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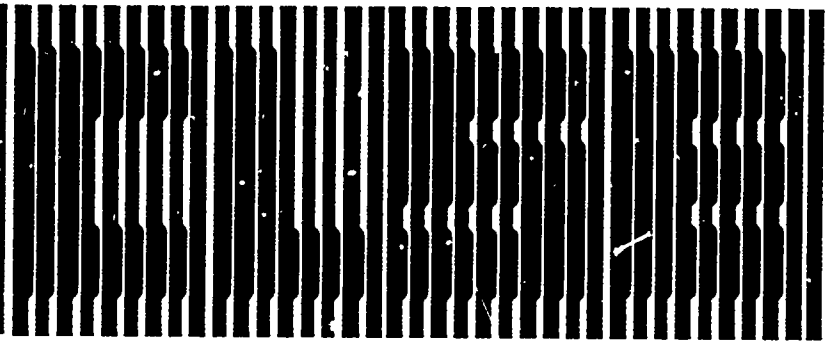
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ABSTRACT

Designed to serve as a means of communication among life science educators who anticipate or are currently using microcomputers as an educational tool, this volume of newsletters provides background information and practical suggestions on computer use. Over 80 articles are included. Topic areas include: (1) using a personal computer in a plant physiology course; (2) annual report and directory; (3) Apple workstations; (4) hypercard; (5) expert systems; (6) computer graphics for simulations; (7) the use of computer simulations to reinforce laboratory experiences; (8) Stella simulation software; (9) software evaluation; and (10) authoring systems. Also provided are guidelines for the preparation and submission of articles to the newsletter, subscription information, indexes for each volume, and lists of meetings and synopses. "Where's the Software?" and "Keeping Abreast of the Literature" are periodic feature articles. (CW)

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# COMPUTERS IN LIFE SCIENCE EDUCATION



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## FIRST STEPS USING PERSONAL COMPUTERS IN A PLANT PHYSIOLOGY COURSE

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I have been enamored of the micro-computer since the days (ancient history!) of the Osborne 1, my first micro. The microcomputer gradually became the indispensable tool for my research and teaching in plant physiology. I used the old ozzie (I called it "Porky" in honor of its oinky-sounding disk drives) for obvious things like writing up lecture notes for class distribution, class handouts, and for keeping grades. I started to write tutorials (not very effective on Porky) and to dream about hooking up Porky to lab equipment, but then came the IBM-PC: goodbye CP/M, hello color.

Microcomputers clearly promise to make instruction in beginning and advanced plant physiology courses more effective and more stimulating to students. All that is needed is the equipment, the time to get it all together and to write programs, and a bit of imagination - a mighty tall order. A NSF grant to a group of several members of our department, including me, made it possible for me to make a start. I now had an XT with a 10 Mb hard disk permanently available in the teaching lab, plus 1 to 3 others that I could borrow, along with an A/D board and a digitizing tablet. Time to quit dreaming and

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get started.

Since we began, the beginning one-quarter course in plant physiology has been given twice. The following sections describe the major modifications made in the course to utilize microcomputers.

#### LABORATORY DATA ACQUISITION AND ANALYSIS

I modified four laboratory exercises to utilize microcomputers for data acquisition (via either the serial port or an internal A/D board) and data analysis. I also wrote simple user interfaces in BASIC. The four laboratories dealt with transpiration, leaf temperatures, CO<sub>2</sub> uptake in photosynthesis and measuring pH changes in chloroplast suspension.

##### Transpiration

In an exercise that I have used for several years, students measure the rate of transpiration of 2-week-old bean seedlings employing a weighing method. Data are collected on the effects of an anti-transpirant chemical and on the effects of light versus darkness. The plants that are used have two expanded leaves and are growing in a small pot. The total weight of each potted plant is less than 100 g (this is important because of the limited capacity of the top-loading mg balance that is used).

Prior to taking transpiration rate measurements, the pots are sealed with tape to prevent water loss from the soil, and any growth above the two expanded leaves is severed. Thus, essentially all water loss is through the two leaves. The students determine the transpiration rate in units of mmol water lost per m<sup>2</sup> of leaf area per second.

In previous years, leaf area was determined by weighing cut-out paper leaf tracings that were then compared to the weight of 0.01 m<sup>2</sup> piece of the same paper. The rate of water loss was determined from weight measurements made at the beginning and end of a 30 minute period.

Because of the time necessary for students to gain an accurate understanding of the written instructions once in the lab setting, and for actually carrying out the required steps of the experiment, it was necessary in the past to

assign different treatments to different student teams. Each team posted its results on a blackboard at the end of the period, and at the next lab period, all students copied all data and prepared a report.

I realized that using microcomputers could improve the efficiency of this exercise in several ways. Data collection and computations could be substantially automated, prompts from the computer could keep the students on track, and final data reports could be accurately prepared. Summary and interpretation of data, done outside of the lab period, would still be the student's responsibility. With the increased efficiency, I found that *each* team could collect a set of data on *all* variables. Class data were still combined for statistical analysis. The latest implementation of this exercise utilizing microcomputers is outlined below.

**Equipment.** Two stations were set up, each with an IBM PC (with color monitor and one floppy disk drive), a printer, and a top-loading mg balance connected to the PC via the serial port. A large cardboard wind deflector was placed around the balance so that air currents would not influence plant weight measurement.

For light conditions, three high intensity lamps were set up under which plants were placed when not being weighed. A small side room was used for dark conditions.

A digitizing tablet, attached to another PC, was set up at another station. The students had learned how to use the digitizing tablet for measuring leaf area in a previous exercise.

**Software.** I wrote a QuickBASIC program to prompt the students for data entry and to print data reports. This program ran continuously on the computers at the weighing stations. SigmaScan, a commercial software package for use with the digitizing tablet, was used for determining leaf areas in m<sup>2</sup>.

**Protocol.** Each team of 2 students used 2 plants, one of which they treated with an anti-transpirant chemical. The area of the 2 leaves on each plant was determined with the digitizer. After a per-

iod of adaptation under the lamps, the plants were taken to a weighing station for the initial check-in. In response to prompts from the computer, the team members entered their team number, names, leaf areas of their plants, and the name of the anti-transpirant they used. They were then instructed to place each plant, in turn, on the balance. The computer recorded the weight from the balance and time from the computer's system clock. The program then stored the data in a disk file, using the students' team number and the day's date (read from the system clock) for the file name (for example, A50187.DAT, for team A5). The students then placed their two plants back under the lamps.

After 10 to 15 minutes in the light following the first weighing, and when their weighing station computer was free, the team returned with the plants for a second weighing. The plants were next dark-adapted for a few minutes, and then weighings were made at the beginning and end of 10-20 minutes in the dark. During each session with the computer at their weighing station, the student team's file was opened and updated with the new weight and time data.

After making the final weighing, the team requested a report of the results from the weighing station computer. The report included all the data they had entered (names, leaf areas, and leaf resistances that they measured with a Li-Cor autoporometer) and the calculated transpiration rates of each plant in light and dark, based on weight, time, and leaf area data. A report of all class data was subsequently prepared (it included a statistical analysis of the data) and given to each student.

**Results.** The use of microcomputers enabled this lab exercise to run more smoothly than in previous years, and each team obtained more data. The students were impressed with the capabilities of the computer and appeared to take the exercise more seriously than in the past.

##### Leaf temperatures

I have run a demonstration exercise on the effects of radiation, wind speed,

and other factors on leaf temperature. One small thermocouple is pressed against the lower surface of a leaf, while another shielded thermocouple is used for air temperature. In the past, readings from both thermocouples were recorded on a two-pen chart recorder. I decided to use the microcomputer for this exercise.

An IBM PC with a Metra-Byte A/D board (model Dash-8/Exp-16) was used in place of the chart recorder. Thermocouples were attached directly to the Exp-16.

In the first implementation of the computerized version of this exercise, I used Labtech-Notebook Software. For the second, I wrote a BASICA program, that was easier to use and set up. In each case, a run-time graph of temperatures was displayed on the computer screen.

Leaf and air temperatures were recorded under various conditions of radiation and wind and saved in an ASCII file. A screen dump of the computer screen on-line graph, or a graph produced after the data was imported into Lotus 1-2-3, was given to each student for inclusion in his or her report on the demonstration. Graphing the data on the screen using Lotus provided an "instant replay" of the temperature records.

The data was easier for the assembled class to see from the computer screen display than from the chart recorder used in the past. The ability to re-run a graph of the temperature versus time data using Lotus made reviewing the results more dynamic. Furthermore, setting up the demonstration is easier using the computer than using the chart recorder.

#### Photosynthesis - CO<sub>2</sub> uptake

For several years, I have used an exercise in which students make measurements of the rate of photosynthesis using an infra-red gas analyzer and the CO<sub>2</sub> depletion method. Briefly, a leaf is placed in a small plastic box under lights. A 10 ml sample of gas is extracted from the box with a syringe, and, after 1 to 4 minutes, another gas sample is extracted. The gas samples are injected into CO<sub>2</sub>-free air passing through the sample port of the infra-red

gas analyzer. The CO<sub>2</sub> content of each sample is determined, and the difference between the two calculated.

Based on the volume of the box, the change in CO<sub>2</sub> content, the area of the leaf, and the elapsed time between sample extractions, the photosynthesis rate can be calculated in micromols of CO<sub>2</sub> taken up per m<sup>2</sup> per second.

I modified the exercise to include data acquisition by a microcomputer. In the past, students took data from a chart recorder tracing, a process that was time consuming, prone to error, and required a calibration and conversion from chart paper units to ppm CO<sub>2</sub>. Use of the microcomputer promised to eliminate these problems. I did decide, however, to leave final calculation of photosynthesis rates up to the students. In the future, I will provide a program that will allow students to confirm their calculations, but still not do it for them!

The infra-red gas analyzer was connected to a Metra-byte A/D board (model Dash-8/Exp 16) on an IBM PC. CO<sub>2</sub>-free air flowed through both the sample and reference ports of the infra-red gas analyzer. Students injected 10 ml samples of air into the air flowing into the sample port.

A digitizing tablet attached to another microcomputer was also set out. This was used for leaf area determination (as noted above, the students had learned to use it in earlier exercises).

I wrote a BASICA program that used a simple user interface to accept data, carry out calibration calculations, and determine the CO<sub>2</sub> content of samples.

One or more students were enlisted to inject calibration gases for the calibration procedure. Each student team was then instructed to obtain a sample of outdoor air for CO<sub>2</sub> content determination. (I had them do this mainly to help reinforce classroom discussions of the CO<sub>2</sub> greenhouse effect, and implications for plants in the future. We're keeping a year to year database of the results.) Each team then obtained CO<sub>2</sub> measurements for photosynthesis determinations at 2 or 3 different light intensities. Finally, the CO<sub>2</sub> compensation point of a particular species was determined by each team.

Use of the A/D board and the microcomputer gave students the CO<sub>2</sub> con-

centration immediately after sample gas injection, helping to maintain their interest.

#### Photosynthesis - pH change in chloroplast suspensions

As part of an exercise on the light reactions of photosynthesis using suspensions of isolated chloroplasts, I conduct a demonstration on change of the bathing solution pH upon illumination (an apparent result of proton uptake into the chloroplast thylakoids). In the past, I used a chart recorder attached to a sensitive pH meter to record changes in pH. I have now switched to using a microcomputer and an A/D board in place of the chart recorder. The benefits of using the microcomputer in this exercise are similar to those mentioned above in the case of the exercise on leaf temperatures.

#### LAB DATABASES

For several of the lab exercises used in this course, I ask the students to gather and analyze all of the data obtained by the whole class. In some cases, this is so that class averages, rather than just one data set, may be used for drawing conclusions. For example, in the case of the effect of auxin on the number of adventitious roots formed on cuttings, there is considerable variation from student team to student team. In other cases, this is done to expand the number of species sampled. For example, in determinations of the frequency (number per area) of stomates on leaf surfaces, different teams are asked to make counts on different species.

In the past, I had the students post their team's data on the board, and when all data have been posted, all class members were asked to copy all of the data for use in preparing reports.

I now use Lotus 1-2-3 custom worksheets for this purpose. Student teams enter their data, and when all class data has been entered, each student has the computer print a summary. Lotus macros were used to make friendlier user interfaces for the worksheets. I will accumulate data from year to year for certain databases, for example, the stomate frequency versus plant species database.



## TUTORIALS

I have written five BASICA microcomputer tutorials to go with this course. They are available on the hard disk of the plant physiology lab PC and also in the computer lab at the University's library. Students may also copy these onto their own personal disks, if they desire.

I plan to write additional tutorials in the future as time allows using BASIC or other formats such as Show Partner. The latter promises to make tutorial writing a much easier and more rewarding task, although I have just begun to explore its use.

The five tutorials include the following.

- Hangperson, a word game designed to foster familiarity with plant physiology terminology.
- Transpiration Rate Tutorial, based on mathematical models of plant transpiration, allows students to enter various parameters (such as wind speed, air humidity, and leaf resistance) to see how they would affect the transpiration rate.
- Spectrum, a tutorial to familiarize students with the spectrum of visible light, providing drill in equating colors with corresponding wavelength ranges.
- Hit the Reaction Center, a graphics representation of photon capture, energy transfer, and fluorescence in the light reactions of photosynthesis.
- The Calvin Cycle, a tutorial designed to give the student a familiarity with the overall process of carbon dioxide fixation in photosynthesis and to gain familiarity with the major intermediate substances involved.

## ADVANCED PLANT PHYSIOLOGY

I used our microcomputers extensively during the most recent offering of my one quarter course called Advanced Plant Physiology. The students, who are generally well motivated, are expected to carry out more involved lab projects with less direction than is the case in the beginning course. I decided to have the students use Lotus 1-2-3 extensively for data handling and graphing, as I have found it very useful

for these purposes in my research.

I introduced the students to this spreadsheet software during the first week. I taught them the rudiments of data entry and graphing and encouraged them to study the software manual to increase their expertise in using the package. In the various exercises they performed in subsequent weeks, they became adept at using Lotus 1-2-3 for data handling and analysis and for preparing tables of data and graphs for their reports. Of course, the students learned how to use the Metra-Byte A/D board and the digitizing tablet for use in their lab investigations (I do all the setting-up in the beginning course).

## CONCLUSIONS

Although the full potential for using microcomputers in plant physiology courses is far from being realized, the experiences described above have convinced me of the value of micros. In evaluating the course, students often mention, in a positive way, their experience with one or another of the computer applications, and they say they would have liked more. With time and the acquisition of additional equipment, the courses will surely be made more stimulating learning experiences through the use of microcomputers. I want to write more tutorials, especially interactive ones. The software tools that are available for programmers of limited sophistication (like me) are certainly adequate for writing valuable, graphics-based tutorials. It just takes a lot of patience and imagination. I also want to extend the number of lab exercises using micros for data collection and analysis. Their use should illustrate how things are done in the real world, as well as be instructional. I would like the students to be more independent in using the micros in the lab (a lot of hand-holding has been necessary so far). At some point in the future, I will require that each student provide a floppy disk on which to store a set of programs and lab data.

To anyone teaching courses like this who hasn't yet gotten into the world of microcomputers, I would heartily recommend that you do so. It will bring a new spark to your enjoyment of teach-

ing (although expect some periods of frustration!), and it will surely enhance the educational experiences of your students. I assure you that the possibilities are so numerous that you will never, in a dozen careers, run out of new things to try in the quest for the best ways of getting your educational messages across.

The tutorials described are available free to anyone who might like to try them. They were written in BASICA for a CGA card and are GW-BASIC compatible. Please send a stamped, self-addressed mailer and a blank 5 1/4" disk to: Larry Blakely, Biological Sciences Dept, California State Polytechnic University, Pomona, CA 91768.

## SUBSCRIPTION INFORMATION

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# DIAGNOSTIC PRACTICE REINFORCES LESSONS FROM LABORATORIES

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In the past, simulations have been widely heralded as powerful learning vehicles. A large number of such models have been created. A common property of these models is interaction among a large number of system properties producing a range of responses in system variables. Instructors using these simulations have used a variety of strategies, ranging from non-directed suggestions that students "play" to prescribed "experiments" with specified data recording, reporting, and analysis. Presumably, those teachers used simulations because they felt models would help students develop an appreciation for system interactions, compensations, and limits of function. The success of that supposition remains largely conjecture. As a modeler, I know that "playing" with a model develops my "feel" for it, but does the average student play as systematically?

The goal in giving students simulations is for them to build a framework for integrating system interactions. To achieve this, the learner must focus on predicting model behavior, not watching passively. One very successful way to use a simulation is in small group discussions where the teacher asks, "If I increase this parameter, can you predict what will happen to that variable? How much? Why?" This is cause to effect reasoning. Certainly, it represents the type of mental dialog we hold with ourselves when exploring a system. Prediction requires more effort and sophistication than watching, and students are less prone to adopt it. Consequently, most students truly play with simulations rather than study the operation predictively.

Previously, we reported a study of veterinary medical students in which, after a "traditional" lecture and wet lab unit on the cardiovascular system, they still held strongly anatomic views of

the cardiovascular system.<sup>3,4</sup> Subsequently we asked them to solve a series of "diagnostic" problems. These problems were cast as games in which one cardiovascular property from a list of 8 was disturbed randomly either up or down. After playing 50 faults, we reassessed their perceptions of the system. Their views had developed a new functional dimension, and that new dimension had shifted in the direction of an expert's understanding.

Our diagnosis game helped students pose questions that are the reciprocal of those described above. They were asking, "If this variable has changed, which properties could explain it?"

This is effect to cause reasoning. Our program is called the Fault Identification Game. Like clinical experiences, the games reinforce both a functional understanding of the specific system and general problem-solving skills.

This paper will show how new games can be written to compliment many other simulations.

The data representations within the game are very general. Problems can be created for any system that can be described in terms of properties (maximum 20), variables (maximum 20) and the interrelationships between them. We define "properties" as those entities that are independent variables, that is, external inputs to the simulation. Our "variables" are dependent values that arise from the interactions between properties. The game is designed to permit non-programmers to develop problem sets easily. Creating a problem requires entering data about a system into a specified format, using either a text editor or a spreadsheet. Required data are shown in Table 1.

## AN ISOLATED HEART EXAMPLE

In the Isolated Heart preparation of the author's Cardiovascular Systems and

Table 1. Data required to create a problem.

*For the problem:*

1. A short "presenting complaint"
2. A short closing discussion

*For each property and variable:*

1. Name, abbreviation, and units
2. An order describing where (and if) each will be displayed

*For each property:*

1. A value
2. The value's deviation from normal (+,-,0)

*For each variable:*

1. A flag describing the variable as either quantitative or qualitative
2. A deviation from normal (+,-,0), or qualitative description (eg, "missing", "open", "elevated")

Dynamics (CVSAD) program<sup>4</sup>, four properties determine the performance of the heart: preload, afterload, heart rate and left heart pump state. If the values of these properties deviate from normal, the new state of the system will be reflected in values of its variables: cardiac output, stroke volume, end systolic volume, end diastolic volume, average arterial pressure, and peak arterial pressure.

In this example, one or more of the properties was changed from normal. Some of the variables are abnormal, others show no change. In Figure 1, the player has requested values of arterial pressure, cardiac output, end diastolic volume and most recently, stroke volume. The game reports SV is 22 ml and that this is normal (for a dog). Some possible faults can be eliminated and some are clearly suggested. At this point, the player begins to enter a tentative hypothesis, (Figure 2, right). A hypothesis is recorded as a

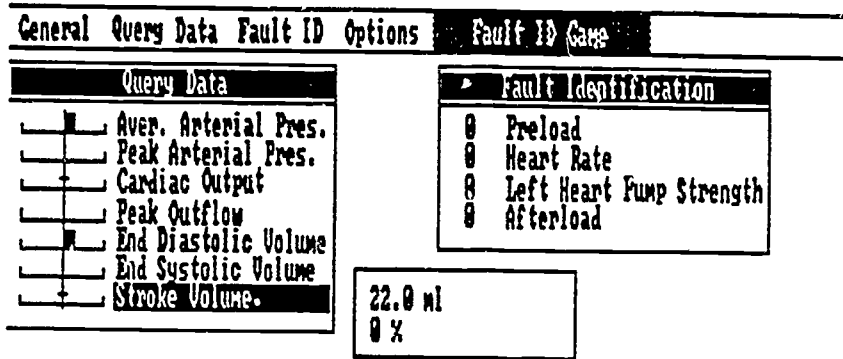


FIGURE 1. Querying data from the Fault Identification Game. Possible data are shown on the left. Possible faults are shown on the right.

setting (-,0,+) for each property. In this case, the diagnosis reads increased preload, decreased left heart strength, and others normal.

When a diagnosis is entered, the computer can be asked to evaluate it. In this case, the answer is partly right, as shown in Figure 2, bottom. The response is reported as a Venn diagram showing fault(s) suspected (S for suspected) by the player and fault(s) actually contained in the problem (D for disturbed). Verbal confirmation is also provided. Correct answers show per-

fect intersection of sets "S" and "D", partial answers show partial intersection, and wrong answers show no intersection. In this case another aspect of cardiac failure remains to be diagnosed.

#### AUTHORING A PROBLEM

The previous example comes from a simulation developed by the authors. It is equally possible to create problems from other simulations. It is equally possible to create problems from other simulations. It is even possible to cre-

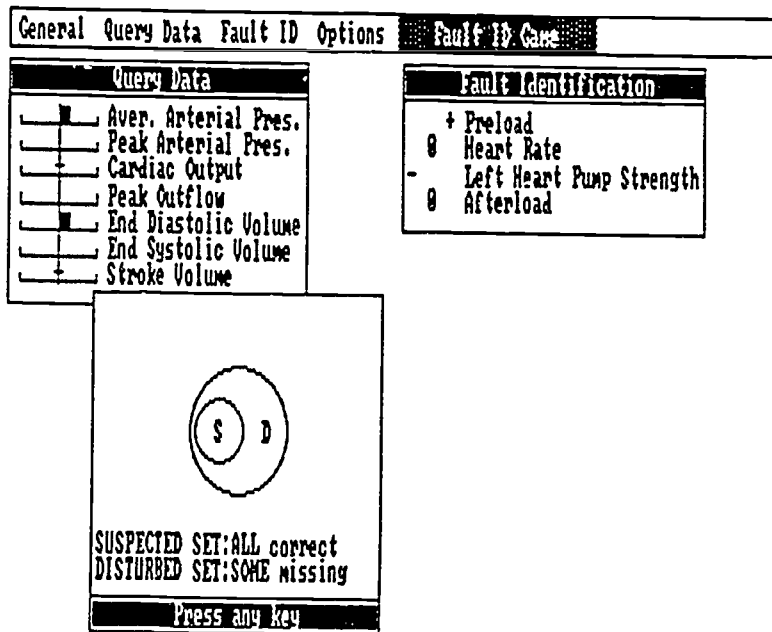


FIGURE 2. Checking a diagnosis. The diagnosis is entered at the top left as high preload, low heart strength, others normal. The computer says the answer is partly correct, but something(s) is still missing.

ate useful student problems from a model that is otherwise too complex for instructional purposes. The central task is phrasing the problem with only those properties and variables that are appropriate to the student's academic level. The following examples show the process of authoring a problem and some of the diversity of material that can be encoded.

The data for a problem are stored in a compacted file format. We have given these files DOS filename extensions .SDF, for System Data File. Problem authors need not concern themselves with the organization of SDF files, rather they may fill in textual forms in either wordprocessor (ASCII) or spreadsheet (DIF) formats. In each case, a utility program converts SDF files to and from the more familiar formats. We chose wordprocessors and spreadsheets as simple interfaces faculty already have experience using. Further, they provide a potential hardware independent link for transferring faults from one computer to another because the SDF format is machine dependent.

#### Encoding a clinical example with a wordprocessor

In *Pulmonary Pathophysiology*, West<sup>5</sup> describes clinical observations for a variety of respiratory diseases. A fault problem can be written from his description of restrictive disease (Chapter 5),

"Spirometry typically reveals a restrictive pattern. The VC [Vital Capacity] is markedly reduced, but the gas is exhaled so rapidly that although the FEV<sub>1.0</sub> [Forced Expiratory Volume in 1 second] is low, the FEV/FVC% [Forced Vital Capacity] may exceed the normal value. The square shape of the forced expiratory spirogram is in striking contrast to the obstructive pattern. The MMFR [Maximum Mid-Expiratory Flow Rate] is normal or high. The flow volume curve does not show the scooped out shape of obstructive disease, and the flow rate is often higher than normal when related to absolute lung volume."<sup>5</sup>

"The arterial PO<sub>2</sub> and PCO<sub>2</sub> are typically reduced and the pH is normal.



Table 2. Data extracted from problem in West's *Pulmonary Pathophysiology*

Variable name	Value	Units	% from normal
Forced Vital Capacity	2.9800	liters	-38
Forced Exp Vol - 1 Sec	2.8800	liters	-30
FEV <sub>1</sub> /FVC	0.9700	unitless	13
Max Midexpiratory Flow	4.7600	l/sec	6
Peak Expir Flow	5.0000	l/sec	-50

The hypoxemia is usually mild at rest until the disease is advanced. However, on exercise the PO<sub>2</sub> often falls dramatically and cyanosis may be evident."<sup>5</sup>

The data in Table 2 were extracted from this description. These constitute the variables for the problem. The "faults" can be encoded in one of two ways. One set would focus on the clinical description such as restrictive disease or obstructive disease. Alternately, the fault could be encoded in terms of defects in functional entities such as lung compliance and airway resistance.

Note that not all the diagnostic in-

formation mentioned by West was used. The data selected are at the discretion of the problem's designer. However, adequate data must be present to distinguish this fault from others in the same problem set.

Table 3 shows how the data from Table 2 are entered into the wordprocessor template generated by the Fault Game's utility. One data item is entered to the left of the descriptions marked in single angle brackets. Double bracketed terms indicate a division within the data record and have no data associated with them. After entering all the data into this file, it is processed back into the compact SDF used by the game.

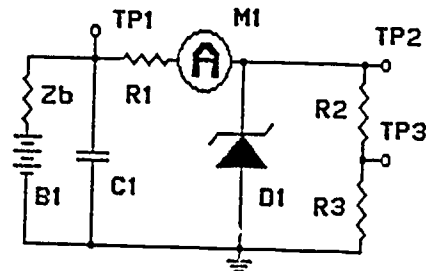
Table 3. Encoding properties and variables in a wordprocessor

<<variable 1>>	
Forced Vital Capacity  <name>	
FVCI <abbr>	
1 <active var?>	1=yes, make active in this game
1 <index>	place data in cell 1 of display
-38 <dev>	
liters <units>	
0 <binary var?>	0=no, its not binary data
1 <display value?>	1=yes, display data in the problem
2.9800 <value>	
<<property 1>>	
Compliance <name>	
Compl <abbr>	
1 <active prop?>	1=yes, show it in this game
1 <index>	place this fault in display position 1
-1 <qual value>	this property is below normal
0 <binary prop?>	0=no, its not binary
0.0000 <value>	actual value is unknown from West's text, but it won't be shown in that game, so its not necessary

Programming a model into a spreadsheet Perhaps the most powerful method of entering data is via a spreadsheet capable of writing DIF (Data Interchange Format) files. Most spreadsheet programs have this capacity. In this example, Twin (Mosaic Software), a "work alike" of Lotus 1-2-3, was used. A file called FIGPLATE.WKS, provided with the game, is a blank template to be filled with author's data. The critical parts of the sheet are protected to prevent inadvertent corruption of the format.

The equations for the model were entered into the bottom of the sheet. All the functions and indirect referencing features of the spreadsheet were used to relate the various properties and variables. This master model can be saved to disk and reloaded. Faults are created by altering desired properties and letting the model in the spreadsheet recalculate the variables. Just a few key strokes are required to modify the sheet and save the data for a new problem.

To augment a course in electronic troubleshooting, a circuit model was developed using the schematic in Figure 3. Nine properties were used: battery internal impedance, capacitor (dc) impedance, resistors R-1, R-2 and R-3, diode



B1 6.3 volts Zb=0.1Ω  
 R1=5Ω R2,R3=1000Ω  
 D1=5.0 volt Zener  
 C1=10 μF  
 M1=0-2 amps

FIGURE 3. Circuit diagram for an electric circuit model that was developed in a spreadsheet and then had its data exported to FIG game datasets.

impedance, D1, battery voltage, and the total impedance of the external circuit. A portion of the spreadsheet (Table 4) shows the layout.

In building the model, some properties, such as resistances and voltages, were direct analogs of the physical component they represented. The functional impedance of the zener diode, however, was quite complex. A reverse biased zener diode exhibits high impedance (nearly an open circuit) until the zener voltage is reached. From then on, it shows a voltage dependent impedance that maintains the voltage drop across the diode at exactly the zener voltage. Using the logical @IF function of the spreadsheet, these two possibilities were modeled. When the fault prevented the voltage at TP2 from exceeding 5.0 volts, the diode impedance was 1010 ohms (used for open circuits), otherwise an impedance was calculated based on current flow in the circuit to provide zener action. Variables presented to the student were limited to the voltages present at the test points (relative to ground) and the current in Meter M1. Resistance measurements could easily have been allowed, but the design goal for the model was to teach active, in-circuit troubleshooting.

The model used rather complex formulas to compute the variables. For



example, the Test Point 1 voltage was computed from the battery voltage (E<sub>batt</sub>), the external system impedance (R<sub>s</sub>), and the internal or battery impedance (R<sub>b</sub>) using the following formulas.

$$\text{Total current (I}_{\text{total}}) = E_{\text{batt}} / (R_s + R_b)$$

$$\text{Voltage of Test Point 1} = I_{\text{total}} * R_s$$

In the normal circuit, the calculated voltage was 6.2743 volts. Faults were either shorts ( $R=0$ ) or opens ( $R=1010$  ohms) in the resistors, except for the battery impedance which was merely increased to simulate a "weak" battery condition. Only single fault sets were used in this case, but multiple faults could be used. A further refinement would be to add ratings, such as power, to some components. The spreadsheet could then evaluate to see if the instructor induced fault caused an overload somewhere else that "took out" additional components.

#### CUSTOM VERSUS GENERIC FAULT GAMES

Boyle and Robinson described an acid-base balance game.<sup>1</sup> In this program the student sees a graphical display showing arterial PCO<sub>2</sub>, plasma bicarbonate (HCO<sub>3</sub><sup>-</sup>), and pH. The normal range, blood buffer line and current patient status are displayed graphically and in tabular form. The player is asked to determine the most appropriate diagnosis for the given acid-base state. The screen then presents a list of diagnoses that represents all combinations of primary and compensatory acid-base states. After a correct diagnosis is selected, the player can select treatment options. These include defining values for respiratory rate and tidal volume and ordering an infusion of acid or basic solution over a 60-minute interval to adjust base excess or deficit. The patient's status is altered on the display during the course of therapy. Tutoring is provided for misdiagnoses and improper therapies.

Boyle and Robinson's fault finding game is a custom application for acid-base balance. The game described here is generic. Acid-base problems could be implemented in the generic engine, but they would lose the graphical interface and the tutorial feedback of the custom version. On the other hand, custom programs require new programming for each application. Our goal was to develop an engine sufficiently flexible that non-programmers could use it to create problem-solving exercises in a wide variety of domains with little instructor time investment.

#### AVAILABILITY OF THE PROGRAM

A preliminary version of the program for the IBM PC is available. The program uses the UCSD P-system and has sample problems along with student and faculty manuals on disk. These may be obtained by sending a blank floppy disk and self addressed envelope (a diskette mailer is best) to the authors. An improved version, used to illustrate this article is being prepared for publication. Problem sets created

for the initial version will be fully compatible with the new one.

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Table 4. Encoding a problem in a spreadsheet

<<PROPERTIES>>			
<name>	<property 1>	<property 2>	<property 3>
<abbr>	Battery	Impedance C1	Limit Resistor
<active prop?>	Z <sub>batt</sub>	Z <sub>cap-1</sub>	R-1
<index>	1	1	1
<qual value>	1	2	3
<binary prop?>	0	0	0
<value>	0	0	0
	0.10000	10000000.0000	5
<<VARIABLES>>			
<name>	<variable 1>	<variable 2>	<variable 3>
<abbr>	Test Point 1	Test Point 2	Test Point 3
<active var?>	E-TP1	TP-2	TP-3
<index>	1	1	1
<dev>	1	2	3
<units>	0	0	0
<binary var?>	Volts	Volts	Volts
<display value?>	1	1	1
<value>	1	1	1
	6.2743	4.9899	2.4950

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**NRCLSE**

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## NRCLSE ANNUAL REPORT FOR 1987

The overall purpose of NRCLSE is to cultivate collaborative efforts among faculty with expertise in using computers in life science education. The broad goal of the Resource is fourfold:

- 1) to educate faculty in effective uses of computers in the curriculum;
- 2) to promote research aimed at evaluating new applications of the computer to life science education;
- 3) to promote development of a critical mass of high quality, versatile software; and
- 4) to serve in a consultant capacity for life science faculty currently active or interested in becoming active in this area.

This year's activities were focused on further establishing the Resource as a viable entity capable of serving the life science community. A statement of NRCLSE's financial status is presented in Table 1.

### GENERAL OVERVIEW OF ACTIVITIES

In 1987, NRCLSE applied for and received non-profit [501(c)(3)] status from the Internal Revenue Service. In addition to making us eligible to receive tax deductible donations, the IRS approval of our non-profit status makes us eligible to apply to various foundations and funding agencies for support of specific projects. We have begun a process to identify foundations whose interest areas include use of computers in life science education as an initial step in developing future projects.

Publication of *Computers in Life Science Education* continues to be our primary activity. However, during the past year, we have continued our software development project in respiratory physiology, provided information to colleagues throughout the world concerning use of computers in life science curricula, participated in regional and national meetings, and

conducted demonstrations and workshops at a number of institutions.

Computers in Life Science Education Subscriptions to CLSE continue at the same level as in previous years, with close to half of the subscriptions being held by libraries. The geographic distribution of subscribers continues to grow. Distribution now includes subscribers in 10 countries representing 4 continents.

In addition to our regular features, "Keeping Abreast of the Literature" and "Where's the Software?", we have begun publishing an annual colleague directory of life science educators interested in using or actively using the computer as a teaching tool. The intent in providing this information is to help life science educators establish links with potential colleagues. Entries for the directory have been drawn primarily from responses to the NRCLSE questionnaires published in CLSE.

#### Software development

Work on conversion of our Apple II package of simulations (Simulations in Physiology - The Respiratory System) to MS-DOS and Macintosh formats continued during this year. All three packages are now complete and available for distribution. Appropriate announcements and purchase information for each set was included in Computers in Life Science Education when each set was completed.

#### Resource Information

During 1987, NRCLSE responded to over 75 requests for information concerning use of the computer as an educational tool. The majority of these concerned suitable software for use in the laboratory setting.

The geographic origin of the requests indicates that we are beginning to accomplish our goal of providing widespread service. Requests originated from institutions in Australia, Belgium, Canada, Denmark, England, Italy, Jamaica, Mexico, Spain, Sweden, and the United States (24 states, the District of Columbia, the U.S. Virgin Islands, and Puerto Rico).

Table 1. NRCLSE Financial Report for year ending December, 1987.

Fund Balance, December 31, 1986		\$14,928
<b>Revenues</b>		
Cash donations		845
Equipment donations		0
Subscriptions		5,080
Software sales		<u>2,008</u>
Total Revenues		7,933
<b>Expenses</b>		
CLSE production		4,182
Operating supplies		2,102
Contractual services		2,258
Travel		567
Software purchases		<u>141</u>
Total Expenses		9,250
Fund balance, December 31, 1987		\$13,612

**Presentations/Consultation**  
Personnel from NRCLSE organized or participated in symposia, panels, and workshops at 5 regional and national meetings during the year. Workshops and demonstrations were also presented at two universities. Topics included teaching problem-solving in physiology, the computer as a teaching aid in biological sciences, the role of the computer in the lecture hall, future trends in physiology teaching, and computer applications other than traditional computer-aided instruction in health science education.

#### Establishing a peer critique mechanism for software

In 1987, we began our effort to establish a peer critique mechanism for reviewing software by initiating a dialog on the topic in Computers in Life Science Education. This effort will continue in 1988 with our first goal being to establish criteria for such a mechanism. To get as much input as possible before establishing specific criteria, we will survey other groups already involved in peer review of software, and the two articles published in CLSE will be reprinted in Science

Software, a quarterly publication serving the science community.

#### SPECIAL THANKS

NRCLSE extends a very special "Thank you" to the following people, organizations, and institutions for their support in 1987.

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# CLSE 1987-88 COLLEAGUE DIRECTORY - PART I

The primary goal of the National Resource for Computers in Life Science Education (NRCLSE) is to cultivate collaborative efforts among life science faculty interested in using the computer as a teaching tool.

The directory that follows is updated from the 1986-87 directory and was drawn primarily from respondents to questionnaires printed in CLSE. It is intended to help readers identify col-

leagues with common interest areas.

The listings are arranged by the content areas identified in response to the question, "What content areas do you teach?" As a result, entries may appear under more than one heading.

Although every attempt has been made to ensure that the information is current and correct, it is likely that some errors appear in this list. We apologize in advance for any inconve-

niences that may arise due to such oversights. Part II of the directory will appear next month.

If you are aware of other colleagues that should be listed, please send their names, addresses, phone numbers, and teaching content areas to NRCLSE, Mail Stop RC-70, Univ. of Washington, Seattle, WA 98195 or let us know via BITNET. NRCLSE's BITNET address is MODELL@UWALOCKE.

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**AIMS AND SCOPE**

The goal of *Computers in Life Science Education* is to provide a means of communication among life science educators who anticipate or are currently employing the computer as an educational tool. The range of content includes, but is not limited to, articles focusing on computer applications and their underlying philosophy, reports on faculty/student experiences with computers in teaching environments, and software/hardware reviews in both basic science and clinical education settings.

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Articles consistent with the goals of *Computers in Life Science Education* are invited for possible publication in the newsletter.

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Articles submitted for publication should not exceed 2000 words and should be typewritten, double spaced, with wide margins. The original and two copies including two sets of figures and tables should be sent to the Editor: Dr. Harold Modell, NRCLSE, Mail Stop RC-70, University of Washington, Seattle, WA 98195.

Title page should include full title, list of authors, academic or professional affiliations, and complete address and phone number of the corresponding author.

Illustrations should be submitted as original drawings in India ink or sharp, unmounted photographs on glossy paper. The lettering should be such that it can be legible after reduction (width of one column = 5.7 cm).

Reference style and form should follow the "number system with references alphabetized" described in the Council of Biology Editors Style Manual. References should be listed in alphabetical order by the first author's last name, numbered consecutively, and cited in the text by these numbers.

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**NRCLSE**

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## CLSE 1987-88 COLLEAGUE DIRECTORY - PART II

The primary goal of the National Resource for Computers in Life Science Education (NRCLSE) is to cultivate collaborative efforts among life science faculty interested in using the computer as a teaching tool.

The listing that follows is a continuation of the 1987-88 directory published last month. It is updated from the 1986-87 directory and was drawn primarily from respondents to questionnaires printed in CLSE. It is intended to help readers identify colleagues with common interest areas.

The listings are arranged by the content areas identified in response to the question, "What content areas do you teach?" As a result, entries may appear

under more than one heading.

Although every attempt has been made to ensure that the information is current and correct, it is likely that some errors appear in this list. We apologize in advance for any inconveniences that may arise due to such oversights. The directory will be completed in next month's issue.

If you are aware of other colleagues that should be listed, please send the pertinent information (name, address, phone number, BITNET address, teaching content area) to NRCLSE, Mail Stop RC-70, Univ. of Washington, Seattle, WA 98195 or let us know via BITNET. NRCLSE's BITNET address is MODELL@UWALOCKE.

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The goal of *Computers in Life Science Education* is to provide a means of communication among life science educators who anticipate or are currently employing the computer as an educational tool. The range of content includes, but is not limited to, articles focusing on computer applications and their underlying philosophy, reports on faculty/student experiences with computers in teaching environments, and software/hardware reviews in both basic science and clinical education settings.

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Articles consistent with the goals of *Computers in Life Science Education* are invited for possible publication in the newsletter.

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Reference style and form should follow the "number system with references alphabetized" described in the Council of Biology Editors Style Manual. References should be listed in alphabetical order by the first author's last name, numbered consecutively, and cited in the text by these numbers.

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## APPLE //e MEDIATED WORKSTATIONS IN THE UNDERGRADUATE PHYSIOLOGY LABORATORY

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Several years ago, the National Science Foundation established the College Science Instrumentation Program (CSIP) to upgrade laboratory instrumentation in undergraduate science education. I was the principle investigator for a CSIP award in 1985 (#CSI-8557398) that permitted me to assemble five Apple //e-mediated workstations for use in my physiology laboratories. The workstation concept exploited the idea that a microcomputer, when linked to an appropriate environmentally sensitive interface, could be transformed into a versatile, data-gathering laboratory instrument.<sup>2,3</sup> I developed software to permit use of the Apple //e

monitor screen as a strip chart recorder and, when the desired screen image was obtained, students could employ a dot-matrix printer to make hard copies of the graphics screen for their lab notebooks. By simply changing the interface or the controlling software, a single workstation could be made to act like many single-function laboratory instruments. For example, my students have used an inexpensive, self-constructed photosensitive interface to transform the Apple //e into a transmission densitometer to study red blood cell hemolysis,<sup>3</sup> a kymograph to investigate frog skeletal muscle contraction,<sup>3</sup> a pneumograph to investigate human

respiratory mechanics and as a device to measure human reaction time in response to audio or visual stimuli. Several types of commercially available interfaces were also employed to allow the Apple //e to act as a pH meter (HRM Software, 175 Tompkins Ave, Pleasantville, New York 10570) and a digital oscilloscope (W.H. Nail Co, 1096 C oro Dam Blvd., Oroville, CA 95965) for electrophysiological studies. For an excellent description of the use of the Apple as a digital storage oscilloscope, I refer you to a report by Mary Coyne.<sup>1</sup>

In this report, I would like to focus on the photosensitive interface and its application in the undergraduate physiology laboratory. The simplicity of the photosensitive interface may be seen in the diagram of the schematics shown in Figure 1. An FPT-100 phototransistor is used to measure light intensity by changes in resistance. A 16 pin Dual Input Plug (DIP) is used to connect the collector pin and the emitter pin of the phototransistor to the +5 Volt and ground pins of the Apple game I/O connector. The DIP and phototransistor plus the "breadboard" (microcircuit board) which hold the components cost under \$20.00 to construct and can be obtained from local or regional electronics suppliers.

#### SOFTWARE DESIGN

Applesoft BASIC was used to write the data sampling and display programs for

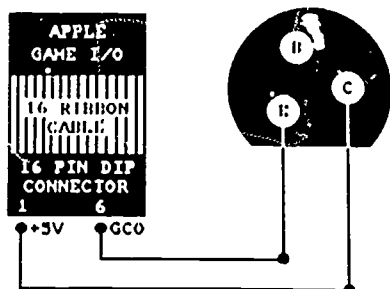


FIGURE 1. Schematic for photosensitive interface. A view of the underside of an FPT-100 phototransistor (upper right) showing placement of the pins and their connection to the DIP connector. E=emitter pin, B=base pin, C=collector pin. Pin 1 of the DIP is the +5 volt source, and pin 6 is the ground (GND) connection.

all experiments in this report. These programs cause the monitor screen to act like a strip chart recorder by drawing a single trace across the high resolution graphics screen from left to right. In order to do this in BASIC, the statement  $Y = PDL(0)$  was used to sample the interface resistance reading and return a value ranging from 0 to 255. The value for Y determines the pixel to be lit on the vertical axis of the graphics screen. The movement of the trace from left to right is driven by FOR-NEXT loop statements that increase the value of X by one each time the loop is executed. When the statement, H PLOT X,Y, is included within the FOR-NEXT loop, pixels are lit in sequence from left to right to simulate a strip chart recording.

A Grappler card in expansion slot 4 connects to a dot-matrix printer and gives students the option to print the high resolution graphics screen in the desired format. Selected screens may also be saved in floppy disk files for review at a later time.

#### STUDENT APPLICATIONS

Students used the photosensitive interface as a transmission densitometer for two types of experiments with red blood cells (RBCs). First, slow hemolysis kinetics of RBCs were followed after introduction of the RBCs into solutions of 0.5 M urea, glycerol or glucose.<sup>3</sup> In the second type of experiment, students examined the effects of tonicity on RBC shape and size. For both experiments, a 3 Volt miniature lamp was positioned so that its light traveled through a test tube containing the RBC suspension before falling on the phototransistor. The results from a typical slow hemolysis experiment in which an RBC suspension was introduced into a solution of 0.5M glycerol can be seen in Figure 2. When RBCs were added, the trace immediately moved upward indicating less light falling on the phototransistor due to an increase in light scattered by the RBCs. Thereafter, the slow downward trace movement to the baseline gave a measure of hemolysis time and indicated that more light reached the phototransistor as light scattering decreased due to RBC swelling and eventual hemoly-

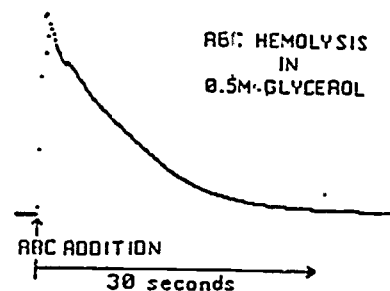


FIGURE 2. Red blood cell slow hemolysis kinetics following addition to 0.5M glycerol. A Digi-View digitizing system (Newtek, Inc., 701 Jackson Suite B3, Topeka KS 66603) was used to transfer the original results from the Apple //e monitor to an Amiga 1000 computer serving as a low cost graphics workstation to permit addition of explanatory text using an Electronic Arts Deluxe Paint program. A photograph was then made using a Polaroid Palette system. All other figures and the single table were produced using the Amiga graphics workstation.

sis. Hemolysis time under these conditions also indicates how fast molecules pass through the RBC membrane. Students find that urea solutions hemolyze RBCs in under 5 seconds, glycerol solutions exhibit hemolysis times around 30 seconds, and in glucose solutions, hemolysis is still incomplete even after 90 seconds. Students may conclude that there is a relationship between molecular size and the ability to pass through a membrane.

The results of tonicity studies in which RBCs were suspended in different NaCl concentrations are shown in Table 1. Transmitted light decreases as the salt concentration increases to 2%. By observing RBC morphology with

Table 1. Use of photosensitive interface to measure light transmitted through RBCs suspended in different salt concentrations. Higher interface readings indicate more light falling on the phototransistor.

TRANSMISSION DENSITOMETRY	
LIGHT TRANSMITTED BY RBC'S IN DIFFERENT CONCENTRATIONS OF SODIUM CHLORIDE	
SOLUTION	INTERFACE READING
DISTILLED H <sub>2</sub> O	237
0.43% NaCl	213
0.85% NaCl	177
2.0% NaCl	128

the light microscope, students can also relate RBC shape with its ability to transmit and scatter light.

The photosensitive interface was also utilized as an updated version of a kymograph to study frog muscle contraction. A typical kymograph arrangement for muscle study was employed except that a phototransistor hung from the tip of the muscle lever instead of an ink writing stylus, and an Apple //e monitor replaced the revolving drum. The phototransistor extended into a short length of PVP (polyvinylpropylene) pipe and was illuminated by a 3 Volt miniature lamp positioned at the bottom. During contraction, the muscle lifted the lever and phototransistor away from the light source, and when the muscle relaxed, the phototransistor returned to its original position. The detected change in light intensity was displayed on the Apple //e monitor screen. The results in Figure 3 were obtained by measuring individual muscle contraction-relaxation responses to a series of electrical stimuli of increasing voltage. The software was designed to wait for the student team to get everything ready for stimulation. When ready, the team signals the program to enter a 15 second period during which they stimulate the muscle with the selected voltage. If the muscle contracts, the monitor screen displays the detected light change as a single, vertical line above the baseline. At the end of 15 seconds, the program moves the baseline to the right a short distance and waits again for the student team to prepare the muscle for the next voltage stimulus.

The strip chart recorder mode was used to study muscle contraction by increasing the frequency of stimulation with supramaximal threshold voltage. Selected results are shown in Figure 4. As with the traditional kymograph, individual muscle contraction-relaxation cycles are observed at low frequencies of stimulation (Figure 4A), and complete tetanus is attained at higher stimulation frequencies (Figure 4C).

The strip chart recorder mode software may also be used with a Marey tambour for pneumographic study of human breathing mechanics. Again, a typical kymographic setup was em-

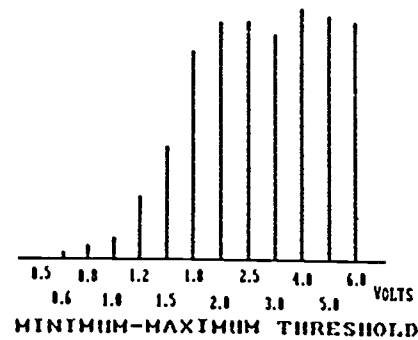


FIGURE 3. Response of frog gastrocnemius muscle to increasing voltage stimuli as measured by the photosensitive interface when used with a kymograph setup.

ployed in which a bellows pneumograph was strapped around the subject's chest wall and attached to the Marey tambour. The major difference was that measurements were being made by a phototransistor hung from the tip of the lever resting on the tambour diaphragm. Typical experimental results may be seen in Figure 5. The upper trace presents a resting breathing rate and is followed by the middle trace which was made while the subject held his breath for over a minute. The lowest trace demonstrates the expected increase in the breathing rate and depth of inspiration that follows a prolonged breath hold. The response of human breathing mechanics under a variety of

conditions such as hyperventilation, rebreathing of exhaled air, or during speech may also be obtained for a more complete study of the physiological conditions governing rate and depth of breathing.

The photosensitive interface may also be utilized to measure human reaction time to audio or visual stimuli. As a visual stimulus, the subject is asked to respond to the disappearance of a square drawn on the graphics screen by pressing a switch that turns on a small light near the photosensitive transistor. The software measures the time from the disappearance of the square to the detection of light by the phototransistor. For the audio cue, the subject is asked to respond at the time a computer generated tone stops. The time before the square disappears or the tone stops is variable and is dependent on a value obtained from a random number generator. Reaction time may be measured for both the hand and the foot and for differences observed in response to either audio or visual cues.

#### TIME AND DISTANCE CONSIDERATIONS

The accuracy of time measurement using FOR-NEXT loops must be considered when using the game I/O as the entry site for signals from the photosensitive interface. I utilized a self-built real time clock interface connect-

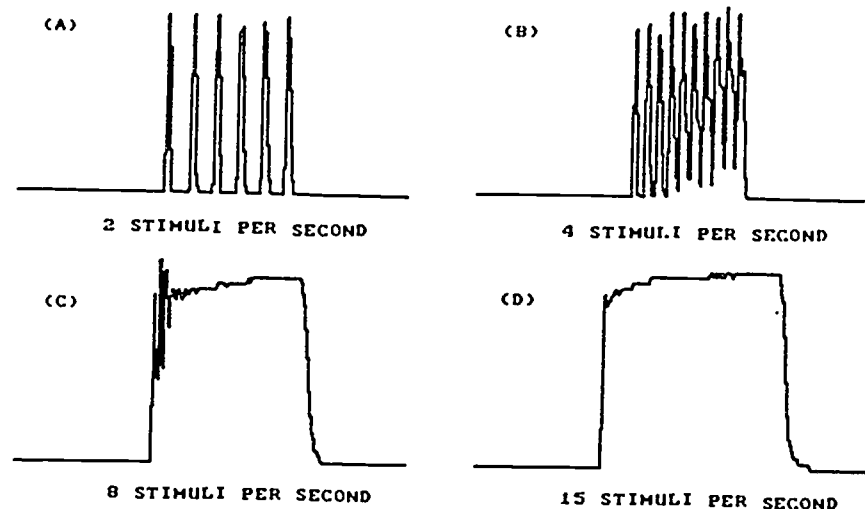


FIGURE 4. Four separate screens (A-D) demonstrating the response of frog gastrocnemius muscle to supramaximal voltage stimuli presented at different frequencies. Strip chart recorder mode software used with the photosensitive interface in the kymograph setup.



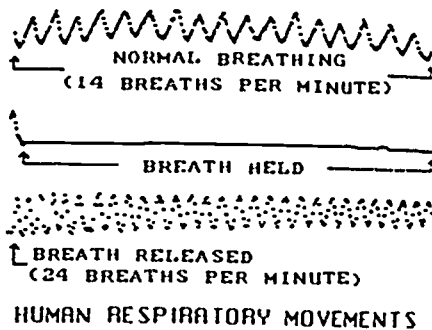


FIGURE 5. Photosensitive interface used as a pneumograph to study human respiratory movements. Strip chart recorder mode software modified to permit the sequential recording of three traces on one screen.

ed to one of the Apple expansion slots to measure, within 0.1 second, the time the trace took to traverse the screen at different light intensities. The results, shown in Figure 6, indicate a linear relationship between trace time and light intensity. At the two extremes of saturating light and total darkness, there is about a 7% difference in trace time indicating that the FOR-NEXT loop timing method collects data in a *near time* rather than *real time* mode. For many experiments at the undergraduate level, near time data collection may be acceptable if it does not compromise student interpretation of results (eg, muscle contraction data in Figures 3 and 4). However, when real time data collection is essential, an external clock interface must be utilized. For a more detailed discussion of analog-digital interfaces and timing considerations, see the article written by Richard Olivo.<sup>4</sup>

For those experiments in which the phototransistor is moved away from the light source, the collected data are not an accurate estimation of the actual distance of the phototransistor from the light source. The error exists because

no correction is made to account for the physical fact that light intensity decreases with the square of the distance from the light source. This is reflected in the results shown in Figure 7 in which light intensity measured via the phototransistor is plotted as a function of the phototransistor distance from the light source. The problem can be corrected, however, by referring to a standard curve as shown in Figure 7 to transform the intensity reading into a true distance measurement.

#### CONCLUSION

The CSIP award supported the assembly of five workstations. Each station serves from three to four students and increases greatly student access to instrumentation. The cost per station was considerably less than the funds that would be needed to purchase each of the single function instruments separately. The multifunctional nature of the Apple //e-mediated workstation coupled with its lower cost when compared to traditional instrumentation of-

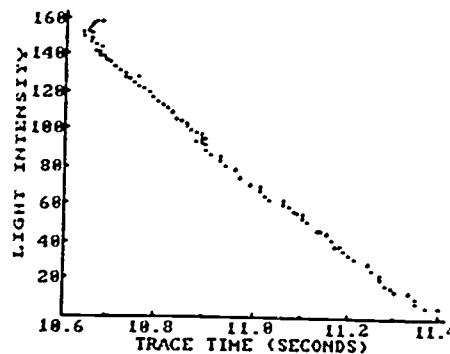


FIGURE 6. Comparison of time for the trace to move from left to right across the monitor screen at different light intensities. Trace time was measured accurately to 0.1 second with a self-built clock interface connected to one of the Apple expansion slots.

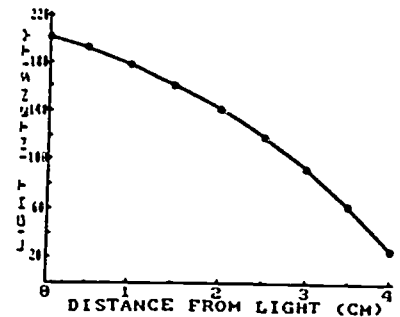


FIGURE 7. Comparison of light intensity reading from the interface with distance of the phototransistor from light source. This particular standard curve was obtained with a phototransistor hanging down from the tip of a kymograph muscle lever into a short length of PVP pipe. A 3 volt miniature lamp was positioned at the bottom of the pipe to serve as a light source.

fers instructors the opportunity to upgrade undergraduate physiology laboratory instrumentation in a cost effective manner without sacrificing the quality of the laboratory learning experience.

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# CLSE 1987-88 COLLEAGUE DIRECTORY - PART III

The primary goal of the National Resource for Computers in Life Science Education (NRCLSE) is to cultivate collaborative efforts among life science faculty interested in using the computer as a teaching tool.

The listing that follows is a continuation of the 1987-88 directory published in the February and March issues of CLSE. It is updated from the 1986-87 directory and was drawn primarily from respondents to questionnaires

printed in CLSE. It is intended to help readers identify colleagues with common interest areas.

The listings are arranged by the content areas identified in response to the question, "What content areas do you teach?" As a result, entries may appear under more than one heading.

Although every attempt has been made to ensure that the information is current and correct, it is likely that some errors appear in this list. We

apologize in advance for any inconveniences that may arise due to such oversights.

If you are aware of other colleagues that should be listed, please send the pertinent information (name, address, phone number, BITNET address, teaching content area) to NRCLSE, Mail Stop RC-70, Univ. of Washington, Seattle, WA 98195 or let us know via BITNET. NRCLSE's BITNET address is MODELL@UWALOCKE.

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**NRCLSE**

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## HYPERCARD — WHAT IS IT?

Dorothy Wooley-McKay

*Department of Biology, Glendale Community College, Glendale, Arizona*

Many individuals familiar with the Apple Macintosh microcomputer are talking about what some believe is a revolutionary new product. The product is called HyperCard, and it was released by Apple Computer at the MacWorld Expo in Boston in August, 1987.

### WHAT IS HYPERCARD?

This is a difficult question to answer because it is many things to different people. The author of HyperCard, Bill Atkinson (also author of MacPaint), has called HyperCard a "software erector set". Presumably, it is a means by which individuals who have never programmed a computer can now do so. However, HyperCard is more than that.

HyperCard is based on the concept of hypertext. Hypertext is a system of transferring information by means of a computer rather than books or other

print media. Some, of course, will not like this manner of information transfer. However, it has some advantages.

What does HyperCard have to do with using computers in the life sciences? Consider the following situation. A student is studying the electrical activity of the heart. In order to understand the principles, he must be familiar with electrophysiology, including the concepts of depolarization and repolarization. Of course, the student has a textbook and lecture notes to use as reference material. However, the sections on electrophysiology were covered in the previous semester, and the student has forgotten much of what was learned in those sections. He must go back and review that material. Unfortunately, the student cannot find the lecture notes from the previous semester. There is still the textbook, but this student never opened the textbook. He

used only lecture notes to study because the exams were based on the lecture notes, and he doesn't know where to find the textbook on electrophysiology. This, of course, is an extreme case. However, using HyperCard or another form of hypertext, we can create the following scenario.

This same student is studying the electrical activity of the heart using HyperCard stacks. HyperCard software consists of stacks. The analogy is a stack of cards. Each stack is divided into cards that are accessed one at a time. On each card, there are fields for text, graphics and buttons (see Figure 1). Part of the heart stack is a tutorial including text (read from the computer monitor) and graphics. When the student gets stuck with the electrophysiological concepts, he clicks (with the mouse) on a button located on one of the cards of the heart stack. When he clicks on this particular button, he is immediately shown a stack on basic electrophysiology. In this stack there may be a simulation of an action potential in a nerve fiber as well as text and graphics explaining the principles of electrophysiology. When the student has finished with electrophysiology, he clicks on another button and goes back to the heart. The same process could be used to connect the anatomy of the heart with the physiology of the heart. For example, when studying the Purkinje system, the student could click on Purkinje system and go to a card illustrating the location of this system. All of the information is linked together in an intuitive way. Instead of information being presented in a serial form, it can be linked together in a way more logical to human thought.

#### EXAMPLE APPLICATIONS

The following applications for teaching life sciences using HyperCard were described in the March 29th issue of MacWeek magazine.

- **Interactive Medical Record** This was developed by Dr. Edward Shultz at Dartmouth Medical School. Using this stack, a student can, among other things, point to a patient's chest and

hear the digitized sound of a heart murmur. In addition, the students can click on buttons to see vital signs, X-Rays, past visits, and blood smears. There is also a glossary, so students can click on any word and look it up in the glossary.

- **Electric Cadaver** This application, developed by Drs. Robert Chase and Steven Freedman of Stanford University Medical School presents anatomical drawings on the monochrome monitor of a Mac II. When a student clicks on a part of the image, that portion is presented on the screen. In addition, color photographs of those structures are shown on a video monitor. The color photographs are taken from the Basset series of 3 dimensional anatomical drawings.

(HyperCard can access videodiscs that contain color photographs of almost anything. The student can thus move easily from a large view of an anatomical structure to more detailed aspects of the same structure.)

- **MACInical** This is a project at Georgetown University Medical School. At the present time, there are two HyperCard stacks being used, HyperRite Up and a stack teaching the anatomy of the hand.

The spectrum of possibilities using HyperCard in education is almost limitless. There is the ability to incorporate digitized sound, such as heart murmurs (realistic sounds, not computer sounds) into HyperCard stacks. HyperCard stacks can be linked to videodiscs. HyperCard stacks can also be linked to real time animations using VideoWorks. Students can easily jump from one subject to a related subject. HyperCard stacks can be made as tutorials, simulations or addenda to laboratory exercises. Eventually, a large data base of any kind of information can be stored on CD ROM players and accessed with HyperCard.

If I wax eloquent on the possibilities of using HyperCard in education, it is because I think the potential of HyperCard has barely been tapped.

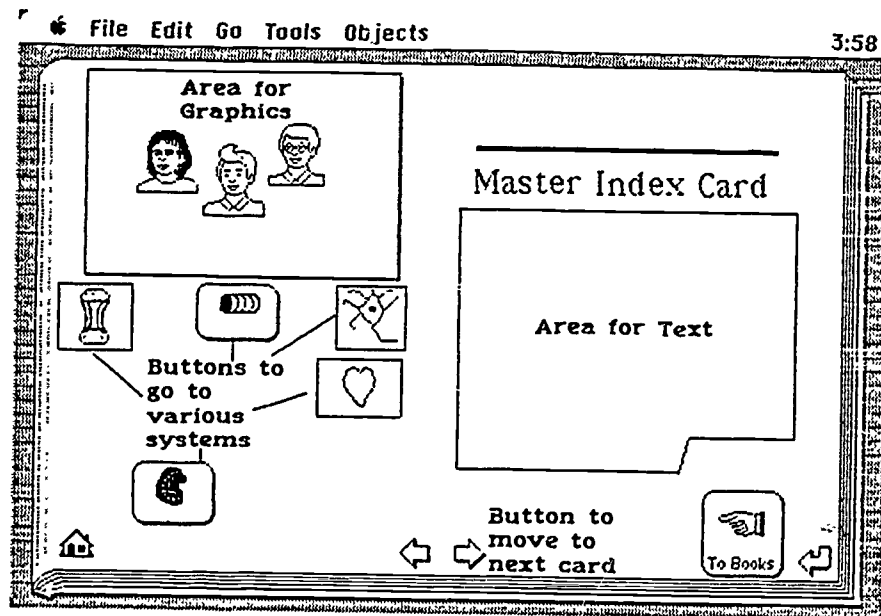


FIGURE 1. Example of fields available on HyperCard cards.

## WHERE'S THE SOFTWARE?

In the past, we have published lists of life science software sources and programs available through them. The following list is presented as the latest in a continuing effort to make colleagues aware of potential resources. As in the past, no attempt has been made by NRCLSE to review these materials.

This month's listings are arranged by content area. Each item includes a vendor code relating the software to the vendors list appearing on page 39.

If you have found specific software helpful in your teaching efforts, please share your good fortune by letting us know about the program(s) and supplier(s) so that we can make this information available through future software lists. Send pertinent information to Dr. Harold Modell, NRCLSE, Mail Stop RC-70, University of Washington, Seattle, WA 98195 or send us a note on BITnet. Our BITnet address is [MODELL@UWALOCKE](mailto:MODELL@UWALOCKE).

### ANATOMY

#### GROSS ANATOMY TUTORIAL

Tutorial for gross anatomy review by region and for self-test in National Board format. Program available for Apple II equipment. L.1

#### HEART LAB

Animated graphics simulation of human heart. Program available for Apple II, TRS-80 Models I and III, PET, and Atari 800/800XL equipment. E.1

#### HUMAN ANATOMY PICTURE FILE

Hi-Res diagrams of heart, brain, eye, ear, respiratory system, kidney, endocrine system, neurons, circulatory system schematic, and digestive system. Program available for Apple II equipment. D.1

#### LOCOMOTION

Tutorial reviewing types and functions of bones and muscles. Program available for Apple II and TRS-80 Model III equipment. J.1

### BIOCHEMISTRY

#### ANIMATIONS

Contains animations for demonstrating DNA structure and synthesis, RNA structure and synthesis, and protein synthesis. Program available for Apple II equipment. E.3

#### BIOCHEMISTRY

Tutorial covering basic atomic structure, balancing equations, and properties of proteins and carbohydrates. Program available for Apple II and TRS-80 Model III equipment. J.1

#### DNA STRUCTURE AND SYNTHESIS

Tutorial dealing with nucleotide structure and linkage between nucleotide, base complementarity and hydrogen bonding. Program available for Apple II equipment. E.3

#### ENZKIN: Enzyme Kinetics

Simulation of enzyme-catalyzed reactions. Program available for Apple II equipment. C.4

#### ENZLAB

Simulations for designing and carrying out enzyme kinetics experiments. Program available for Apple II and IBM-PC compatible equipment. B.1

#### ENZPACK

An enzyme kinetics teaching and calculation program. Program available for Apple II and IBM-PC compatible equipment. B.1

#### ENZYME ACTION

Tutorial on the basic nature and function of enzymes. Program available for Apple II equipment. B.1

#### ENZYME-SIMULATION, ENZYME ACTION

Simulation of enzyme action and the effects of inhibitors using acetylcholinesterase. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

#### GENE MACHINE

Tutorial/Simulation dealing with DNA replication and protein synthesis. Program available for Apple II equipment. H.1

#### MOLECULAR BIOLOGY SERIES

Programs demonstrating central processes of RNA and protein synthesis and

DNA synthesis and repair. Program available for Apple II and IBM-PC compatible equipment. B.1

#### MOLGRAF

Molecular graphics package. Program available for Apple II and IBM-PC compatible equipment. B.1

#### THE NUCLEIC ACIDS

Tutorial/Simulation dealing with principal nucleotides and synthesis of RNA. Program available for Apple II equipment. E.2

#### PROTEIN SYNTHESIS

Tutorial dealing with the general structure of the amino acids and formation of peptide bonds. Program available for Apple II equipment. E.3

#### RNA STRUCTURE AND SYNTHESIS

Tutorial extending the concept of hydrogen bonding between complementary bases to show the synthesis of RNA on the DNA template and the analogies in structure between DNA and RNA. Program available for Apple II equipment. E.3

### BIOLOGY

#### ADAPTION AND IDENTIFICATION

Tutorial covering animal adaptation to different environments and animal identification. Program available for Apple II equipment. S.1

#### ANIMAL REPRODUCTION

Tutorial reviewing sperm development, egg and fertilized egg. Program available for Apple II and TRS-80 Model III equipment. J.1

#### ASEXUAL REPRODUCTION

Tutorial reviewing cell division. Program available for Apple II and TRS-80 Model III equipment. J.1

#### BIOLOGY COMPUTER

#### STIMULATIONS

Tutorial covering various biological concepts and experimental areas including enzymes, photosynthesis, respiration, diffusion, meiosis, muscles, nerves, and genetics. Program available for Apple II and TRS-80 Models I and III equipment. L.1

#### DIFFUSION AND ACTIVE TRANSPORT



Tutorial covering diffusion, osmosis, and active transport in biological systems. Program available for Apple II equipment. S.2

**ENERGETICS AND METABOLISM, GARDEN OF BIOLOGY: VOLUME 1**

Data base illustrating reactions of metabolism and interactions between the several metabolic compartments of a cell. Program available for Macintosh equipment. K.1

**EVOLUTION, GARDEN OF BIOLOGY: VOLUME 2**

Data base illustrating relations among organisms of many kinds, emphasizing the history and mechanics of their evolutionary change. Program available for Macintosh equipment. K.1

**KNOWLEDGE MASTER - BIOLOGY 2**

Test-item database for test generation. Content covers coelenterates, arthropods, insects, fish, amphibians, and reptiles. Part of a 5-program Biology series for Apple II equipment. A.1

**KNOWLEDGE MASTER - BIOLOGY 3**

Test-item database for test generation. Content covers birds, mammals, primates, protists, bacteria and taxonomic zoology. Part of a 5-program Biology series for Apple II equipment. A.1

**OSMO- OSMOSIS IN RED BLOOD CELLS**

Simulation of red blood cells in hypertonic, hypotonic, and isotonic solutions. Program available for Apple II, TRS-80 Model III, IBM-PC, and Commodore 64/128 equipment. D.2

Tutorial/Simulation covering effects of temperature, concentration, solubility, molecule size and charge, and membrane pore size on flow of matter across semi-permeable membranes.

Program available for Apple II, TRS-80 Models I and III equipment. E.2

Simulation of thistle tube experiments and animation of a molecular model for osmosis. Program available for Apple II equipment. C.4

**PASSIVE TRANSPORT**

Tutorial-simulation covering diffusion and osmosis. Program for MS-DOS compatible equipment. C.1

**SIMULATION OF HEMOGLOBIN FUNCTION**

Simulations of hemoglobin and myoglobin functions. Program for

Apple II equipment. C.2

**BOTANY**

**ALGAL GROWTH**

Simulation of the effects of eight variables on growth of algae. Program available for Apple II and IBM-PC compatible equipment. O.1

**BIOLOGY FRUIT KEY**

Identifies 125 trees and shrubs. Program available for Atari 400/800 equipment. D.3

**COMPETE:Plant competition**

Simulation of experiments involving interaction between flowering plants. Program available for Apple II equipment. C.4

**FAMILY IDENTIFICATION**

Data retrieval program to review the characteristics of 74 North American flowering plant families. Program available for Apple II equipment. C.4

**LEAF: STRUCTURE AND FUNCTION**

Tutorial-simulation covering the anatomy and physiology of the leaf with respect to its role as the "chemical factory" of the plant. Program for IBM-PC (PC-DOS). C.1

**PHOTOSYNTHESIS AND LIGHT ENERGY**

Simulation focuses on characteristics of light and its role as an energy source. Program for IBM-PC (PC-DOS). C.1

**PHOTOSYNTHESIS AND RESPIRATION**

Demonstration and simulations focusing with light and dark reactions of photosynthesis, respiration, and ATP cycle. Program available for Apple II equipment. S.1

**PHOTOSYNTHESIS & TRANSPORT**

Tutorial dealing with photosynthesis and transport in plants. Program available for Apple II and TRS-80 Model III equipment. J.1

**PLANT ANATOMY PICTURE FILE**

Hi-Res diagrams of roots, stem cross-section, leaf cross-section, photosynthesis, flowers, seeds, and germination. Program available for Apple II equipment. D.1

**PLANT GROWTH**

Tutorial-simulation covering physiology of growth beginning with the seed. Covers hormone control, feedback mechanisms, transport, and differentia-

tion. Program for IBM-PC (PC-DOS). C.1

**PLANT-PLANT GROWTH SIMULATION**

Simulation of the effects of light intensity and duration on growth and development of green plants. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

**REPRODUCTION IN PLANTS**

Tutorial reviewing asexual and sexual reproduction in plants. Program available for Apple II and TRS-80 Model III equipment. J.1

**SOLAR FOOD**

Tutorial/Simulation dealing with photosynthesis. Program available for Apple II equipment. H.1

**CYTOLOGY**

**CELL CHEMISTRY I**

Tutorial covering various chemical structures. Program available for Apple II and IBM-PC compatible equipment. S.3

**CELL CHEMISTRY II**

Tutorial covering the chemical and physical processes that occur within cells. Program available for Apple II and IBM-PC compatible equipment. S.3

**CELLGROW**

Simulation of cell kinetics. Program available for Apple II equipment. U.1

**CELL GROWTH AND MITOSIS**

Interactive simulation covering surface area-volume ratio, chromosome number, chromosome replication, and cytoplasmic division. Program for IBM-PC (PC-DOS). C.1

**CELLS: STRUCTURE AND FUNCTION**

Simulation reinforces basic concepts of cell structure, cell functions, water movement and concentration gradients, and diffusion and active transport. Program available for Apple II equipment. S.1

**ECOLOGY**

**AIR POLLUTION**

Simulation of carbon monoxide pollution in an urban environment. Program available for Apple II and TRS-80 Model I and III equipment. E.2

**AQUATIC ECOLOGY**

Utilities to perform many of the cal-

culations common to aquatic ecology. Program available for Apple II and IBM-PC equipment. O.1

#### **AQUATIC ECOLOGY DATA SIMULATION**

25 simulations covering aquatic systems. Program available for Apple II and IBM-PC equipment. O.1

#### **ECOLOGICAL DATA SIMULATION**

25 simulations covering ecological systems. Program available for Apple II and IBM-PC equipment. O.1

#### **ECOLOGY**

Rote drill reviews and reinforces concepts of general terrestrial, and aquatic ecology. Available for Apple II and IBM-PC compatible equipment. S.3

#### **ECOLOGY**

Simulation dealing with plant population sizes and growth pattern. Available for Apple II equipment. S.1

#### **ECOLOGICAL ANALYSIS - PC**

Utilities that perform life table analysis, interspecific association indices, community similarity, diversity indices, descriptive statistics, mark-recapture analysis, plus regression and correlation analysis. Program available for IBM-PC compatible equipment. O.1

#### **ECOLOGICAL ANALYSIS VOL. 2 - PC**

Utilities that perform community similarity analysis, indices of dispersion, species-area curve, and step-wise multiple regression. Program available for IBM-PC compatible equipment. O.1

#### **ECOLOGICAL ANALYSIS PROGRAMS PLUS**

Utilities that perform life table analysis, community similarity indices, diversity indices, predator-prey modeling, mark-recapture analysis, descriptive statistics, plus regression and correlation analysis. Program available for Apple II equipment. O.1

#### **ECOLOGICAL MODELING**

Series of 7 programs dealing with a variety of techniques for modeling ecological systems and processes. Program available for Apple II and IBM-PC compatible equipment. C.4

#### **NICHE-ECOLOGICAL GAME/ SIMULATION**

Game in which students attempt to place an organism in its proper ecological niche correctly by specifying environment, range, and competitor. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

#### **POLLUTE**

Simulation of factors affecting water quality. Includes temperature, amount and type of pollutant, and water treatment. Program available for Apple II, PET/CBM, and TRS-80 Model III equipment. C.3

#### **POLLUTE/IMPACT/WATER POLLUTANTS**

Simulation of the impact of various pollutants on typical bodies of water. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

#### **WATER POLLUTION**

Simulation of the effects of temperature, type of waste, dumping rate, and method of treatment on the impact of pollution on aquatic life. Program available for Apple II, TRS-80 Models I and III equipment. E.2

#### **EVOLUTION**

##### **EVOLUT: Evolution and Natural Selection**

Simulation of fluctuations in gene frequencies of wild populations. Program available for Apple II equipment. C.4

##### **EVOLUTION**

Simulations covering mutation, gene flow, natural selection, and genetic drift on populations. Program available for Apple II and IBM-PC compatible equipment. O.1

#### **GENETICS**

##### **ADVANCED GENETICS**

Tutorial/Simulation presented as a nine-part program covering dominance and recessiveness, partial dominance, lethality, mechanism of inheritance, multiple alleles, sex linkage, multi-trait inheritance, crossing over, and gene mapping. Program available for Apple II equipment. E.2

##### **CATGEN**

Simulation allowing students to mate domestic cats of known genotypes. Program available for Apple II and IBM-PC compatible equipment. C.4

##### **CATLAB (Second Edition)**

Simulation in introductory genetics. Program available for Apple II and IBM-PC compatible equipment. C.4

#### **CELLS AND GENETICS PICTURE FILE**

Hi-Res diagrams of animal cell, plant cell, mitosis, meiosis, Punnett Square, sex linked traits, DNA replication, protozoa, energy reactions, and pedigree. Program available for Apple II equipment. D.1

#### **DICROSS-DIHYBRID CROSSES**

Simulation of various types of dihybrid crosses. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

#### **DNA-THE MASTER MOLECULE**

Simulation dealing with DNA structure. Programs available for Apple II equipment. E.2

#### **DNAGEN-DNA/GENETIC CODE SIMULATION**

Simulation of genetic code to produce protein sequences. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

#### **FLYGEN**

Simulation of monohybrid or dihybrid crosses with 25 varieties of *Drosophila*. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

#### **GENESIM**

Simulations of experiments in bacterial and molecular genetics. Program available for Apple II and IBM-PC compatible equipment. B.1

#### **GENETIC DRIFT**

Tutorial-simulation focusing on random changes with time in the distribution of individuals in small populations. Program for Apple II equipment. C.2

#### **GENETICS**

Tutorial covering various crosses in plants and fruit fly populations. Program available for Apple II and TRS-80 Model III and IV equipment. J.1

#### **GENETICS**

Tutorial that allows students to explore Mendel's experiments, Punnett Squares, sex linkage in fruit flies, and multiple alleles. Program available for Apple II equipment. S.1

#### **GENETICS**

Tutorial examines DNA molecule and progresses to applied genetics. Program available for Apple II and IBM-PC compatible equipment. S.3

**HEREDITY DOG**

Tutorial covering various genetic topics. Program available for Apple II and Commodore 64/128 equipment. H.1

**HUMAN GENETIC DISORDERS**

Simulation investigating inherited disorders. Program available for Apple II equipment. H.1

**INTRODUCTORY GENETICS**

Three part tutorial covering a variety of topics. Program available for Apple II, TRS-80 Models I and III equipment. E.2

**LIFE**

Educational game dealing with changing distributions of individuals. Program for Apple II equipment. C.2

**LINKOVER: Genetic Mapping**

Simulation of genetic mapping experiments. Program available for Apple II equipment. C.4

**MEIOSIS**

Tutorial/Simulation providing an interactive portrayal of gamete formation. Program available for Apple II and IBM-PC equipment. E.2

**MEIOSIS, MITOSIS, PROTEIN****SYNTHESIS**

Simulation demonstrating mitosis, meiosis, DNA replication, and protein synthesis. Program available for Apple II equipment. S.1

**MENDELIAN GENETICS**

Simulation covering dominance, partial dominance, lethality, linkage, and sex linkage. Program for Apple II equipment. C.2

**MONOCROS-MONOHYBRID****CROSSES**

Simulation of various monohybrid genetic crosses. Program available for Apple II, TRS-80 models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

**NATURAL SELECTION**

Tutorial/Simulation dealing with genetics and evolution to populations. Program available for Apple II equipment. E.2

**POPGEN-POPULATION GENETICS**

Simulation of the effects of Hardy-Weinberg Law conditions on gene, genotype, and phenotype frequencies of a population over time. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

**MICROBIOLOGY****DILUTE-MICROBIAL DILUTION SERIES**

Simulation covering design and testing of microbial dilution series to determine concentration of a bacterial solution. Program available for Apple II, TRS-80 Model III, IBM-PC, Commodore 64/128 equipment. D.2

**MICROBIOLOGY TECHNIQUES**

Tutorial/Simulation covering various laboratory procedures. Program available for Apple II equipment. E.2

**NEUROSCIENCE****NEUROMUSCULAR CONCEPTS**

Tutorial covering muscle action potentials, use of electromyograph, contraction, muscle action and movement disorders. Program for Apple II equipment. B.2

**PHARMACOLOGY****ILEUM**

Simulates laboratory experiments investigating effects of drugs on the in vitro guinea pig ileum. Program available for Apple II and IBM-PC compatible equipment. B.1

**PRINCIPLES OF PHARMACOLOGY**

Tutorial covering history, drug absorption and distribution, biotransformation and elimination, mechanisms of action, and drug safety and efficacy. Program for Apple II equipment. B.2

**PHYSIOLOGY****ABGAME**

Tutorial and game providing practice in acid-base principles. Program available for IBM-PC compatible equipment. N.3

**ACID-BASE PHYSIOLOGY****SIMULATION**

Simulation of acid-base disturbances based on Davenport Diagram. Program available for IBM-PC compatible equipment. I.1

**BASIC HUMAN**

Integrated systems model of human physiology. Program available for IBM-PC compatible equipment. R.1

**BIOFEEDBACK**

Part of 10 program package Experiments in Human Physiology. Experiments include biofeedback, condi-

tioning, and perception measurements. Program available for Apple II equipment. H.1

**BIOFEEDBACK MICROLAB**

Package includes a pulse rate sensor that measures EMG, a thermistor probe to measure skin temperature, and an interface circuit that enables student to connect the sensors to the computer. Program available for Apple II and Commodore 64/128 equipment. H.1

**THE BODY IN FOCUS**

Tutorial for investigating body systems including skeletal, muscular, respiratory, cardiovascular, gastrointestinal, endocrine, and integumentary. Available for Apple II and IBM-PC compatible equipment. N.2

**CALIBRATION**

Part of 10 program package Experiments in Human Physiology. Temperature and timing functions are calibrated against standards. Program available for Apple II equipment. H.1

**CAPEXCH**

Simulation dealing with exchange at the capillary level. Available for IBM-PC compatible equipment. N.3

**CARDIOVASCULAR FITNESS LAB**

Provides students with everything they need in order to use the microcomputer to monitor cardiovascular activity. Program available for Apple II and Commodore 64/128 equipment. H.1

**CARDIOVASCULAR INTERACTIONS**

Cardiovascular Physiology simulation. Program available for IBM-PC compatible equipment. I.1

**CARDIOVASCULAR PHYSIOLOGY****PART I: PRESURE/FLOW****RELATIONS**

Tutorial dealing with a variety of calculations in the area of hemostatics/hemodynamics. Program available for IBM-PC compatible equipment. R.2

**CARDIOVASCULAR PHYSIOLOGY****PART II: REFLEX**

Tutorial dealing with carotid sinus regulation of blood pressure, and reflex responses in hemorrhage and exercise. Program available for IBM-PC compatible equipment. R.2

**CIRCSIM: A TEACHING EXERCISE****ON BLOOD PRESSURE****REGULATION**

Simulated experiment based on a model of the baroreceptor reflex loop.



Program available for IBM-PC compatible equipment. R.2

**CIRCSYST**

Simulation of hemodynamics. Available for IBM-PC compatible equipment. N.3

**DIGESTION**

Tutorial covering digestion in simple organisms and humans.

Program available for Apple II and TRS-80 Model III equipment. J.1

**ENDOCRINE SYSTEM**

Tutorial covering hormones, effects and problems. Program available for Apple II and TRS-80 Model III equipment. J.1

**EXCRETION**

Tutorial reviewing metabolic wastes, waste removal, and kidney function. Program available for Apple II and TRS-80 Model III equipment. J.1

**VENDORS**

**A.1 Academic Hallmarks**

P.O. Box 998  
Durango, CO 81301  
(303) 247-8738

**B.1 BIOSOFT**

22 Hills Road  
Cambridge CB2 1JP, United Kingdom

P.O. Box 580  
Milltown, NJ 08850

**B.2 Biosource Software**

2105 S. Franklin, Suite B  
Kirksville, MO 63501  
(816) 665-3678

**C.1 Classroom Consortia Media, Inc.**

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Staten Island, NY 10301  
(800) 237-1113  
(800) 522-2210

**C.2 COMpress**

P.O. Box 102  
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(603) 764-5831

**C.3 Compuware**

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Randolph, NJ 07869  
(201) 366-8540

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Iowa City, IA 52242  
(319) 335-4100

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Aurora, CO 90013

**D.2 Diversified Education Enterprises**  
725 Main Street  
Lafayette, IN 47901  
(317) 742-2690

**D.3 DYNACOMP, Inc.**  
1064 Gravel Road  
Webster, NY 14580  
(716) 671-6160  
(800) 828-6772

**E.1 Educational Activities, Inc.**  
P.O. Box 392  
Freeport, NY 11520  
(800) 645-3739  
(516) 223-4666

**E.2 Educational Materials and Equipment Co.**  
P.O. Box 17  
Pelham, NY 10803  
(914) 576-1121

**E.3 EduTech, Inc.**  
303 Lamatine Street  
Jamaica Plain, MA 02130  
(617) 524-1774

**H.1 HRM Software**  
175 Tompkins Avenue  
Pleasantville, NY 10570  
(914) 769-7496  
(800) 431-2050

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**AIMS AND SCOPE**

The goal of *Computers in Life Science Education* is to provide a means of communication among life science educators who anticipate or are currently employing the computer as an educational tool. The range of content includes, but is not limited to, articles focusing on computer applications and their underlying philosophy, reports on faculty, student experiences with computers in teaching environments, and software/hardware reviews in both basic science and clinical education settings.

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Articles consistent with the goals of *Computers in Life Science Education* are invited for possible publication in the newsletter.

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Articles submitted for publication should not exceed 2000 words and should be typewritten, double spaced, with wide margins. The original and two copies including two sets of figures and tables should be sent to the Editor: Dr. Harold Modell, NRCLSE, Mail Stop RC-70, University of Washington, Seattle, WA 98195.

Title page should include full title, list of authors, academic or professional affiliations, and complete address and phone number of the corresponding author.

Illustrations should be submitted as original drawings in India ink or sharp, unmounted photographs on glossy paper. The lettering should be such that it can be legible after reduction (width of one column = 5.7 cm).

Reference style and form should follow the "number system with references alphabetized" described in the Council of Biology Editors Style Manual. References should be listed in alphabetical order by the first author's last name, numbered consecutively, and cited in the text by these numbers.

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## AN OVERVIEW OF EXPERT SYSTEMS

Monica C. Nagy

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Expert systems is a branch of artificial intelligence that has close ties to the life sciences. One of the first successful expert systems was developed by a team of researchers headed by Edward Shortliffe at Stanford University. The system, known as Mycin, gave advice on the diagnosis and treatment of infectious diseases and could perform as well as human experts or even outperform human experts in the field.<sup>2</sup> A recent bibliographic search revealed that twenty-four articles on expert systems have been published by health science journals in the last three years. This paper will examine the characteristics of an expert system, explain its functioning parts, and give a working

example of an actual expert systems development project.

An expert system is a computer program that performs the actions of a human expert consultant. The best way to understand an expert system is to examine the characteristics of a human expert consultant. Consider the following scenario.

Jane is the office manager of a small company. Being a high prestige organization, the appearance of the client waiting area is of the utmost importance. Several months ago, the company invested a large sum of money in indoor plants for the client waiting area. The plants gave the waiting area a friendly and relaxed atmosphere that

many clients have since commented on. In the last week, however, the plants have started to look limp. Some of them have developed yellow spots on their leaves. Since Jane's bachelor degree in business administration did not prepare her for dealing with sick plants, she decides to hire an indoor plant consultant.

Jane's first step in hiring the plant consultant is to draw up a list of criteria. She decides that the plant consultant must have the following five qualities:

- **KNOWLEDGE**

The consultant must be very knowledgeable about the problem at hand. Someone with several years of experience would be preferred. This person does not need to have a Ph. D. in botany, but they do need to know a good deal about the care of indoor plants.

- **RELIABILITY**

The consultant should be reliable and capable of judging whether he or she is able to give advice on the problem. Far too often has Jane been stung by high-priced consultants who gave the wrong answer or would not admit that they could not solve the problem.

- **ABILITY TO EXPLAIN WHY**

The consultant should also be able to explain why certain factors are important. The company cannot afford to hire a consultant every time the plants get sick. The consultant, therefore, should be able to explain what's wrong and give advice on how to prevent the problem from happening again.

- **ABILITY TO EXPLAIN HOW**

Most of all, the consultant should be able to explain how he or she arrived at the conclusions and be able to justify the results. This provides a backup in case the results are unusual or do not seem logical.

- **FRIENDLINESS**

Finally, an ideal consultant should be friendly and easy to work with. Although this criteria is often hard to meet, a friendly consultant makes for

a much more pleasant experience and helps encourage further consultations in the future.

### KEY FEATURES OF AN EXPERT SYSTEM PROGRAM

Let's examine each of Jane's five criteria for an expert human consultant and see how it applies to an expert system program. Jane's first requirement was that the consultant be knowledgeable and experienced in the problem area. An expert system program achieves this by containing all the rules and facts about a certain subject area. This collection of rules and facts is called the knowledge base.

Jane's second criterion, reliability, is another crucial factor in an expert system. An expert system program, just like its human counterpart, would be of little use if it were not reliable. To ensure reliability, expert system programs have a very narrow scope of expertise. At the present time, you would be hard pressed to find an expert system that could simulate the actions and experience of Ph.D. level botanist. You could, however, find one that gave advice on common houseplants. A colleague, Janise Richards, is developing one that diagnoses common houseplant ailments. When an expert system can give advice, the advice is usually in terms of certainty factors. A certainty factor is the degree to which an expert system thinks its advice is correct for that particular case. The certainty factors are reported either as percentages: "50% certainty of outcome #1," or as range: "On a scale of one to ten, your chances of #1 are 5."

Jane's next two criteria are the most important characteristics of an expert system program. The ability to answer how and why are, in my opinion, the two features that distinguish expert systems from other advice-giving programs. This ability also makes expert system programs a good learning tool. Rather than being simply a passive device, an expert system program can show you the logic behind its questions and its decisions.

Jane's final criterion, that the expert consultant be friendly and easy to work with, also applies to expert system pro-

grams. Most people would be reluctant to reuse any computer program with which they had a great deal of difficulty. For this reason, expert systems should strive to be user friendly.

To summarize, the key features of an expert system program are very similar to those of a good human expert consultant. An expert system contains a large amount of knowledge about a specific subject area. In order to ensure reliability, it tells the user when it can and cannot give advice. When it can give advice, it often uses certainty factors to indicate its degree of confidence in that advice. Finally expert systems help their users learn about the subject area by explaining their reasoning and their advice. Ideally, they should be friendly and easy to use.

### COMPONENTS OF AN EXPERT SYSTEM

In order to achieve these goals, an expert system is constructed of three parts, the rule base, the database and the inference engine. The rule base holds all the expert information. It typically stores this information in the form of if-then rules. For example, a rule used in a plant diagnosis expert system might read:

```
If
    white, wooly spots appear on
    stem, leaves, or base of stem
and leaves develop sticky patches
and growth is stunted
```

then diagnosis is mealy bugs

This type of if-then rule is also known as production rule.

The next part of an expert system is called the database. The database holds all the facts about the specific case. These facts are supplied by the user. In the case of Jane and her sick plants, the facts might be "plant appears to be limp" and "yellow patches on leaves." The database changes with each new case. The database and the rule base together are often termed the knowledge base of the expert system because these are the two parts that store the knowledge.

The final, and most important part is

the inference engine. The inference engine is the driver of the expert system. It is where all the actual "thinking" takes place. It uses the rules in the rule base and facts in the database to decide on the advice to give.

#### BUILDING AN EXPERT SYSTEM

Expert systems are easiest to build when the subject area is narrow and the rules are straight-forward and non-conflicting. For example, an expert system could be written to give advice on simple operations on a computer such as changing the default disk drive or displaying a directory.<sup>1</sup> The rules for performing these operations are usually listed in the computer's guide to operations. When the rules have already been written down, as in the case of a manual or quick reference guide, the creation of an expert system is made much easier. Many excellent expert systems have been created by first interviewing a number of human experts and then translating their expertise into rules. This process can be very time consuming and can also produce conflicting rules.

When there is no general consensus on the rules, or there exists a multitude of special exceptions, building an expert system becomes difficult. In college, I once gave a presentation on expert systems to a group of my fellow computer science majors. A short example program on how to identify a science fiction movie seemed to be appropriate since science fiction is of interest to most aspiring computer scientists. It quickly became apparent that science fiction movies were difficult to classify. As soon as one set of rules was finished, a conflicting example would arise. Almost everyone consulted had a different definition of a science fiction movie. Few could agree on even ten common factors. How would you classify *The Twilight Zone*? What is the dividing line between science fiction and fantasy? Everyone agreed that a futuristic setting would be a good indicator. How far in the future the movie would have to be set could not be agreed upon. The presence of new technology was another factor commonly agreed upon. No consensus

could be reached on how new the technology would have to be. I later came to find out that this debate has been going on for some time and shows no signs of being settled any time soon. A general consensus on the rules is necessary for building an expert system.

Let's see how all this information applies to an actual expert system development project. For the past year, I have worked on an expert system with Ernst Schelb, DMD, at the Dental School of the University of Texas Health Science Center at San Antonio. Our goal was to design and implement an expert system in the area of endodontics, to be used by second year dental students. The system was to be used for practice in the student lab. Endodontics is the area of dentistry that deals with the nerves and blood vessels on the inside of the tooth. Our expert system, known as "The Hurting Tooth Advisor," deals with diagnosing pain in a single suspect tooth. It does not deal with pain from other teeth or pain caused by problems outside of the mouth. Dr. Schelb felt that endodontics, as compared to other areas of dentistry, lent itself most to an expert system because of the small number of possible diagnoses. Also, the rules for

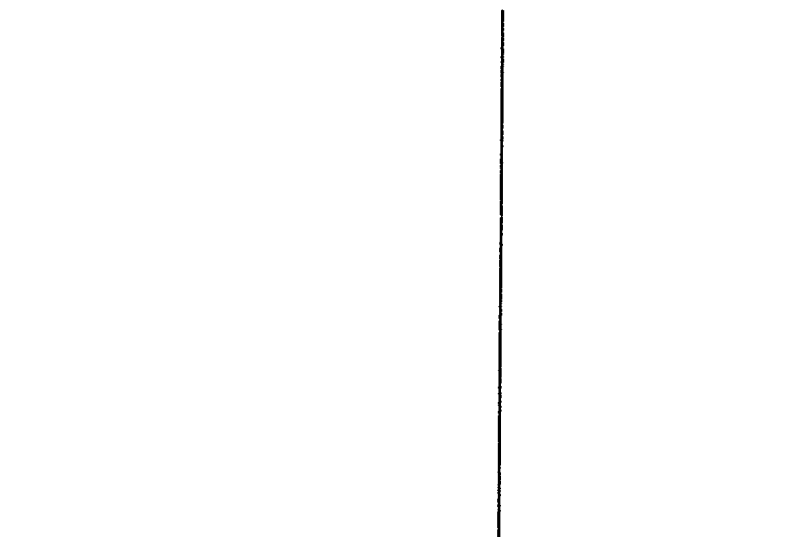
diagnosis in endodontics are more straight-forward and less conflicting.

We began with a matrix that listed signs, symptoms and test outcomes across the top and possible diagnoses along one side. We then narrowed the signs, symptoms and tests down to three categories, description of the pain, clinical findings, and radiographic findings. We used the information in the matrix to write the rules. An expert system is much easier to create if the rules have already been formulated.

#### PROGRAM EXAMPLE

When it came time to transfer the rules to the computer, we chose to use the EXSYS expert system generator (EXSYS Inc., P.O. Box 75158, Station 14, Albuquerque, NM 87194). Generator programs greatly ease and accelerate the creation of an expert system. They take care of all the details. They usually provide the inference engine and a facility for displaying questions to the end user. They also allow the end user to ask why a question is being asked and, if they so desire, to see the rules that were used. Best of all, most generator programs are designed to be used by non-programmers.

Besides the features mentioned above,



Add rule <A> or <ENTER>, Edit rule <E>, Delete rule <D>, Move rule <M>, Print <P>, Store/exit <S>, Run <R>, Options <O>, DOS <Ctrl-X>, Help <H>

FIGURE 1. The main editor screen.



EXSYS has a specialized editor for entering rules. This means that it prompts the person editing the rule for each of the rule's components (see Figure 1). Once the rules are entered, they can be tested right away by pressing "R" for run. The antecedent, or "if" part, of each rule is translated into a question for the student to answer. The expert system uses the rules themselves and the answers that the student has given thus far to determine which question will be asked next. For instance, in Figure 2, if the student answers that the tooth shows signs of caries, the next question would ask how extensive the caries were. If the student did not indicate that caries could be seen, the expert system would go on to the next category of questions. The student would not see the extra question on caries.

While using the expert system, a student can ask why a question is being asked by entering "WHY" as the response to a question. The student also has the option of seeing the rules as they are used. Once the student has answered all the questions, the possible diagnosis and treatment plan are displayed. At this point, the student can either print the advice, change some of

THE SUSPECT TOOTH SHOWS SIGNS OF

1. CARIES
2. PREVIOUS RESTORATIONS
3. WEAR ON THE OCCLUSAL ENAMEL SURFACE
4. CERVICAL ABRASION

---

Enter number(s) of value(s), WHY for information on the rule, <?> for more details, QUIT to save data entered or <H> for help.

FIGURE 2. A question screen. At this point the student could enter the number corresponding to a selected choice, the word "WHY" to see the current rule, a "?" for more details, "QUIT" to save the information entered so far or "H" for help on using the expert system.

their answers, and run the system again, or start over from the beginning (see Figure 3).

The Hurting Tooth Advisor is now in a prototype stage, meaning that it gives

accurate results only for a few chosen cases. We hope to develop this into a fully operational system.

In conclusion, an expert system is a computer program that simulates the human capabilities of knowledge, reliability, the ability to explain the importance of key factors, the ability to explain how a conclusion is reached, and the ability to be friendly and easy to use. Expert systems attain these abilities by organizing information into a rule base and a database. They also contain an inference engine that drives the whole process. Finally, an expert system is easiest to build in an area that has well codified rules and guidelines that are straight-forward and non-conflicting.

#### REFERENCES

1. Nagy, T, D Gault, and M Nagy. Building Your First Expert System. Aston-Tate, Culver City, CA, 1985.
2. Winston, PH and KA Predergast (eds). The AI Business: The Commercial Uses of Artificial Intelligence. MIT Press, Cambridge, MA, 1984.

Values based on 0/1 system	VALUE
1. CAN GIVE ADVICE	1
2. INFERRED TREATMENT IS APPLY SODIUM FLUORIDE	1
3. INFERRED DIAGNOSIS IS CERVICAL ABRASION	

---

All choices <A>, only if value>1 <G>, Print <P>, Change : rd rerun <C>, rules used <line number>, Quit/save <Q>, Help <H>, Done <D>:

FIGURE 3. The conclusions are displayed in terms of certainty factors. In this case, the certainty factor will either be zero, for false, or one, for true.

## WHERE'S THE SOFTWARE?

In the past, we have published lists of life science software sources and programs or program areas available through them. The following list is presented as the latest in a continuing effort to make colleagues aware of potential resources. As in the past, no attempt has been made by NRCLSE to review these materials.

This month's listings continue last month's and are arranged by content area. Each item includes a vendor code relating the software to the vendors appearing at the end of the software lists.

If you have found specific software helpful in your teaching efforts, please share your good fortune by letting us know about the program(s) and supplier(s) so that we can make this information available through future Where's the Software lists. Send pertinent information to Dr. Harold Modell, NRCLSE, Mail Stop K.C.-70, University of Washington, Seattle, WA 98195 or send us a note on BITnet. Our BITnet address is MODEL@UWALOCKE.

### PHYSIOLOGY

#### CONCEPTS IN THERMOGRAPHY

Tutorial covering basic DC concepts, peripheral vascular physiology, detecting skin temperature, amplifiers, and processing DC signals. Program for Apple II equipment. B.2

#### EXERCISE EXPERIMENTS

Part of 10 program package Experiments in Human Physiology. The effect of exercise and physical condition on heart rate, breathing rate, and skin temperature is investigated. Program available for Apple II equipment. H.1

#### GAS DIFFUSION IN THE LUNG

Simulation of oxygen and CO<sub>2</sub> transfer between alveolar air and blood. Program available for IBM-PC compatible equipment. I.1

#### HEART RATE

Part of 10 program package Experiments in Human Physiology. Light and light sensor for measuring and recording heart rate. Program available for Apple II equipment. H.1

#### HOMEOSTASIS-THERMOREGULATION

Part of 10 program package Experiments in Human Physiology. Students investigate the body's ability to maintain a constant

internal temperature by subjecting a volunteer to mild temperature excursion while recording and displaying skin and body temperature. Program available for Apple II equipment. H.1

#### HUMAN BODY-STRUCTURE AND FUNCTION

Simulation covering joint movement, movement of food through digestive system, and enzyme activity. Program available for Apple II equipment. S.1

#### MODEL NEURON

Simulation of the behavior of an isolated neuron. Program available for Macintosh equipment. K.1

#### MUSCLE MECHANICS: A COMPUTER-SIMULATED EXPERIMENT

Simulated experiment that permits the user to determine either the length-tension or the force-velocity relationship of a skeletal muscle. Program available for IBM-PC compatible equipment. R.2

#### NERVOUS SYSTEM

Tutorial covering nerves, reflexes, and chemical transfer of impulses. Program available for Apple II and TRS-80 Model III equipment. J.1

#### PHYSIOLOGICAL DATA SIMULATION

25 simulations covering aspects of physiology. Program available for Apple II and IBM-PC compatible equipment. 0.1

#### PROBLEMS IN FLUID COMPARTMENT RE-DISTRIBUTION

Tutorial covering solution of simple problems of fluid compartment changes in the face of perturbations. Program available for IBM-PC compatible equipment. R.2

#### PSYCHOLOGICAL STRESS-LIE DETECTOR

Part of 10 program package Experiments in Human Physiology. The physiological response to the stress of a frustrating and abusive quiz is measured. Program available for Apple II equipment. H.1

#### PULMONARY MECHANICS

Tutorial and simulation dealing with pulmonary mechanics. Available for IBM-PC compatible equipment. N.3

#### RESPIRATION RATE

Part of 10 program package Experiments in Human Physiology. A napping subject is monitored for heart and breathing rate. Results are compared to the data acquired when the subject is awake. Program available for Apple II equipment. H.1

#### RESPONSE-TIME

Part of 10 program package Experiments in Human Physiology. Users measure finger reaction times with a bright light stimulus (sensor included). Program

available for Apple II equipment. H.1

**RESPONSE-TIME INVESTIGATIONS**  
Part of a 10 program package Experiments in Human Physiology. The effects on reaction times of stimulus type and response location are studied. Program available for Apple II equipment. H.1

#### RESPSYST, GASEXCH

Simulations dealing with pulmonary gas exchange. Available for IBM-PC compatible equipment. N.3

#### SIMULATIONS IN PHYSIOLOGY - THE RESPIRATORY SYSTEM

Series of 12 simulations dealing with respiratory mechanics, gas exchange, chemoregulation and acid-base balance. Program available for Apple II, IBM-PC compatible, and Macintosh equipment. N.1

#### SKELETAL MUSCLE ANATOMY/PHYSIOLOGY

Tutorial covering three muscle categories, skeletal muscle microstructure, sliding filament theory, motor units, and lever systems. Program for Apple II equipment. B.2

#### SKELETAL MUSCLE MECHANICS

Set of six simulations dealing with muscle physiology. Program available for IBM-PC compatible equipment. I.1

#### SKILLS IN ELECTROMYOGRAPHY

Tutorial covering skin preparation, reducing EMG artifact, testing a myograph's operation, electrode location, and preventing shock hazards. Program for Apple II equipment. B.2

#### SKIN TEMPERATURE

Part of a 10 program package Experiments in Human Physiology. Temperature probe (included) senses body and skin temperatures. Program available for Apple II equipment. H.1

### POPULATION DYNAMICS

#### COEXIST:Population Dynamics

Simulation of the growth of two populations either independently or in competition for the same limited resources. Program available for Apple II equipment. C.4

#### ISLAND BIOGEOGRAPHY

Three simulations of island communities dealing with the relationship between island area and number of species, colonization of a new island, and island immigration and extinction. Program available for Apple II equipment. C.4

#### LIMITS

Simulation of the effects of growth on world population, pollution, food supply, industrial output, and natural resources. Program available for Apple II, PET/CBM and TRS-80 Model III equipment. C.3

**MARK & RECAPTURE**

Simulation of mark and recapture experiments to explore three models for estimating population sizes. Program available for Apple II equipment. C.4

**POP**

Simulation of three growth models (exponential, logistical, and logistical with low density). Program available for Apple II, PET/CBM, and TRS-80 Model III equipment. C.3

**POPGRO-POPULATION GROWTH SIMULATION**

Simulation of unlimited growth (J-curve), limited growth (S-curve) and limited growth with response lag time (S-curve with oscillations) models of population growth. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

**POPULATION FLUCTUATIONS**

Tutorial covering factors influencing population growth. Program available for Apple II, TRS-80 Models I and III equipment. E.2

**POPULATION GROWTH**

Simulation dealing with exponential and density-dependent growth. Program for Apple II equipment. C.2

**POPULATION GROWTH**

Simulation of population growth. The package compares and contrasts the geometric or exponential growth model with the logistic or Verhulst-Pearl growth model. Program available for Apple II equipment. C.4

**POPULATION SIZES**

Simulation dealing with a dynamic population. Program for Apple II equipment. C.2

**SUBSTANCE ABUSE****DRINKING AND NOT DRINKING**

Tutorial designed to augment strategies for the prevention of substance abuse. Includes facts about drinking and the effects of alcohol. Program available for Apple II equipment. K.1

**INTRODUCTION TO PSYCHOACTIVE DRUGS**

Tutorial designed to augment strategies for the prevention of psychoactive drug abuse. Program available for Apple II equipment. K.1

**KEEP OFF THE GRASS**

Tutorial designed to augment strategies for the prevention of marijuana abuse. Program available for Apple II equipment. K.1

**SIX CLASSES OF PSYCHOACTIVE DRUGS**

Tutorial designed to augment strategies for the prevention of psychoactive drug abuse. Program available for Apple II equipment. K.1

**SUBSTANCE ABUSE DATA BASE**

Database containing contact information on substance abuse organizations. Program available for Apple II equipment. K.1

**ZOOLOGY****ZOOLOGY I**

Tutorial covering the general characteristics, structures, and functions that define the major invertebrate phyla. Program available for Apple II and IBM-PC compatible equipment. S.3

**ZOOLOGY II**

Tutorial covering physiology in the Phylum Chordata. Program available for Apple II and IBM-PC compatible equipment. S.3

**MISCELLANEOUS****BAFFLES, BAFFLES II**

Game to help students develop deductive reasoning and problem solving skills. Program available for Apple II (BAFFLES) and IBM-PC compatible (BAFFLES II) equipment. C.4

**BALANCE-PREDATOR-PREY SIMULATION**

Simulation of the effects of food supply, carrying capacity, environmental conditions, and external pressures or predator/prey relationships. Program available for Apple II, TRS-80 Models I and III, IBM-PC, and Commodore 64/128 equipment. D.2

**BLANCHAER CLINICAL CASE STUDIES**

Eight simulations of clinical syndromes. Program available for Apple II equipment. B.1

**CLASSIFY-CLASSIFICATION KEY PROGRAM**

Presents an unclassified set of characteristics and labels for classification at various levels. Program available for Apple II, TRS-80 Model III, IBM-PC, and Commodore 64/128 equipment. D.2

**GRADE KEEPER - PC**

Grade book manager that handles classes up to 300 students, up to 25 grades per student. Program available for IBM-PC compatible equipment. O.1

**GRADEBOOK**

Program for the analysis of a large set of grades. Program available for IBM-PC compatible equipment. I.1

**LABPLOT**

Allows the Apple II with any A/D converter card to be used as a multipen chart recorder or as an X/Y plotter. B.1

**LIFE TABLES AND THE LESLIE MATRIX**

Tutorial-simulation dealing with the basic life table and Leslie Matrix. Program available for Apple II equipment. C.4

**MALARIA**

Simulation of the effects of various types of malaria epidemic controls. Program

available for Apple II, PET/CBM, and TRS-80 Model III equipment. C.3

**MULTI-Q**

A general purpose question creation and presentation system. Program available for Apple II and IBM-PC compatible equipment. B.1

**PREDATION**

Simulation of predator-prey interactions. Program available for Apple II equipment. C.4

**PREDATION EQUILIBRIA**

Simulations of equilibrium models of predator-prey interaction. Program available for Apple II equipment. C.4

**"Q" EDUCATIONAL AUTHORIZING SYSTEM**

Authoring system for tutorial and assessment material. Allows incorporation of graphics and videodisc material. Program available for IBM-PC compatible equipment. B.1

**RATS**

Simulation of rat control in city or apartment by sanitation and various poisons. Program for Apple II, PET/CBM, and TRS-80 Model III equipment. C.3

**STERL**

Simulation exploring effectiveness of pest control methods. Program available for Apple II, PET/CBM, and TRS-80 Model III equipment. C.3

**TRIBBLES, TRIBBLES Revisited**

Simulation to introduce students to the scientific method. Programs available for Apple II (Tribbles) and IBM-PC compatible (Tribbles revisited) equipment. C.4

**VENDORS**

A.1 Academic Hallmarks  
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## A GRAPHIC COMPUTER LANGUAGE FOR PHYSIOLOGY SIMULATIONS

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Physiologists and other life scientists can use mathematical models profitably to create and test new scientific theories as well as to demonstrate to their students the behavior of complex biological systems. Unfortunately, creating such models often requires more computer programming expertise or more time and effort than the investigator can afford to invest in a model whether it is a research tool or a didactic device. Therefore, we were intrigued by LabView\*, a graphic pro-

gramming language that recently became available for the Macintosh computer.

Because physiologists generally think of organ systems in terms of block diagrams and feedback loops, we thought perhaps a graphic programming language would make constructing a mathematical model easier, less time-consuming, and more intuitive than conventional computer languages.

Therefore, the purpose of this communication is to report our experience using LabView to construct mathematical models of physiological systems.

Because LabView was originally developed to control laboratory instruments and to collect and analyze data, a LabView program can be thought of metaphorically as a "virtual instrument." A virtual instrument consists of a front panel and an executable block diagram. In one window on the com-

\* The graphic programming language is actually called G, but it is better known by the trade name LabView, an acronym for laboratory virtual instrument engineering workbench. The LabView package consists of four 3.5 inch disks and a two-volume manual. It is available from National Instruments, 12109 Technology Blvd., Austin, Texas, 78727-9112, and it runs on a 1Mb Mac Plus, SE, or II. An external floppy or hard disk is required.

puter screen, the front panel displays the various input and output controls available to the user or programmer: graphic representations of switches, dials, knobs, digital or analog meters, strip-charts, etc. The appearance of these controls or display devices changes to illustrate the data going into or out of the virtual instrument.

A second window contains the block diagram that constitutes the program that the computer executes. In the block diagram, numeric variables, display devices, program control structures such as For-Next loops, and arithmetic operations are nodes represented graphically by icons. The programmer uses various menus to obtain the desired icons and then places them in the block diagram. "Wires" connecting the icons define the path of data flow from one node to the next. The execution environment is inherently parallel and data-driven, (ie, each icon or node executes its particular function when all necessary input data are available to it).<sup>5</sup> The output of that particular node is then passed on to the next. The block diagram of one virtual instrument can even contain other virtual instruments within it. Thus, previously programmed routines can easily be incorporated into a more elaborate model. According to a recent report, "the intended user base for LabView includes engineers and scientists with no programming experience or limited experience with a simple language like BASIC."<sup>5</sup> Therefore, being only moderately proficient in BASIC, we felt appropriately qualified to evaluate LabView as a programming language for physiological simulations. To do so, we programmed two well-known cardiovascular models in LabView. In the rest of this report, we describe the programs themselves, our experience in creating them, and our perceptions of the strengths and weaknesses of LabView as a method for modeling physiological systems.

#### A SIMPLE CARDIOVASCULAR MODEL

The following model of the cardiovascular system was originally proposed by Guyton et al.<sup>1,2</sup> Rothe<sup>4</sup> later developed it into a teaching tool for courses

in cardiovascular physiology. The model was designed to simulate the interaction between the heart and the peripheral vascular system and to demonstrate homeostatic responses to a variety of perturbations to the cardiovascular system (eg, exercise, cardiac failure, and hemorrhage).

The model consists of three compartments: a single-chambered heart, an artery, and a vein. Under steady-state conditions, each compartment contains a specific volume of blood. The pressure in each compartment is determined by the compartment's compliance and its blood volume. Blood flow between compartments is a function of the pressure gradients and resistances between compartments. To simulate the Frank-Starling mechanism crudely, cardiac output is computed simply as a linear function of heart volume. Although there are no "reflexes" *per se*, the appropriate cardiovascular effectors (ie, arterial resistance and compliance, venous resistance and compliance, and cardiac contractility) can be modified

by the user. In addition, there is also a provision for simulating hemorrhage or transfusion by increasing or decreasing the total blood volume.

Figure 1 shows the "front panel" of the cardiovascular model as it appears on the Macintosh screen. The slide switches permit the user to change the resistances (Art.R and Ven.R) and compliances (Art.C and Ven.C) of the two vascular compartments, cardiac contractility (Cirt), and the total blood volume ( $\Delta$ Vol). The two strip-charts continuously display the cardiac output and arterial pressure. The digital indicators for each compartment show the current numerical values for the blood volumes (Art.V, Ven.V, and Hrt.V) and pressures (Art.P, Ven.P, and Hrt.P) as well as the total blood volume (Tot.V). In Figure 1, the input and output values shown on the slide switches and other indicators are the initial default values specified by Rothe.<sup>4</sup>

Figure 2 shows the block diagram for the simple cardiovascular model. All of the actual calculations are performed

Simple Cardiovascular Model Panel

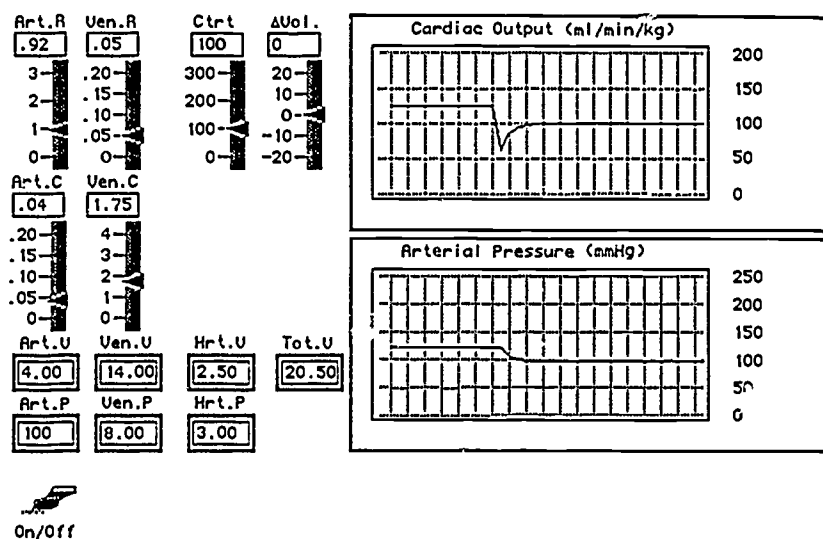


FIGURE 1. Front panel of a simple cardiovascular model. A window on the computer screen shows the four types of control and display devices used in this model. Slide switches select values for cardiac contractility (Cirt), the rate of hemorrhage or infusion ( $\Delta$ Vol), and arterial and venous resistances (Art.R and Ven.R) and compliances (Art.C and Ven.C). Digital indicators show blood volumes in the three compartments, and strip-chart simulators provide a continuous record of cardiac output and arterial pressure. A binary on-off switch stops and starts program execution.



within a While-Wend Loop activated by an On/Off switch located on the front panel. In the diagram, the On/Off switch is "wired" to the "recirculation terminal" (counterclockwise arrow) at the bottom right-hand corner. The While-Wend Loop is used to provide continuous operation and to store the compartmental volumes for use in the next iteration. These values are stored from one iteration to the next by the three shift-registers (boxed arrows) located on the left and right borders of the While-Wend Loop structure. The three numeric constants wired to the shift-registers on the left border of the While-Wend Loop initialize the shift-registers and provide the starting values of the blood volumes in the three compartments.

Within the While-Wend Loop, the program proceeds from left to right. The first step in the program is to calculate the pressures in the three compartments by dividing the volumes specified in the three shift-registers by the corresponding compliance values selected on the front panel. These calculations are performed by three division icons. For example, to calculate the pressure in the arterial compartment, the arterial shift-register is wired to the upper terminal (the numerator) of one division icon, and the arterial compliance slide switch control is wired to the bottom terminal (the denominator). The output terminal of the division icon (the quotient) is wired to both a strip-chart indicator and a numeric indicator to display the arterial pressure on the front panel. The same procedure is used to calculate the pressures in the venous and cardiac compartments.

The next step in the program is to calculate the pressure gradients between the arterial and venous compartments and between the venous and cardiac compartments. These pressure differences are calculated by wiring the three division output terminals into two subtraction icons. The pressure differences are then divided by the arterial and venous resistances to determine the blood flow from the arterial to the venous compartment and from the venous compartment to the heart.

As shown in the lower portion of Figure 2, cardiac output is simply a linear

Simple Cardiovascular Model Diagram

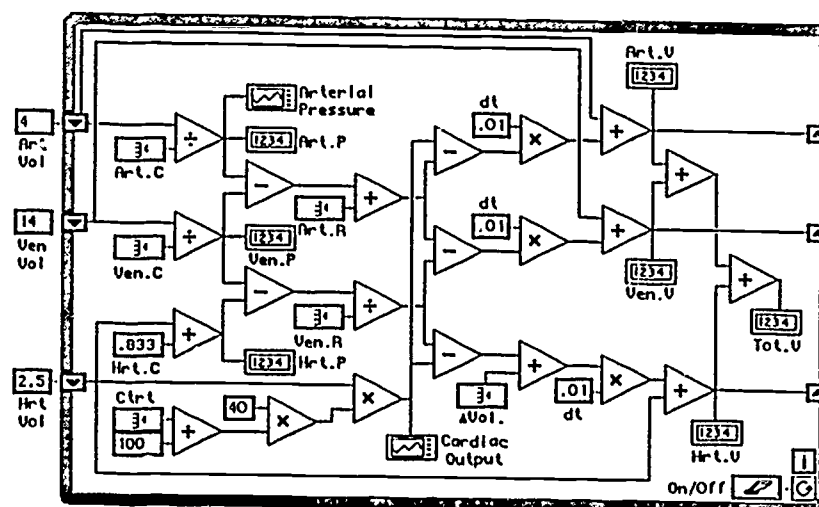


FIGURE 2. Block diagram of a simple cardiovascular model. A LabView diagram is the program that the computer executes. Each icon represents the arithmetic operation denoted by its symbol or corresponds to a switch or display device on the front panel shown in Figure 1. "Wires" define the path of data flow. For details of operation, see text.

function of heart volume and is calculated by multiplying the heart volume by the slope of the relationship between heart volume and cardiac output. However, note that the slope of this relationship depends on the cardiac contractility (Hrt. Ctrt) and that cardiac contractility can be changed from the default valve (100%) by using the slide switch on the front panel.

Under steady-state conditions, blood flow through the three compartments will be equal. However, following any perturbation, a transient imbalance in blood flow between any two compartments will redistribute the blood volume. Therefore, the next step in the program is to calculate the volume shifts among the three compartments (ie, cardiac output minus arterial runoff, arterial runoff minus venous return, and venous return minus cardiac output). These differences between each compartment's inflow and outflow are then multiplied by an increment in time (dt). The result of this integration is added to the old volume from the previous iteration (from the shift-register on the left) to yield the new instantaneous volume in each compartment. The new instantaneous volume is then

passed to the shift register on the right and stored until the next iteration. Finally, the slide switch  $\Delta$ Vol allows the user to select a rate at which blood volume is gained or lost through transfusion or hemorrhage.

#### A MODEL OF PULSE PRESSURE

To show a slightly more elaborate model programmed in LabView and to demonstrate other capabilities of LabView not illustrated in the previous example, we present a model of the arterial pressure pulse that was originally published by Coleman and Sias<sup>1</sup> and later used by Randall for didactic purposes.<sup>3</sup> The model was designed to demonstrate the physics of the arterial pressure pulse and to show how the amplitude of the arterial pressure wave is affected by stroke volume, arterial compliance, and the ventricular ejection rate. The model consists of a single ventricle and an elastic arterial compartment. We modified the Coleman-Sias-Randall model so that the ventricle receives a variable venous return and has an adjustable heart rate. These two factors determine the stroke volume and the rate of ventricular ejection into the arterial compartment. The



instantaneous volume in the arterial compartment is set by the difference between ventricular ejection and the peripheral runoff. The instantaneous volume and the arterial compliance determine the instantaneous pressure.

Figure 3 shows the front panel of the pulse pressure model. The slide switches select values for the venous return (V.R.), heart rate (H.R.), compliance (C.), and total peripheral resistance (T.P.R.). In addition, two numeric controls specify values for the fraction of the cardiac cycle devoted to systole (systolic fraction) and the integration interval (dt). The two strip-charts display the ventricular ejection rate and arterial pressure. The four digital displays show the current values of the stroke volume, systolic and diastolic pressures, and the amplitude of the pressure pulse. The initial values and constants shown in Figure 3 were taken from Randall.<sup>3</sup>

Figure 4 shows the block diagram for the pressure pulse model. Like the previous model of the cardiovascular system, all the calculations for the model are performed within a While-Wend loop to provide continuous operation and to store current values in shift-

registers for use in the subsequent iteration. However, unlike the previous model, the pressure pulse model contains a number of virtual instruments that perform the more complicated program operations, such as timing the cardiac cycle and calculating the rate of

ventricular ejection, discussed in detail below.

As in the previous model, the program execution proceeds from left to right and starts with the initialization of the shift-registers. The complex icon wired to the top shift-register on the left border is an "array builder." The two numeric constants specifying the initial systolic (S) and diastolic (D) pressures are wired to the left side of the array builder which generates a two-element array that is passed to the shift-register. Note that in LabView, the appearance of a wire depends on the type of data it carries (ie, numeric, array, string, or boolean data) and that the array wire is thicker than the wire used for numeric data. The other three shift-registers are initialized by numeric constants specifying the starting value for the timer (n) and the default values for arterial volume (V) and pressure (P).

Inside the While-Wend loop, using values specified on the front panel for venous return, heart rate, and systolic fraction of the cardiac cycle, the program begins by calculating the stroke volume, the period of the complete cardiac cycle, and the duration of systole. The cardiac period and the duration of systole as well as the current values of n and dt are wired to the timer sym-

### Pressure Pulse Model Diagram

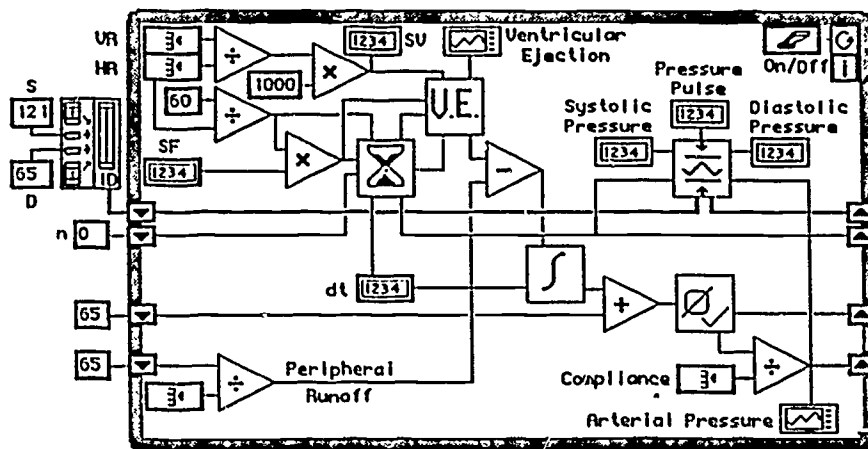


FIGURE 4. Executable diagram of pulse pressure model. The main diagram contains five "virtual instruments" also programmed by the authors: timer (hourglass icon), ventricular ejection (V.E.), integrator (integral sign), error-trapping routine (zero check), and wave analyzer (connected to pressure pulse display). For details, see text.

### Pressure Pulse Model Panel

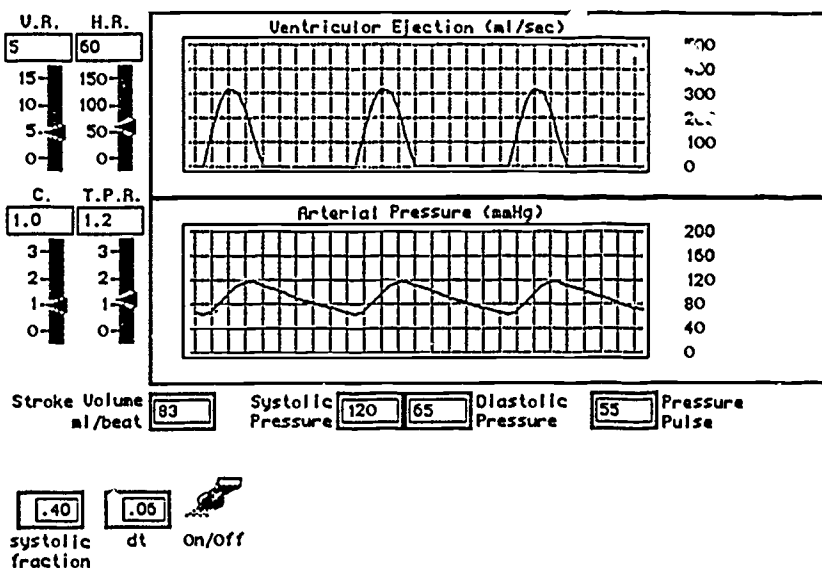


FIGURE 3. Front panel of pulse pressure model. Slide switches select values for venous return (V.R.), heart rate (H.R.), arterial compliance (C.), and total peripheral resistance (T.P.R.). Constants are the systolic fraction of the cardiac cycle and the integration interval (dt). Computed variables are shown on simulated stripcharts or digital displays.

bolized by the hourglass icon. The timer is the virtual instrument responsible for the timing of each cardiac cycle and for signaling the beginning and end of systole.

The diagram for the timer is shown in Figure 5. The timer operates by setting  $n$  equal to zero at the beginning of each cardiac cycle and incrementing  $n$  by one during each iteration until the cardiac cycle is completed. The number of iterations required for a cycle is the cardiac period divided by  $dt$ . During each iteration, by using the "greater than, equal to, or less than" function,  $n$  is compared to the calculated number of iterations per cycle. The output of this comparator is boolean (true or false) as indicated by the dotted line connecting the "less than" terminal to the case structure selector. So long as  $n$  is less than the calculated number of iterations per cycle, the program executes the operations specified in the TRUE form of the case structure, and  $n$  is incremented by one. At the end of a cycle, when  $n$  is no longer less than the calculated number of iterations per cycle, the FALSE form of the case structure is executed, and  $n$  is reset to zero. The time since the beginning of each cycle is simply the product of  $n$  multiplied by  $dt$ . A second comparator icon is used to compare the elapsed cycle time with the calculated systolic duration and to provide a boolean signal for the beginning and end of systole.

The instantaneous rate of ventricular ejection is calculated by the virtual instrument labelled V.E. in Figure 4. The block diagram for the V.E. virtual instrument is shown in Figure 6. From an equation given by Randall,<sup>3</sup> ventricular ejection (VE) is calculated as the positive half of a sine wave:

$$VE = (SV/SD) \cdot \sin(\pi \cdot \text{time}/SD) / 0.636$$

Here, SV is the stroke volume, SD is the systolic duration, and time is the time elapsed since the beginning of systole. The constant, 0.636, simply converts the average systolic ejection rate (SV/SD) to its maximum systolic value. In the diagram (Figure 6), this calculation is performed within a TRUE-FALSE case structure. During systole, while the boolean signal from

## Pressure Pulse Model Timer Diagram

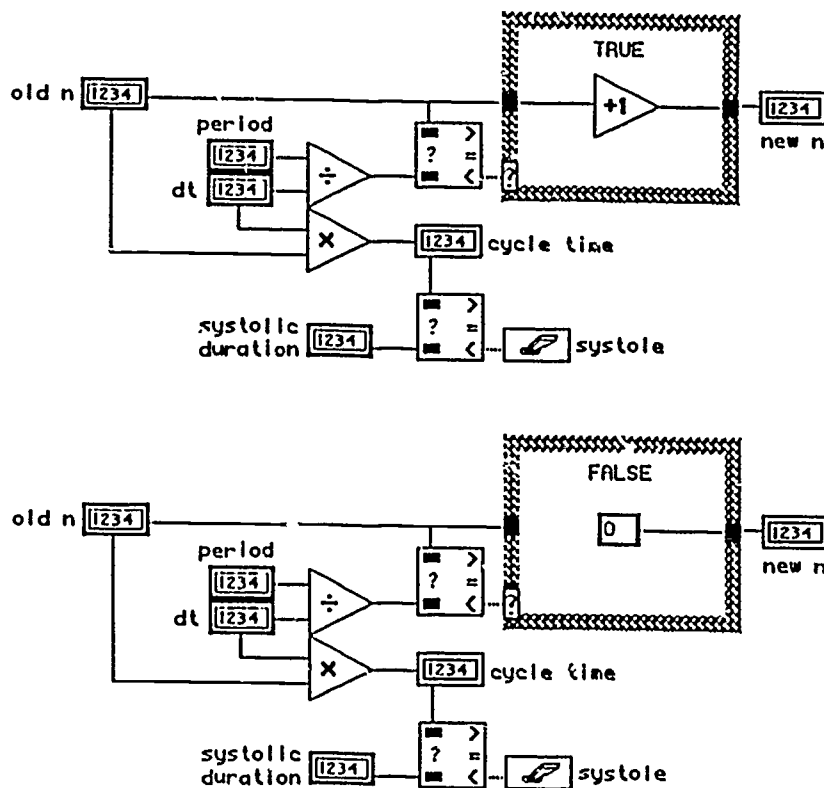


FIGURE 5. Diagram of the virtual instrument "timer." Shown are contents of timer that appears as the hourglass icon in Figure 4. From the period of total cardiac cycle and the integration interval ( $dt$ ), this instrument calculates the number of iterations per cardiac cycle, computes the time elapsed since the beginning of the current cardiac cycle, and provides a boolean (true-false) signal of whether or not the heart is in systole. Both the true and false forms of the case structure are shown.

the timer is true, the calculations specified in the TRUE form of the case structure yield the instantaneous ventricular ejection. During diastole, while the boolean signal from the timer is false, the FALSE form of the case structure simply generates a zero for ventricular ejection. The instantaneous volume in the arterial compartment is determined by the difference between ventricular ejection and peripheral runoff. On the first iteration, peripheral runoff is calculated by dividing the initial diastolic pressure (65 mmHg) by the total peripheral resistance set on the front panel slide switch. At the center of the main diagram (Figure 4), peripheral runoff is subtracted from ventricular ejection. The difference between arterial inflow and outflow is multiplied by  $dt$  within the virtual instrument

symbolized by the homemade integral sign. The result of this integration is the volume increment or decrement that is added to the arterial volume from the previous iteration.

Although it is impossible for an arterial system to have a negative volume, a computer model is under no such restriction. Indeed, depending on the values selected on the front panel, it is quite possible for the present model to develop negative volumes in the arterial compartment or to generate invalid numbers that require the user to restart the program (eg, by dividing by zero). Therefore, the next step in the program is an error checking routine performed by the virtual instrument symbolized by the zero check icon.

Figure 7 shows the diagram for the zero checker. To trap negative or in-

### Pressure Pulse Model V.E. Diagram

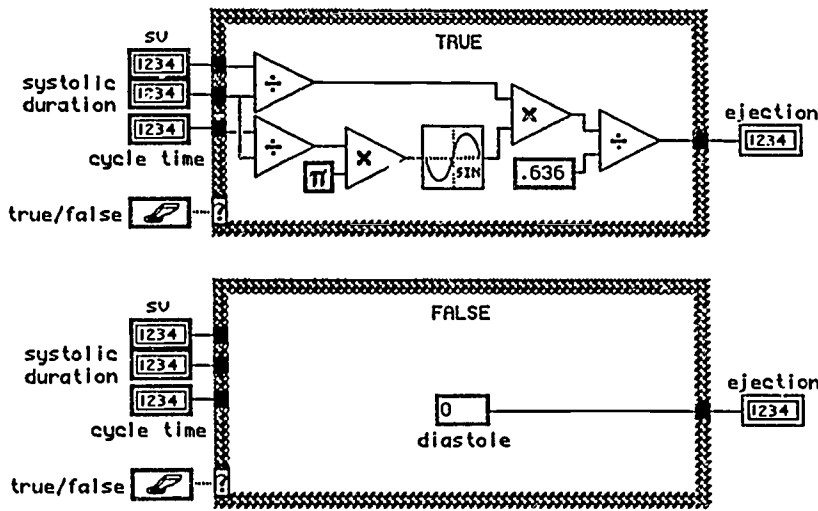


FIGURE 6. Diagram of the virtual instrument for ventricular ejection. Shown are details of the ventricular ejection calculation symbolized in Figure 4 by the V.E. icon. This instrument generates the positive half of a sine wave during systole (true case) and sets ventricular ejection to zero during diastole (false case).

valid numbers, the numbers are tested against zero. All numbers greater than zero are passed through the TRUE form of the case structure, whereas numbers less than or equal to zero are reset to one.

After passing through the zero checker, the arterial volume is simply divid-

ed by the arterial compliance to determine the instantaneous arterial pressure which is then registered on the strip-chart and temporarily stored in the shift-register (lower right border of Figure 4). To determine the systolic and diastolic pressures and the amplitude of the pressure pulse, the instanta-

### Pressure Pulse Model Zero Checker Diagram

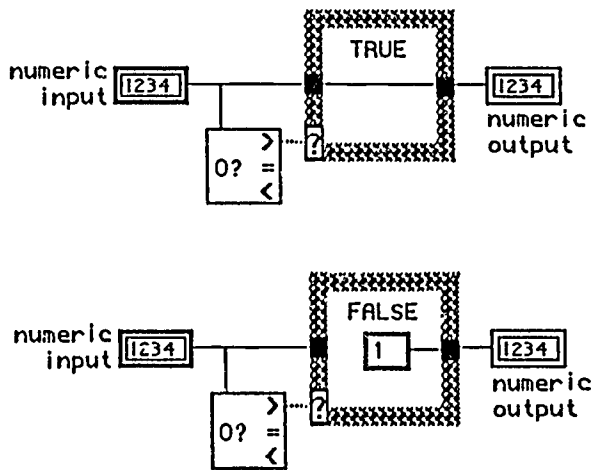


FIGURE 7. Diagram of the zero checker. This is an error-trapping routine that eliminates invalid numbers such as negative arterial volumes and division by zero errors.

neous arterial pressures for each cardiac cycle are accumulated in an array by the virtual instrument symbolized by the wave analyzer icon.

The diagram for the wave analyzer is shown in Figure 8. Recall that the timer sets  $n$  equal to zero at the beginning of each cardiac cycle. The wave analyzer consists of a TRUE-FALSE case structure driven by the current value of  $n$ . At the beginning of each cardiac cycle (when  $n$  equals zero), the array of instantaneous pressures accumulated by the array builder during the previous cycle is processed by the "max-min" function (a menu-available function in LabView) thus retrieving the systolic and diastolic pressures. The amplitude of the pressure pulse is simply the systolic minus the diastolic pressure. The current instantaneous arterial pressure is wired to an empty array builder to become the first element of the new cycle's array which is then passed out of the TRUE form of the case structure to the shift-register on the right border of the While-Wend loop. When  $n$  is greater than zero during each subsequent iteration of the cycle, the FALSE form of the case structure uses a single array builder to add each new instantaneous pressure to the accumulating array until  $n$  is again set to zero by the timer.

### EVALUATION

For the life scientist, even one reasonably adept at computer programming, constructing a mathematical model has usually been tedious and time consuming because he had to develop specific routines for data input and output, for approximating nonlinear functions, and for integrating time-dependent variables. Simulations, particularly those on mainframe computers, were generally written in compiled languages that did not have graphics and that often prevented interactive use of the model. Even with the advent of microcomputers, the investigator had to expend more time and effort on supporting software for graphics and interactive utilities than on the model itself. Therefore, when we became aware of LabView, our hope was that a graphic programming language would make modeling more intuitive and much easier

for the life scientist than it had been in the past.

In evaluating LabView, we used several criteria. First, we felt that the user should only have to build his model, not develop graphics or interactive routines. Second, the user should be able to run the simulation continuously and obtain results in a readily comprehensible form such as strip-charts or X-Y plots. Third, the simulation should be executed in an acceptable amount of time. Fourth, the model should be interactive to the extent that the modeler can easily interrupt the simulation to change a variable's value and resume the simulation. Similarly, altering the model itself and beginning a new simulation should be simple and straight forward. Finally, the simulation language should have all the arithmetic, transcendental, and matrix math routines that high-level languages usually offer. LabView satisfies many of these criteria.

### Programming Environment

The metaphor of a virtual instrument, with controls and indicators on a front panel and an executable block diagram, is a concept readily appreciated by most physiologists. Indeed, the ease of presenting both input and output data in graphic, easily assimilated forms is perhaps the best feature of LabView. In fact, largely because of LabView's virtual instrument approach to programming, it was surprisingly easy to exploit LabView's ability to display data graphically and to implement the two models we described. Starting from a short list of values for constants and the inputs and outputs needed for the front panel, only three hours of programming were required to create an operational version of the simple cardiovascular model. To develop a similar model in BASIC with the same capabilities for user interaction and the same high quality graphics would have required much more time and programming expertise.

### Front Panel

Although the two models we presented make limited use of the many available front panel controls and indicators, it should be noted that all of the controls

### Pressure Pulse Model Wave Analyzer Diagram

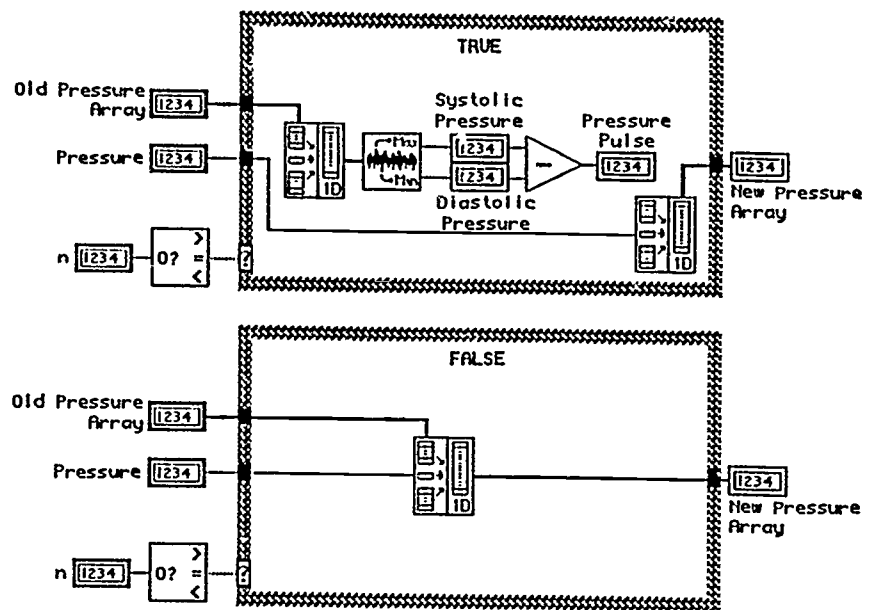


FIGURE 8. Diagram of wave analyzer. At the beginning of each cardiac cycle (true case), this instrument uses the "min-max" function to select the diastolic and systolic pressures from an array of pressure values stored during the previous cardiac cycle (false case).

and indicators in LabView are menu-selected with specified ranges and scales that can easily be changed by the user (though not while a program is running). Range checking is done automatically, and out-of-range errors can be ignored, coerced within the specified range, or allowed to stop the program (as determined by the user). Positioning and changing the size of the controls and indicators to achieve a particular layout on the front panel are relatively simple operations that require some familiarity with the Macintosh mouse. However, there is no provision for customizing controls or indicators or for adding additional graphics to the front panel (eg, it is not possible to add a schematic illustration of a model to the front panel).

### Block Diagram

The block diagram is analogous to the program code in other high-level languages. In a small program such as the simple cardiovascular model (Figure 2), the diagram is truly self-explanatory except for one or two LabView idiosyncrasies like the shift-registers (the only way to use variable values recur-

sively). However, even slightly more complicated programs like the pulse pressure model (Figure 4) can be difficult to decipher and debug. Their indecipherability stems partly from the inclusion of virtual instruments within the main block diagram. The block diagrams of such instruments cannot be viewed at the same time that the overall diagram is examined. Furthermore, case structures and other program control devices are symbolized by several layers of figures, only one of which can be viewed on the screen at a time.

Constructing a comprehensible LabView block diagram requires a sense of spatial organization and some artistry. In LabView block diagrams, data flow occurs most naturally from left to right because the input terminals of function icons are on their left, and their outputs are on the right. The orientation of the icons cannot be changed. A further constraint is the small size of the Mac Plus screen. Therefore, some forethought is necessary to anticipate the overall layout of the diagram and the amount of space that the various function icons will occupy on the screen.



Failing to plan ahead inevitably leads to spaghetti-like tangles of wire that cannot easily be debugged. Even worse, editing such poorly planned diagrams requires completely "un-wiring" the entire diagram just to make a few simple changes.

In LabView block diagrams, four different types of data are depicted by wires with different appearances. Fortunately, LabView does not make the programmer specify the type he needs but instead makes the appropriate selection automatically depending, for example, on whether the selected function requires a simple variable or an array. Logical errors and connections between inappropriate devices are immediately indicated by "broken wires," an error message that often fails to pinpoint the problem.

#### Execution Speed

We have not quantified the execution speed of LabView, but our impression is that it runs at about the same speed as an interpreted BASIC. Thus, execution speed is a potential problem. On the Mac Plus, the pulse pressure model executed at an acceptable rate, but each iteration produced a perceptible jitter as the strip-chart simulator scrolled across the monitor. By contrast, on the faster Macintosh II, the arterial waveform glided smoothly across the screen "like the pulse of a perfect heart." Therefore, to maintain acceptable speed with larger models than those presented here, it may be necessary to run the simulations on a Macintosh II. Alternatively, the program could be compiled. The manufacturer claims to have a compiler for LabView, although it is not yet available commercially.

#### Conclusions

Before using it, we had the impression that LabView would make mathematical modeling possible for all life scientists who are not computer programmers. In most respects, LabView lived up to this expectation, but to some extent it did not. Certainly, the many useful features of LabView, such as the math routines and the graphic displays, can easily be used by the unsophisticated programmer. Similarly, simple diagrams are indeed self-explanatory

programs. In our opinion, an inexperienced programmer could construct a much more sophisticated model in LabView than in other high-level languages. However, more elaborate LabView programs can be as difficult to decipher and debug as those in other languages. Similarly, like other computer languages, LabView can demand arcane programming tricks, and the methods that have to be used to store variable values and to branch on a particular condition are often not straightforward. Nevertheless, even the accomplished programmer will find much to admire and use in LabView: arithmetic and logical functions, array arithmetic, matrix and vector algebra, complex numbers, statistical functions, and signal processing routines including fast Fourier transforms. All of these routines appear as simple icons that naive and sophisticated programmers alike can easily incorporate into their models. Therefore, considering these and the previously mentioned advantages of LabView, we recommend that LabView be seriously considered as a novel modeling methodology with the potential of enabling the average life scientist to simulate complex biological systems.

To obtain a copy of the LabView programs presented here, send the authors an 800k, 3.5 inch disk and an appropriate, stamped, self-addressed mailing container.

This work was supported by USPHS grants HL-36080 and AM-33024.

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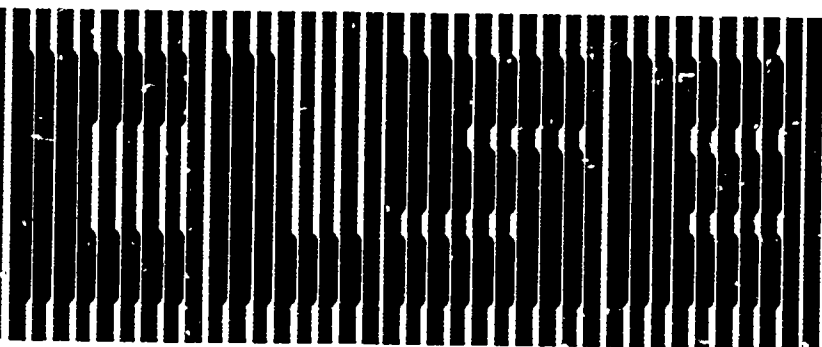
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## USE OF A COMPUTER SIMULATION TO REINFORCE WET LABS IN NUCLEAR MEDICINE

Harold I. Modell and Michael M. Graham  
*Department of Radiology, University of Washington, Seattle, Washington*

Many procedures commonly encountered in Nuclear Medicine are indicator dilution determinations. A known quantity of a radionuclide, for example, is introduced into an unknown volume. By comparing the activity within an aliquot of the unknown to that of a standard, the unknown volume can be calculated. The samples are generally counted in a well counter. If these determinations are to be carried out accurately and interpreted properly, it is essential that Nuclear Medicine technologists and physicians understand the

conservation of mass principles underlying indicator dilution techniques as well as the errors that can arise as a result of well counter characteristics. The American Board of Nuclear Medicine recommends that all resident training programs include principles and limitations of both indicator dilution techniques and well counters.

In our resident program, these issues have been addressed through a series of laboratory exercises conducted once each year. To ensure that the lab sessions do not interfere with Nuclear

Medicine clinic activities, a finite time commitment must be made to deal with various logistical problems related to equipment, space, and scheduling. Although some residents wish to explore these topics further during the remainder of the year, they have been unable to do so because of time and space constraints. To overcome this limitation, we have developed a simulation of a well counter, designed for use currently with Apple II computers, that allows the user to explore indicator dilution principles and the effects of counter characteristics on counting efficiency.

#### WELL COUNTER DESIGN

The "standard" well counter currently used in Nuclear Medicine environments consists of a sodium iodide (NaI) crystal with a hole in it, a photomultiplier tube, a high voltage source, an amplifier, and a multi-channel analyzer. The gamma rays from a sample in the hole interact with the crystal, causing flashes of light. The brightness of each flash of light is proportional to the energy of the gamma ray. The light is sensed by the photomultiplier tube resulting in a small electrical pulse that is proportional to the brightness of the flash of light. The small pulse is amplified and sent to the multi-channel analyzer, which, along with its electronics, displays the data.

Counts obtained with the NaI crystal well counter are subject to a variety of factors. Some of these are related to

THE FOLLOWING ISOTOPE STANDARDS ARE CURRENTLY AVAILABLE FOR YOUR USE:

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- 2) - COBALT-60
- 3) - CHROMIUM-51
- 4) - GALLIUM-67
- 5) - INDIUM-111
- 6) - IODINE-125
- 7) - IODINE-131
- 8) - TECHNETIUM-99M
- 9) - YTTERBIUM-169

CHOICE?

FIGURE 1. Initial screen of well counter simulation listing the isotopes that can be counted.

the NaI crystal and to the electronics processing the detected events, some depend on the sample size and the geometry of the counting system, and others are dependent upon the random nature of radioactive disintegrations. The counting efficiency reflects the interaction of these factors. To gain meaningful information when using such an instrument, it is necessary to understand the factors contributing to the counting efficiency.

The wet labs in our curriculum are designed to help the residents examine a number of these factors including position of the sample within the detector chamber, sample volume, influence of the container in which the sample is placed, and setting the counting window.

#### WELL COUNTER SIMULATION

The well counter simulation provides a means by which students can extend their wet lab experience to further explore indicator dilution principles and the effects of counter characteristics on counting efficiency. Dead time losses and Poisson statistics are incorporated into the counting scheme. Counting characteristics of the simulated counter were drawn from data described by Hine<sup>1</sup> and Sorenson and Phelps.<sup>2</sup>

#### Setting up the counter

When using the simulation, the student is presented first with a list of standards, shown in Figure 1, that are available for setting the counting window. The digitized energy spectrum

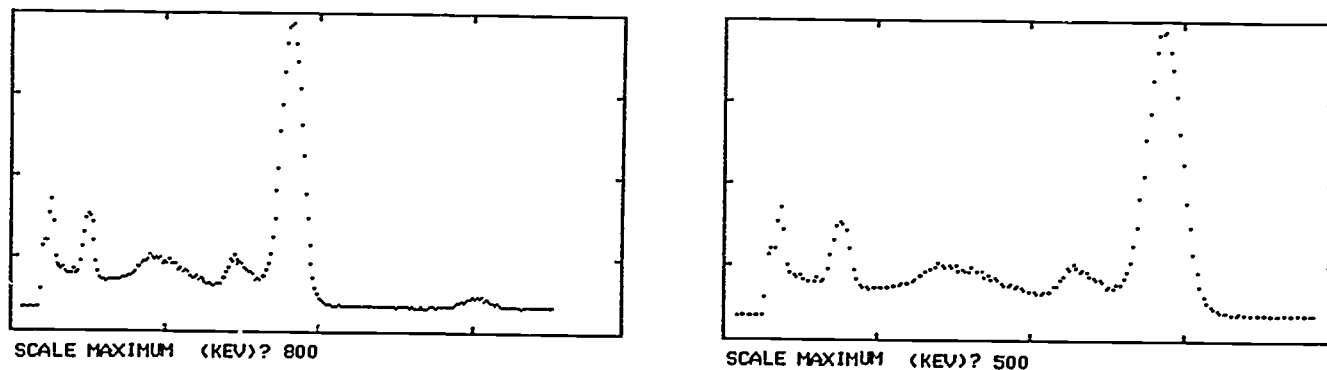


FIGURE 2. Energy spectra for Iodine-131. In the left panel, the student has chosen 800 KeV as the scale limit. In the right panel, the spectrum is shown with 500 KeV as the scale limit. The process can be repeated as many times as desired.

for each of the standards is stored in a separate file and accessed by the program as needed. After choosing the desired standard, the student is presented with an energy scale on which the energy spectrum of the chosen standard will be displayed. The student can adjust the energy scale (Figure 2) to best display the spectrum for setting the desired counting window.

The counting window is then set by using the arrow keys to move a hairline to the desired positions on the spectrum (Figure 3). If the chosen window is unsatisfactory, it can be reset.

### Sample preparation

Having defined the counter set-up, the student must prepare a sample to count (Figure 4). In doing so, the amount of isotope to be used as the indicator and the volume in which it is to be diluted are defined. At this point, a message is presented reminding the student that the basic underlying assumption in all indicator dilution determinations is that the volume in which the indicator resides is well mixed.

The next step is to define the sample volume that is to be put into the counter and define the counting time.

### Counter data

When sample preparation is complete, and counting parameters have been defined, the program reviews the sample definition and counter parameters and

## SAMPLE PREPARATION

AMOUNT OF I131 TO USE AS INDICATOR (MCI)?5

VOLUME FOR DILUTION (L)?1

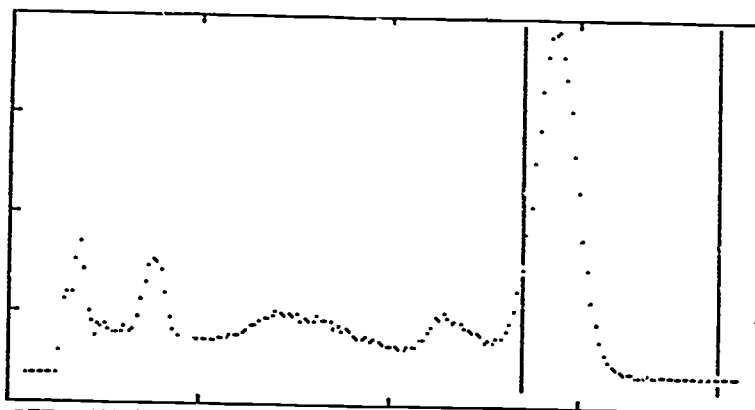
ASSUMPTION

VOLUME WITH INDICATOR IS WELL MIXED!

SAMPLE VOLUME FOR COUNTER (ML)?1

COUNTING TIME (SEC)?30

FIGURE 4. Interactive screen on which the student has defined how the sample is to be prepared.



SET WINDOW UPPER LIMIT  
USE ← → KEYS TO ADJUST HAIRLINE  
PRESS 'A' TO ACCEPT POSITION

FIGURE 3. Spectrum of Iodine-131 with hairlines indicating the energy limits chosen for the counting window. The lower limit has been defined.

presents the results of the counting process (Figure 5). This screen also contains information about the counting efficiency. It presents the apparent counting efficiency (total counts detected relative to the total theoretical counts that should have been detected) as well as the actual overall efficiency (apparent counting efficiency corrected for background).

### Options

After the counting data have been presented, the student is presented with an options menu (Figure 6) that allows redefinition of the tracer, counter parameters, and sample parameters. At

this point, the choice to count the same sample again is also provided. In this way, the random variation in counts seen with actual well counters can be examined. For example, Table 1 lists the counts recorded when the same sample of Iodine-131 was counted five times.

### Error messages

Error messages presented to the student are designed to promote an additional learning experience by providing additional information relevant to the error. In some cases, the error relates to a specific choice made by the student. For example, if the student chooses to count the sample for less than 10 seconds, a message is presented indicating that a longer count time will provide better statistical data. In other cases, the error results from the counter characteristics or from a combination of choices. For example, if the count rate from the sample is too high for the counter, a message appears indicating that the counter is paralyzed.

### MODEL SOLUTION

Data used for incorporating the effects of the various counting factors such as the effects of counter deadtime and sample size were obtained from chapters on well counters presented by Hine<sup>1</sup> and Sorenson and Phelps<sup>2</sup>. Sev-



ISOTOPE: I131

COUNTER SETTINGS:

WINDOW = 342 TO 384 KEV  
COUNTING TIME = 30 SEC

SAMPLE PREPARATION:

5 MCI IN 1 L

SAMPLE = 1 ML

TOTAL COUNTS DETECTED = 402858

APPARENT COUNTING EFFICIENCY = 19 %

ACTUAL OVERALL EFFICIENCY = 7 %

<PRESS ANY KEY TO CONTINUE>

FIGURE 5. Output showing the results of the well counter count.

eral factors enter in to calculating the counts detected for a given sample. Initially, the actual activity (mCi) per ml of sample is calculated. Using this value, the theoretical counts/sec for each ml of sample are calculated according to the following equation.

$$\text{Counts/sec} = \text{mCi} \times (3.7 \times 10^7 \text{ disintegrations/sec}) \times \text{Counting efficiency of the isotope}$$

To obtain the counting efficiency of each isotope, the gamma ray efficiency/disintegration is multiplied by the quantum detection efficiency for each gamma ray produced by the isotope. The resultant values are then summed to obtain the counting efficiency.<sup>1</sup>

The value obtained for counts/sec is then adjusted for the sample volume if the sample volume is greater than 1 ml and for samples that have been diluted.<sup>1</sup>

The influence of deadtime is then incorporated into the calculation according to Sorenson and Phelps.<sup>2</sup>

Background activity, calculated as .35 counts/sec per Kev in the counting window, is the next factor added to the counts detected.

The total counts reported are calculated from the following equation.

$$\text{Total counts} = ((\text{counts/sec} \times \% \text{ spectrum that represents}$$

OPTIONS:

- 1) CHANGE ISOTOPES
- 2) RESET WINDOW
- 3) CHANGE AMOUNT OF INDICATOR
- 4) CHANGE VOLUME FOR DILUTION
- 5) CHANGE SAMPLE VOLUME
- 6) DILUTE SAMPLE VOLUME
- 7) CHANGE COUNTING TIME
- 8) COUNT SAMPLE
- 9) QUIT

CHOICE?

FIGURE 6. Options menu from well counter program.

Table 1. Counts obtained from repeated counting trials of sample described in Figure 4.

Trial	Counts detected
1	403,070
2	402,228
3	402,567
4	402,905
5	403,205

the counting window) + background) x total counting time.

This value is then adjusted by adding a random gaussian variation to it.

The total theoretical counts that should have been detected (used to present the actual overall efficiency to the student) is calculated from the actual activity per ml of sample, the definition of a mCi of activity, the counting time, the fraction of the energy spectrum within the counting window, and the sample size being counted.

#### EVALUATION

A systematic evaluation of the effectiveness of this program has been difficult to perform because the number of new residents exposed to the material each year is very small, normally less than five, and the program is available on an optional basis. Nevertheless, residents who have used the program have indicated that they have found it helpful in reinforcing the material presented in the wet labs.

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<i>O<sub>2</sub> and CO<sub>2</sub> Dissociation Curves</i>	Interrelationships between the O <sub>2</sub> and CO <sub>2</sub> dissociation curves
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<i>Gas Exchange in a Single Alveolus</i>	Effects of ventilator-perfusion ratio and inspired gas composition on exchange in a single exchange unit
<i>The Non-Uniform Lung</i>	Gas exchange from atmosphere to tissues with V <sub>A</sub> /Q mismatching in the lung
<i>Overall Gas Exchange</i>	Gas exchange from atmosphere to tissues with V <sub>A</sub> /Q mismatching and a true shunt
<b>ACID-BASE BALANCE</b>	
<i>Acid-Base Balance: Fundamental Relationships</i>	Acid-Base balance from a Base Excess viewpoint

## CAN YOU HELP?

Among the major goals of NRCLSE are to help life science educators identify colleagues who share a common interest in the use of the computer as an educational tool (see Annual Colleague Directory, February - April, 1988 CLSE) and to help life science educators identify appropriate software for use in their curricula.

In an attempt to achieve the latter, we routinely publish software lists (see Where's the Software, May - June, 1988 CLSE). However, these lists do not offer any information regarding the applicability of programs to specific curricular needs. To provide this type of information, *we need your help*. Please take a few moments to let us know *what software works for you*. Providing information in the format presented below will help us to organize this information to share with colleagues.

Mail information to NRCLSE, Mail Stop RC-70, University of Washington, Seattle, WA 98195. Please include your name, address, phone number, and BITnet address (if applicable) with your response.

### SOFTWARE INFORMATION REPORT

Name of software:

Source of software:

Type of program:

Tutorial (Q & A)

Simulation

Combination

Equipment needed to run program:

Apple II

IBM-PC

Macintosh

Mainframe

Other

Student population using software:

Undergraduate

Graduate

Medical/Veterinary

Dental

Nursing

Content area covered by this software:

How do you use this software:

Independent study

Classroom instruction

Student laboratory

Would you be willing to discuss your use of this program with colleagues?

Yes

No

What do you like *best* about this software?

What do you like *least* about this software?



**AIMS AND SCOPE**

The goal of *Computers in Life Science Education* is to provide a means of communication among life science educators who anticipate or are currently employing the computer as an educational tool. The range of content includes, but is not limited to, articles focusing on computer applications and their underlying philosophy, reports on faculty/student experiences with computers in teaching environments, and software/hardware reviews in both basic science and clinical education settings.

**INVITATION TO CONTRIBUTORS**

Articles consistent with the goals of *Computers in Life Science Education* are invited for possible publication in the newsletter.

**PREPARATION AND SUBMISSION OF MATERIAL**

Articles submitted for publication should not exceed 2000 words and should be typewritten, double spaced, with wide margins. The original and two copies including two sets of figures and tables should be sent to the Editor: Dr. Harold Modell, NRCLSE, Mail Stop RC-70, University of Washington, Seattle, WA 98195.

Title page should include full title, list of authors, academic or professional affiliations, and complete address and phone number of the corresponding author.

Illustrations should be submitted as original drawings in India ink or sharp, unmounted photographs on glossy paper. The lettering should be such that it can be legible after reduction (width of one column = 5.7 cm).

Reference style and form should follow the "number system with references alphabetized" described in the Council of Biology Editors Style Manual. References should be listed in alphabetical order by the first author's last name, numbered consecutively, and cited in the text by these numbers.

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## USING *STELLA* SIMULATION SOFTWARE IN LIFE SCIENCE EDUCATION

Thomas G. Coon

*School of Forestry, Fisheries and Wildlife, University of Missouri, Columbia, Missouri*

Mathematical modeling has a rich potential for use in teaching the life sciences. While traditional approaches may emphasize memorization of structures and functions, simulation allows a student to develop his or her understanding and analysis of living systems by testing hypotheses about how the systems work. In the past, simulation has been unapproachable in other than advanced undergraduate or graduate level courses, mainly for logistical reasons. The use of modeling in teaching is dependent on students 1) having access to computer facilities adequate to

handle many iterative calculations and large data sets and 2) being facile programmers. Before the development of personal computers with large memory and high speed, students were limited to the use of mainframe computers and standard multi-purpose programming languages such as FORTRAN. While the newer personal computers have sufficient speed and memory to serve even the sloppiest of modelers, they did not eliminate the need for students to be programmers before becoming modelers. It was nearly inevitable that a simulation class would devolve into a

### EDITOR'S NOTE

Due to space limitations, *Keeping Abreast of the Literature*, CLSE's regular quarterly feature due to appear in this issue will appear in next month's issue.

programming class; the students would develop their programming skills, but rarely to the level at which they could use those skills to build, test and use their own models.

Several solutions have developed to alleviate the need for students in biological modeling to be expert programmers. One solution is custom-made software packages that are programmed to run simulations on particular subjects, such as those described by Eckblad.<sup>1</sup> Another, more general solution is a programming language designed for modeling that students can learn and use on personal computers. One package designed to meet this need is STELLA, developed by Barry Richmond and others at High Performance Systems, Inc.<sup>3</sup> STELLA was designed by Richmond et al<sup>4</sup> to make modeling available as a learning tool for students of all disciplines. The name STELLA is an acronym for Structural Thinking, Experiential Learning Laboratory with Animation, a cumbersome explanation that is less prominently expressed in later versions of the user's guide. The creators of this software state that their motivation for developing STELLA was to make the discovery approach to any discipline of study more accessible and more fruitful. Although this article does not address all disciplines, STELLA is a resounding success for teaching the dynamic aspects of the life sciences. It is important to keep in mind that the reason for using modeling as a teaching tool is not to make it easier for students to make predictions about the outcome of life processes, but to make it possible for students to develop their understanding of life processes. Students can effectively accomplish this goal if they are challenged to 1) state their understanding of the dynamics of a living system in a structured format (ie, a model), and 2) test their understanding by asking questions of their model using a structured, experimental procedure. STELLA allows students to carry out both of these steps.

STELLA is designed to operate on Macintosh computers (512K memory or greater), and it takes full advantage of the distinctive Macintosh user inter-

face. It allows a user to begin with a conceptual model of a system and move through the standard steps of model building and testing until the model is ready to use as an experimental instrument of study. It reinforces the standard modeling process of building a conceptual model, making the model quantitative, validating the model, and then using the model as a study tool.<sup>2</sup> Furthermore, it keeps the user honest in that the model will not work unless every parameter is defined and connected to the rest of the model. It is friendly enough to inform the errant user of the parameters that remain to be defined.

As is true for any good software, STELLA has been modified and updated several times since it became commercially available in late 1985. The initial version used some home-grown substitutes for standard Macintosh menu selections (eg, icons representing sticks of dynamite substituted for the Cut editing command) and offered no convenient means of transferring model output to another file electronically. Similarly, data could be entered into the model only by typing in changes to the initial conditions. These and other shortcomings have been eliminated in Version 2.0, released in early 1988. The STELLA developers have divided the initial version into two packages designed for different user groups, STELLA for Business and STELLA for Education. While the basic architecture is the same for both packages, they differ in the selection of built-in functions and other subtleties. For this review, I have confined my attention to the latter.

#### HOW IT WORKS

As in any modeling exercise, the STELLA modeler begins by constructing a conceptual model that identifies the *essential* components of the model and the *important* relationships between them. I emphasize essential and important because a modeler must be able to sift through the multitude of components and factors inherent to a complex system and focus on those that are crucial to understanding the particular dynamics of the system that interest

the modeler. The advantage of modeling is that it allows the user to simplify the system sufficiently to understand and study it, but it challenges the modeler not to simplify to the extent that the model no longer behaves in the same way as the system. For example, in studying the redfish fishery in the Gulf of Mexico, it may be interesting and true that the culinary and marketing skills of a New Orleans chef had a major impact on the fishery. Yet, for a basic understanding of how redfish populations sustain themselves in an environment that includes fishing nets, it is more important to include processes such as redfish reproductive rates, mortality rates, and fishing rates than Yuppie economics. By simplifying the model, the user can develop an understanding of the system sufficient to later ask the question, "What happens if the demand for redfish quadruples in one year, perhaps due to the success of some Cajun marketing?"

STELLA uses a hydraulic analogy for representing the conceptual model on the monitor screen. The basic elements of the model are stocks, flows, converters and input links. The icons for these are illustrated in Figure 1. Stocks are defined as components that can accumulate or lose their basic elements. In a

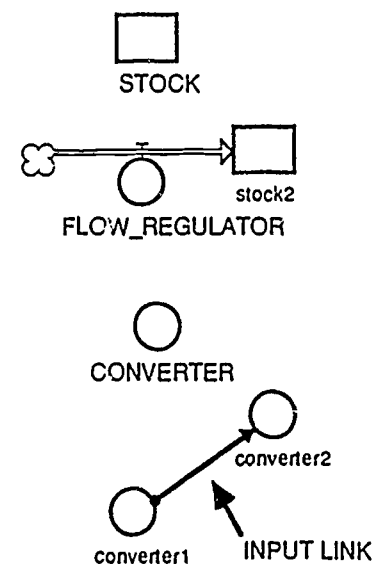


FIGURE 1. Icons used to build conceptual model diagrams in STELLA.

hydraulic analogy, any vessel that can contain water (eg, cup, water tower or ocean) is a stock. In the redfish analogy, a population of redfish could be a stock, or the fleet of fishing boats may be a stock. The stock analogy is a natural for fishery population models, however a little imagination can extend it to any biological system. For the physiologist, the accumulation of acetylcholine in a pre-synaptic neuron or of free oxygen in erythrocytes may be stocks of interest. In constructing a model with STELLA, the modeler begins by placing all of the pertinent stocks on the screen and naming them.

For a stock to accumulate or lose the articles it holds, it must have a means of passage for the articles into and out of the stock. STELLA uses "flows", hydraulic pipes with control valves ("regulators"), to represent this function. The stock representing a redfish population might have flows coming into it that represent births and immigration and flows going out of it that represent natural mortality and fishing mortality (Figure 2). While the flow icons represent the concept of fish moving into and out of the population, they do not show either where the fish come from or go to, or what factors determine how many fish enter or leave the population. The sources and sinks of fish may be extraneous to the goal of the modeler, in which case, STELLA allows the modeler to be vague ("clouds" are at the appropriate end of such flows). If the modeler does want to designate the source of inflows or outflows, she can do so by planting the appropriate source stock at the appropriate end of the flow pipe. Of course this new stock may need some inflow/outflow designations, as well.

All varying flows, such as births or fishing deaths, are regulated by some additional factors. They may be dynamic factors of critical importance to the model (eg, water temperatures during the time of larval development) or other factors of unknown importance to the model. STELLA incorporates the influence of these factors, using icons for "converters", the factors, and "input links" that specify the direction of action. In the redfish model, the

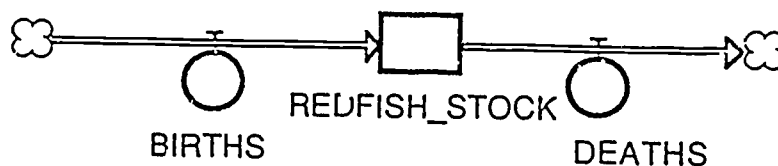


FIGURE 2. Conceptual model of a simple population model showing the model stock, flows, and converters using STELLA.

temperature converter acts on the regulator of the birth flow (Figure 3). Other factors that influence birth rates may be added to the model (eg, plankton abundance, larval abundance) as converters, or taken from other parts of the model already represented (eg, the population stock). Furthermore, converters can be used to influence other converters rather than flow regulators. All other stocks, flows, converters, and input links can and should be added to the diagram on the monitor screen before the modeler goes on to define the quantitative relationships between the converters and their objects of action.

STELLA allows the modeler to define the quantitative aspects of the model in either of two ways, by typing in an equation or by drawing a graphical representation of the function. For example, the modeler may not have a reliable data set on the relationship between the number of adult redfish and the number of larval redfish they produce each year, but she may have a general model for such a relationship based on studies of other species. Rather than worry about the details of unknown regression coefficients, the modeler can simply draw a graphical

representation of such a relationship using the command "Become Graph" (Figure 4). On the other hand, the modeler may have derived a regression equation for the relationship between temperature and plankton abundance and the survival of redfish larvae to the end of their first year, based on years of data collection. The modeler can simply type in the equation that expresses this relationship to complete the inflow side of the model. A similar approach can be used to complete the outflow side of the model. In defining equations, a number of mathematical, logical, and trigonometric functions are available as "Built-ins" to the software.

Once the quantitative aspects of the model have been completely defined, the modeler can view the structure of the model in one of two ways. The Diagram window remains as the conceptual representation of the model, and the quantitative expression for each relationship can be viewed by opening the converter or regulator of interest. For a complete list of the equations and graphical relationships, the user can open the Equation window. Both windows can be printed by a simple command.

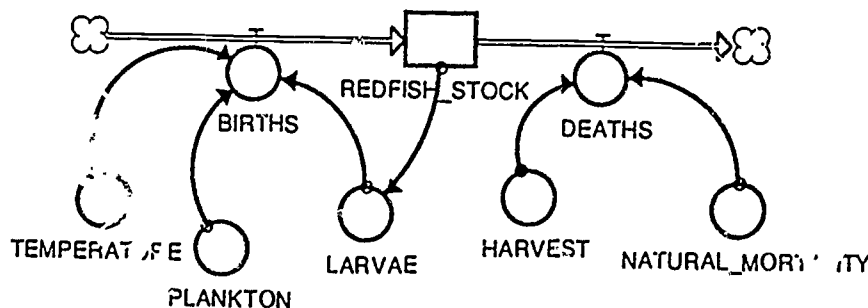


FIGURE 3. Complete conceptual diagram of a population model showing the converters and input links added to the diagram.



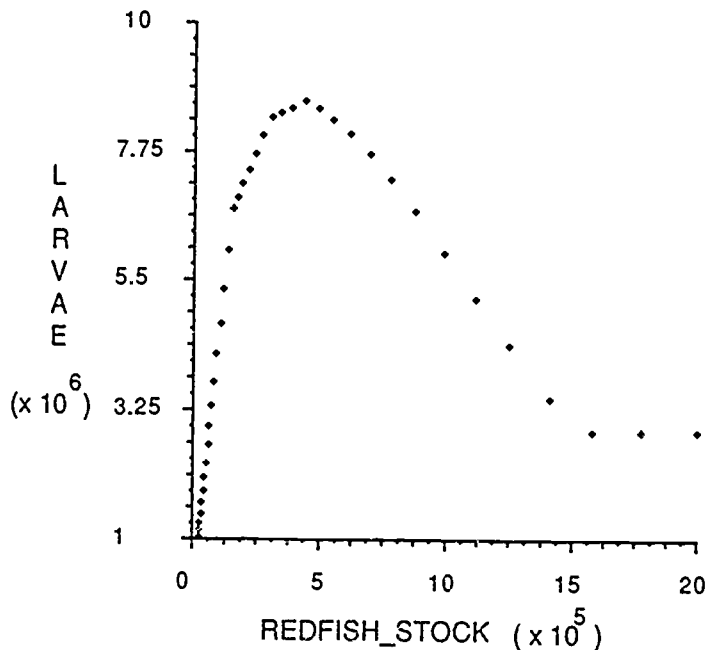


FIGURE 4. Graphical definition of the relationship between the converter "LARVAE" and the stock "REDFISH\_STOCK" using the operation "BECOME GRAPH" in STELLA.

At this point, the model is ready to be run. Before running the model, the user has the opportunity to designate several aspects of the simulation technique. All models are run as time-dependent systems. Thus, the model begins with the initial conditions and makes iterative calculations to determine how the flows and stocks will change over time. The modeler can designate one of three integration methods (Euler's, 2nd-order Runge-Kutta, and 4th-order Runge-Kutta) as well as the increments of time to use in calculations and the starting and ending time for the simulation run.

The value of any of the model stocks or converters can be produced as a model output, and these can be presented in user-defined graphic and tabular forms. The graphic output can be presented as a time-series plot (Figure 5a) or as a plot of one parameter relative to another (Figure 5b). The user defines which parameters to include in the output table and at what time intervals they should be included (Table 1). Output graphs and tables can be viewed on the monitor and can be printed to an external device.

Version 2.0 allows the user to save a

Diagram window or an output graph as a bit-map file accessible by "paint"-type software or as a file of PICT elements accessible by "draw"-type software. Tables and the Equation window can be saved as text files, accessible by many applications, including word processors and spreadsheets (tables only).

After a model is operative, the user may need to make some adjustments to assure that the model behaves like the system it represents. Adjustments to equations can be made from the Equation window or by opening the stock, converter, or regulator of interest. Additional converters and input links can be added and defined quickly, without redefining the rest of the model. The different output options and the ability to stop a simulation run before it is completed allow the user to evaluate the effects of these adjustments on the model's performance. Further tests, such as sensitivity analyses, can be performed by use of some built-in functions. Adding the PULSE function to an equation will cause the model to make a one-time and large scale change in the value of a parameter used in the simulation. Thus a pulse of plankton

could be included halfway through a simulation run to determine how the model responds to a one-time alteration in the regulating conditions. Other functions allow for continuous change in the value of a parameter over time (RAMP) or discrete step-wise changes (STEP).

One of the most valuable features of STELLA is the availability of two functions that introduce user-prescribed stochastic behavior in what would otherwise be deterministic models. The RANDOM function will generate a randomly selected value between 0 and 1 to use in describing random behavior of a variable or function. The RANDOM-generated value can be modified by arithmetic transformation to obtain values drawn from any continuous probability function desired. For normally-distributed variation, the NORMAL function can be used. The default normal distribution has a mean of 0 and standard deviation of 1, however these parameters can be modified by adjusting the arguments of the NORMAL function. These functions not only allow the user to observe the sensitivity of the model to variation in the model components, but they can be used to study the influence of natural variation on the behavior of the system being modeled. This is invaluable in helping students to learn what levels of variation are important in modifying system behavior, as well as the particular parameters that are most sensitive to variation.

#### EXAMPLES OF STUDENT PROJECTS

In the process of developing an advanced graduate course in fishery management, I was faced with the dilemma of providing students an opportunity to plan and evaluate fishery management plans without having the luxury of replicable lakes, ponds, and streams or years to carry out the experiments. Simulation seemed a viable means for the students to develop their plans and to evaluate their consequences within the reality of a classroom. A variety of mainframe and microcomputer lab facilities are available at no additional expense to the students on our campus. However, I anticipated that few of the

students would have the computer skills necessary to convert their models into program code and feared that lab sessions would be little more than debugging sessions. With some frustration, I made my course plans based on a traditional review of case histories, structured discussion sessions, and required term papers.

I received a copy of the original version of STELLA the week before classes began and reviewed it immediately. It was obvious immediately that STELLA would allow me to teach the course as I wanted and within the constraints (eg, limited student programming ability) that I faced. With some trepidation, I cast aside my syllabus and outlines and embarked on an adventure of experiential learning of my own as I tried to stay ahead of the students.

My goal was simple. I wanted the students to examine and develop their understanding of how fish populations respond to environmental and human influences and how humans adjust their fishing behavior in response to the same factors. Rather than learning simply from others' mistakes and successes, they were challenged to state their own understandings in the form of models. They could then test the predictions based on these principles against the data of past fishery management practices. In the process of carrying out this exercise, they could refine and deepen their understanding of how fishery systems function, rather than simply find that their ideas were correct or incorrect.

I began the course with an introduction to the kinds of problems faced by fishery managers (relatively small data bases and relatively poor understanding of how the systems function) and then introduced the basic methods of model development and use. The first assignment was to develop a model of a catfish fishery in which fish entered the population by stocking (no natural reproduction) and were removed only by angler harvest or natural mortality. The class worked together to construct a conceptual model agreeable to everyone. They used data available in published literature and agency reports to establish the quantitative relationships

between "converters" (determining factors) and "flow regulators" (processes). In the absence of documented data, they relied upon data on related species and fisheries and expert advice from fishery managers.

Once the class had their quantitative model designed on paper, I gave them a thirty minute introduction to the use of STELLA and then turned them loose in a laboratory with Macintosh computers. None of the students had used a Macintosh computer before. Within one hour, most had their model entered onto the computer and operating.

My reason for describing this se-

quence is to highlight how easily and how quickly students can learn to use STELLA. They need some prior instruction in the practice of modeling, some practice constructing and quantifying a simple model, and less than one hour of instruction to be able to use STELLA. As with any programming language, students become more accomplished and can create more complex models with practice. Prior programming experience certainly facilitates a more advanced approach initially. Even so, students become functionally literate in STELLA extremely quickly.

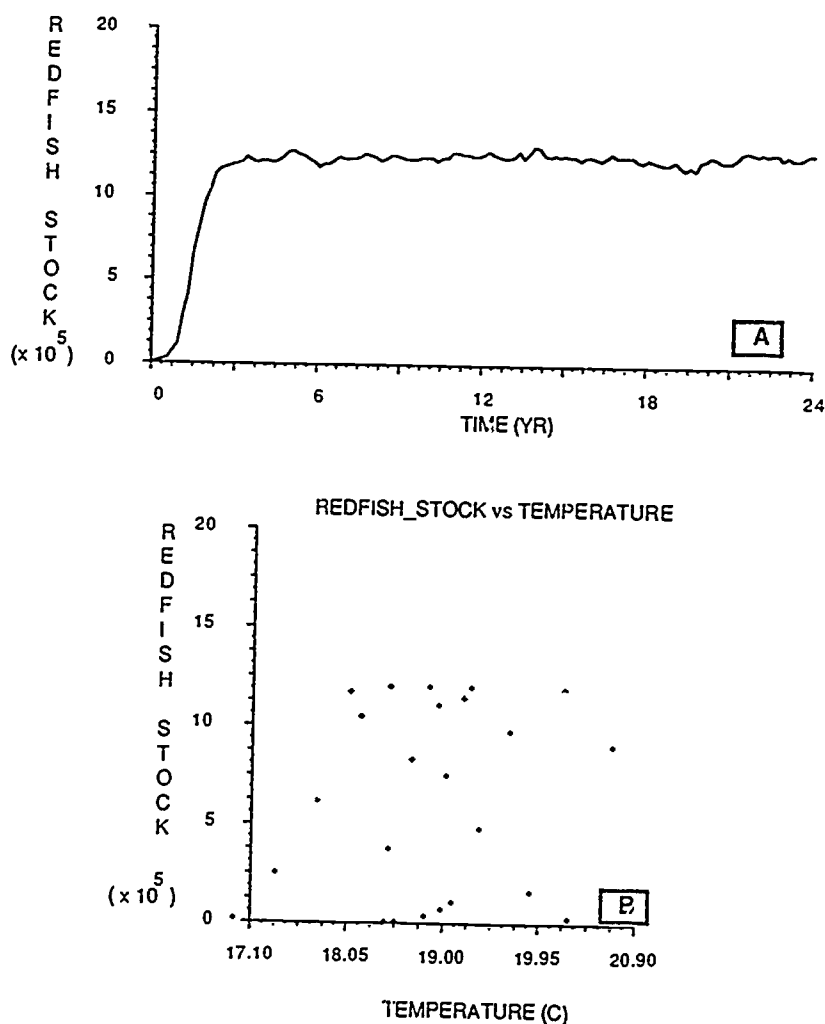


FIGURE 5. Graphical output of the population model illustrated in Figure 3 using STELLA. Graphs can be presented as time series plots (A) or as scatter plots of two variables (B). The function NORMAL was applied to converters "Temperature" and "Plankton" to generate the variation in (A).

TABLE 1. Tabular output of data generated by a simulation run of the REDFISH model using STELLA. Conditions of variation for TEMPERATURE and PLANKTON are the same as in Figure 5.

Time	REDFISH S...	TEMPERAT...	PLANKTON	HARVEST	NATURAL ...
0.0	5000.00	20.76	243.22	950.00	1900.00
1.00	249840.59	16.25	223.74	47469.71	94939.42
2.00	1008724.6	18.66	192.74	191657.67	383315.34
3.00	1194601.3	19.35	214.39	226974.25	453948.50
4.00	1221407.9	19.90	229.57	232067.48	464134.97
5.00	1249427.1	18.15	212.31	237391.16	474782.31
6.00	1247460.6	20.51	207.53	237017.52	474035.03
7.00	1231574.4	18.41	223.52	233999.14	467998.28
8.00	1230035.9	19.45	268.25	233706.83	467413.66
9.00	1242133.0	17.65	225.50	236005.27	472010.53
10.00	1209216.9	19.34	208.97	229751.19	459502.38
11.00	1228582.4	19.41	242.13	233430.66	466861.31
12.00	1207441.1	18.90	206.85	229413.81	458827.63
13.00	1213822.5	18.51	286.60	230626.28	461252.56
14.00	1210243.1	18.38	306.84	229946.19	459892.38
15.00	1264946.0	17.00	194.24	240339.73	480679.47

After the students had used their initial model to ask some basic questions about stocking rates and harvest rates, they quickly found that their model was extremely simple to use; in fact, it was too simple. In the second version of the model, they incorporated individual growth, so that fish could be stocked at a sub-harvestable size and would grow into the harvestable size class. They could have used a variety of model structures to accomplish this, but they settled on breaking the single catfish stock of the original model into stocks of catfish of different sizes. Growth rates (ie, age of passage into the next larger size class) were determined by feeding rates and by the total density of all catfish. They increased the complexity of the outflow side of the model by incorporating length restrictions on fish harvest such that each size class had different harvest mortality rates.

Successive assignments built more demanding features into the modeling problem, yet all could be accomplished with relatively simple models. As with any modeling assignment, students tended to make models more complex than were necessary. STELLA became

unwieldy and slow as model complexity increased, serving as a negative reinforcement against excessive complexity. To incorporate a more biological basis for fish somatic growth into population models, the students designed a parallel model that used millimeters of fish length as a stock that could be increased by environmentally- and density-regulated growth equations. By linking a series of these length stocks, each representing an age class of fish, they could model fish growth with a von Bertalanffy model.<sup>5</sup> The students used the NORMAL function in an equation for recruitment of young fish into the adult stock to study the effects of random variation in environmental conditions (lake level) on the success of fish spawning and juvenile survival. The class developed all of these models as a group project which allowed them to learn from each other as well as from their own experiences.

For the final project however, I required each student to develop his or her own research problem, a model to use in studying the research problem, and a report on his or her findings from the model. This assignment was the most difficult for the students, perhaps

because they had been too accustomed to the group dynamic in finding modeling tactics to accomplish their assigned tasks. The fact that they had been given the problem in previous assignments and had to define the problem in the final assignment may have also limited their success in this exercise.

#### LIMITATIONS TO THE USE OF STELLA

It should be obvious that I consider STELLA an asset in my teaching. Although my initial use was for a graduate course, it is quite adaptable to undergraduate courses. It is simple enough that high school students (or even junior high students) could learn to use it, as long as they are given an appropriate introduction to the methods of modeling. Yet it is powerful enough to be useful in research applications.

One of our Ph.D. graduates used STELLA to model the physiological response of ruffed grouse to various winter habitat conditions. In spite of this breadth of usefulness, STELLA does have some limitations.

Perhaps the first limitation is one that some may consider a trivial one. Educational software that is usable only on a Macintosh computer is less accessible than that which is produced for all microcomputer formats. STELLA is so closely tied to the Macintosh interface that it seems unlikely that it will be available in PC or Apple II format. We are fortunate that our campus has recognized the need for PC and Mac uses and has established computer labs equipped with both types of machines. Other educators may be more limited. This is a fault of educational planners, not of the software.

My only frustrations with STELLA have come when I have tried to push it beyond a pedagogical use. STELLA can be quite useful in research applications, however its major limitation is in the size of models it can handle efficiently and its inability to link separate models in joint or successive simulation runs.

Simulation time slows considerably as the integration time steps are reduced

and as the number of stocks and flows increases beyond 8 or 10. Obviously, one solution is to keep models simple, yet some applications require more detail to produce results usable in a research application. Another solution is to break cumbersome models into subunits that can be run in sequence. STELLA can be used in this way, but it requires continuous involvement of the user to move it from one step to the next. It is not designed to allow the user to automatically link these subunits into a master run, feeding output from one run as input to the next. The only way to accomplish this is to run the first subunit in the series, save the output table in a file, import this data into the next subunit as initial data, run the next subunit, feed the output to the successive subunit, and repeat these steps for each subunit. In defense of the software, this use is extending the package beyond its designed use (to teach modeling), yet it is so close to being useful in other research applications that it seems a desirable and justifiable improvement (perhaps as a "STELLA

for Research").

One other limitation that concerned me involved the random number generating functions (RANDOM and NORMAL). These are extremely useful and efficiently designed functions built into the software. Yet they suffer from an inconvenient limitation. When using stochastic models, it is important to make multiple runs of the model to generate enough sets of output to determine the statistical characteristics of the model results. STELLA requires the user to initiate each simulation run directly, rather than programming the machine to run the simulation 10 or 100 times automatically and to save the appropriate output from each run.

Against the strengths of all that STELLA can do, particularly as a teaching tool, these complaints seem minor groushings. A more canny user might be capable of programming the machine to accomplish what I cannot. If these uses of the software were the goals of the designers, it should not be necessary to be a canny programmer. But the developers knew what they

wanted,<sup>3</sup> and they succeeded remarkably well in meeting their goals. I encourage anyone interested in teaching with simulations to try STELLA. You will not turn it away.

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5. Ricker, WE. *Computation and interpretation of biological statistics of fish populations*. Bulletin 191, Fisheries Research Board of Canada, 1975.

## ANNOUNCEMENT

A Physiology teaching software demonstration will take place at the XXXI International Congress of Physiological Sciences in Helsinki, Finland, July 9-14, 1989. Computers will be available on-site to allow participants a hands-on experience with the programs. Authors wishing to display their software are urged to submit abstracts describing the topic, educational goals, and hardware requirements of their programs. Copies of the programs and minimal documentation (operating instructions) must be forwarded to Helsinki by June 1, 1989. Information about the software demonstration can be obtained from:

Dr. Joel Michael  
Department of Physiology  
Rush Medical College  
Chicago, IL 60612  
(312) 937-6426

Information about the Congress and the required registration material can be obtained from:  
The Secretariat, XXXI International Congress of Physiological Sciences, P.O. Box 722, SF-00101 Helsinki, Finland.



**AIMS AND SCOPE**

The goal of *Computers in Life Science Education* is to provide a means of communication among life science educators who anticipate or are currently employing the computer as an educational tool. The range of content includes, but is not limited to, articles focusing on computer applications and their underlying philosophy, reports on faculty/student experiences with computers in teaching environments, and software/hardware reviews in both basic science and clinical education settings.

**INVITATION TO CONTRIBUTORS**

Articles consistent with the goals of *Computers in Life Science Education* are invited for possible publication in the newsletter.

**PREPARATION AND SUBMISSION OF MATERIAL**

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Title page should include full title, list of authors, academic or professional affiliations, and complete address and phone number of the corresponding author.

Illustrations should be submitted as original drawings in India ink or sharp, unmounted photographs on glossy paper. The lettering should be such that it can be legible after reduction (width of one column = 5.7 cm).

Reference style and form should follow the "number system with references alphabetized" described in the Council of Biology Editors Style Manual. References should be listed in alphabetical order by the first author's last name, numbered consecutively, and cited in the text by these numbers.

**RESPONSIBILITY AND COPYRIGHT**

Authors are responsible for accuracy of statements and opinions expressed in articles. All authors submitting manuscripts will be sent a copyright transfer form to complete. The completed form must be returned before the work will be published.

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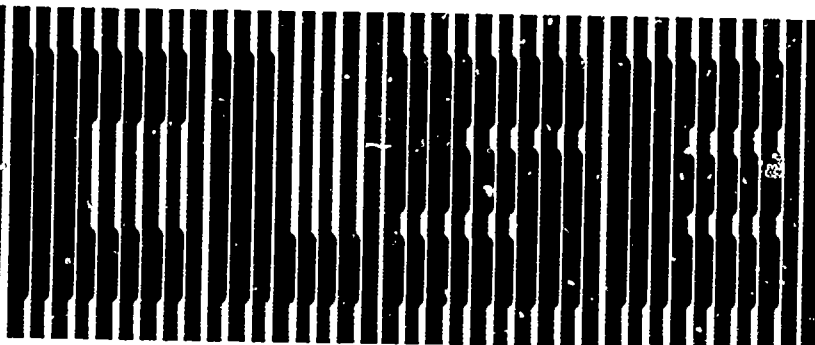
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**NRCLSE**

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## NRCLSE BEGINS SOFTWARE EVALUATION PROGRAM

Harold I. Modell

*Department of Radiology, University of Washington, Seattle, Washington*

One of the goals of NRCLSE has been to establish a peer critique mechanism for software related to life science education.<sup>4</sup> Over the past two years, broad issues related to the overall focus of such a program have been highlighted in CLSE, and reader input was encouraged to help define evaluation criteria.<sup>3,5</sup> The underlying issues, however, appear to be too complex to define simple criteria.

Recently, EDUCOM has also tried to address these issues through two evaluation efforts. The National Center for Research to Improve Postsecondary Teaching and Learning (NCRIPAL) established an annual higher education software awards program, and EDUCOM initiated a scholarly review of academic software project. In its first year, the latter has focused on computer based learning modules in three

general curricular areas, only one of which is life science related. These programs tend to illustrate aspects of the dilemma facing any evaluation process in this area.

Entries in the NCRIPAL program undergo evaluation by instructional design experts as well as content experts. As a result, finalists in the competition tend to be at the "cutting edge" in terms of instructional design criteria and utilization of available computer capability. Thus, although some older software may be very effective in presenting material and hence potentially very helpful to educators, they may fall short of ratings that would convey this to potential users.

The guidelines for reviewers in EDUCOM's software review project reflect an assumption that the software has been designed primarily to "teach" a particular topic. Thus, programs that have been designed to augment classroom interaction or provide animated illustration material in a group setting<sup>1,2</sup> may not receive appropriate reviews.

Two approaches to the overall problem may be taken. The first is to delay implementing a critique mechanism until clear criteria are evident. The second is to implement a mechanism that will not be ideal but that will provide a framework in which evaluation criteria may be tested and allowed to evolve as more experience is gained. After taking the first approach for several years, it has become clear to NRCLSE that significant progress will not occur until the latter approach is attempted.

#### THE NRCLSE PEER CRITIQUE PROGRAM

The NRCLSE peer critique program being initiated with this month's issue of Computers in Life Science Education will serve as the first step in this process. Its initial focus will be micro-computer software (Apple II, MS-DOS, and Macintosh), and its goal will be

two-fold: to provide software authors with feedback and to provide users with information. The feedback will include a critique of the software and suggestions for ways to make it more versatile. Each piece of software will be critiqued from content and instructional design standpoints. Because computer memory and mass storage are no longer limiting factors as they were several years ago, program efficiency does not seem to be as important an issue as it once was. Thus, the programs will not be examined from a computer programming standpoint.

Each author will be required to submit a written narrative addressing the following questions to give reviewers a basis for initial evaluation.

- What need prompted the development of the software?
- For what student population was the software intended?
- In what environment is the software intended to be used (eg, group discussion, independent study)?
- Why was the specific format of the software chosen? (ie, What is the underlying philosophy of the software?)
- How long has the software been used with students?
- What attempts have been made to evaluate the impact of the software, and what were the results of those evaluations?

The software (copies supplied by the author) along with copies of the narrative will be sent to at least two content reviewers and one reviewer with expertise in instructional design. The software will be returned to the author with the reviewers' comments unless the author designates that reviewers may keep the software.

#### FOCUS OF EVALUATIONS

Although development of specific evaluation criteria will require some experi-

ence with the critique process, guidelines, in the form of a series of questions, will be provided for the reviewers. The content reviewers will be asked to focus initially on the following questions.

- Is the content accurate?
- Does the software appear to meet the design criteria specified by the author?
- Are error messages provided where appropriate?
- Do the form of error messages (if appropriate) provide an additional learning experience for students?
- Is the software suitable for student populations or educational settings other than those specified by the author? If not, what modifications would be necessary to make the software more versatile?
- Are you aware of other software that covers the same material? If so, who are the authors, and how does this program differ from the others?
- What features did you like best about the program?
- What features did you like least about the program?

Reviewers with expertise in instructional design will be asked to examine the program with the following questions in mind.

- Are the input and output schemes followed in this program consistent with sound instructional design principles (not necessarily "state of the art")?
- How could the input and output screens be improved and still meet the program needs as stated by the author?

#### FUNDING FOR THE PROGRAM

Because NRCLSE has been unable to identify potential sources of funding for such a program, the program must

be self funding. Each author will be required to submit a nominal fee of \$25 to cover handling, mailing, and follow-up costs associated with the review process.

#### SOFTWARE REPORTS

To meet the goal of providing software users with information concerning available software, each complete program received will be included in the CLSE Where's the Software listing. In addition, each complete package will

be considered for an in-depth review article in CLSE.

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## A STUDENT'S VIEW OF COMPUTER-BASED CURRICULUM INNOVATIONS

Susan Erdman

*Division of Comparative Medicine, Massachusetts Institute of Technology, Cambridge, Massachusetts*

Unlike the classes before them, each student of the Class of '88 of Mississippi State University College of Veterinary Medicine was required to purchase a computer upon matriculation.<sup>1</sup> Having been one of these students, I had the opportunity to witness, first hand, computer-based innovations in the curriculum. After four years, some of us fantasized about nationwide computerized networking systems while others imagined a 128K Macintosh computer falling twenty stories and crashing to the concrete while David Letterman's audience cheered. No doubt, the Class of '88 were pioneers, and, no doubt, the introduction of computers into the curriculum was progressive. The progress, however, was not without controversy.

The computers were added to our curriculum primarily to facilitate information management. The computer applications in the curriculum included

word-processing, diagnostic tools, data organization aids, spreadsheets, and graphics. MacWrite data disks replaced the traditional hard-copy sophomore and junior course objectives. PKC diagnostic and management software was briefly introduced. Think-tank was applied to organize projects, and Jazz was applied to hypothetical practice management exercises. Portions of our freshman anatomy course utilized Filevision to computerize lessons so that information leaves coordinated with anatomical illustrations presented logically on the computer screen.<sup>2</sup> Today's classes also use computerized case analyses that unfold uniquely with each management decision throughout a hypothetical case.

As freshmen, most of us sat staring blankly at the computer screens. Most of us had little previous computer experience, and some had no interest in learning. Fortunately, the curriculum

planners chose Macintosh computers, which are generally easy to learn to use. This standardization also created computer compatibility that facilitated exchange of information and skills. Word processing with MacWrite quickly became popular for vocational and avocational computer use from editing course objectives to polishing resumes. Only computer games were used more frequently than MacWrite. The voluminous course objectives that were presented on a computer disk were easier to store (for example, in a pocket) and access than hard-copies. Moreover, students now had the option of personalizing the course outlines for future reference. (Course objectives are currently presented with Hypercard, which further increases flexibility and accessibility of information storage by establishing software links among related data.) Although we had few opportunities to apply diagnostic software in



the didactic portion of the curriculum, some students applied diagnostic software to clinical cases during the senior year. On several occasions, computer-generated differential diagnoses demonstrated their utility in the clinics.

Interestingly, on a more negative note, my classmates frequently chose to study and work from hard-copies instead of from the computer screen, perhaps because they were more comfortable with the familiar format or, perhaps, because computer screens were harder on the eyes during long hours of study. Even more serious, opponents of the computerization suggested that computers might compromise mastery of the "veterinary essentials."

Considering the burgeoning advances within the veterinary disciplines, it seems difficult to construct an appropriate and manageable knowledge base using traditional approaches. Yet, it is important that veterinarians have access to pertinent and current information. Our veterinary curriculum demonstrated that computers facilitate access to a dynamic information source that can grow with the field. Indeed, computers can help veterinarians harness some of our expanding knowledge resources to improve future veterinary care.

Who knows how much professional impact our computer experience had on the Class of '88? When the future

demonstrates that computers improve veterinary services, no doubt, computers will be further integrated and utilized within our profession. The Class of '88 and their successors will be well prepared.

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## KEEPING ABREAST OF THE LITERATURE

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# NRCLSE SOFTWARE EVALUATION SUBMISSION FORM

The NRCLSE software evaluation program has been initiated to promote development of high quality, versatile educational software in the life sciences by providing authors with feedback from critiques by life science educators and instructional designers. To ensure that software is reviewed in an appropriate fashion, it is essential that reviewers fully understand the rationale underlying the design criteria chosen by the author and the environment for which the software is intended. Please provide the following information concerning each software package to be evaluated.

Submit 3 complete copies of the software and all supporting documentation along with \$25 (to cover handling, mailing, and follow-up costs) to Software Evaluation Program, NRCLSE, Mail Stop RC-70, University of Washington, Seattle, WA 98195.

Author's name:

Author's address:

Title of software:

Content area:

Minimum hardware requirements:

Optimal hardware configuration:

Software requirements (operating system, etc):

Student population for which software was written:

Environment for which software is primarily intended (independent study, classroom discussion, lecture enhancement, etc):

How long has this software been used by students?

Can reviewers keep the review copy of this software?

Describe the underlying philosophy of this software. What need prompted the development of this software? Why was the specific format of the software chosen? What goals did the input/output scheme (or screen design) address? Is documentation an integral part of the package? How is it intended to be used? (Use additional pages if necessary.)

Has the software been evaluated by students?

Briefly describe the format and results of any student evaluation.

What attempts have been made to evaluate the impact of the software on student progress?

What were the results of the impact evaluation?



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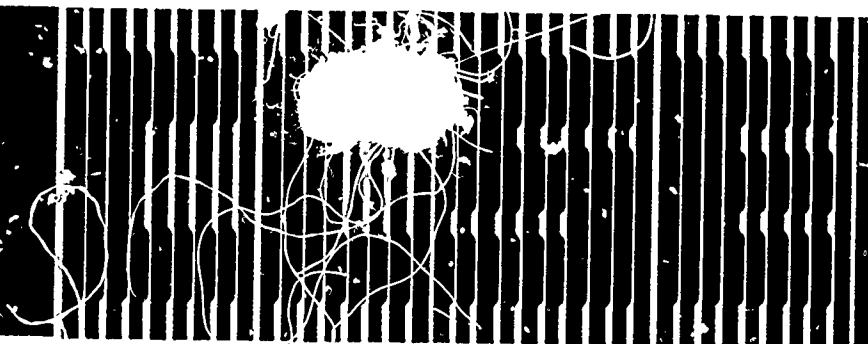
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## SELECTING AN AUTHORIZING SYSTEM 1988: HOW THE VENDORS MAKE IT MORE DIFFICULT

Mary S. Trainor and P. Alexandra Schultejaan

*Los Alamos National Laboratory, Los Alamos, New Mexico*

Selecting an authoring system can be an intimidating prospect, considering the number and diversity of systems on the market today. Most organizations contemplating an authoring system purchase have neither the time nor personnel to devote to lengthy benchmark

testing.<sup>5,8,9</sup> Instead, these organizations are forced to make an evaluation of numerous authoring systems based solely on the information and demonstration packages provided by the vendors. The major problem posed by this approach is that these packages do not

### *Editor's Note*

*The papers in this month's CLSE were presented at the 30th International Conference of the Association for the Development of Computer-Based Instructional Systems held in Philadelphia, PA, November 7-10, 1988 and are reprinted from the proceedings of that conference.*

include the information necessary to evaluate the usefulness and applicability of an authoring system for the organization's purposes.

The Cognitive Systems Engineering Group at Los Alamos National Laboratory has been active in the computer-based training (CBT) field for more than three years. We have developed several prototype computer-based systems for various agencies and are well acquainted with the constraints of programming languages versus authoring systems.

This paper describes our experiences in attempting to recommend an authoring system that would best suit the purposes of our client. Our client had very specific requirements, and we were limited by both time and resources in the evaluation process. We encountered numerous difficulties in our attempt to conduct a thorough evaluation under our time and funding constraints. The information and demonstration packages provided by the vendors contained insufficient information for us to thoroughly evaluate the authoring systems. We present some suggestions to vendors on the information that should be provided to allow potential customers adequate data for evaluation. In addition, this same information can be used by potential authoring systems customers (eg training/education managers and designers) in their evaluation of authoring systems. The goals of the paper are

- 1) to suggest to vendors that they provide more consistent and thorough information, and
- 2) to encourage potential customers to be more demanding of the vendors and more educated in their evaluations.

#### METHODS

Selecting an authoring system should be done with some care to ensure that the system is compatible with the customer's needs and desires. We divided the selection process into four major steps:

##### Step 1. Define requirements

It is important to define the needs of the

group that will be developing the CBT, both now and in the future. These requirements not only include such factors as the skills of the person(s) who are going to be doing the authoring but also the desired features of the courseware. A tradeoff analysis needs to be done, determining priorities for in the development of each authoring system, priorities were set. Your priorities should match those of the system developers. For example, it is difficult to obtain an easy-to-learn interface which also allows for sophisticated instructional strategy development (eg, simulation). It also needs to be taken into account that as a CBT program grows, the level of sophistication desired increases. Thus, features which seem unnecessary now (eg, ability to jump out to a programming language, digital audio) may become highly desirable a year from now. Unless these features are anticipated selection of a highly constraining system may occur. The specific needs and requirements for the evaluation we undertook were:

- An authoring system that is compatible with Zenith Z248's for authoring and the EIDS system for delivery.
- A system capable of handling a variety of instructional strategies, including drill and practice, tutorial, gaming and simulation.
- A computer-managed instruction (CMI) capability which allows for such features as customized record keeping, report generation, a variety of test modes, and learning prescription generation.
- The capability to incorporate other graphics and routines developed in a programming language into the authored courseware. This was desired because of future flexibility, allowing for addition of features not possible through the authoring system.
- A system that is easy-to-learn for nonprogrammers with minimal time spent on documentation.
- Versatility so that authors who become familiar with the system will be able to customize the interface to

their own preferences.

- The capability to customize the feedback relative to the course and to include sophisticated answer-judging routines.
- A system that provides a variety of questioning formats.
- A system that does not charge royalties or include a separate presentation system.
- Minimum maintenance costs.
- The ability to add videodisc capability, and perhaps CD-ROM, in the future.

##### Step 2. Select authoring systems for in-depth evaluation

Because of the number of authoring systems on the market, it was not possible to evaluate them all. We started the narrowing down process by doing background research on authoring systems<sup>1-4,6,7,10,11</sup> and making a preliminary selection from the *CBT Guide to Computer-Based Training Systems and Courseware* published by Data Training and Weingarten Publications, Inc.<sup>12</sup> The Guide covers the basic attributes of 98 authoring systems, and Data Training provides a matrix of system features versus authoring system. The Guide, literature, and our own experience allowed us to select eleven systems as candidates for evaluation.

##### Step 3. Contact vendors for information and demonstration packets

All eleven vendors were contacted by phone for information about their products and for a demonstration of the system. Four vendors were not able to send demonstration discs for evaluation because they had not produced one. One vendor subsequently sent the entire authoring system for us to preview, but we did not have the 8 to 12 hours required to adequately evaluate the system. Of those having a demonstration disc, most desired a small payment for the disc (\$15-25). We explained that we felt, since this was a main marketing tool of theirs and we were potential customers, we should be provided the disc free of charge. In addition, in our institution it would have cost far more than \$15 to \$25 to get the necessary purchasing paperwork

through the system (not to mention time). All vendors cooperated, but in several instances, many phone calls were required in order to obtain the disc. We had to describe in detail who we were and the nature of the evaluation.

The printed information we received on the authoring systems consisted primarily of sales literature; many vendors did not send the pertinent technical details we required for a thorough evaluation. When queried on the phone about this, several vendors said that we were more inquiring than most of their potential customers, and thus they did not have literature prepared at that level of technical detail.

These factors contributed to this being a most time consuming and unpleasant step in the evaluation process.

#### Step 4. In-depth evaluation of the authoring systems

The next step was to analyze each authoring system relative to our criteria and based upon the information and demonstration packages received. Our criteria, shown in Table 1, were developed from our experience, the literature, and from the evaluation requirements. We had thought that the Data Training Guide would provide us the necessary criteria, but after reading it through, we discovered that Data Training only set the features list, and the vendors had great liberty in what they submitted under each feature. None of the information was parallel, making comparisons difficult, if not impossible. We are not faulting Data Training in this discussion, as they were providing a very useful data clearinghouse function compiling the system/vendor information.

#### Step 5. Recommendation of an authoring system for purchase

The last step was to examine our evaluation information and make a final selection. Unfortunately, our results were quite inconclusive because of inadequate information (see discussion of this under RESULTS below). We were able to narrow final candidates to three, however. Because we needed to make a recommendation, we selected

Table 1. Questions to ask when choosing an authoring system

1. What is the cost per instructor station? per student station? What kind of agreement do I need? If number of student stations is large and undefined at this point, then an agreement for unlimited usage and no special presentation system is probably desired.
2. What is the royalty agreement? Am I willing to pay the vendor for each external usage of my courseware? If unlimited distribution is desired, ask about cost of this up front.
3. How easy-to-learn and easy-to-use is the system? Many vendors say their system is easy, and yet they require extensive training delivered by them. Call current system users and ask them and/or try the system out for yourself to answer this question.
4. What are the hardware requirements? Does the system require some specialized hardware you do not have access to or cannot afford? Some vendors developed their system to run on one set of hardware and then retrofit for others. If your hardware is retrofitted, it may be awkward to use.
5. What support does the vendor supply? Does the vendor supply telephone consulting (you will need it as you get started)? Are software updates provided? Are the updates designed to be compatible with earlier versions?
6. What graphics, video, input devices and audio are supported? The power of the technology comes through in the use of high quality graphics (including overlay on video), digital audio (not just audio on videodisc), the ability to use touch or mouse as well as keyboard, and the videodisc. You may not want these now, but you probably will later. Don't bind yourself to an authoring system which is constrained and not keeping up with new technologies.
7. Which instructional strategies are supported? Many authoring systems constrain you to tutorial and drill and practice formats, yet the simulation and gaming strategies may be needed for a particular application. How does the vendor define a particular strategy? The vendor may define it differently than you do as well.
8. How does the computer-managed instruction (CMI) work? Most systems have CMI, but it may not match your needs. What record keeping and reporting do you require? Will the system require a central file server? Do you have to pay extra for CMI?
9. What response and questioning strategies can be used? You will want to be able to ask some open-ended questions, as well as multiple choice questions. Will the system allow for this? Does it have a built-in spelling corrector? Can several specific hints be given for different wrong answers, to aid the student in reasoning to the correct answer?
10. How flexible is the interface for different screen designs? Can you have different types of menus: located at different screen locations? of different sizes? with icons/graphics?
11. Does the system support team development? Can one person be working on graphics while another working on audio and text for the same lesson at the same time?
12. How easy is it to change and update a lesson? Maintenance costs can be high and maintenance is always necessary. What features does the system have to accommodate maintenance?
13. Can it talk to programming languages? If an authoring system has an efficient way to jump out to code written in a programming language, then it provides much greater flexibility than a system which does not have this capability. This is desirable whether you need this flexibility now or not.
14. Can it import courseware written using other systems? You may need to author or write scripts using other software systems, and if you can import those files into the authoring system directly, you can save a great deal of time.



one of the final three systems/vendors on the basis of the degree of cooperation they gave us on the phone, their reputation, and the apparent expansibility of the system.

## RESULTS

Some of our evaluation criteria were readily answerable either through the demonstration packages or in the information sheets (numbers 1,2,6,8,16). Compatibilities and system features were fairly easily determined. In fact, the authoring systems we were evaluating were remarkably similar in their features with little apparent differences to use for discrimination purposes.

In the area of cost, it became apparent that the advertised cost was very negotiable depending upon the size and nature of the contract, as well as the business of the contracting institution (eg, education or government). For example, if an institution were to purchase several authoring workstations and many student presentation systems, the price would be reduced significantly. One could only obtain this type of information on the phone through probing, and therefore many potential customers would not obtain this information. Another cost issue resulting in confusion was royalties. In our case, we did not want to have to pay royalties on internally developed and distributed courseware, but rather have the freedom of distribution. Several vendors seemed ill-prepared to cope with this requirement in terms of cost estimate, even though it seems to be a commonly held and desirable requirement in the community.

We were unable to make any judgment on several key criteria because of inadequate information. These criteria are listed and described below.

### Ease-of-use

Only two of the demonstration packages included enough of the program so that a limited course could be built, giving us a true "hands-on" experience with the authoring system. Without actually trying to use the authoring system, it was impossible to evaluate ease-of-use. Similarly, without an example of documentation, we were unable to

evaluate it or how much the program depended upon it. We discussed this drawback of the demonstration discs with several vendors, and they agreed with our criticism. The vendor claimed that most people evaluating the systems do not request evaluating ease-of-use and ease-of-learning (even though these are very relative) and that they would not be producing demo discs to make this possible. The only alternative they suggested was trying to use the complete authoring system package (some offered to send one on a trial basis), yet such use requires a larger time investment than we (and most potential customers) have for evaluation.

### External interface

We were unable to determine the extent the authoring system was capable of interfacing with other software and programming languages. Although all of the systems we were evaluating claimed this capability, it became apparent through review of the literature that some systems were more capable than others. In one case, for example, jumping out to some external code was quite easy, but quickly jumping back into the authoring system courseware was not possible.

### Computer managed instruction (CMI)

Only three of the systems evaluated included sufficient information on the CMI capabilities. The others informed you that they did have the capability of testing and record keeping, but did not provide information on the nature of the CMI interface, the number and types of records kept routinely, the method of access for students versus instructors, the learning prescription feature availability, or the types of testing possible.

### Question format

Only two of the systems evaluated discussed the subject of possible question formats adequately and, then, only on the demonstration disc. Most of the systems mentioned multiple choice and fill-in as questioning modes, and some included matching and true/false. For most of the systems, we could not determine if short answer or long answer

questions were possible. In addition, there are a variety of ways of designing questioning screens, and yet we had no way of knowing from the demonstration discs which types of interfaces were available for which systems and how easy they were to generate.

### Customized feedback

We were able to determine the extent of customized feedback and answer judging only from the two demonstration discs that allowed us to actually build a program. These discs represented a wide range in feedback treatments possible, thus the complexity of the criterion was evident. We were interested in information such as how long the feedback could be, how many customized responses could be programmed per course, and how the answer judging was handled. This parameter was important to us, because we wanted to aid our students in reasoning to the correct answer for a problem instead of just being told "try again" or being told the right answer after only two incorrect answers. The majority of systems we evaluated did not fully discuss the subject, thus our evaluation was incomplete for this criterion.

### Instructional strategies

Instructional strategy, or the method used to aid the student in achieving the objectives, is critical to a good match between the subject matter and the student needs. For example, a drill and practice strategy is adequate for learning multiplication tables but is very inadequate in assessing a student's ability to operate a particular control panel. Since most installations need to provide courseware on a variety of applications requiring a variety of instructional strategies, flexibility and robustness here was necessary. None of the systems we evaluated adequately treated the instructional strategies issue. It was also not possible to deduce this information based on the data provided in the information and demonstration packages.

### Compilation

One of the systems indicated that the resultant courseware was an uncom-

piled program written in a programming language. The other seven products we evaluated did not indicate if the courseware generated by the authoring language was compiled or co-compiled.

#### DISCUSSION AND CONCLUSIONS

The information provided by the vendors on their authoring systems was not adequate to make a thorough evaluation of the products. We found that most of the information provided by the vendors was sales- and feature-oriented rather than oriented toward the performance of the system and the modes of instruction it supports. Most of the demonstration discs we looked at were computer-based sales pitches that demonstrated samples of courses and graphics created by the authoring system. They did not indicate how the program was achieved and the amount of time one could expect to spend generating a similar course, nor did they indicate the types of instructional designs that could be generated beyond a linear set of questions and answers.

When questioned by us, the vendors agreed to the limitations of the information packages and demonstration discs. It seems, however, that we as customers are not demanding that they produce alternatives. Potential customers of the authoring systems need to be more thorough in their investigations and evaluations and more demanding of the vendors' materials. Similarly, vendors need to lend greater assistance to their prospective customers by providing the necessary information so that users can make an intelligent selection. Based on our experience, we recommend that potential customers generate a list of criteria prior to evaluation and be firm with the vendors in obtaining answers. We recommend that vendors include more detailed technical information in their sales and demonstration packages. The specifics of the additional information needed is described below.

1. Our foremost recommendation is for the vendors to allow the potential customer to actually use the authoring system in the demonstra-

tion disc. Proprietary protection can be achieved by limiting the number of screens that can be built or by not enabling a "save file" routine. It is virtually impossible to get an indication of system responses, capabilities, performance, and ease-of-use without actually using the authoring system itself. This type of demonstration disc is widely available in the "expert system shell" business, so vendors of expert systems can be contacted for details of logistics and design. The expert system shell demo discs do enable you to create and save your own mini-expert system knowledge base, and yet they are constraining enough so that you could not actually produce a fully operational expert system with them. Through the use of three of these discs we were able to make a very intelligent evaluation of the different interfaces relative to our needs.

2. Provide examples of different instructional strategies in both the sales literature and on the demonstration discs that can be achieved with the authoring system. Define what you mean by "tutorial" and "simulation", as these terms have a variety of meanings in the field. Show the types and varieties of questioning formats that are allowed within the constraints of the system.
3. Offer demonstration discs to potential customers, if only on a "borrowed" basis. Potential customers should not have to pay to obtain essential information to adequately evaluate a vendor's product.
4. Provide more technical information on the performance of the authoring system.
5. Include more information on the system's ability to interface with other software and programming languages. Also, discuss the capabilities to build library files of graphics and subroutines.
6. Include more information on the CMI package and its capabilities and applications. Give examples of the types of records possible and

identify how scorekeeping is accomplished.

7. Discuss if and how feedback can be customized to fit the needs of the user. Include the constraints on customizing feedback as well as capabilities.

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## FROM CAI TO ICAI: TWO EXAMPLES ALONG THE WAY

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There is a continuum of computer-based exercises,<sup>14</sup> each with its own advantages and limitations, that ranges from the "conventional" (CAI) to the "intelligent" (ICAI). Application of significant concepts from the field of artificial intelligence (AI) to the development of computer-based exercises, even in the absence of expensive AI "tools", can result in innovative teaching programs.

We have already written several CAI exercises in physiology.<sup>6,8,9,10</sup> For the past two years, we have focused on developing two tutoring programs in the domain of the cardiovascular (CV) physiology. Neither program functions at the level of an ICAI exercise; they are not yet "smart tutors". But, we have produced sophisticated programs that are of use with our (JAM and AAR) students, and at the same time, are pointing the way to our eventual development of truly "smart" programs.

### WHAT IS A "SMART TUTOR"

A "smart tutor" (sometimes referred to as an intelligent tutoring system or ITS) is a computer program able to assist a student in mastering some subject matter domain.<sup>4,11</sup> The essence of such a program is that its tutoring — its interaction with the student — is context dependent.<sup>13</sup> This means that the particular action it takes is determined by the current state of the student's knowledge or understanding. To produce such a behavior, a "smart tutor" must be able to:

- 1) model the student's understanding of, or misconceptions about, the domain being studied;<sup>1</sup>
- 2) solve the problems being presented to the student;
- 3) tutor the student, selecting a sequence of actions (tactics) accord-

ing to some strategy that depends on the current state of the student;<sup>13</sup> and

- 4) communicate with the student in a way that reveals what the student understands; in many, but not all, cases this requirement can best be met by the ability to generate a two-way natural language conversation.<sup>13</sup>

As interim steps along the way to the development of our two "smart tutors", we have employed approaches to partially realize some of the above functions that do not require expensive or complex AI resources. Before describing these solutions, however, we need to briefly describe the particular programs we are working on.

### "SMART TUTORS" IN CV PHYSIOLOGY

As teachers of medical physiology, we (JAM and AAR) routinely ask students to learn the behavior of complex, multi-component systems with feedback, and to do this in a way that facilitates their ability to "solve problems". Our programs are intended to assist students in this task.

The first program assists students in solving pathophysiology problems. Here, the student is asked to first determine the pathophysiology giving rise to a constellation of signs and symptoms presented by a patient, and then to explain the origin of these features from the physiological cause they propose.<sup>5</sup> The general educational objective of this exercise is the development of those skills needed to apply a hypothetico-deductive problem-solving approach<sup>2</sup> to a particular class of problems.

A second program coaches students working with a computer simulation of

the system that maintains a constant blood pressure. Here, the student predicts the changes that will occur in seven parameters following selected experimental procedures (the tutor is based on a teaching program called CIRCSIM<sup>10</sup>). The general educational objective of this exercise is the development of the ability to analyze the behavior of complex systems possessing feedback.<sup>6,8</sup>

### INCORPORATING "INTELLIGENCE" INTO CAI EXERCISES

While the present versions of these two programs are not as "smart" as they will eventually be, they effectively perform some of the functions necessary for an ITS *without using sophisticated AI techniques*.

#### Student - computer interface

Effective tutoring will require that the student be able to communicate with the program using natural language. However, the development of a robust natural language capability, one able to understand ill-formed inputs generated by the student, and produce complex, context-dependent outputs, is a major undertaking which we have only started. In the absence of these capabilities we have developed two alternative ways for students to provide inputs that contain enough information to drive a tutorial interaction.

In the pathophysiology tutor<sup>3</sup> the student must:

- 1) identify patient signs and symptoms,
- 2) generate hypotheses,
- 3) request data,
- 4) validate hypotheses, and
- 5) provide explanations.

A menu-driven approach provides these



functions at this time.

To identify pertinent patient findings, the student uses a cursor to highlight text in the description of the patient; the selected text is then saved to a file for later use by the student and by the tutor.

To generate an hypothesis to account for the findings he has selected, the student "pages" through a set of structured menus until he finds an hypothesis that he identifies as the source of the patient's problem. Data about the patient is requested by the student in a similar way. With the set of possible hypotheses or data is clearly finite, it is sufficiently large that it places no significant constraint on the students' selection — it does not suggest "answers" that the student might not have arrived at if he was working independently.

In our simulation-based program, the student makes predictions about the direction of change of specified parameters using a predictions table in which the student can move a cursor and toggle the desired entry into each cell.

This set of inputs provides the tutor with an assemblage of data from which it can deduce a cognitive model of the student and then decide on the appropriate response to generate. There is clearly more information available here than in a conventional CAI situation where a student answers a single question whose correctness is assessed and used to select feedback.

#### Expert problem-solvers

A "smart tutor" must itself "understand" the subject matter so that it can solve the problems presented to the students and interpret students' responses.

For example, in a simulation-based tutor, the simulation itself is an expert problem-solver in that it calculates the behavior of the system for each input.<sup>7</sup> Thus, whether the student is asked to make qualitative predictions about the system, or to calculate the quantitative responses he expects, the correct answer is internally available.

Alternatively, in many domains it is possible to develop rule-based qualitative models with which to generate predictions and explanations about system behavior.<sup>12</sup> This is the approach that

we have taken with the preliminary version of our CV simulation tutor. An algorithm has been written with which the qualitative behavior of the system to any input can be determined. Such an approach enables one to model complex systems where an analytical model may be too difficult to generate, or where sufficient data is not available upon which to base such a model.

Where the nature of the exercise does not permit a model-based approach, one may use a stored data-base of solutions to the problems to be presented. This is the approach we have taken with the pathophysiology tutor where we are dealing with a necessarily small number of problems (both because there are a limited number of patient problems that are relevant to our educational objectives, and because the time required to solve each problem is sufficiently long that the student can only do one or two problems at a sitting). Solutions to each of the three problems in our exercise are stored using an approach that resembles a data-base program.

#### Student-modelling

Conventional CAI generally utilizes conditional branching from individual answers to determine the particular feedback to be provided or the next step in the exercise to be pursued. While this "individualizes" the students' experience in some limited sense, it provides a fixed response to any wrong answer *regardless of the reason that the student answered incorrectly*. However, the same incorrect response can result from great differences in students' general background, available knowledge about the domain, or skill at problem-solving.

A "smart tutor" attempts to provide a deeper level of assistance to the student by determining the specific basis for the wrong answer. This process of modeling the "cognitive state" of the student is amongst the most difficult confronting the development of ICAI programs.

Our approach to student modeling in our simulation tutor, one with general applicability, is based on a concept map of the relevant domain and an experi-

mentally derived catalog of student "bugs". Predictions for a minimum of seven parameters are available as each experiment is carried out. Particular patterns of errors then allow the modeler to pick from the list of "bugs" the most probable source of the students' incorrect responses. For example, errors in predicting the changes that will occur in CC, HR, and TPR and the failure to have entered these predictions sequentially suggests that the student does not understand the common neural control of these parameters. With such information available to the "tutor", the most helpful intervention can then be selected.

#### SUMMARY

The development of sophisticated tutoring programs, programs able to tailor their interactions with the student to meet the students' specific needs, will go a long way towards overcoming many of the limitations of current CAI. There is much to be learned from the fields of artificial intelligence and cognitive science that is relevant to this effort, even if the technical resources to implement these ideas are unavailable. It is ultimately the ideas that are most relevant, not the use of any particular AI programming language or AI workstation.

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#### AIMS AND SCOPE

The goal of *Computers in Life Science Education* is to provide a means of communication among life science educators who anticipate or are currently employing the computer as an educational tool. The range of content includes, but is not limited to, articles focusing on computer applications and their underlying philosophy, reports on faculty/student experiences with computers in teaching environments, and software/hardware reviews in both basic science and clinical education settings.

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