

BMJ Open Development of guidance on the timeliness in response to acute kidney injury warning stage test results for adults in primary care: an appropriateness ratings evaluation

Tom Blakeman,¹ Kathryn Griffith,² Dan Lasserson,³ Berenice Lopez,⁴ Jung Y Tsang,¹ Stephen Campbell,⁵ Charles Tomson⁶

To cite: Blakeman T, Griffith K, Lasserson D, *et al.* Development of guidance on the timeliness in response to acute kidney injury warning stage test results for adults in primary care: an appropriateness ratings evaluation. *BMJ Open* 2016;**6**:e012865. doi:10.1136/bmjopen-2016-012865

► Prepublication history and additional material is available. To view please visit the journal (<http://dx.doi.org/10.1136/bmjopen-2016-012865>).

Received 8 June 2016

Revised 19 August 2016

Accepted 12 September 2016



CrossMark

For numbered affiliations see end of article.

Correspondence to

Dr Tom Blakeman;
tom.blakeman@manchester.ac.uk

ABSTRACT

Objectives: Tackling the harm associated with acute kidney injury (AKI) is a global priority. In England, a national computerised AKI algorithm is being introduced across the National Health Service (NHS) to drive this change. The study sought to maximise its clinical utility and minimise the potential for burden on clinicians and patients in primary care.

Design: An appropriateness ratings evaluation using the RAND/UCLA Appropriateness Method.

Setting: Clinical scenarios were developed to test the timeliness in (1) communication of AKI warning stage test results from clinical pathology services to primary care, and (2) primary care clinician response to an AKI warning stage test result.

Participants: A 10-person panel was purposively sampled with representation from clinical biochemistry, acute and emergency medicine and general practice. General practitioners (GPs) represented typical practice in relation to rural and urban practice, out of hours care, GP commissioning and those interested in reducing the impact of medicalisation and 'overdiagnosis'.

Results: There was agreement that delivery of AKI warning stage test results through interruptive methods of communication (ie, telephone) from laboratories to primary care was the appropriate next step for patients with an AKI warning stage 3 test result. In the context of acute illness, waiting up to 72 hours to respond to an AKI warning stage test result was deemed an inappropriate action in 62 out of the 65 (94.5%) cases. There was agreement that a clinician response was required within 6 hours, or less, in 39 out of 40 (97.5%) clinical cases relating AKI warning stage test results in the presence of moderate hyperkalaemia.

Conclusions: The study has informed national guidance to support a timely and calibrated response to AKI warning stage test results for adults in primary care. Further research is needed to support effective implementation, with a view to examine the effect on health outcomes and costs.

Strengths and limitations of this study

- The study used systematic RAND/UCLA consensus methods to optimise clinician response to acute kidney injury (AKI) warning stage test results for adults in primary care.
- Efforts to maximise the utility of AKI alerts and minimise the potential for burden included engagement with general practitioners who represented concerns about medicalisation and overdiagnosis.
- Although a range of clinically relevant scenarios were considered, in terms of burden on RAND/UCLA panellists, it was not feasible to test all major comorbidity groups (eg, patients with type 2 diabetes).
- The study sought clarity on 'necessary' actions required under ideal conditions and further studies are needed to understand the implementation of AKI alerts and guidance in routine clinical practice.

INTRODUCTION

Maintenance of kidney health is central to addressing two global health priorities: the prevention of cardiovascular disease and support for people vulnerable to complications during episodes of acute illness.¹⁻⁴ The international Oby25 initiative (<http://www.Oby25.org/>) aims to prevent avoidable harm associated with acute kidney injury (AKI).² In a similar vein, the National Health Service (NHS) Five Year Forward View emphasises tackling AKI as a priority to improve patient safety and health outcomes across the NHS.⁵

AKI is a clinical syndrome characterised by a sudden reduction in kidney function.⁴ The diagnosis of AKI and its staging is based on acute changes in serum creatinine and/or a

reduction in urine output.⁴ There are many causes for AKI though it is most commonly associated with episodes of acute illness on the background of increased risk.^{4 6–8} AKI is associated with frailty and is more common in older people living with complex comorbidities.^{4 6 8–12} These include people with pre-existing chronic kidney disease (CKD), diabetes and heart failure as well as those with underlying cognitive decline, who may be reliant on a carer for adequate fluid intake during episodes of acute illness.⁶

AKI is associated with ~1 in 5 unplanned hospital admissions, with almost two-thirds of these patients with AKI having developed it in the community.¹³ It is associated with increased length of hospital admission and need for renal replacement therapy.⁴ AKI is also associated with poorer health outcomes including increased risk of progression of CKD and increased mortality both in the short and long terms.⁴ Care of people with AKI costs around 1% of the NHS budget, reflecting the fact that AKI is a strong marker for severity of acute illness.¹⁴

In a major NHS England patient safety initiative, a computerised AKI algorithm has been established, based on the Kidney Disease Improving Global Outcomes (KDIGO) classification system for AKI, to automatically detect biochemical changes in kidney function and identify potential cases of AKI.^{4 15 16} The test result is named an 'AKI warning stage' and is categorised according to severity (ie, AKI stage 1, 2 or 3).¹⁵ A level 3 NHS England Patient Safety Notice made it mandatory that all NHS Acute Trusts and Foundation Trusts implement the use of the algorithm within laboratory information systems, providing warning stage results to users in secondary care, by March 2015. This is combined with a mandatory submission of a data set to a national registry allowing audit of the frequency of alerts.¹⁶ The next phase of AKI alerting involves direct communication of AKI warning stage test results to primary care in England, from April 2016.^{15 16}

National clinical guidelines for AKI recommend 'identifying AKI in patients with acute illness' who are deemed at risk of AKI.⁶ At present, while a significant proportion of episodes of illness complicated by AKI start in the community, AKI tends to be identified on hospital admission.¹⁷ As a potential marker of the 'sick patient', this new test could support earlier detection and prompt management of acute illness in primary care.^{15 18} However, trade-offs exist between their clinical utility and the potential for burden on clinical care.^{19 20}

First, kidney function tests are taken in primary care for many reasons both as part of routine disease management monitoring as well as assessment of urgent care. Current data suggest that only a small proportion of all primary care serum creatinine requests generate an AKI alert.²¹ In primary care, a decision on whether or not to check kidney function needs to be tailored to the individual circumstance.⁶ Consideration needs to be given as to whether this will support clinical management. For example, blood tests are not necessary when

immediate admission is required (eg, there is evidence of sepsis).^{22 23} At the other end of the spectrum, checking kidney function tests may also not be necessary for patients presenting with a minor self-limiting acute illness, such as a single episode of diarrhoea or vomiting.

Second, 'false-positive' test results may occur. For example, a change in serum creatinine may be due to progression of CKD rather than represent an acute deterioration in kidney function.^{15 19 20} Third, there is also need to address the risks associated with a test result being communicated from laboratories to out of hours services, requiring a clinician who may have limited access to clinical information to act on the result. Fourth, there is evidence to suggest that the introduction of 'e-alerts' in isolation are unlikely to improve health outcomes.^{24 25} In particular, 'alert fatigue' is a recognised problem limiting implementation and effectiveness.^{26 27} If implemented well, the release of AKI warning stage test results to primary care has the potential to generate major improvements in outcomes. However, experience of other 'top-down' initiatives intended to improve kidney outcomes in primary care has been mixed.²⁸

In order to maximise their clinical utility and minimise burden on health professionals and patients, guidance is required to ensure a timely response and for test results to be considered with an understanding of the clinical context in which the blood test was taken.

METHODS

Study design

The RAND/UCLA Appropriateness Method (RAM) was used to help develop guidance on the response to AKI warning stage test results in primary care.²⁹ RAM is an established consensus method to generate appropriateness ratings.²⁹ It is a systematic approach to address specific dilemmas in clinical practice in which clinical decisions are required but where 'robust scientific evidence' about the benefits is lacking.^{29–31} The methodology combines 'best available scientific evidence with the collective judgement of experts' in order to provide a statement regarding the appropriateness of undertaking a particular action 'at the level of patient-specific symptoms, medical history and tests results'.^{29–31} As such, it is a rigorous methodology congruent with a need to place response to AKI warning stage test results in clinical context. Findings from the process are deemed to have face validity and provide a platform for prospective development and evaluation of guidance to support clinical decision-making.³²

Developing clinical scenarios for appropriateness rating

Through NHS England's Think Kidneys Programme, a working group was established to develop clinical scenarios for appropriateness rating.³³ In order to ensure scenarios were informed from a range of perspectives,

the group comprised general practitioners (GPs), a nephrologist, a chemical pathologist with a GP background and a methodologist with expertise in the RAM as well as patient safety in primary care. The group identified national AKI clinical guidelines and invited hospital trusts, clinical commissioning groups and strategic clinical networks throughout the country to send guidance documents in use, or being developed, to support primary care in responding to AKI warning stage test results. These documents were reviewed to identify key elements of care relating to AKI, which through group discussion led to the generation of a list of clinical cases, with a focus on rating the timeliness in response to each case.

Clinical scenarios were categorised according to the following clinical characteristics: level of AKI warning stage test result (AKI warning stage 1, 2 and 3); medical history (episode of acute illness, history of chronic heart failure, CKD stage 4 or 5, or history of renal transplant, recent change in dose of a diuretic, ACE inhibitor or angiotensin receptor blocker); and evidence of complicating factors (poor fluid intake/urine output, hyperkalaemia, both mild ($K^+ 5.5\text{--}5.9\text{ mmol/L}$) or moderate ($K^+ 6.0\text{--}6.4\text{ mmol/L}$), evidence of risk of urinary tract obstruction or intrinsic renal disease). Scenarios were not tested for severe hyperkalaemia as UK Renal Association Clinical Practice Guidelines recommend emergency assessment and treatment of severe hyperkalaemia ($K^+ \geq 6.5\text{ mmol/L}$).³⁴ In order to reduce research burden on panel members and keep the process salient, a decision was made to restrict the number of ratings for AKI warning stage 3 test results in which timeliness in management was deemed less controversial.⁴ Also, the clinical scenarios considered during the consensus process were not specific to either in hours or out of hours care. Other categories considered in scenarios presented were: age (adult, or children and young person); and whether a patient was receiving end of life care. These clinical characteristics were then used to develop 163 clinical cases, with four responses (next steps) requiring rating for each clinical case (ie, $163 \times 4 = 652$ clinical scenarios).

Each clinical case represented a separate testing indication consisting of variable combinations of the clinical characteristics. Clinical cases were independently linked to two main areas of decision-making:

- 1 Based on knowledge of an AKI warning stage test result and potassium level, clinical pathology service/laboratory staff need to make one of four decisions:
 - Send an AKI warning stage result via the data transfer service without comment (non-interruptive communication).
 - Send an AKI warning stage result via the data transfer service with comment (non-interruptive communication).
 - Send an AKI warning stage result to an NHS email address that is known to be monitored regularly during working hours (non-interruptive communication).

- Send an AKI warning stage result by telephone call to GP/practice/out of hours service provider (interruptive communication).

- 2 Based on the knowledge of the presenting AKI warning stage test result, clinical history including complications, GPs (responsible primary care team, ie, either in hours or out of hours service provider) will need to make one of four decisions:

- Seek immediate admission.
- Respond to the AKI warning stage result within 6 hours.
- Respond to the AKI warning stage result within 24 hours.
- Respond to the AKI warning stage result within 72 hours.

An established rating scale was used to rate each potential decision for each clinical scenario on a nine-point integer scale, with 1 indicating an 'extremely inappropriate next step' and 9 indicating an 'extremely appropriate next step'.^{29 31}

- Scores 1–3: inappropriate next step (ie, no benefit, possible harms).
- Scores 4–6: uncertainty about next step (ie, when harms and benefits are judged as approximately equal, or when the best available evidence does not support a judgement either way).
- Scores 7–9: Appropriate next step (ie, benefits judged to outweigh harms).

Panel membership and rating

Panel members were purposively sampled and recruited to ensure that a range of perspectives were considered and ensure that the appropriateness ratings were valid for routine clinical general practice. A 10-person panel was identified with representation from clinical biochemistry ($n=2$), acute and emergency medicine ($n=2$) and general practice (GP principals, $n=5$; salaried GP, $n=1$). Individuals with an expressed interest and expertise in AKI through membership of the NHS England Think Kidneys Programme were invited as well as GPs who represented typical practice in relation to out of hours care, rural and urban practice, GP commissioning and those with an interest in reducing medicalisation and 'overdiagnosis'.^{33 35} The panel was co-chaired by one researcher (SC), who is expert in the RAM, and a nephrologist (CT) who is expert in AKI, neither of whom contributed to the RAND/UCLA decision-making scores.

The RAND process entailed two rounds of rating:

Round 1 was conducted by email with the panel members rating the clinical scenarios individually at home, with no discussion between panellists. In order to help rate the appropriateness of each decision for every scenario, panel members were sent an 'instructions' document and a 'context' document that provided an overview of the literature with a focus on the rationale for the clinical characteristics chosen. The evaluation aimed to gain clarity on 'necessary' processes of care

under ideal conditions. When scoring each scenario, panellists were instructed to assume that they had available all the clinical information presented.

Round 2 entailed a 1-day face-to-face meeting held in September 2015. Panel members were presented with their own data from round 1 as well as data outlining the distribution of all panel members' ratings. This summary was used to stimulate discussion with a focus on areas of disagreement. Panel members then rated each scenario on their own individual-blinded rating sheets. Panel members were not required to reach consensus.

Data entry and analysis

RAMs were used to conduct analysis.²⁹ Data from round 2 were collated and analysed to determine the level of agreement within the panel for a proposed action for each clinical scenario. Agreement was defined by 8 out of the 10 (80%) panel members rating the same three-point region on the nine-point integer scale (ie, 1–3, 4–6, 7–9). Disagreement was defined to exist where $\geq 30\%$ of panel members rated a scenario in the 1–3 range and where $\geq 30\%$ rated the same scenario in the 7–9 range on the nine-point integer scale. A proposed action was then categorised as an 'appropriate' next step if a scenario rated 7–9 without disagreement and a rating of 1–3 without disagreement was deemed to be an 'inappropriate' next step. Ratings of clinical scenarios without consensus (either 'agreement' or 'disagreement') were considered as equivocal.

RESULTS

The results are presented in tables 1–4 for round 2 ratings. Table 1 summarises the ratings across the 652 clinical scenarios, with 415 (63.7%) achieving agreement; 32 (4.9%) rated with disagreement and 205 (31.4%) with an equivocal rating. A proposed action was deemed the appropriate next step in 137 (21.0%) out of the 652 scenarios. Two hundred and ten (32.2%) of the proposed actions were deemed inappropriate, and in 68

Table 1 Summary of round 2 appropriateness ratings

Total number of scenarios	652
Agreement (8 out of 10 (80%) of panel members rating in the same 3-point region)	415 (63.7%)
Appropriate (80% of panel members rating 7–9)	137 (21.0%)
Inappropriate (80% of panel members rating 1–3)	210 (32.2%)
Agreement with uncertain benefit (80% of panel members rating the same consecutive 3-point region—but NOT 1–3 or 7–9)	68 (10.4%)
Disagreement ($\geq 30\%$ of scores in 1–3 AND $\geq 30\%$ in 7–9 for same scenario)	32 (4.9%)
Equivocal	205 (31.4%)

Table 2 Communication of AKI warning stage test results for adults by clinical pathology services to primary care

	AKI warning stage 1			AKI warning stage 2			AKI warning stage 3		
	Response	Result	Median	Response	Result	Median	Response	Result	Median
Potassium not raised	Non-IC and no C	E	5	Non-IC and no C	A	1	Non-IC and no C	A	1
	Non-IC and C	U	7	Non-IC and C	U	5.5	Non-IC and C	A	2
	Non-IC and M	E	5	Non-IC and M	E	6	Non-IC and M	A	2
	IC (Telephone)	E	5	IC (telephone)	E	7	IC (telephone)	A	9
Mild hyperkalaemia (5.5–5.9 mmol/L)	Non-IC and no C	A	3	Non-IC and no C	A	1	Non-IC and no C	A	1
	Non-IC and C	D	7	Non-IC and C	E	3	Non-IC and C	A	1
	Non-IC and M	E	5	Non-IC and M	E	7	Non-IC and M	A	1.5
	IC (telephone)	E	5	IC (telephone)	A	8	IC (telephone)	A	9
Moderate hyperkalaemia (6.0–6.4 mmol/L)	Non-IC and no C	A	1	Non-IC and no C	A	1	Non-IC and no C	A	1
	Non-IC and C	E	3.5	Non-IC and C	A	2	Non-IC and C	A	1
	Non-IC and M	E	5.5	Non-IC and M	E	6	Non-IC and M	A	1
	IC (telephone)	A	9	IC (telephone)	A	9	IC (telephone)	A	9

Colour key: red: agreement for appropriate communication as an appropriate action; no colour: no agreement on an appropriate action.

A, agreement; AKI, acute kidney injury; C, comment; D, disagreement; E, equivocal; IC, interruptive communication; M, email; U, agreement with uncertain benefit.

Table 3 Timeliness in response to AKI warning stage test results for adults in primary care—in the context of acute illness

AKI in the context of acute illness	AKI warning stage 1				AKI warning stage 2				AKI warning stage 3			
	No comorbidities	Chronic heart failure	CKD 4/5 or renal transplant	CKD 4/5 and chronic heart failure	No comorbidities	Chronic heart failure	CKD 4/5 or renal transplant	CKD 4/5 and chronic heart failure	No comorbidities	Chronic heart failure	CKD 4/5 or renal transplant	CKD 4/5 and chronic heart failure
Potassium not raised	SIA=A(2) <6=D(5.5) <24=E(7) <72=E(5)	SIA=E(3) <6=D(6) <24=A(7) <72=E(2.5)	SIA=(4.5) <6=U(7) <24=A(7.5) <72=A(1)	SIA=(4.5) <6=U(7) <24=E(5.5) <72=A(1)	SIA=E(4.5) <6=A(7) <24=E(7) <72=A(1)	SIA=E(5) <6=U(7.5) <24=U(6) <72=A(1)	SIA=E(5) <6=A(7) <24=D(5.5) <72=A(1)	SIA=E(7) <6=A(8) <24=E(4) <72=A(1)	SIA=A(8) <6=U(7) <24=A(1) <72=A(1)	Not tested	Not tested	Not tested
Mild hyperkalaemia (5.5–5.9 mmol/L)	SIA=E(3.5) <6=E(7) <24=A(7) <72=E(3)	SIA=E(4.5) <6=U(7) <24=U(6.5) <72=A(1.5)	SIA=E(5) <6=A(7.5) <24=E(6) <72=A(1)	SIA=U(5.5) <6=A(7) <24=E(4) <72=A(1)	SIA=E(5) <6=A(8) <24=U(7) <72=A(1)	SIA=U(6) <6=A(7.5) <24=E(5.5) <72=A(1)	SIA=U(6) <6=A(8) <24=E(3) <72=A(1)	SIA=A(7) <6=A(7) <24=U(3) <72=A(1)	Not tested	Not tested	Not tested	Not tested
Moderate hyperkalaemia (6.0–6.4 mmol/L)	SIA=E(6) <6=A(8) <24=E(5) <72=A(1)	SIA=D(5.5) <6=A(7) <24=E(5) <72=A(1)	SIA=E(7) <6=A(7.5) <24=E(2.5) <72=A(1)	SIA=A(8) <6=U(6.5) <24=A(2) <72=A(1)	SIA=A(7.5) <6=E(8) <24=A(3) <72=A(1)	SIA=A(7) <6=A(8) <24=U(3) <72=A(1)	SIA=A(7.5) <6=U(7) <24=A(3) <72=A(1)	SIA=A(8) <6=U(7) <24=A(2) <72=A(1)	Not tested	Not tested	Not tested	Not tested
Risk of urinary tract obstruction or intrinsic renal disease	SIA=E(7) <6=A(7.5) <24=E(5) <72=A(1)	SIA=E(6.5) <6=E(6.5) <24=D(7) <72=A(1)	SIA=E(7.5) <6=E(7) <24=E(6) <72=A(1)	SIA=A(8) <6=E(7) <24=E(2) <72=A(1)	SIA=A(7) <6=E(7) <24=E(4.5) <72=A(1)	SIA=E(7) <6=U(7) <24=E(6) <72=A(1)	SIA=U(7) <6=E(7) <24=A(3) <72=A(1)	SIA=A(7.5) <6=E(7) <24=A(2) <72=A(1)	Not tested	Not tested	Not tested	Not tested
Poor oral intake/urine output	SIA=E(4) <6=A(7) <24=E(7) <72=A(1.5)	SIA=E(4) <6=U(7) <24=E(7) <72=A(2)	SIA=E(4.5) <6=A(7) <24=U(7) <72=A(1.5)	SIA=E(6) <6=U(7) <24=E(5) <72=A(1)	SIA=U(7) <6=A(7) <24=E(4) <72=A(1)	SIA=E(6) <6=A(7) <24=E(5.5) <72=A(1)	SIA=U(6) <6=A(8) <24=U(3) <72=A(1)	SIA=U(7) <6=A(8) <24=E(3.5) <72=A(1)	Not tested	Not tested	Not tested	Not tested
Mild hyperkalaemia +poor oral intake/urine output	SIA=U(5) <6=A(7) <24=D(6) <72=A(1)	SIA=U(5) <6=E(7) <24=E(7) <72=A(1)	SIA=U(6) <6=A(7) <24=D(5) <72=A(1)	SIA=U(6.5) <6=U(7) <24=A(2) <72=A(1)	SIA=U(7) <6=A(8) <24=E(4.5) <72=A(1)	SIA=E(6.5) <6=A(8) <24=U(3.5) <72=A(1)	SIA=U(7) <6=A(8) <24=E(3) <72=A(1)	SIA=A(7) <6=A(8) <24=A(2) <72=A(1)	Not tested	Not tested	Not tested	Not tested
Moderate hyperkalaemia+poor oral intake/urine output	SIA=A(7) <6=A(8) <24=A(2.5) <72=A(1)	SIA=E(7) <6=E(7) <24=E(3) <72=A(1)	SIA=A(8) <6=E(7) <24=E(3) <72=A(1)	SIA=A(9) <6=D(5.5) <24=A(2) <72=A(1)	SIA=A(9) <6=U(7) <24=A(2.5) <72=A(1)	SIA=A(8) <6=U(7) <24=A(2.5) <72=A(1)	SIA=A(8.5) <6=E(6.5) <24=A(2) <72=A(1)	SIA=A(9) <6=E(6.5) <24=A(1.5) <72=A(1)	Not tested	Not tested	Not tested	Not tested
Risk urinary tract obstruction or intrinsic renal disease+poor oral intake/urine output	SIA=E(7.5) <6=E(7.5) <24=E(3) <72=A(1)	SIA=E(6.5) <6=E(5.5) <24=D(3.5) <72=A(1)	SIA=E(7.5) <6=E(6.5) <24=D(1.5) <72=A(1)	SIA=A(8) <6=A(7) <24=A(2) <72=A(1)	SIA=E(8) <6=E(6.5) <24=E(3) <72=A(1)	SIA=A(7.5) <6=E(6.5) <24=E(3) <72=A(1)	SIA=A(8) <6=E(7) <24=A(2) <72=A(1)	SIA=A(8) <6=E(7) <24=A(2) <72=A(1)	Not tested	Not tested	Not tested	Not tested

Colour key: red: agreement that seeking immediate admission is an appropriate action; orange: agreement that responding in <6 hours is an appropriate action; brown: agreement that responding in <24 hours is an appropriate action; no colour: no agreement on an appropriate action. Numerical value in brackets: median rating. A, agreement; AKI, acute kidney injury; D, disagreement; E, equivocal; SIA, seek immediate admission; U, agreement with uncertain benefit; CKD 4/5, chronic kidney disease stage 4 or 5.

Table 4 Timeliness in response to AKI warning stage test results for adults in primary care—no acute illness

AKI with no history of acute illness	AKI warning stage 1				AKI warning stage 2				AKI warning stage 3			
	No comorbidities	Chronic heart failure	CKD 4/5 or renal transplant	CKD 4/5 and chronic heart failure	No comorbidities	Chronic heart failure	CKD 4/5 or renal transplant	CKD 4/5 and chronic heart failure	No comorbidities	Chronic heart failure	CKD 4/5 or renal transplant	CKD 4/5 and chronic heart failure
Potassium not raised	SIA=A(1) <6=A(1) <24=E(6.5) <72=E(7.5)	SIA=A(1) <6=E(4) <24=A(7) <72=E(7)	SIA=A(1) <6=E(4.5) <24=A(8) <72=E(4.5)	SIA=A(1) <6=E(5) <24=A(8) <72=E(5)	SIA=E(1.5) <6=E(6.5) <24=E(6) <72=E(1.5)	SIA=E(2.5) <6=E(7) <24=U(7) <72=A(1)	SIA=E(5.5) <6=E(7) <24=D(6) <72=A(1)	SIA=D(5) <6=U(7) <24=E(5) <72=A(1)	SIA=E(7) <6=A(7.5) <24=E(2.5) <72=A(1)	SIA=U(7) <6=A(8) <24=E(4) <72=A(1)	SIA=A(7) <6=U(7) <24=A(1.5) <72=A(1)	SIA=A(8) <6=E(6.5) <24=A(1) <72=A(1)
Mild hyperkalaemia (5.5–5.9mmol/L)	SIA=A(1) <6=E(3.5) <24=A(8) <72=E(6)	SIA=A(1) <6=E(5) <24=A(8) <72=D(3)	SIA=A(1.5) <6=E(5) <24=A(8) <72=E(1)	SIA=A(1) <6=D(5.5) <24=A(8) <72=A(1)	SIA=E(2.5) <6=E(6.5) <24=E(7) <72=E(2.5)	SIA=E(2.5) <6=U(7) <24=E(7) <72=A(1)	SIA=D(5.5) <6=U(7) <24=D(5.5) <72=A(1)	SIA=D(5.5) <6=A(7.5) <24=D(5.5) <72=A(1)	SIA=U(7) <6=A(7.5) <24=E(1.5) <72=A(1)	SIA=A(7) <6=A(7.5) <24=E(3) <72=A(1)	SIA=A(8) <6=E(7) <24=A(1.5) <72=A(1)	SIA=A(8) <6=E(7.5) <24=A(1) <72=A(1)
Moderate hyperkalaemia (6.0–6.4 mmol/L)	SIA=E(3) <6=A(7) <24=E(7) <72=A(1.5)	SIA=E(3.5) <6=A(7) <24=E(6) <72=A(1)	SIA=E(6) <6=A(8) <24=D(5.5) <72=A(1)	SIA=E(6) <6=A(7.5) <24=D(5.5) <72=A(1)	SIA=E(4) <6=A(8) <24=D(6) <72=A(1)	SIA=E(5.5) <6=A(8) <24=E(6) <72=A(1)	SIA=E(6.5) <6=A(8) <24=E(3.5) <72=A(1)	SIA=A(8) <6=A(8) <24=E(3) <72=A(1)	SIA=A(8) <6=A(7.5) <24=A(1.5) <72=A(1)	SIA=A(8) <6=A(7.5) <24=A(1.5) <72=A(1)	SIA=A(9) <6=E(6) <24=A(1) <72=A(1)	SIA=A(9) <6=D(4.5) <24=A(1) <72=A(1)
Recent increase diuretics/ACEI+potassium not raised	SIA=A(1) <6=A(2) <24=A(8) <72=U(6)	SIA=A(1) <6=E(3.5) <24=U(6.5) <72=D(6)	SIA=A(2) <6=U(5.5) <24=A(8) <72=E(3)	SIA=A(2.5) <6=E(5) <24=A(8) <72=A(2.5)	SIA=E(3) <6=E(7) <24=E(7) <72=E(1)	SIA=E(3) <6=U(7) <24=U(7.5) <72=E(1)	SIA=U(3) <6=U(7) <24=E(6.5) <72=A(1)	SIA=U(4) <6=A(7) <24=D(5.5) <72=A(1)	SIA=E(7) <6=A(7.5) <24=E(2) <72=A(1)	SIA=A(7) <6=A(7) <24=E(3) <72=A(1)	SIA=A(7) <6=E(7) <24=E(1) <72=A(1)	SIA=A(7.5) <6=U(7.5) <24=A(1.5) <72=A(1)
Recent increase diuretics/ACEI+mild hyperkalaemia	SIA=A(1.5) <6=E(4) <24=A(8) <72=E(5.5)	SIA=A(1.5) <6=E(5) <24=A(8) <72=A(2.5)	SIA=A(2.5) <6=E(7) <24=A(7.5) <72=A(2)	SIA=E(3) <6=E(6) <24=A(8) <72=A(2)	SIA=A(3) <6=U(7) <24=E(6.5) <72=A(1.5)	SIA=E(3) <6=U(7) <24=E(6.5) <72=E(1.5)	SIA=U(4) <6=A(7) <24=E(5.5) <72=A(1)	SIA=U(5) <6=A(8) <24=D(4) <72=A(1)	SIA=A(7) <6=A(7.5) <24=E(2) <72=A(1)	SIA=A(7) <6=A(7) <24=E(1.5) <72=A(1)	SIA=A(7) <6=E(7.5) <24=A(1) <72=A(1)	SIA=A(7.5) <6=U(7.5) <24=A(1.5) <72=A(1)
Recent increase diuretics/ACEI+moderate hyperkalaemia	SIA=E(3) <6=A(7) <24=E(7) <72=A(1)	SIA=U(4) <6=A(7.5) <24=D(6.5) <72=A(1)	SIA=U(5.5) <6=A(8) <24=D(6.5) <72=A(1)	SIA=E(6) <6=A(7) <24=D(5) <72=A(1)	SIA=E(3.5) <6=A(8) <24=D(6) <72=A(1)	SIA=E(6.5) <6=A(8) <24=E(6) <72=A(1)	SIA=U(7) <6=A(8) <24=E(3.5) <72=A(1)	SIA=A(7) <6=A(7.5) <24=U(4) <72=A(1)	SIA=A(8) <6=U(7.5) <24=A(1.5) <72=A(1)	SIA=A(8) <6=E(7) <24=A(1) <72=A(1)	SIA=A(8.5) <6=E(7) <24=A(1) <72=A(1)	SIA=A(8) <6=E(7) <24=A(1) <72=A(1)

Colour key: red: agreement that seeking immediate admission is an appropriate action; orange: agreement that responding in <6 hours is an appropriate action; brown: agreement that responding in <24 hours is an appropriate action; no colour: no agreement on an appropriate action. Numerical value in brackets: median rating.

A, agreement; ACEI, ACE inhibitors; AKI, acute kidney injury; CKD 4/5, chronic kidney disease stage 4 or 5; D, disagreement; E, equivocal; SIA, seek immediate admission; U, agreement with uncertain benefit.

(10.4%) of the proposed actions for a scenario, there was agreement of uncertain benefit.

The study was primarily undertaken to support the development of new guidance on the response to AKI warning stage test results for adults in primary care. Tables 2–4 and the Results section focus on presenting data for adults. Supplementary tables present data for a child or young person as well as for adults with chronic heart failure on an end of life care pathway.

Communication of an AKI warning stage test result to primary care

Table 2 presents the data for the timeliness in communication of AKI warning stage test results for adults by clinical pathology services to primary care. There was agreement that delivery of AKI warning stage test results through interruptive methods of communication (ie, telephone result) from laboratories to primary care was the appropriate next step for patients with: an AKI warning stage 3 test result irrespective of potassium level; an AKI warning stage 2 test result in the presence of mild (K^+ 5.5–5.9 mmol/L) or moderate hyperkalaemia (K^+ 6.0–6.4 mmol/L); and an AKI warning stage 1 test result in the presence of moderate hyperkalaemia (K^+ 6.0–6.4 mmol/L).

Timeliness in clinician response to an AKI warning stage test result

Data on the timeliness in clinician response to AKI warning stage test results for adults in primary care are presented in table 3 (in the context of acute illness) and table 4 (no history of acute illness). Agreement was reached on an appropriate next step for the majority of clinical cases presented to panel members: 48 out of the 65 (73.8%) clinical cases relating to AKI test results in the context of acute illness and 58 out of 72 (80.6%) clinical cases for AKI test results with no history of acute illness.

In the context of acute illness, there was a greater level of uncertainty on the appropriate action for clinical cases relating to AKI warning stage 1 test results, particularly in the context of patients with chronic heart failure. In the absence of acute illness, there was a greater level of uncertainty on the appropriate next step for clinical cases relating to AKI warning stage 2 test results.

In the context of acute illness, waiting up to 72 hours to respond to an AKI warning stage test result was deemed an inappropriate action in 62 out of 65 (94.5%) clinical cases tested. In the absence of acute illness, waiting up to 72 hours was deemed inappropriate in 56 out of the 72 (77.8%) clinical cases, with uncertainty for this timeframe tending to relate to AKI warning stage 1 test results.

There was agreement that a clinician response was required within 6 hours, or less, in 39 out of 40 (97.5%) clinical cases relating AKI warning stage test results in the presence of moderate hyperkalaemia (K^+ 6.0–6.4 mmol/L).

These findings included agreement to seek immediate admission if moderate hyperkalaemia was present in the context of AKI warning stage 3 results, AKI warning stage 2 results in the context of acute illness and AKI warning stage 1 results in the context of acute illness associated with poor oral intake/urine output.

There was uncertainty on the appropriate response to AKI warning stage test results generated for patients with chronic heart failure on an end of life care pathway, though waiting up to 72 hours to respond to these test results was deemed an inappropriate next step.

DISCUSSION

Principal findings

The introduction of warning stage test results based on a national AKI algorithm is a major patient safety initiative across the NHS in England.¹⁵ However, its implementation in primary care is at a period in time where clinical workload is deemed to be reaching saturation point.³⁶ The study is an important starting point in terms of navigating the challenges of addressing the harm and costs associated with AKI while avoiding the creation of additional workload for professionals and unnecessary burden on patients.³⁷ Through use of rigorous consensus methods, the findings highlight clinical scenarios where interruptive methods of communication (ie, telephoning a test result) by clinical pathology services to primary care are considered to be appropriate. There was also agreement in the majority of clinical cases in terms of determining the timeliness in response to an AKI warning stage test result by a clinician in primary care. In the main, waiting up to 72 hours was deemed an inappropriate response. The findings from the study have informed the development and recent publication of NHS England Think Kidneys guidance (<https://http://www.thinkkidneys.nhs.uk/aki/resources/primary-care/>).³³

Strengths and weaknesses of this study

The study used systematic RAND/UCLA consensus methods to optimise the development of guidance for clinicians on how to respond to AKI warning stage test results for adults in primary care.²⁹ Investment in ‘overcoming questions about the legitimacy of new guidelines’ is important to ensure successful implementation.³⁸ A key feature of the study was to maximise the utility of AKI alerts and minimise the potential for burden on patients and health professionals. Efforts to achieve this included engaging primary care clinicians who represented concerns about medicalisation and overdiagnosis.³⁵

Multiple scenarios were tested through use of RAMs.²⁹ Key features of these scenarios were identified through identification and comparison of existing relevant guidance documents. Cross-referencing and integrating different guidelines may address challenges of clinical decision-making in primary care, particularly for patients with multimorbidity.³⁹ For example, rather than treating

results in isolation, the methodological approach enabled AKI warning stage test results to be considered in the context of hyperkalaemia. The results offer a platform to inform alignment in guidance developed by NHS England Think Kidneys Programme, the Royal College of Pathologists and the UK Renal Association.^{33 34}

AKI is a clinical syndrome and AKI warning stage test results need to be considered with an understanding of the clinical context in which they were taken.⁴ The UCLA/RAND consensus process aimed to provide face validity regarding the timeliness in response to AKI warning stage test results by clinicians based on the assumption that all the relevant clinical information was available. There is evidence of the predictive validity of quality indicators developed through the RAM.^{32 40} In terms of developing 'timeliness in response' to AKI test results as an indicator of patient safety in primary care, future research needs to provide evidence of acceptability, feasibility, reliability, sensitivity to change and predictive validity.³²

Although a range of clinically relevant scenarios were considered, in terms of burden on RAND/UCLA panelists, it was not feasible to test all major comorbidity groups (eg, patients with type 2 diabetes, patients receiving cancer treatment and not on an end of life care pathway). In addition, developing scenarios for working hours and out of hours may have produced greater clarity on timeliness in response though would have also lengthened the ratings process for panel members with the potential for 'cognitive strain'.⁴¹ On balance, the chosen response times for rating (ie, seek immediate admission; respond within 6 hours; respond within 24 hours; respond within 72 hours) sought to evaluate appropriate actions taking into account key time frames in current clinical practice (eg, respond within 6 hours in order to determine whether appropriate to be seen within working hours, or before next day if knowledge of test result during out of hours; respond within 72 hours to determine if appropriate to wait until Monday if knowledge of a test result on Friday afternoon/evening).

Though the study sought clarity on 'necessary' next steps conducted under ideal conditions, it is hoped that the subsequent published guidance will seek to drive system change to better support clinician decision-making, both for in and out of hours care. Developing robust systems for managing laboratory results is deemed key to ensuring patient safety in general practice.⁴²

Comparison with other studies

The introduction of AKI as a new classification system has both the potential to structure and constrain the delivery of acute illness care.^{4 43} Although AKI alerts are a potential driver for improving patient safety, AKI is a clinical diagnosis and diagnostic accuracy is dependent on the clinician interpreting the test result in the context of an individual patient. This includes

recognising the potential for 'false positives' such as cases of misclassified CKD.^{15 19 20} An assessment of the sensitivity of the NHS England AKI algorithm has been carried out in secondary care but so far, there remain little data on either its specificity or its real-time use in primary care.¹⁹ A retrospective review of electronic medical records would help estimate the diagnostic AKI error rate as well as determine the incidence and factors associated with missed diagnostic and management opportunities.⁴⁴

In the majority of UCLA/RAND clinical cases tested through a multidisciplinary professional panel, a wait of up to 72 hours to respond to an AKI warning stage test result was considered to be inappropriate. This is a potentially important marker of patient safety and requires further evaluation. Caution is required interpreting an AKI warning stage test result in primary care as AKI diagnostic and staging criteria are based on changes in serum creatinine by $>26 \mu\text{mol/L}$ within the previous 48 hours, or an increase in serum creatinine by ≥ 1.5 times baseline that is known or presumed to have occurred within the prior 7 days.^{4 15} Currently, unlike the hospital setting, 'baseline' creatinine levels for comparison are less likely to have taken place within these time periods, reiterating the importance that AKI is a clinical (not merely a biochemical) diagnosis that requires clinician review in order to place a test result in context.

Though there is evidence that AKI is associated with adverse clinical outcomes, the majority of the studies informing the development of the KDIGO classification system for AKI were based on critical care databases.⁴⁵ As such, its application may not necessarily be generalisable to the primary care setting.⁴⁵ However, irrespective of whether an AKI warning stage test result leads to confirmation of AKI or alternatively flags CKD progression, there is evidence to indicate that the e-alerts are still likely to 'identify patients at risk of poor outcomes' in terms of non-recovery, chronic renal replacement therapy and long-term mortality.^{17 46}

Implications for clinicians, policymakers and future research

There was greater uncertainty in how to respond to AKI warning stage test results for people with chronic heart failure, a population who have an increased morbidity and mortality and who account for ~5% of emergency hospital admissions.⁴⁷ This highlights a need for better evidence to support response to changes in serum creatinine for these patients.^{48 49} As indicated by the European Society of Cardiology Heart Failure (ESC) guidelines (2016), increases in serum creatinine are not always the main clinical priority, 'especially when they are accompanied by appropriate decongestion, diuresis and haemoconcentration'⁴⁹—although this statement was in the context of acute heart failure in hospital; data on the prognostic importance of changes in kidney

function during treatment of heart failure in primary care are lacking. The ESC guidelines recommend that 'when large increases in serum creatinine occur, care should be taken to evaluate the patient thoroughly and should include assessment of a possible renal artery stenosis, excessive hypervolaemia or hypovolaemia, concomitant medication and hyperkalaemia, which frequently coincides with worsening renal function'.⁴⁹ This again highlights the importance of 'treating the patient and not the test result' and of comparing the test result with current and previous creatinine levels.^{33 48} Knowing the reason for why kidney function tests were taken is of primary importance. Were they taken in the context of: routine chronic disease monitoring, pharmacotherapy monitoring (eg, dose adjustment) or to support decision-making for patients presenting with a history of acute illness?³³

Effective minimally disruptive medicine is a key principle underpinning care delivery.³⁷ Resourcing initiatives to address the growth in demand for urgent care services is a priority, particularly with an ageing population with increasing care needs.^{50 51} Rather than being an additional burden, targeting AKI may provide an approach to better assessment and management of acute illness across the primary/secondary care interface.⁵² As stated by Guthrie *et al*,³⁹ people with complex comorbidities 'have patterns of illness that do not neatly fit disease categories, in which case syndrome focused guidance or tools will often be relevant'.³⁹ Aligned with the implementation of the national AKI algorithm and professional guidance, resourcing patient and public involvement is critical to the development of a more responsive urgent care service.^{50 51} However, while dialogue about the importance of kidney health may provide a platform to support self-management, this currently remains limited.^{33 53–56} Evidence exists that only one in two people are aware that the kidneys make urine and that only 12% of the population have knowledge that the kidneys have a role in processing medicines.⁵⁷

CONCLUSIONS

Tackling the harm associated with care complicated by AKI is a global priority.² In England, a national AKI algorithm is being introduced across the to drive this change.¹⁵ However, in a context where clinical workload in English general practice is reaching saturation point,³⁶ there is a need to maximise its clinical utility and minimise the potential for burden on clinicians and patients.³⁷ The study sought to address this challenge and the findings have informed the development of guidance to support a timely and calibrated response to AKI warning stage test results for adults in primary care. Further research is needed to support effective implementation, with a view to examine the effect on health outcomes and costs.

Author affiliations

¹National Institute for Health Research Collaboration for Leadership in Applied Health Research and Care (NIHR CLAHRC) Greater Manchester, Centre for Primary Care, Institute of Population Health, The University of Manchester, Manchester, UK

²RCGP Clinical champion for Kidney Care, Royal College of General Practitioners, London, UK

³Department of Geratology, Oxford University Hospitals NHS Foundation Trust, Oxford, UK

⁴Department of Clinical Biochemistry, Norfolk and Norwich University Hospitals Foundation Trust, Norwich, UK

⁵NIHR Greater Manchester Primary Care Patient Safety, Translational Research Centre, Institute of Population Health, Centre for Primary Care, University of Manchester, Manchester, UK

⁶Department of Renal Medicine, Newcastle upon Tyne Hospitals NHS Foundation Trust, Newcastle upon Tyne, UK

Twitter Follow Tom Blakeman at @TomBlakeman

Acknowledgements The following colleagues are from the NHS England Think Kidneys Programme: Richard Fluck, Ron Cullen, Mike Jones, Karen Thomas, Julie Slevin, Annie Taylor, Teresa Wallace and Melanie Dillon. TB is funded by the National Institute for Health Research Collaboration for Leadership in Applied Health Research and Care (NIHR CLAHRC) Greater Manchester. The work outlined in this document may be considered to be affiliated to the work of the NIHR CLAHRC Greater Manchester. DL is funded by the NIHR Oxford Biomedical Research Centre.

Contributors TB, CT and SC designed the study. TB, KG, DL, BL, SC and CT developed the clinical scenarios for appropriateness rating and organised panel membership for round 1 and 2 ratings. SC, CT and TB conducted the round 2 meeting. TB, JYT and CT conducted the analysis and SC provided methodological expertise. All authors contributed to the interpretation of the results, commented on draft manuscripts and have given their approval for publication.

Funding This work was supported by Think Kidneys. Think Kidneys is a national programme led by NHS England in partnership with UK Renal Registry.

Disclaimer The views expressed in this document are those of the author(s) and not necessarily those of the NHS, NIHR or the Department of Health.

Competing interests None declared.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

REFERENCES

1. Matsushita K, van der Velde M, Astor BC, *et al*. Association of estimated glomerular filtration rate and albuminuria with all-cause and cardiovascular mortality in general population cohorts: a collaborative meta-analysis. *Lancet* 2010;375:2073–81.
2. Mehta RL, Cerda J, Burdmann EA, *et al*. International Society of Nephrology's Oby25 initiative for acute kidney injury (zero preventable deaths by 2025): a human rights case for nephrology. *Lancet* 2015;385:2616–43.
3. Jha V, Garcia-Garcia G, Iseki K, *et al*. Chronic kidney disease: global dimension and perspectives. *Lancet* 2013;382:260–72.
4. Kidney Disease Improving Global Outcomes (KDIGO) Acute Kidney Injury Work Group. KDIGO clinical practice guideline for acute kidney injury. *Kidney Int Suppl* 2012;2:1–138.
5. NHS England. *The forward view into action: planning for 2015/16*. NHS England Publications, 2014.
6. National Institute for Health and Care Excellence. *Acute kidney injury: prevention, detection and management of acute kidney injury*

- up to the point of renal replacement therapy. CG169 London: NICE, 2013.
7. Liao F, Pascual J. Epidemiology of acute renal failure: a prospective, multicenter, community-based study. *Kidney Int* 1996;50:811–18.
 8. Wonnacott A, Meran S, Amphlett B, *et al.* Epidemiology and outcomes in community-acquired versus hospital-acquired AKI. *Clin J Am Soc Nephrol* 2014;9:1007–14.
 9. Liangos O, Wald R, O'Bell JW, *et al.* Epidemiology and outcomes of acute renal failure in hospitalized patients: a national survey. *Clin J Am Soc Nephrol* 2006;1:43–51.
 10. Hsu CY, McCulloch CE, Fan D, *et al.* Community-based incidence of acute renal failure. *Kidney Int* 2007;72:208–12.
 11. Kolhe NV, Muirhead AW, Wilkes SR, *et al.* National trends in acute kidney injury requiring dialysis in England between 1998 and 2013. *Kidney Int* 2015;88:1161–9.
 12. Abraham KA, Thompson EB, Bodger K, *et al.* Inequalities in outcomes of acute kidney injury in England. *QJM* 2012;105:729–40.
 13. Selby NM, Kolhe NV, McIntyre CW, *et al.* Defining the cause of death in hospitalised patients with acute kidney injury. *PLoS ONE* 2012;7:e48580.
 14. Kerr M, Bedford M, Matthews B, *et al.* The economic impact of acute kidney injury in England. *Nephrol Dial Transplant* 2014;29:1362–8.
 15. NHS England. *UK renal registry, think kidneys. Acute kidney injury warning alert best Practice.* Think Kidneys, 2014.
 16. NHS England. *Patient safety alert: stage 3 directive: standardising the early identification of acute kidney injury.* NHS England, 2014.
 17. Sawhney S, Fluck N, Fraser SD, *et al.* KDIGO-based acute kidney injury criteria operate differently in hospitals and the community—findings from a large population cohort. *Nephrol Dial Transplant* 2016;31:922–9.
 18. Kolhe NV, Staples D, Reilly T, *et al.* Impact of compliance with a care bundle on acute kidney injury outcomes: a prospective observational study. *PLoS ONE* 2015;10:e0132279.
 19. Sawhney S, Fluck N, Marks A, *et al.* Acute kidney injury—how does automated detection perform? *Nephrol Dial Transplant* 2015;30:1853–61.
 20. Sawhney S. Automated alerts for acute kidney injury warrant caution. *BMJ* 2015;350:h19.
 21. Barton AL, Mallard AS, Parry RG. One year's observational study of acute kidney injury incidence in primary care; frequency of follow-up serum creatinine and mortality risk. *Nephron* 2015;130:175–81.
 22. National Institute for Health and Care Excellence. *Sepsis: recognition, diagnosis and early management.* NICE, 2016.
 23. Singer M, Deutschman CS, Seymour C, *et al.* The third international consensus definitions for sepsis and septic shock (sepsis-3). *JAMA* 2016;315:801–10.
 24. Wilson FP, Shashaty M, Testani J, *et al.* Automated, electronic alerts for acute kidney injury: a single-blind, parallel-group, randomised controlled trial. *Lancet* 2015;385:1966–74.
 25. Laing C. On the alert for outcome improvement in acute kidney injury. *Lancet* 2015;385:1924–6.
 26. Thomas ME, Sitch A, Baharani J, *et al.* Earlier intervention for acute kidney injury: evaluation of an outreach service and a long-term follow-up. *Nephrol Dial Transplant* 2015;30:239–44.
 27. McCoy AB, Waitman LR, Gadd CS, *et al.* A computerized provider order entry intervention for medication safety during acute kidney injury: a quality improvement report. *Am J Kidney Dis* 2010;56:832–41.
 28. Blakeman T, Protheroe J, Chew-Graham C, *et al.* Understanding the management of early-stage chronic kidney disease in primary care: a qualitative study. *Br J Gen Pract* 2012;62:e233–42.
 29. Fitch K, Bernstein SJ, Aguilar MD, *et al.* *The RAND/UCLA appropriateness method user's manual.* Santa Monica: RAND, 2001.
 30. Avery AJ, Dex GM, Mulvaney C, *et al.* Development of prescribing-safety indicators for GPs using the RAND Appropriateness Method. *Br J Gen Pract* 2011;61:e526–36.
 31. Campbell SM, Fuat A, Summerton N, *et al.* Diagnostic triage and the role of natriuretic peptide testing and echocardiography for suspected heart failure: an appropriateness ratings evaluation by UK GPs. *Br J Gen Pract* 2011;61:e427–35.
 32. Campbell SM, Braspenning J, Hutchinson A, *et al.* Research methods used in developing and applying quality indicators in primary care. *Qual Saf Health Care* 2002;11:358–64.
 33. NHS England, UK Renal Registry. Think Kidneys. *Acute kidney injury best practice guidance: responding to AKI warning stage test results for adults in primary care.* Think Kidneys, 2016. <http://www.thinkkidneys.nhs.uk/>
 34. UK Renal Association. *Clinical practice guidelines: treatment of acute hyperkalaemia in adults.* UK Renal Association, 2014.
 35. McCartney M, Treadwell J. The RCGP's new standing group on overdiagnosis. *BMJ* 2014;349:g4454.
 36. Hobbs FDR, Bankhead C, Mukhtar T, *et al.* Clinical workload in UK primary care: a retrospective analysis of 100 million consultations in England, 2007–14. *Lancet* 2016;387:2323–30.
 37. May C, Montori VM, Mair FS. We need minimally disruptive medicine. *BMJ* 2009;339:b2803.
 38. Johnson MJ, May CR. Promoting professional behaviour change in healthcare: what interventions work, and why? A theory-led overview of systematic reviews. *BMJ Open* 2015;5:e008592.
 39. Guthrie B, Payne K, Alderson P, *et al.* Adapting clinical guidelines to take account of multimorbidity. *BMJ* 2012;345:e6341.
 40. Kravitz RL, Laouri M, Kahan JP, *et al.* Validity of criteria used for detecting underuse of coronary revascularization. *JAMA* 1995;274:632–8.
 41. Kravitz RL, Park RE, Kahan JP. Measuring the clinical consistency of panelists' appropriateness ratings: the case of coronary artery bypass surgery. *Health Policy* 1997;42:135–43.
 42. Bell BG, Spencer R, Avery AJ, *et al.* Tools for measuring patient safety in primary care settings using the RAND/UCLA appropriateness method. *BMC Fam Pract* 2014;15:1–7.
 43. Armstrong D. Diagnosis and nosology in primary care. *Soc Sci Med* 2011;73:801–7.
 44. Cheraghi-Sohi S, Singh H, Reeves D, *et al.* Missed diagnostic opportunities and English general practice: a study to determine their incidence, confounding and contributing factors and potential impact on patients through retrospective review of electronic medical records. *Implement Sci* 2015;10:105.
 45. Thomas ME, Blaine C, Dawney A, *et al.* The definition of acute kidney injury and its use in practice. *Kidney Int* 2015;87:62–73.
 46. Hobbs H, Bassett P, Wheeler T, *et al.* Do acute elevations of serum creatinine in primary care engender an increased mortality risk? *BMC Nephrol* 2014;15:1–10.
 47. National Institute for Health and Care Excellence. *Management of chronic heart failure in adults in primary and secondary care.* Manchester: NICE, 2010.
 48. Perazella MA, Coca SG. Three feasible strategies to minimize kidney injury in 'incipient AKI'. *Nat Rev Nephrol* 2013;9:484–90.
 49. Ponikowski P, Voors AA, Anker SD, *et al.* 2016 ESC guidelines for the diagnosis and treatment of acute and chronic heart failure: the Task Force for the diagnosis and treatment of acute and chronic heart failure of the European Society of Cardiology (ESC) developed with the special contribution of the Heart Failure Association (HFA) of the ESC. *Eur Heart J* 2016;37:2129–200.
 50. NHS England. *High quality care for all, now and for future generations: transforming urgent and emergency care services in England—Urgent and Emergency Care Review End of Phase 1 Report.* Leeds: NHS England, 2013.
 51. NHS England. *General Practice: Forward View.* NHS England, 2016.
 52. Blakeman T, Harding S, O'Donoghue D. Acute kidney injury in the community: why primary care has an important role. *Br J Gen Pract* 2013;63:173–4.
 53. Blakeman T, Blickem C, Kennedy A, *et al.* Effect of information and telephone-guided access to community support for people with chronic kidney disease: randomised controlled trial. *PLoS ONE* 2014;9:e109135.
 54. Daker-White G, Rogers A, Kennedy A, *et al.* Non-disclosure of chronic kidney disease in primary care and the limits of instrumental rationality in chronic illness self-management. *Soc Sci Med* 2015;131:31–9.
 55. Greer RC, Cooper LA, Crews DC, *et al.* Quality of patient-physician discussions about CKD in primary care: a cross-sectional study. *Am J Kidney Dis* 2011;57:583–91.
 56. Gaffney H, Blakeman T, Blickem C, *et al.* Predictors of patient self-report of chronic kidney disease: baseline analysis of a randomised controlled trial. *BMC Fam Pract* 2014;15:196.
 57. NHS England, UK Renal Registry, Think Kidneys. Understanding what the public know about their kidneys and what they do. Findings from Ipsos MORI survey—July 2014: Think Kidneys, 2014.