

BMJ Best Practice

Dupuytren's contracture

Straight to the point of care



Last updated: Jul 14, 2023

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Summary

Dupuytren's contracture is an inherited disease of progressive fibrous tissue contracture of the palmar fascia.

Predominantly affects men of northern European descent aged >40 years old who smoke, drink alcohol, or have diabetes.

Patients present with a small lump or multiple lumps with pits in the palm of the hand, progressing to contractures of the fingers.

Intralesional corticosteroid injections have been shown to reduce the need for surgery.

Surgical referral should be made when metacarpophalangeal joint contractures reach 30 degrees, or if any degree of proximal interphalangeal joint contracture is present.

Percutaneous needle fasciotomy and collagenase injections are significant therapeutic alternatives to surgery.

Definition

An inherited disease of progressive fibrous tissue contracture of the palmar fascia, Dupuytren's contracture is seen predominantly in men of northern European descent aged >40 years. Its inheritance pattern is believed to be autosomal dominant with variable penetrance.[1]

Epidemiology

There are differing reports on the prevalence of Dupuytren's contracture depending on region and study design, however worldwide disease prevalence is 8.2%, with the highest rates per continent observed in Africa at 17.2%, and the lowest in America at 2.3%.^[4] Other studies report higher prevalence among those of European descent than in other races, and within Europe it is higher in northern than in Mediterranean countries.^[5] Prevalence in Western countries ranges from 0.6-31.6%.^[6] For instance, in the same age group, prevalence has been reported as 22.0% (with progression to contractures in 4.2%), 2.6%, and 0.67% in Dutch, Swedish and UK cohorts respectively.^[7] ^[8] ^[9] In Japan, disease prevalence is estimated to be 7.0%.^[10]

Dupuytren's disease is twice as common in men as in women, and incidence increases with age.^[6] ^[7] ^[8] ^[9] ^[10]

Aetiology

Dupuytren's contracture is a progressive fibroproliferative disease that is believed to show autosomal dominant inheritance with variable penetrance.^[1] The higher prevalence of the disease in men is probably due to the fact that androgen receptors are expressed in Dupuytren nodules.

The familial occurrence and its presence in identical twins suggest a genetic basis for the disease. The sibling recurrence risk ratio equals 2.9 (95% CI 2.6 to 3.3). Although the genetic factors involved have not been fully elucidated, in one family the gene has been mapped to chromosome 16q.^[1] DNA microarray has demonstrated that >30 unique genes are upregulated and 6 unique genes downregulated by a factor of 4 or greater in affected patients.^[11] A novel gene, MafB (musculoaponeurotic fibrosarcoma oncogene homolog B), is overexpressed in patients with Dupuytren's contracture but not in those with normal fascia.^[11]

A high number of mitochondria have been demonstrated in fibroblasts derived from diseased tissue, and a mitochondrial defect in the 16S ribosome RNA has been noted in 90% of affected patients compared with none in the control patients.^[12] Thus, a mitochondrial aetiology may be significant in some families, whereas a chromosomal defect may be important in others.

A number of growth factors, immunological mediators, and free radicals are also implicated as causative factors, and it seems that an inciting factor triggers these mediators to stimulate myofibroblast proliferation.^[13]

Diabetes mellitus has a strong association with Dupuytren's contracture.^[4] ^[10] ^[14] ^[15] ^[16] The prevalence of the disease in patients with diabetes ranges from 3% to 33% and increases with the duration of the diabetes. It is hypothesised that diabetes causes microvascular changes that produce local hypoxic tissue damage, inducing Dupuytren's contracture. Patients with diabetes tend to have a mild form of the disease, with slow progression.

Greater alcohol intake is associated with an increased likelihood of Dupuytren's contracture.^[4] ^[7] ^[10] ^[17] However, it should be noted that most patients with the disease do not misuse alcohol, and not all studies have found a significant association.^[18] For unclear reasons, smoking increases risk, although the microvascular changes associated with smoking and the effect of carbon monoxide on mitochondrial cytochrome c, released by mitochondria in response to pro-apoptotic stimuli, may play a role.^[17]

Trauma was first proposed to be a cause by Dupuytren himself. Since the first report that linked trauma to the disease, the strength of this association has been controversial. It is unclear why the disease would be caused by heavy labour, and further studies are required to control for smoking and alcohol use.^[19]

There have been reports of a connection between epilepsy and Dupuytren's contracture. When discontinuing anticonvulsant drugs, findings such as palmar cords and knuckle pad thickening have been reported to regress, although large cohort studies have not found an association with epilepsy or epilepsy medication and the disease. It would seem that epilepsy is not a risk for the disease, but anticonvulsant drugs (e.g., phenytoin and carbamazepine) may trigger it.^[20]

Pathophysiology

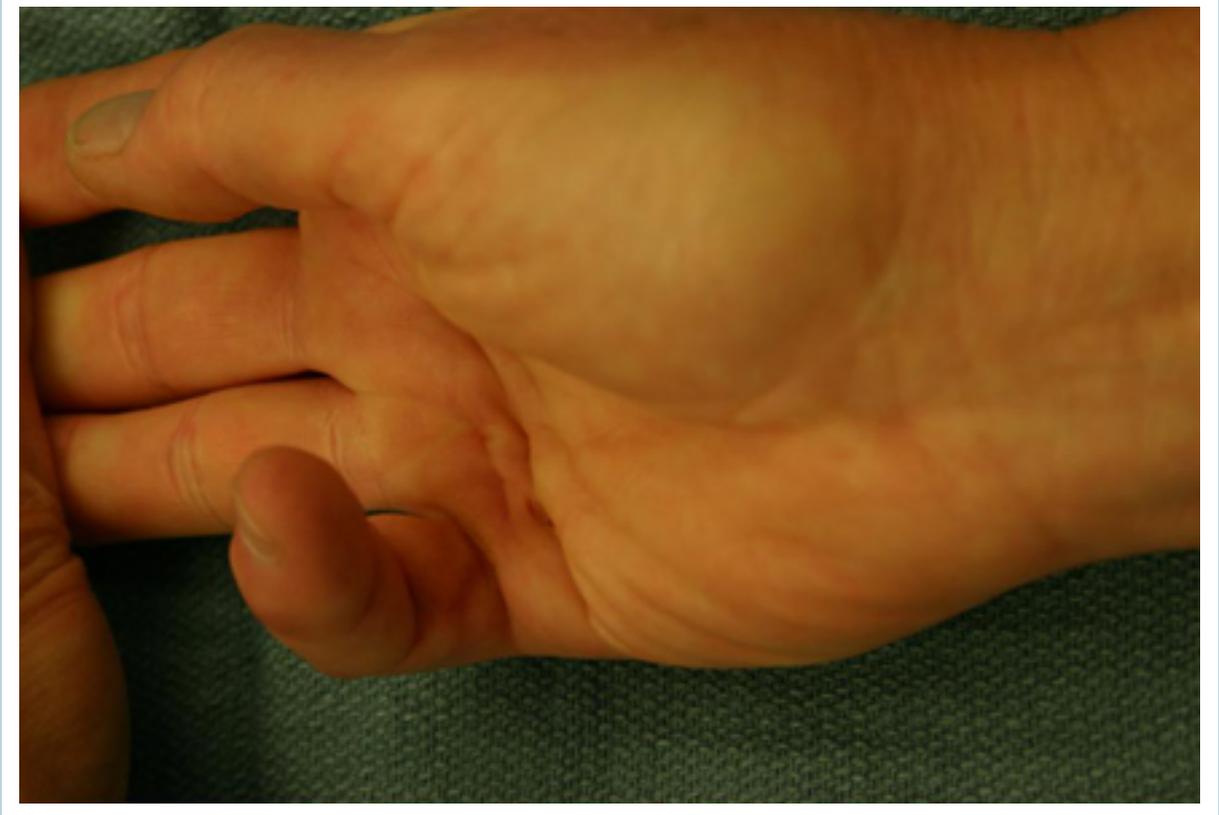
Several hypotheses have been proposed for the underlying pathophysiological mechanisms of Dupuytren's contracture. Localised ischaemia from diabetes, trauma, or other aetiologies is thought to produce xanthine oxidase free radicals that may then damage the perivascular connective tissue and bring on a reparative response by surrounding fibroblasts or triggering myofibroblasts. Fibroblast growth factor (FGF), platelet-derived growth factor (PDGF), and transforming growth factor-beta (TGF- β) may signal an overproduction of myofibroblasts, leading to the formation of nodules and contractures. It has also been demonstrated in Dupuytren's contracture that the proportions of the different types of collagen are altered, with collagen type 1 replaced by collagen type 3, similar to the proliferative phase of wound healing.

Classification

Characteristics of fibrous tissue deformity and contracture^[2]

Three grades of Dupuytren's contracture have been described, based on the characteristics of the fibrous tissue deformity and the presence of a contracture.

- Grade 1: thickened nodule and a band in the palmar aponeurosis that may progress to skin tethering, puckering, or pitting.
- Grade 2: peritendinous band with limited extension of the affected finger.
- Grade 3: presence of flexion contracture.



Preoperative view of a small finger flexion contracture with surgical indications

From the collection of Dr C.M. Rodner; used with permission



*Preoperative view of a small finger flexion contracture with surgical indications
From the collection of Dr C.M. Rodner; used with permission*

Presence of nodules and degree of contracture^[3]

The 7 stages of Dupuytren's contracture have been described, based on the presence of nodules and the severity of the contracture.

- Stage 0: no contracture
- Stage N: no contracture, palpable nodule
- Stage N/1: 0° to 5° contracture, palpable nodule
- Stage 1: 6° to 45° contracture
- Stage 2: 46° to 90° contracture
- Stage 3: 91° to 135° contracture
- Stage 4: >135° contracture

Case history

Case history #1

A 60-year-old man of northern European descent presents with a several-year history of a mass on the ulnar side of his palm. He has no significant past medical history. The patient states that this mass has increased in size over the past year and is now causing his small finger to bend down. He has no pain but is increasingly bothered by the flexion deformity of his small finger, as it is hard for him to put on a glove, and his finger often gets caught on things. On physical examination there is a cord palpable over the small finger and ring finger rays, with extension across the metacarpophalangeal joint of the small finger producing a contracture of 50 degrees in this joint. There is no proximal interphalangeal joint involvement. Plain x-rays and laboratory studies are normal.

Approach

The diagnosis of Dupuytren's contracture is clinical and based on a thorough physical examination of the hands, supported by an indicative clinical history.

Clinical history

Dupuytren's contracture is typically seen in men of northern European descent aged >40 years. As it is believed to show autosomal dominant inheritance with variable penetrance, there may be a family history of the disease.^[1] Medical history may be positive for diabetes mellitus or epilepsy, and social history may reveal that the patient is a smoker or a heavy drinker.

Patients describe difficulties with face washing, combing their hair, and putting their hands in their pockets or fitting them into gloves.

Physical examination

A detailed physical examination of the affected hand(s) reveals a number of characteristic findings, depending on the disease progression. Bilateral hand involvement is common, with one hand usually more severely affected than the other, although the handedness of the patient is not a predictor of severity.

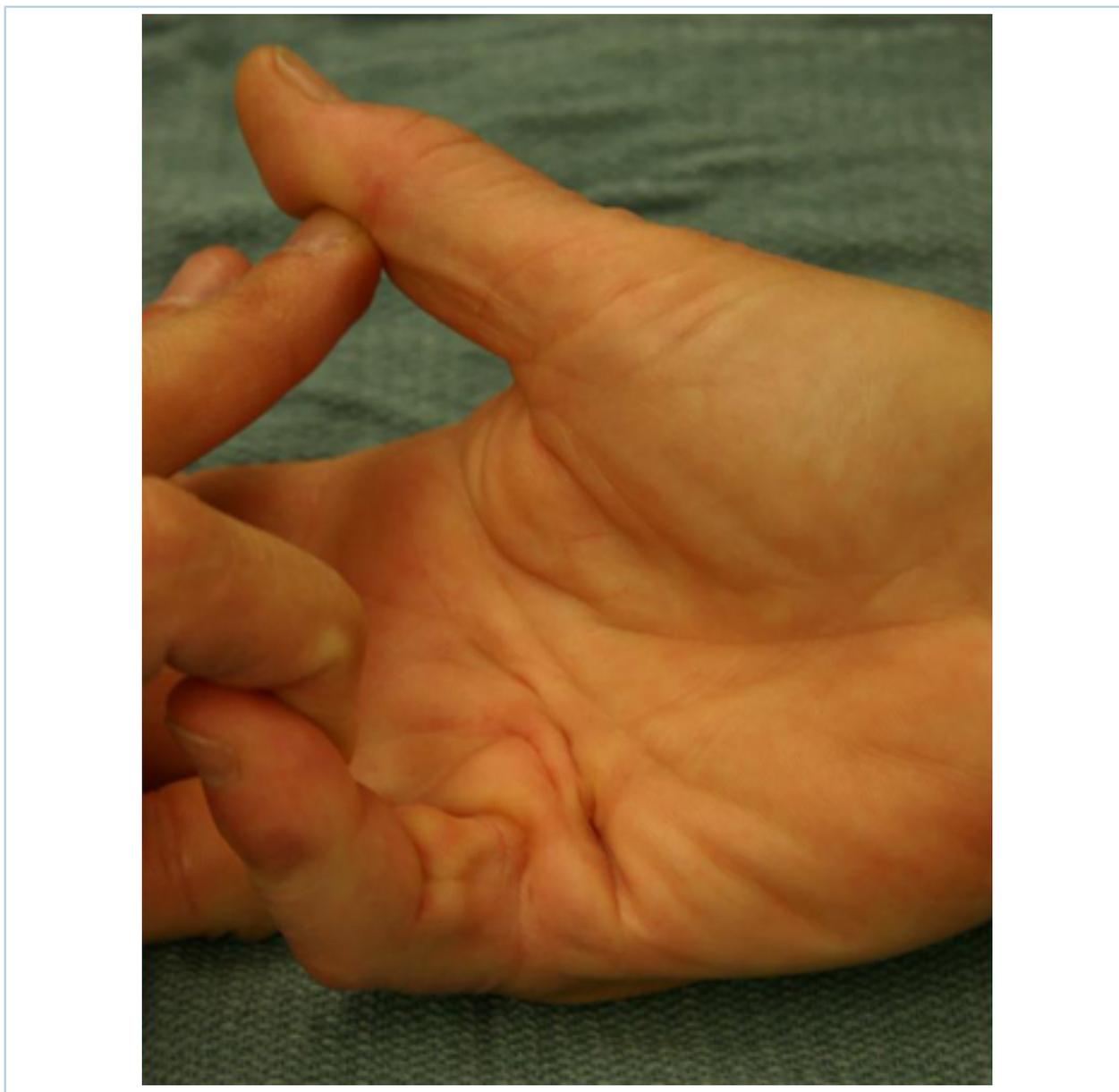
A firm and thickened palmar nodule over the metacarpal head at the level of the distal palmar crease, proximal to the metacarpophalangeal (MCP) joint, is often the earliest sign. This nodule may be associated with a band in the palmar aponeurosis. After nodule formation, palmar skin changes occur, with skin thickening, tethering, puckering, or pitting, as well as subcutaneous fat fibrosis.

The formation of pretendinous cords usually occurs as isolated nodules coalesce, although nodules and cords can be present simultaneously. The most commonly affected digit is the ring finger, followed by the small finger, thumb, middle finger, and index finger. As pretendinous cords in the palm progress they may travel across the MCP joint and, over time, produce MCP joint flexion contractures, leading to limited extension of the affected finger.



Preoperative view of a small finger flexion contracture with surgical indications

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The degree of contracture is dependent on the severity of the disease. Digital cords that cross the proximal interphalangeal (PIP) joint may cause PIP joint contractures.

Examination of the dorsal aspect of the PIP joints may reveal areas of subcutaneous fibrosis, known as Garrod's nodes or knuckle pads, which are indicative of systemic fascial disease and predictive of bilateral involvement. Garrod's nodes are found in about one half of patients.

The term 'Dupuytren's diathesis' refers to patients with severe disease. These patients are usually younger, with very rapid disease progression involving both hands, and are more likely to have systemic fascial disease, including Ledderhose's disease, affecting the plantar surface of the feet and, in men, Peyronie's disease, which affects the penis.[19]

The Hueston table-top test aids diagnosis and involves the patient attempting to lay the palm of the hand flat on a table surface. The test is positive if the patient is unable to flatten the hand on the table.

Investigations

As the diagnosis of Dupuytren's contracture is predominantly clinical, ultrasound of the hand has limited usefulness in the diagnosis of the disease. It shows a mass lying between the flexor tendon below and the skin above. MRI and radiographs are not indicated in the diagnosis of the disease.

History and exam

Key diagnostic factors

presence of risk factors (common)

- Risk factors for Dupuytren's contracture include a family history of the disease, diabetes mellitus, anticonvulsant therapy, smoking, alcohol, and trauma.

male >40 years of age (common)

- Dupuytren's contracture is predominantly seen in men of northern European descent aged >40 years.^{[6] [7] [8] [9]}

difficulties with manual activities (common)

- Patients describe difficulties with face washing, combing their hair, and putting their hands in their pockets or fitting them into gloves.^[19]

palmar nodule (common)

- A firm and thickened palmar nodule over the metacarpal head at the level of the distal palmar crease, proximal to the metacarpophalangeal (MCP) joint, is often the earliest sign.
- Nodules may be associated with a band in the palmar aponeurosis.

palmar skin changes (common)

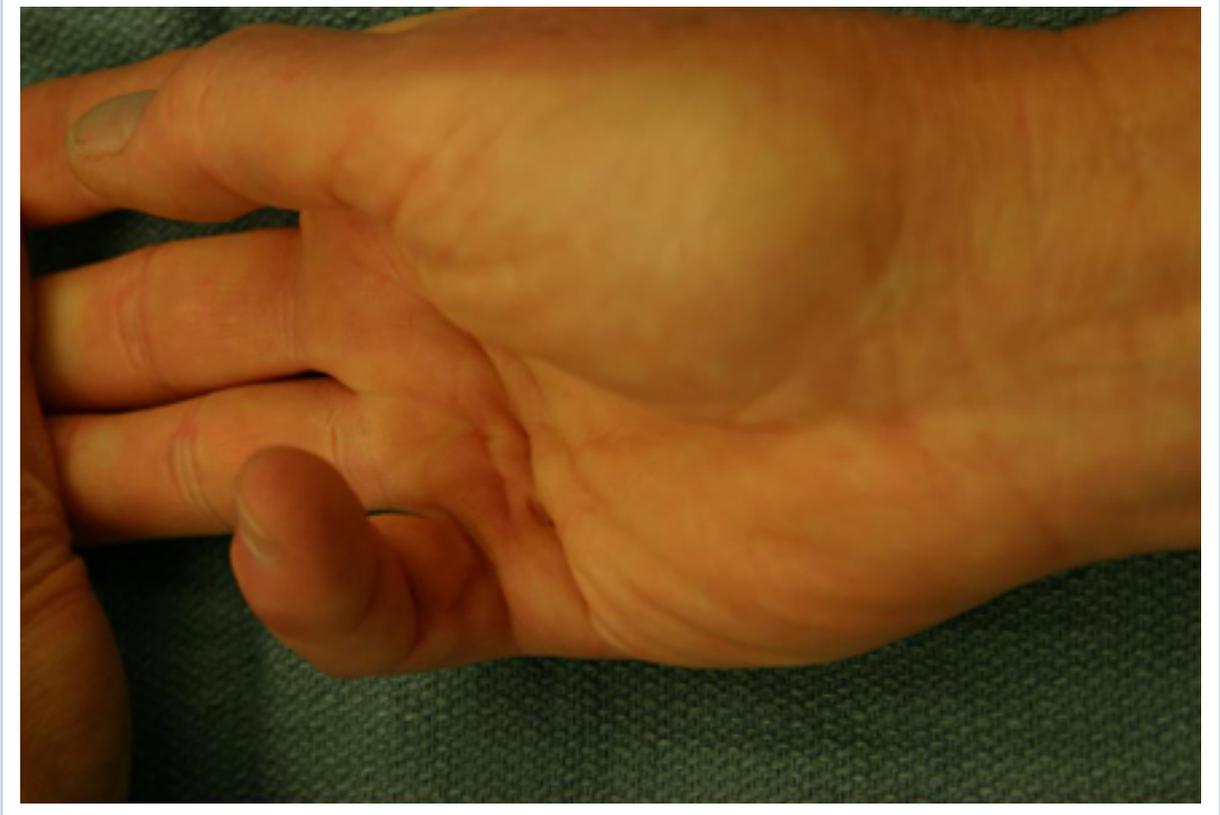
- After nodule formation, skin changes occur, with skin thickening, tethering, puckering, or pitting, as well as subcutaneous fat fibrosis.

pretendinous cords (common)

- The formation of pretendinous cords usually occurs as isolated nodules coalesce, although nodules and cords can be present simultaneously.
- The most commonly affected digit is the ring finger, followed by the small finger, thumb, middle finger, and index finger.

MCP joint contracture (common)

- As pretendinous cords in the palm progress they may travel across the MCP joint and, over time, produce MCP joint flexion contractures, leading to limited extension of the affected finger.



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- The degree of contracture is dependent on the severity of the disease.

proximal interphalangeal (PIP) joint contracture (common)

- Digital cords that cross the PIP joint may cause PIP joint contractures.

positive Hueston table-top test (common)

- Involves the patient attempting to lay the palm of the hand flat on a table surface.
- The result is positive if the patient is unable to flatten the hand on the table.

Other diagnostic factors

bilateral involvement (common)

- Bilateral hand involvement is common, with one hand usually more severely affected than the other, although the handedness of the patient is not a predictor of severity.
- Patients with Dupuytren's diathesis have rapid disease progression involving both hands.

Garrod's nodes (uncommon)

- Examination of the dorsal aspect of the PIP joints may reveal areas of subcutaneous fibrosis, known as Garrod's nodes, or knuckle pads, which are indicative of systemic fascial disease and predictive of bilateral involvement.
- Garrod's nodes are found in about one half of patients.

involvement of plantar surface of the feet (uncommon)

- Patients with Dupuytren's diathesis are more likely to have systemic fascial disease, including Ledderhose's disease, which affects the plantar surface of the feet.

involvement of penis (uncommon)

- Patients with Dupuytren's diathesis are more likely to have systemic fascial disease, including Peyronie's disease in men, which affects the penis.

Risk factors

Strong

male sex

- Dupuytren's contracture is twice as common in men as in women.[8] [9] It predominantly affects men of northern European descent aged >40 years.

age >40 years

- The incidence of Dupuytren's contracture increases with age and it predominantly affects men of northern European descent aged >40 years.[6] [7] [8] [9]

family history

- Dupuytren's contracture is believed to show autosomal dominant inheritance with variable penetrance.[1]
- The familial occurrence and its presence in identical twins suggest a genetic basis for the disease. The sibling recurrence risk ratio equals 2.9 (95% CI 2.6 to 3.3).
- Although the genetic factors involved have not been fully elucidated, in one family the gene has been mapped to chromosome 16q.[1]
- DNA microarray has demonstrated that >30 unique genes are upregulated and 6 unique genes downregulated by a factor of 4 or greater in affected patients.[11]
- A novel gene, MafB (musculoaponeurotic fibrosarcoma oncogene homolog B), is overexpressed in patients with Dupuytren's contracture but not in those with normal fascia.[11]

diabetes mellitus

- Diabetes mellitus has a strong association with Dupuytren's contracture.[4] [10] [14] [15] [16] The prevalence of Dupuytren's contracture in patients with diabetes ranges from 3% to 33% and increases with the duration of the diabetes.
- It is hypothesised that diabetes causes microvascular changes that produce local hypoxic tissue damage, inducing Dupuytren's contracture. Patients with diabetes tend to have a mild form of the disease with slow progression.

Weak

high alcohol intake

- Greater alcohol intake is associated with an increased likelihood of Dupuytren's contracture.[4] [7] [10] [17] However, it should be noted that most patients with the disease do not misuse alcohol, and not all studies have found a significant association.[18]

smoking

- For unclear reasons, smoking increases the risk of Dupuytren's contracture, although the microvascular changes associated with smoking and the effect of carbon monoxide on mitochondrial cytochrome c, released by mitochondria in response to pro-apoptotic stimuli, may play a role.[17]

trauma

- Trauma was first proposed to be a cause by Dupuytren himself. Since the first report that linked trauma to the disease, the strength of this association has been controversial. It is unclear why the disease would be caused by heavy labour or vibrational exposure, and further studies are required to control for smoking and alcohol use.
- One theoretical explanation for the connection of trauma and Dupuytren's contracture is the induction of transforming growth factor-beta (TGF- β) with trauma and the stimulation of myofibroblasts from the increased TGF- β . [21]

anticonvulsant medication

- There have been reports of a connection between epilepsy and Dupuytren's contracture. When discontinuing anticonvulsant drugs, findings such as palmar cords and knuckle pad thickening have been reported to regress, although large cohort studies have not found an association with epilepsy or epilepsy medication and the disease. It would seem that epilepsy is not a risk for the disease, but anticonvulsant drugs (phenytoin and carbamazepine) may trigger it.[20]

Investigations

Other tests to consider

Test	Result
ultrasound of hand <ul style="list-style-type: none"> • As the diagnosis is predominantly clinical, ultrasound of the hand has limited usefulness. 	mass lying between the flexor tendon below and the skin above

Differentials

Condition	Differentiating signs / Differentiating tests symptoms	
Trigger finger	<ul style="list-style-type: none"> The finger can be fully extended with a notable click. 	<ul style="list-style-type: none"> Ultrasound: thickening and hypoechogenicity, with possible increased vascularity of the A1 pulley.
Epithelioid sarcoma	<ul style="list-style-type: none"> Tends to be progressive and extend beyond localised digits. 	<ul style="list-style-type: none"> Biopsy: results can be difficult to determine but are often seen as well-defined eosinophilic cytoplasm and one or more atypical, eccentrically located nuclei, resulting in a plasmacytoid appearance.
Camptodactyly	<ul style="list-style-type: none"> Presence of little finger contracture from an early age. 	<ul style="list-style-type: none"> No differentiating tests.
Traumatic finger contracture	<ul style="list-style-type: none"> History of significant preceding trauma leading to injury of the proximal interphalangeal joint. 	<ul style="list-style-type: none"> X-ray of hand may show associated fracture.

Approach

Patients with early Dupuytren's contracture can be managed expectantly, although the injection of nodules with a corticosteroid may be carried out if the Dupuytren lesions are bothersome. Those with metacarpophalangeal (MCP) joint contractures of 30 degrees or less and with no proximal interphalangeal (PIP) joint contractures can be treated with needle aponeurotomy, percutaneous fasciotomy, or corticosteroid injections. When function is impaired or a severe and disabling deformity is present, surgery is recommended, and patients with MCP joint contractures >30 degrees or PIP joint contractures can be treated with either open partial fasciectomy, segmental aponeurotomy, or percutaneous fasciotomy.[19] [22] Partial fasciectomy has become the favoured technique of hand surgeons in the surgical treatment of Dupuytren's contracture, due to the relatively low rate of recurrence seen with this procedure.[23] It is difficult, however, to compare the efficacy of the many different modalities used to treat Dupuytren's contracture. A reason for this is the inconsistency in reporting of outcomes across studies evaluating treatments for Dupuytren's contracture.[24] [25] Only 3 Level I studies have compared the different surgical techniques for the treatment of primary Dupuytren's contracture, and current evidence does not support one procedure as being better than the other, apart from a particularly high recurrence rate after needle fasciotomy. More research will be needed in the future.[26]

All patients with contractures should receive hand therapy post-procedure, and postoperative splinting following open partial fasciectomy is used at the discretion of the surgeon. The effects of stretching on prevention of contracture development for up to 7 months have shown no added benefit, and no studies have been done for more than 7 months. There is therefore no current evidence to support the use of stretching to prevent contractures.[27]

Corticosteroid injections

Patients with early Dupuytren's contracture who have evidence of the disease but have not yet developed contractures can be treated with corticosteroid injections if they are experiencing bothersome symptoms. Those with MCP joint contractures of 30 degrees or less with no PIP joint contractures who wish to avoid a more invasive procedure may also benefit from corticosteroid injections.

The injection of Dupuytren nodules with triamcinolone acetonide monthly for up to 5 months, or every 6 weeks for 3 injections, has been shown to produce significant regression of the disease, with an average of 3.2 injections per nodule required for improvement of function.[28] After corticosteroid injection, fewer patients progress to surgery than would be predicted with expectant management alone.[28]

Collagenase injection

Collagenase clostridium histolyticum is used to treat adult patients with a palpable cord along with MCP or PIP contracture, with a corresponding decrease in both fasciotomies and fasciectomyes.[29] The efficacy and tolerability of injectable mixed collagenase subtypes as an alternative to surgical intervention in Dupuytren's contracture has been examined in a phase 3, double-blind study, which found that collagenase safely and effectively restored normal finger extension in the majority (87%) of patients.[30] A mean of 1.4 injections was required to normalise affected joints, and clinical success was achieved within 29 days. Contracture recurrence was relatively low, occurring in just 5 joints (1 MCP, 4 PIP) between 6 and 24 months after treatment, and the recurrence severity ranged from 20 degrees to 40 degrees. Adverse events were localised to the injection site, generally of mild-to-moderate severity, and transient in nature. In another study, more cords that were injected with collagenase met criteria of a reduction in contracture to 0 to 5 degrees of full extension 30 days after the last injection (64.0% vs.

6.8%, $P < 0.001$) than cords injected with placebo.[31] The most commonly reported adverse events were localised swelling in the hand, pain, bruising, pruritus, and transient regional lymph-node enlargement and tenderness. Serious adverse events were seen in 2% of collagenase recipients, including tendon ruptures and complex regional pain syndrome.[31] Although there is some literature with small numbers of patients to suggest that recurrence is more common in PIP joint contractures than in MCP joint contractures, some guidelines recommend that collagenase treatment is limited only to clinical trials.[32][33]

One systematic review has found that having previous surgery did not affect the efficacy and safety of collagenase injections, making this an option in patients with recurrent Dupuytren's contracture.[34]

The technique for collagenase injection is relatively straightforward. First, the skin is prepared with antiseptic solution. Without the use of any local anaesthesia, an insulin needle (28-gauge) is inserted to a depth of 5 mm for MCP joint lesions and 3 mm for PIP joint lesions. The gristly structure of the cord is easily palpated with the end of the needle. A gentle amount of passive motion ensures that the needle is not in the flexor tendon before the injection of collagenase into the MCP or PIP lesion. The injections are administered proximally-to-distally, repositioning the needle prior to injection of the next dose. After the injection is given, a dressing is placed on the hand. The patient is then instructed to return to the surgery the next day for the finger extension manoeuvre, in which the finger is extended to break the cord. This may be done with or without a local anaesthetic. After the manipulation is successfully completed, there are no restrictions and early movement is encouraged. Although wearing a night-time splint afterwards has not been proven to decrease recurrence, it should be offered to the patient. The patient is followed to assess the need for up to 2 further injections every 4 to 6 weeks.[30] Collagenase clostridium histolyticum may not be available in some countries.

Needle aponeurotomy

Needle aponeurotomy, also called percutaneous needle fasciotomy (PNF), is a minimally invasive technique that can be undertaken in the office. While needle aponeurotomy is typically used for early Dupuytren's contracture, it has also been described for more advanced stages of the disease and can be used in those with MCP joint contractures of 30 degrees or less with no PIP joint contractures.

Needle aponeurotomy is usually successful in correcting the Dupuytren's contracture, takes very little time to perform (usually 20 to 30 minutes), requires only local anaesthesia, and is not very painful. In comparison with open surgical procedures, it results in similar resolution,[35] [36] minimal scarring, faster recovery, and can be repeated easily if the contracture recurs.[37] Needle aponeurotomy is thus an attractive option for patients with less aggressive and early disease.[38] One systematic review reported a tendency to greater patient satisfaction with needle aponeurotomy, with fewer adverse effects, compared with other procedures.[39] Recurrence rates of up to 58% have been reported during 3 to 5 years of follow-up, but long-term outcomes are not well reported.[40]

Randomised studies report no significant difference in treatment outcome between needle aponeurotomy and collagenase injection.[41] [42] In one trial, however, collagenase treatment led to more, mainly transient, complications than needle aponeurotomy.[42]

Percutaneous fasciotomy

Percutaneous fasciotomy is a similar procedure to needle aponeurotomy but uses a scalpel to cut and release the band causing the digital contracture instead of a needle to weaken it.[43][44] It is thus performed by a hand surgeon in the operating room. This technique is normally used in patients with

MCP joint contractures of 30 degrees or less with no PIP joint contractures, but can also be used in more advanced stages of Dupuytren's contracture.[22]

As with needle aponeurotomy, the finger is brought into full extension with a characteristic snap, and the goal of percutaneous fasciotomy is to promote greater extension and function of the affected digit.[22]

The major advantages of percutaneous fasciotomy are that it causes less pain and allows a faster recovery than traditional open fasciectomy interventions. However, as with needle aponeurotomy, this procedure is associated with a risk of recurrence of up to 43%, as diseased fascia is unavoidably left behind. There is concomitant risk of flexor tendon or nerve injury.[43][44][45]

Segmental aponeurotomy

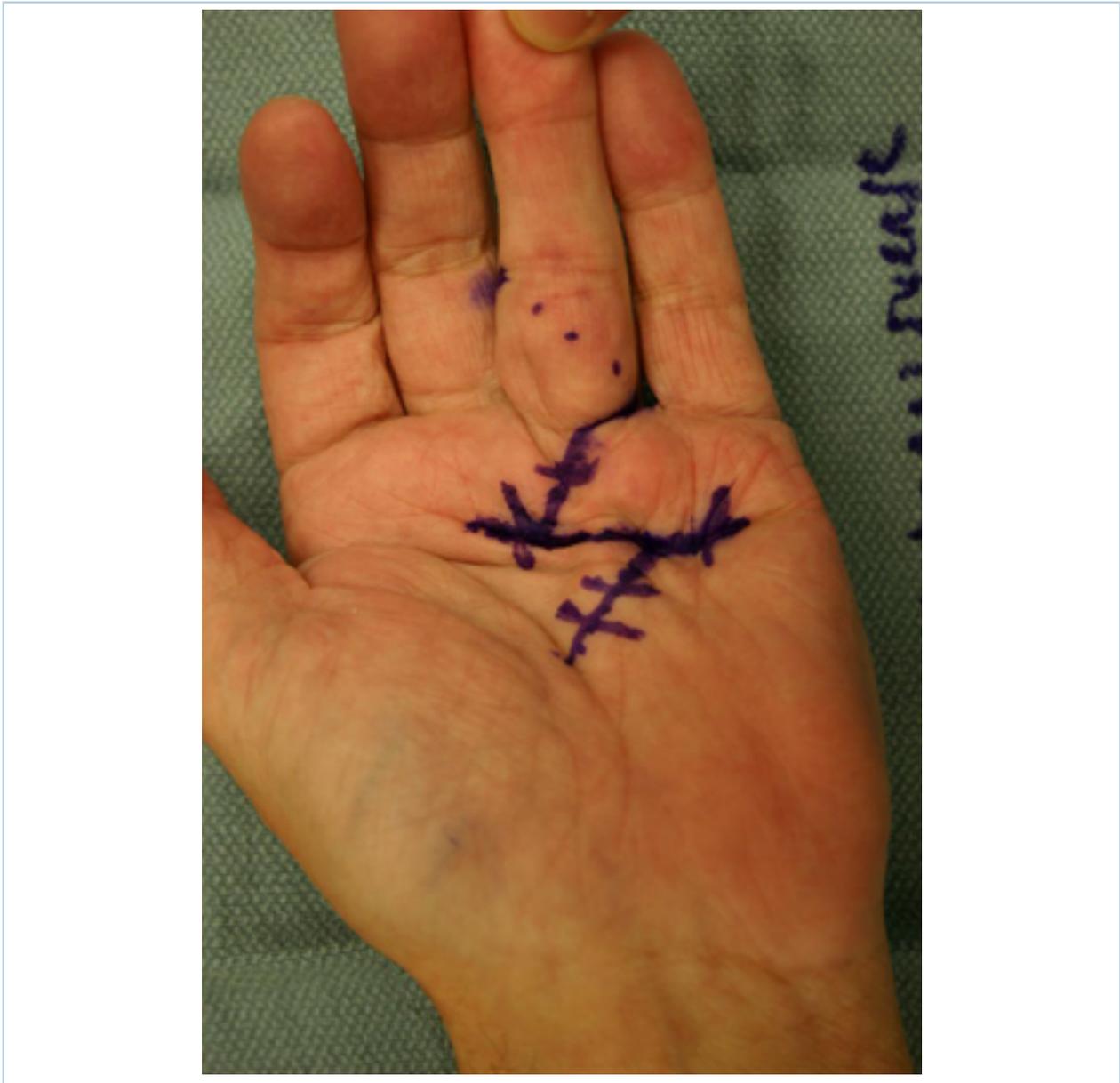
Segmental aponeurotomy is a compromise between percutaneous techniques and open fasciectomy, in which multiple small incisions are made in the palm and the digits to remove segments of the Dupuytren cord and achieve discontinuity between the segments of diseased tissue, with no effort made to remove all of the pathological tissue. It is typically used in patients with MCP joint contractures >30 degrees, or in those with PIP joint contractures. The clinical results of this technique compare fairly well with traditional open fasciectomy techniques, with a recurrence rate ranging from 20% to 35%.[46][47] [48]

Open partial fasciectomy

Although the timing of surgical intervention varies, the decision to operate is generally taken if the MCP joint contracture exceeds 30 degrees or a PIP joint contracture develops.[49] The presence of any PIP joint contracture alone is a relative indication for surgery.[50] Open partial fasciectomy is a successful treatment and is associated with a postoperative recurrence rate of 15%.[49] The risk of postoperative recurrence increases with severity of the contracture, and the longer a deformity is present, the greater the chance of the joint contracture becoming irreversible.[49] [50]

Partial fasciectomy, also known as subtotal, limited, or regional fasciectomy, was first described by Goyrand in 1834 and involves opening up the hand and, under direct visualisation, excising the diseased Dupuytren tissue. Excising only the pathological cord(s) with a partial fasciectomy is now more commonly employed than total fasciectomy or dermatofasciectomy. Postoperative infection is countered with the use of perioperative antibiotics and careful soft tissue handling.

An open partial fasciectomy remains the most common procedure used in the surgical management of Dupuytren's contracture and is the preferred surgical procedure for advanced disease in patients with functional impairment who are surgical candidates. Using this technique, a small amount of potentially diseased fascia could theoretically be left behind, as a radical fasciectomy of the entire palmar fascia is not typically performed. Through a transverse palmar incision, usually overlying the distal palmar crease, the fascia that has formed pathological cords is excised in a proximal-to-distal direction.



Preoperative view of the ring finger of a patient with a flexion contracture with surgical indications, showing the incision marking, demonstrating a transverse incision overlying the distal palmar crease, and oblique Brunner incisions coursing from it proximally and distally

From the collection of Dr C.M. Rodner; used with permission

The choice of digital incision is surgeon dependent and is most often between a Brunner incision (angled 'zigzag' incisions made from the ulnar aspect of the MCP joint crease to the radial aspect of the PIP joint or vice versa) and a longitudinal incision (often combined with multiple Z-plasties). Brunner or longitudinal incisions are carried out along the middle of each affected digit, and skin flaps are carefully raised from the pathological cord to avoid skin necrosis. Brunner 'zigzag' incisions are made methodically in a proximal-to-distal direction from the transverse palmar incision, following the path of the palpable Dupuytren cord.

In the palm the neurovascular structures that lie deep to the involved fascia at this level are identified and, with particular attention paid to retracting these neurovascular bundles, the diseased fascia is excised and elevated in a proximal-to-distal direction.



Intraoperative view of the ring finger of a patient with a flexion contracture, with the radial digital neurovascular bundle identified and isolated coursing volar over the Dupuytren cord, which is being held up by forceps as it is excised in a proximal-to-distal direction

From the collection of Dr C.M. Rodner; used with permission

As the Dupuytren cord moves into the finger, great care is taken to identify the radial and ulnar digital arteries and nerves, as they bifurcate from the parent common artery and nerve. As the proper digital neurovascular bundles may be displaced towards the midline in PIP joint contractures, careful dissection is required to identify them in such cases. The fascia must be cut only if it is directly visualised and the artery and nerve are protected. If a spiral cord is present, which is often the case with PIP joint contractures, dissection proceeds from both a proximal-to-distal and a distal-to-proximal direction so as to safely free the neurovascular bundle from the cord that is displacing it centrally.[51] When the small finger is involved, the insertion of the abductor digiti quinti muscle should be identified and excised in order to fully release the digit.

After all of the diseased tissue is excised, the joints are inspected clinically for the persistence of a contracture. Although the MCP joint contracture usually resolves with the excision of the cord alone, PIP joint contractures often do not. Residual contracture of the PIP joint after the fasciectomy is completed is addressed initially with a release of the volar plate, followed by release of the collateral ligaments if necessary.

Currently, most hand surgeons endeavour to dissect the normal skin away from the underlying diseased tissue.^[46] If the palmar skin is adherent to the cord so that it cannot be saved and the defect is small, primary closure is often performed. Such defects may also occur as the result of straightening out a previously contracted digit. As the finger is extended, primary skin coverage in the palm may prove to be difficult. If the palmar defect is too large for primary closure, skin grafting or the McCash 'open palm' technique have both shown good clinical results.^[52] When possible, direct primary closure of the palmar skin is performed over a Penrose drain to prevent haematoma, as this method of closure allows for early motion and good skin sensibility, avoiding the meticulous wound care required with an open wound.



Postoperative view of the ring finger of a patient with a flexion contracture, showing the closed wound over a Penrose drain, which is used to minimise subsequent haematoma formation

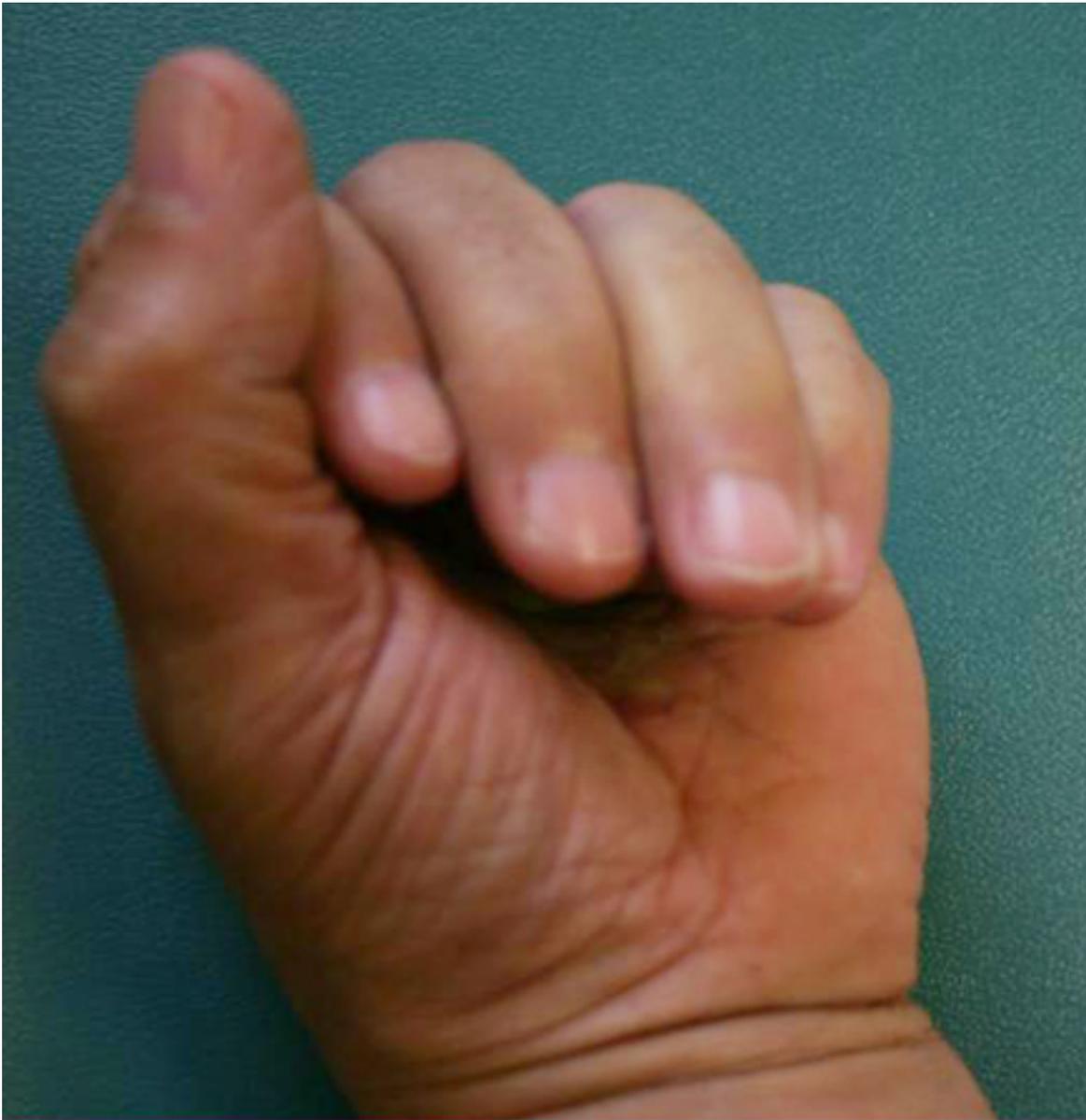
From the collection of Dr C.M. Rodner; used with permission

The McCash 'open palm' technique, used by many surgeons, is an alternative to direct palmar wound closure and is associated with a greater active range of motion without any increased risk of infection, although this technique has the disadvantage of leaving the wound open for about 1 month.^[52]

The fingers are then splinted in full extension, and the patient is followed up within a few days to pull out the drain and assess the wound. By the fifth postoperative day, patients are sent to the hand therapist for a forearm-based digital extension splint that is worn full time between therapy visits. Flexion exercises begin after the wound has stabilised. Regaining digital flexion often proves more difficult than maintaining extension after fasciectomy, due to the postoperative extension splinting required.



*One-month postoperative view of the ring finger of a patient with a flexion contracture, demonstrating full active digital extension
From the collection of Dr C.M. Rodner; used with permission*



*One-month postoperative view of the ring finger of a patient with a flexion contracture, demonstrating active digital flexion
From the collection of Dr C.M. Rodner; used with permission*

By the third postoperative week, the splint is weaned, to be worn at night only, and night-time extension splinting can continue for as long as 6 months. Some surgeons have abandoned postoperative splinting, favouring earlier mobilisation in order to minimise difficulties with flexion. There is also evidence to suggest that splinting (including night-time extension splinting) after surgery provides no additional benefit to standard hand therapy in maintaining finger extension, except perhaps for cases in which extension loss occurs postoperatively, whereby night-time extension splinting may provide some benefit.^{[53] [54]} Postoperative splinting may not be justified in all patients.

Other surgical techniques

Total fasciectomy and dermatofasciectomy are now seldom used but are described here for completeness.

The technique of total (or radical) fasciectomy via a transverse palmar incision and multiple digital Z-plasties in which all of the palmar fascia is excised has not been found to have any clinical advantage over a more limited fasciectomy and is associated with a higher complication rate.^[55] ^[56] Although total fasciectomy should theoretically reduce recurrence rates by excising the entire palmar fascia responsible for Dupuytren's contracture, this has not been supported by the literature.

Dermatofasciectomy, involving a simultaneous excision of the skin, which is subsequently replaced with full-thickness skin grafts, has produced fairly good results, although morbidity associated with skin grafting, such as poor graft sensibility, has made this technique less popular than partial fasciectomy.^[57] ^[58]^[59]^[60]

Radiotherapy

Although its use in Dupuytren's contracture is extremely limited, radiotherapy is described here for completeness, as it has been reported to be successful in the treatment of the disease. One study showed that the majority (77%) of lesions did not progress following a total radiotherapy dose of 30 Gy. Long-term studies have not found differences in disease progression 7 years after radiotherapy, compared with controls.^[61]

Treatment algorithm overview

Please note that formulations/routes and doses may differ between drug names and brands, drug formularies, or locations. Treatment recommendations are specific to patient groups: [see disclaimer](#)

Ongoing		(summary)
no MCP joint or PIP joint contracture		
	1st	expectant management
	adjunct	corticosteroid injections
≤30 degrees MCP joint contracture with no PIP joint contracture		
	1st	collagenase injection
	1st	needle aponeurotomy
	1st	percutaneous fasciotomy
	1st	corticosteroid injections
>30 degrees MCP joint contracture and/or PIP joint contracture		
	1st	open partial fasciectomy + perioperative antibiotics
	adjunct	postoperative splinting
	1st	segmental aponeurotomy
	1st	percutaneous fasciotomy
	1st	collagenase injection

Treatment algorithm

Please note that formulations/routes and doses may differ between drug names and brands, drug formularies, or locations. Treatment recommendations are specific to patient groups: [see disclaimer](#)

Ongoing

no MCP joint or PIP joint contracture

1st expectant management

» Patients with early Dupuytren's contracture who have evidence of the disease but have not yet developed contractures may be managed expectantly with regular follow-up every 6 months to evaluate for disease progression. Use of the Hueston table-top test and appropriate staging of Dupuytren's contracture are essential during follow-up sessions.[49] [50]

adjunct corticosteroid injections

Treatment recommended for SOME patients in selected patient group

Primary options

» [triamcinolone acetonide](#): consult specialist for guidance on dose

» Patients with early Dupuytren's contracture who have evidence of the disease but have not yet developed contractures can be treated with corticosteroid injections if they are experiencing bothersome symptoms.

» The injection of Dupuytren nodules with triamcinolone acetonide monthly for up to 5 months or every 6 weeks for 3 injections has been shown to produce significant regression of the disease, with an average of 3.2 injections per nodule required for improvement of function.[28] After corticosteroid injection, fewer patients progress to surgery than would be predicted with expectant management alone.[28]

≤30 degrees MCP joint contracture with no PIP joint contracture

1st collagenase injection

Primary options

» [collagenase clostridium histolyticum](#): 0.58 mg intralesionally as a single dose, maximum 3 doses/cord
May repeat 2 doses at intervals of 4 weeks if the contracture remains.

» Collagenase clostridium histolyticum is used to treat adult patients with a palpable cord

Ongoing

along with metacarpophalangeal (MCP) joint contracture or proximal interphalangeal (PIP) joint contracture, with a corresponding decrease in both fasciotomies and fasciectomyes.[29] A mean of 1.4 injections is required to normalise affected joints, and clinical success is achieved within 29 days.[30] In one study, reductions in contracture to 0 to 5 degrees of full extension, 30 days after the last injection, were achieved in cords injected with collagenase as compared with those injected with placebo.[31] Adverse events are localised to the injection site, generally of mild-to-moderate severity, and transient in nature.[30] The most commonly reported adverse events were localised swelling in the hand, pain, bruising, pruritus, and transient regional lymph-node enlargement and tenderness. Serious adverse events were seen in 2% of collagenase recipients, including tendon ruptures and complex regional pain syndrome.[31] Although there is some literature with small numbers of patients to suggest that recurrence is more common in PIP joint contractures than in MCP joint contractures, additional long-term studies involving larger numbers of patients are warranted to verify the long-term effectiveness and recurrence rates of collagenase treatment, as well as its long-term safety.[32]

» One systematic review has found that having previous surgery did not affect the efficacy and safety of collagenase injections, making this an option in patients with recurrent Dupuytren's contracture.[34]

» After the injection is given, a dressing is placed on the hand. The patient is then instructed to return to the surgery the next day for the finger extension manoeuvre, in which the finger is extended to break the cord. This may be done with or without a local anaesthetic. After the manipulation is successfully completed, there are no restrictions and early movement is encouraged. Although wearing a night-time splint afterwards has not been proven to decrease recurrence, it should be offered to the patient. The patient is followed to assess the need for up to 2 further injections every 4 to 6 weeks.[30]

» Collagenase clostridium histolyticum may not be available in some countries.

1st

needle aponeurotomy

» Under sterile conditions, the area is anaesthetised with lidocaine, and an 18-gauge needle is used to puncture the aponeurotic

Ongoing

band causing the digital contracture. This weakens the contracture until it can be broken by mechanical force, typically with a characteristic snap. The needle is introduced volar (palmar side) to the tendon at various sites, progressing from proximal to distal. Care must be taken not to insert the needle into the tendon, to avoid iatrogenic injury. Use of ultrasound to guide the procedure may reduce the risk of accidental lesions.[62]

» This procedure is usually successful in correcting the contracture, takes very little time to perform (20-30 minutes), requires only local anaesthesia, and is not very painful. In comparison with open surgical procedures, it results in similar resolution, minimal scarring, faster recovery, and can be repeated easily if the contracture recurs.[35] [36] [37] Needle aponeurotomy is thus an attractive option for patients with less aggressive and early disease.[38] Recurrence rates of up to 58% have been reported during 3 to 5 years of follow-up, but long-term outcomes are not well reported.[40]

» All patients with contractures should receive hand therapy post-procedure.

1st percutaneous fasciotomy

» A similar procedure to needle aponeurotomy but uses a scalpel to cut and release the band causing the digital contracture and is thus performed by a hand surgeon in the operating room.[43] [44] The finger is brought into full extension with a characteristic snap.

» This procedure causes less pain and allows a faster recovery than traditional open fasciectomy interventions. However, it is associated with a risk of recurrence of up to 43%, as diseased fascia is unavoidably left behind, and there is risk of flexor tendon or nerve injury.[43][44][45]

» All patients with contractures should receive hand therapy post-procedure.

1st corticosteroid injections

Primary options

» **triamcinolone acetonide**: consult specialist for guidance on dose

» Patients who wish to avoid a more invasive procedure may benefit from corticosteroid injections.

Ongoing

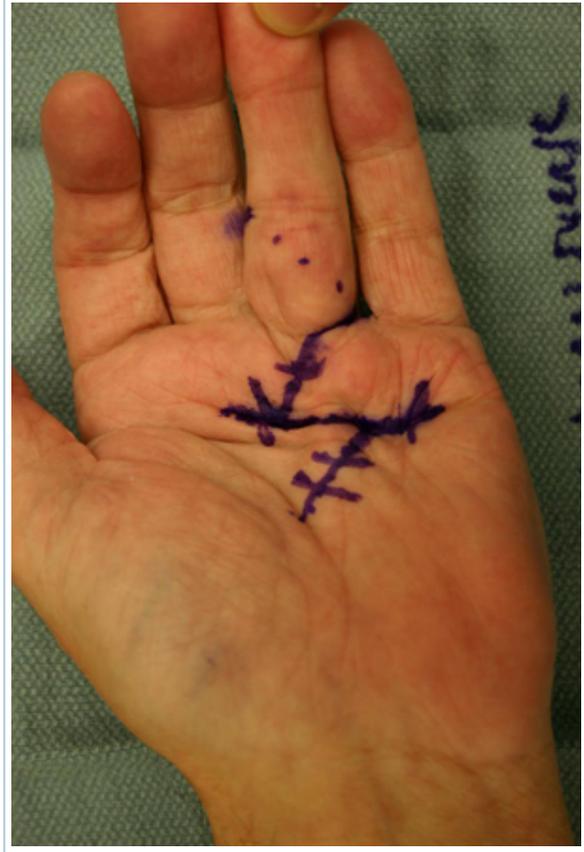
- » The injection of Dupuytren nodules with triamcinolone acetonide monthly for up to 5 months or every 6 weeks for 3 injections has been shown to produce significant regression of the disease, with an average of 3.2 injections per nodule required for improvement of function.^[28] After corticosteroid injection, fewer patients progress to surgery than would be predicted with expectant management alone.^[28]
- » All patients with contractures should receive hand therapy post-procedure.

>30 degrees MCP joint contracture and/or PIP joint contracture

1st open partial fasciectomy + perioperative antibiotics

- » The most common procedure used in the surgical management of Dupuytren's contracture, as it is associated with a postoperative recurrence rate of 15%.^[49]
- » Through a transverse palmar incision overlying the distal palmar crease, the fascia forming pathological cords is excised in a proximal-to-distal direction.

Ongoing



Preoperative view of the ring finger of a patient with a flexion contracture with surgical indications, showing the incision marking, demonstrating a transverse incision overlying the distal palmar crease, and oblique Brunner incisions coursing from it proximally and distally
 From the collection of Dr C.M. Rodner; used with permission

The choice of digital incision is surgeon dependent and is most often between a Brunner and a longitudinal incision (often combined with multiple Z-plasties).

» In the palm, the neurovascular structures deep to the involved fascia are identified and retracted.

Ongoing



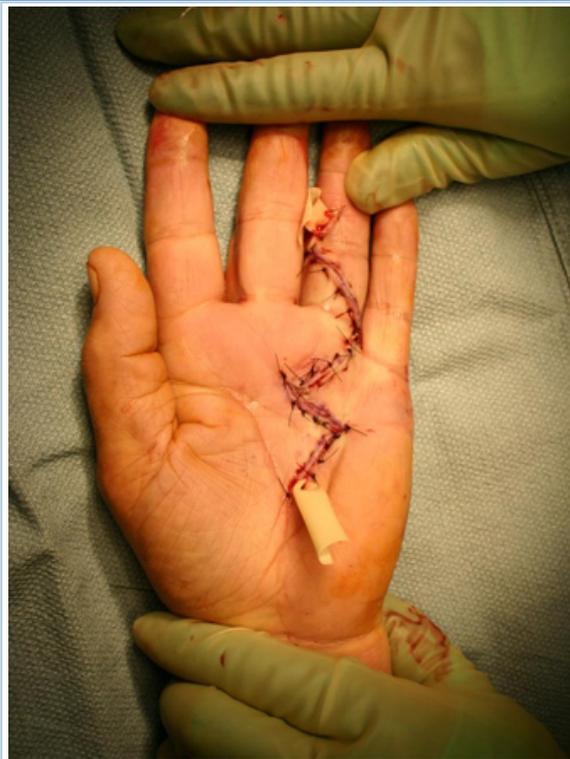
Intraoperative view of the ring finger of a patient with a flexion contracture, with the radial digital neurovascular bundle identified and isolated coursing volar over the Dupuytren cord, which is being held up by forceps as it is excised in a proximal-to-distal direction

From the collection of Dr C.M. Rodner; used with permission

The diseased fascia is excised and elevated in a proximal-to-distal direction. As the Dupuytren cord moves into the finger, care is taken to identify the radial and ulnar digital arteries and nerves, particularly in PIP joint contractures. The joints are inspected for persistent contractures, and residual contracture of the PIP joint is addressed with a release of the volar plate, followed by release of the collateral ligaments if necessary.

» When possible, direct primary closure of the palmar skin is performed over a Penrose drain to prevent haematoma, as this method of closure allows for early motion and good skin sensibility, avoiding the meticulous wound care required with an open wound.

Ongoing



Postoperative view of the ring finger of a patient with a flexion contracture, showing the closed wound over a Penrose drain, which is used to minimise subsequent haematoma formation

From the collection of Dr C.M. Rodner; used with permission

If the palmar defect is too large for primary closure, skin grafting or the McCash 'open palm' technique are used. The patient is followed up within a few days to pull out the drain and assess the wound.

- » All patients with contractures should receive hand therapy post-procedure. After an open partial fasciectomy, flexion exercises begin once the wound has stabilised.
- » Postoperative infection is countered with the use of perioperative antibiotics and careful soft tissue handling. First- or second-generation cephalosporins are usually given. Clindamycin can be used as an alternative in patients with penicillin allergy, to provide gram-positive cover. However, practice varies between centres, and local protocols should be consulted and followed.

adjunct postoperative splinting

Treatment recommended for SOME patients in selected patient group

- » After an open partial fasciectomy, the fingers can be splinted in full extension.

Ongoing

» By the fifth postoperative day, patients are sent to the hand therapist for a forearm-based digital extension splint that is worn full time between therapy visits. Flexion exercises begin once the wound has stabilised. Regaining digital flexion often proves more difficult than maintaining extension after fasciectomy, due to the postoperative extension splinting required.



One-month postoperative view of the ring finger of a patient with a flexion contracture, demonstrating full active digital extension

From the collection of Dr C.M. Rodner; used with permission

Ongoing



*One-month postoperative view of the ring finger of a patient with a flexion contracture, demonstrating active digital flexion
From the collection of Dr C.M. Rodner; used with permission*

By the third postoperative week the splint is weaned, to be worn at night only, and night-time extension splinting can continue for as long as 6 months.

» Some surgeons have abandoned postoperative splinting, favouring earlier mobilisation in order to minimise difficulties with flexion. There is also evidence to suggest that splinting (including night-time extension splinting) after surgery provides no additional benefit to standard hand therapy in maintaining finger extension, except perhaps for cases in which extension loss occurs postoperatively, whereby night-time extension splinting may provide some benefit.^{[53] [54]} Postoperative splinting may not be justified in all patients.

1st segmental aponeurotomy

» A compromise between percutaneous techniques and open fasciectomy, in which multiple small incisions are made in the palm and the digits to remove segments of the Dupuytren cord and achieve discontinuity between the segments of diseased tissue, with no effort made to remove all of the pathological tissue.

» The clinical results of this technique compare quite well with traditional open fasciectomy techniques, with a recurrence rate ranging from 20% to 35%.^{[46][47] [48]}

Ongoing

- » All patients with contractures should receive hand therapy post-procedure.
- 1st** **percutaneous fasciotomy**
- » A similar procedure to needle aponeurotomy but uses a scalpel to cut and release the band causing the digital contracture and is thus performed by a hand surgeon in the operating room.[43] [44] The finger is brought into full extension with a characteristic snap.
- » This procedure causes less pain and allows a faster recovery than traditional open fasciectomy interventions. However, it is associated with a risk of recurrence of up to 43%, as diseased fascia is unavoidably left behind, and there is a risk of flexor tendon or nerve injury.[43][44][45]
- » All patients with contractures should receive hand therapy post-procedure.
- 1st** **collagenase injection**
- Primary options**
- » **collagenase clostridium histolyticum**: 0.58 mg intralesionally as a single dose, maximum 3 doses/cord
May repeat 2 doses at intervals of 4 weeks if the contracture remains.
- » Collagenase clostridium histolyticum is used to treat adult patients with a palpable cord along with metacarpophalangeal (MCP) joint contracture or proximal interphalangeal (PIP) joint contracture, with a corresponding decrease in both fasciotomies and fasciectomyes.[29] A mean of 1.4 injections is required to normalise affected joints, and clinical success is achieved within 29 days.[30] In one study, reductions in contracture to 0 to 5 degrees of full extension, 30 days after the last injection, were achieved in cords injected with collagenase as compared with those injected with placebo.[31] Adverse events are localised to the injection site, generally of mild-to-moderate severity, and transient in nature.[30] The most commonly reported adverse events were localised swelling in the hand, pain, bruising, pruritus, and transient regional lymph-node enlargement and tenderness. Serious adverse events were seen in 2% of collagenase recipients, including tendon ruptures and complex regional pain syndrome.[31] Although there is some literature with small numbers of patients to suggest that recurrence is more common in PIP joint contractures than in MCP joint contractures, additional long-term studies involving larger

Ongoing

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» Collagenase clostridium histolyticum may not be available in some countries.

Emerging

Tumor necrosis factor (TNF)-alpha inhibitors

There is emerging evidence supporting the use of TNF-alpha inhibitors (e.g., adalimumab) for early-stage Dupuytren's disease. One randomised, double-blind, placebo-controlled trial on a UK cohort found that the use of intranodular adalimumab injections were effective in softening and reducing the size of nodules.^[63]

Primary prevention

No primary prevention studies exist, and it remains to be seen whether management of diabetes or reduction in smoking and alcohol will reduce the prevalence of the disease. As occupational trauma is thought to be associated with Dupuytren's disease, proper protection of the hand from trauma at work is indicated.

Patient discussions

Minimisation of the risk factors for Dupuytren's contracture may help slow progression of the disease. Patients should therefore be instructed to stop smoking, reduce their alcohol intake, and use proper protection of the hand from trauma at work if appropriate. Adequate control of diabetes is also important. Patient information is available. [NHS choices: Dupuytren's contracture] (<http://www.nhs.uk/conditions/Dupuytren's-contracture/Pages/Introduction.aspx>)

Monitoring

Monitoring

Follow-up every 6 months is useful to evaluate for disease progression. Use of the Hueston table-top test and appropriate staging of Dupuytren's contracture are essential during follow-up sessions.^{[49] [50]}

The risk of postoperative recurrence increases with severity of the contracture. The longer a deformity is present, the greater the chance of the joint contracture becoming irreversible.

Complications

Complications	Timeframe	Likelihood
intraoperative neurovascular injury	short term	low
<p>Neurovascular injury can be minimised by meticulous intraoperative dissection, identifying the nerves and arteries, and by cutting only tissue that is well visualised after the neurovascular bundles for a particular digit have been identified and retracted.</p> <p>Digital neurapraxia is sometimes unavoidable when a previously contracted digit is straightened. Occasionally, digital ischaemia may occur after extension of a previously contracted digit, due to arterial stretch and vasospasm. If this occurs intraoperatively, the finger should be passively flexed and treated with warm saline. If this fails to restore blood flow, the local application of lidocaine should be considered, and if this is unsuccessful, intravenous heparin may be administered.</p>		
postoperative haematoma	short term	low
<p>The risk of haematoma formation is minimised by tourniquet deflation when the fasciectomy is completed, by achieving adequate haemostasis prior to wound closure, and/or by closing the wound over a drain.[65]</p>		
postoperative infection	short term	low
<p>Infection is countered with the use of perioperative antibiotics and careful soft tissue handling.</p> <p>Complications associated with wound healing can be minimised by careful flap planning, meticulous elevation of flaps, and haematoma prevention. Reported incidence of postoperative surgical site infection is 0.04% at 90 days.[66]</p>		
postoperative stiffness	variable	high
<p>Prophylaxis is difficult due to the need for a prolonged period of postoperative immobilisation after Dupuytren surgery.</p> <p>Supervised hand therapy exercises should be instituted as soon as the state of the wound allows exercising of the hand.</p>		
post-procedural recurrence	variable	medium
<p>The post-procedural recurrences reported may represent both true recurrence (disease at the operative site) and disease extension (disease outside the prior surgical area).</p> <p>Recurrence is significantly more common in patients who present with proximal interphalangeal (PIP) joint contractures, a diseased little finger, or multiple affected digits, and the risk of recurrence increases with severity of the contracture.[50]</p> <p>Due to the fact that diseased fascia is left behind, needle aponeurotomy is associated with a recurrence rate of at least 50% and percutaneous fasciotomy is associated with a recurrence rate of at least 43%, whereas open partial fasciectomy is associated with a postoperative recurrence rate of 15%, and segmental aponeurotomy is associated with a recurrence rate of 20% to 35%.[46][47] [48]</p>		

Complications	Timeframe	Likelihood
postoperative reflex sympathetic dystrophy	variable	low
<p>Early warning signs of reflex sympathetic dystrophy, such as excessive postoperative pain, should be responded to with diligence and a high index of suspicion.</p> <p>Postoperative dressings should be loosened or changed in the presence of severe hand or digit swelling.</p>		

Prognosis

Dupuytren's contracture is progressive, with 75% of patients developing features of more advanced stages of the disease. Men seem to progress more rapidly than women, and patients aged <50 years tend to progress more rapidly than older patients. Although most patients will progress in severity, about 10% will regress. Many patients with nodules or cords do not progress to contracture, as evidenced by a Scandinavian study where only 35% of patients developed contracture during an 18-year follow-up period.^[64] Smoking and alcohol use increase the likelihood of progression to surgery. There is currently no proven preventative treatment that can interrupt the progression of contracture.^[22]

Treatment guidelines

United Kingdom

Radiation therapy for early Dupuytren's disease (<https://www.nice.org.uk/guidance/ipg573>)

Published by: National Institute for Health and Care Excellence

Last published: 2016

Europe

Dutch multidisciplinary guideline on Dupuytren disease (<https://www.jhsgo.org>)

Published by: Netherlands Society of Plastic Surgery

Last published: 2022

Online resources

1. NHS choices: Dupuytren's contracture (<http://www.nhs.uk/conditions/Dupuytren's-contracture/Pages/Introduction.aspx>) (*external link*)
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Key articles

- Karbowski M, Holme T, Khan K, et al. Dupuytren's disease. *BMJ*. 2021 Jun 4;373:n1308. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/34088710?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/34088710?tool=bestpractice.bmj.com)
- Boe C, Blazar P, Iannuzzi N. Dupuytren contractures: an update of recent literature. *J Hand Surg Am*. 2021 Oct;46(10):896-906. [Full text \(https://www.jhandsurg.org/article/S0363-5023\(21\)00427-5/fulltext\)](https://www.jhandsurg.org/article/S0363-5023(21)00427-5/fulltext) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/34452797?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/34452797?tool=bestpractice.bmj.com)
- Ketchum LD, Donahue TK. The injection of nodules of Dupuytren's disease with triamcinolone acetonide. *J Hand Surg Am*. 2000 Nov;25(6):1157-62. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/11119679?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/11119679?tool=bestpractice.bmj.com)
- Badalamente MA, Hurst LC. Efficacy and safety of injectable mixed collagenase subtypes in the treatment of Dupuytren's contracture. *J Hand Surg Am*. 2007 Jul-Aug;32(6):767-74. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/17606053?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/17606053?tool=bestpractice.bmj.com)
- Hurst LC, Badalamente MA, Hentz VR, et al. Injectable collagenase clostridium histolyticum for Dupuytren's contracture. *N Engl J Med*. 2009 Sep 3;361(10):968-79. [Full text \(https://www.nejm.org/doi/full/10.1056/NEJMoa0810866\)](https://www.nejm.org/doi/full/10.1056/NEJMoa0810866) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/19726771?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/19726771?tool=bestpractice.bmj.com)

References

1. Hu FZ, Nystrom A, Ahmed A, et al. Mapping of an autosomal dominant gene for Dupuytren's contracture to chromosome 16q in a Swedish family. *Clin Genet*. 2005 Nov;68(5):424-9. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/16207209?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/16207209?tool=bestpractice.bmj.com)
2. Hart MG, Hooper G. Clinical associations of Dupuytren's disease. *Postgrad Med J*. 2005 Jul;81(957):425-8. [Full text \(https://academic.oup.com/pmj/article/81/957/425/7032223\)](https://academic.oup.com/pmj/article/81/957/425/7032223) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/15998816?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/15998816?tool=bestpractice.bmj.com)
3. Tubiana R. Evaluation of deformities in Dupuytren's disease. *Ann Chir Main*. 1986;5:5-11. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/3963905?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/3963905?tool=bestpractice.bmj.com)
4. Salari N, Heydari M, Hassanabadi M, et al. The worldwide prevalence of the Dupuytren disease: a comprehensive systematic review and meta-analysis. *J Orthop Surg Res*. 2020 Oct 28;15(1):495. [Full text \(https://josr-online.biomedcentral.com/articles/10.1186/s13018-020-01999-7\)](https://josr-online.biomedcentral.com/articles/10.1186/s13018-020-01999-7) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/33115483?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/33115483?tool=bestpractice.bmj.com)
5. Ross DC. Epidemiology of Dupuytren's disease. *Hand Clin*. 1999;15:53-62. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/10050242?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/10050242?tool=bestpractice.bmj.com)
6. Lanting R, Broekstra DC, Werker PMN, et al. A systematic review and meta-analysis on the prevalence of Dupuytren disease in the general population of Western countries. *Plast Reconstr*

- Surg. 2014 Mar;133(3):593-603. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/24263394?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/24263394?tool=bestpractice.bmj.com)
-
7. Lanting R, van den Heuvel ER, Westerink B, et al. Prevalence of Dupuytren disease in The Netherlands. *Plast Reconstr Surg*. 2013 Aug;132(2):394-403. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/23897337?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/23897337?tool=bestpractice.bmj.com)
-
8. Nordenskjöld J, Englund M, Zhou C, et al. Prevalence and incidence of doctor-diagnosed Dupuytren's disease: a population-based study. *J Hand Surg Eur Vol*. 2017 Sep;42(7):673-7. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/28093015?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/28093015?tool=bestpractice.bmj.com)
-
9. Broekstra DC, Kuo RYL, Burn E, et al. Dupuytren disease: prevalence, incidence, and lifetime risk of surgical intervention. A population-based cohort analysis. *Plast Reconstr Surg*. 2023 Mar 1;151(3):581-91. [Full text \(https://journals.lww.com/plasreconsurg/Fulltext/2023/03000/Dupuytren_Disease__Prevalence,_Incidence,_and.24.aspx\)](https://journals.lww.com/plasreconsurg/Fulltext/2023/03000/Dupuytren_Disease__Prevalence,_Incidence,_and.24.aspx) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/36730480?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/36730480?tool=bestpractice.bmj.com)
-
10. Tajika T, Kobayashi T, Kaneko T, et al. Epidemiological study for personal risk factors and quality of life related to Dupuytren's disease in a mountain village of Japan. *J Orthop Sci*. 2014 Jan;19(1):64-70. [Full text \(https://www.doi.org/10.1007/s00776-013-0478-y\)](https://www.doi.org/10.1007/s00776-013-0478-y) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/24129389?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/24129389?tool=bestpractice.bmj.com)
-
11. Lee LC, Zhang AY, Chong AK, et al. Expression of a novel gene, MafB, in Dupuytren's disease. *J Hand Surg Am*. 2006 Feb;31(2):211-8. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/16473681?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/16473681?tool=bestpractice.bmj.com)
-
12. Bayat A, Walter J, Lambe H, et al. Identification of a novel mitochondrial mutation in Dupuytren's disease using multiplex DHPLC. *Plast Reconstr Surg*. 2005 Jan;115(1):134-41. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/15622243?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/15622243?tool=bestpractice.bmj.com)
-
13. Thurston AJ. Dupuytren's disease. *J Bone Joint Surg Br*. 2003 May;85(4):469-77. [Full text \(https://boneandjoint.org.uk/Article/10.1302/0301-620X.85B4.14215\)](https://boneandjoint.org.uk/Article/10.1302/0301-620X.85B4.14215) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/12793547?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/12793547?tool=bestpractice.bmj.com)
-
14. Chammas M, Bousquet P, Renard E, et al. Dupuytren's disease, carpal tunnel syndrome, trigger finger, and diabetes mellitus. *J Hand Surg Am*. 1995 Jan;20(1):109-14. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/7722249?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/7722249?tool=bestpractice.bmj.com)
-
15. Geoghegan JM, Forbes J, Clark DI, et al. Dupuytren's disease risk factors. *J Hand Surg Br*. 2004 Oct;29(5):423-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/15336742?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/15336742?tool=bestpractice.bmj.com)
-
16. Rydberg M, Zimmerman M, Löfgren JP, et al. Metabolic factors and the risk of Dupuytren's disease: data from 30,000 individuals followed for over 20 years. *Sci Rep*. 2021 Jul 19;11(1):14669. [Full text \(https://www.doi.org/10.1038/s41598-021-94025-7\)](https://www.doi.org/10.1038/s41598-021-94025-7) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/34282190?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/34282190?tool=bestpractice.bmj.com)
-

17. Godtfredsen NS, Lucht H, Prescott E, et al. A prospective study linked both alcohol and tobacco to Dupuytren's disease. *J Clin Epidemiol*. 2004 Aug;57(8):858-63. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/15485739?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/15485739?tool=bestpractice.bmj.com)
18. Loos B, Puschkin V, Horch RE. 50 years experience with Dupuytren's contracture in the Erlangen University Hospital - a retrospective analysis of 2919 operated hands from 1956 to 2006. *BMC Musculoskelet Disord*. 2007;8:60. [Full text \(http://www.biomedcentral.com/1471-2474/8/60\)](http://www.biomedcentral.com/1471-2474/8/60) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/17610744?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/17610744?tool=bestpractice.bmj.com)
19. Karbowski M, Holme T, Khan K, et al. Dupuytren's disease. *BMJ*. 2021 Jun 4;373:n1308. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/34088710?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/34088710?tool=bestpractice.bmj.com)
20. Arafa M, Noble J, Royle SG, et al. Dupuytren's and epilepsy revisited. *J Hand Surg Br*. 1992 Apr;17(2):221-4. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/1588209?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/1588209?tool=bestpractice.bmj.com)
21. Bisson MA, McGrouther DA, Mudera V, et al. The different characteristics of Dupuytren's disease fibroblasts derived from either nodule or cord: expression of alpha-smooth muscle actin and the response to stimulation by TGF-beta1. *J Hand Surg Br*. 2003 Aug;28(4):351-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/12849947?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/12849947?tool=bestpractice.bmj.com)
22. Boe C, Blazar P, Iannuzzi N. Dupuytren contractures: an update of recent literature. *J Hand Surg Am*. 2021 Oct;46(10):896-906. [Full text \(https://www.jhandsurg.org/article/S0363-5023\(21\)00427-5/fulltext\)](https://www.jhandsurg.org/article/S0363-5023(21)00427-5/fulltext) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/34452797?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/34452797?tool=bestpractice.bmj.com)
23. Freehafer AA, Strong JM. The treatment of Dupuytren's contracture by partial fasciectomy. *J Bone Joint Surg Am*. 1963;45:1207-16. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/14077984?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/14077984?tool=bestpractice.bmj.com)
24. Ball C, Pratt AL, Nanchahal J. Optimal functional outcome measures for assessing treatment for Dupuytren's disease: a systematic review and recommendations for future practice. *BMC Musculoskelet Disord*. 2013;14:131. [Full text \(http://www.biomedcentral.com/1471-2474/14/131\)](http://www.biomedcentral.com/1471-2474/14/131) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/23575442?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/23575442?tool=bestpractice.bmj.com)
25. Rodrigues JN, Becker GW, Ball C, et al. Surgery for Dupuytren's contracture of the fingers. *Cochrane Database Syst Rev*. 2015;(12):CD010143. [Full text \(http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD010143.pub2/full\)](http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD010143.pub2/full) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/26648251?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/26648251?tool=bestpractice.bmj.com)
26. Becker GW, Davis TR. The outcome of surgical treatments for primary Dupuytren's disease--a systematic review. *J Hand Surg Eur Vol*. 2010 Oct;35(8):623-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/20621942?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/20621942?tool=bestpractice.bmj.com)
27. Harvey LA, Katalinic OM, Herbert RD, et al. Stretch for the treatment and prevention of contractures. *Cochrane Database Syst Rev*. 2017;(1):CD007455. [Full text \(http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD007455.pub3/full\)](http://onlinelibrary.wiley.com/doi/10.1002/14651858.CD007455.pub3/full) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/28146605?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/28146605?tool=bestpractice.bmj.com)

28. Ketchum LD, Donahue TK. The injection of nodules of Dupuytren's disease with triamcinolone acetonide. *J Hand Surg Am.* 2000 Nov;25(6):1157-62. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/11119679?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/11119679?tool=bestpractice.bmj.com)

29. Lipman MD, Carstensen SE, Deal DN. Trends in the treatment of Dupuytren disease in the United States between 2007 and 2014. *Hand (N Y).* 2017 Jan;12(1):13-20. [Full text \(https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5207289\)](https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5207289) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/28082837?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/28082837?tool=bestpractice.bmj.com)

30. Badalamente MA, Hurst LC. Efficacy and safety of injectable mixed collagenase subtypes in the treatment of Dupuytren's contracture. *J Hand Surg Am.* 2007 Jul-Aug;32(6):767-74. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/17606053?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/17606053?tool=bestpractice.bmj.com)

31. Hurst LC, Badalamente MA, Hentz VR, et al. Injectable collagenase clostridium histolyticum for Dupuytren's contracture. *N Engl J Med.* 2009 Sep 3;361(10):968-79. [Full text \(https://www.nejm.org/doi/full/10.1056/NEJMoa0810866\)](https://www.nejm.org/doi/full/10.1056/NEJMoa0810866) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/19726771?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/19726771?tool=bestpractice.bmj.com)

32. Watt AJ, Curtin CM, Hentz VR. Collagenase injection as nonsurgical treatment of Dupuytren's disease: 8-year follow-up. *J Hand Surg Am.* 2010 Apr;35(4):534-9. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/20353858?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/20353858?tool=bestpractice.bmj.com)

33. Kemler MA, de Wijn RS, van Rijssen AL, et al. Dutch multidisciplinary guideline on Dupuytren disease. *J Hand Surg Glob Online.* 2023 Mar;5(2):178-83. [Full text \(https://www.jhsgo.org/article/S2589-5141\(22\)00177-3/fulltext\)](https://www.jhsgo.org/article/S2589-5141(22)00177-3/fulltext) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/36974283?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/36974283?tool=bestpractice.bmj.com)

34. Bainbridge C, Gerber RA, Szczypa PP, et al. Efficacy of collagenase in patients who did and did not have previous hand surgery for Dupuytren's contracture. *J Plast Surg Hand Surg.* 2012 Sep;46(3-4):177-83. [Full text \(http://informahealthcare.com/doi/pdf/10.3109/2000656X.2012.683795\)](http://informahealthcare.com/doi/pdf/10.3109/2000656X.2012.683795) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/22670890?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/22670890?tool=bestpractice.bmj.com)

35. Toppi JT, Trompf L, Smoll NR, et al. Dupuytren's contracture: an analysis of outcomes of percutaneous needle fasciotomy versus open fasciectomy. *ANZ J Surg.* 2015 Sep;85(9):639-43. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/24438029?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/24438029?tool=bestpractice.bmj.com)

36. Zhou C, Selles RW, Slijper HP, et al. Comparative effectiveness of percutaneous needle aponeurotomy and limited fasciectomy for Dupuytren's contracture: a multicenter observational study. *Plast Reconstr Surg.* 2016 Oct;138(4):837-46. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/27307334?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/27307334?tool=bestpractice.bmj.com)

37. van Rijssen AL, Gerbrandy FS, Ter Linden H, et al. A comparison of the direct outcomes of percutaneous needle fasciotomy and limited fasciectomy for Dupuytren's disease: a 6-week follow-up study. *J Hand Surg Am.* 2006 May-Jun;31(5):717-25. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/16713831?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/16713831?tool=bestpractice.bmj.com)

38. Foucher G, Medina J, Navarro R. Percutaneous needle aponeurotomy: complications and results. *J Hand Surg Br.* 2003 Oct;28(5):427-31. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/12954251?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/12954251?tool=bestpractice.bmj.com)
39. Soreide E, Murad MH, Denbeigh JM, et al. Treatment of Dupuytren's contracture: a systematic review. *Bone Joint J.* 2018 Sep;100-B(9):1138-45. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/30168768?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/30168768?tool=bestpractice.bmj.com)
40. Chen NC, Srinivasan RC, Shauver MJ, et al. A systematic review of outcomes of fasciotomy, aponeurotomy, and collagenase treatments for Dupuytren's contracture. *Hand (N Y).* 2011 Sep;6(3):250-5. [Full text \(https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3153627\)](https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3153627) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/22942847?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/22942847?tool=bestpractice.bmj.com)
41. Strömberg J, Ibsen-Sörensen A, Fridén J. Comparison of treatment outcome after collagenase and needle fasciotomy for Dupuytren contracture: a randomized, single-blinded, clinical trial with a 1-year follow-up. *J Hand Surg Am.* 2016 Sep;41(9):873-80. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/27473921?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/27473921?tool=bestpractice.bmj.com)
42. Skov ST, Bisgaard T, Søndergaard P, et al. Injectable collagenase versus percutaneous needle fasciotomy for Dupuytren contracture in proximal interphalangeal joints: a randomized controlled trial. *J Hand Surg Am.* 2017 May;42(5):321-8. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/28473158?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/28473158?tool=bestpractice.bmj.com)
43. Rowley DI, Couch M, Chesney RB, et al. Assessment of percutaneous fasciotomy in the management of Dupuytren's contracture. *J Hand Surg Br.* 1984 Jun;9(2):163-4. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/6747419?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/6747419?tool=bestpractice.bmj.com)
44. Colville J. Dupuytren's contracture - the role of fasciotomy. *Hand.* 1983 Jun;15(2):162-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/6884846?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/6884846?tool=bestpractice.bmj.com)
45. Bryan AS, Ghorbal MS. The long-term results of closed palmar fasciotomy in the management of Dupuytren's contracture. *J Hand Surg Br.* 1988 Aug;13(3):254-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/3171286?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/3171286?tool=bestpractice.bmj.com)
46. Rayan GM. Dupuytren disease: Anatomy, pathology, presentation, and treatment. *J Bone Joint Surg Am.* 2007 Jan;89(1):189-98. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/17256226?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/17256226?tool=bestpractice.bmj.com)
47. Moermans JP. Long-term results after segmental aponeurectomy for Dupuytren's disease. *J Hand Surg Br.* 1996;21:797-800. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/8982932?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/8982932?tool=bestpractice.bmj.com)
48. Moermans JP. Segmental aponeurectomy in Dupuytren's disease. *J Hand Surg Br.* 1991 Aug;16(3):243-54. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/1960487?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/1960487?tool=bestpractice.bmj.com)
49. Townley WA, Baker R, Sheppard N, et al. Dupuytren's contracture unfolded. *BMJ.* 2006;332:397-400. [Full text \(http://www.ncbi.nlm.nih.gov/pmc/articles/PMC1370973\)](http://www.ncbi.nlm.nih.gov/pmc/articles/PMC1370973) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/16484265?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/16484265?tool=bestpractice.bmj.com)

50. Au-Yong IT, Wildin CJ, Dias JJ, et al. A review of common practice in Dupuytren surgery. *Tech Hand Up Extrem Surg.* 2005 Dec;9(4):178-87. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/16340578?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/16340578?tool=bestpractice.bmj.com)
51. Umlas ME, Bischoff RJ, Gelberman RH. Predictors of neurovascular displacement in hands with Dupuytren's contracture. *J Hand Surg Br.* 1994 Oct;19(5):664-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/7822934?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/7822934?tool=bestpractice.bmj.com)
52. McCash CR. The open palm technique in Dupuytren's contracture. *Br J Plast Surg.* 1964;17:271-80 [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/14191131?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/14191131?tool=bestpractice.bmj.com)
53. Kemler MA, Houtp P, van der Horst CM. A pilot study assessing the effectiveness of postoperative splinting after limited fasciectomy for Dupuytren's disease. *J Hand Surg Eur Vol.* 2012 Oct;37(8):733-7. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/22311918?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/22311918?tool=bestpractice.bmj.com)
54. Collis J, Collocott S, Hing W, et al. The effect of night extension orthoses following surgical release of Dupuytren contracture: a single-center, randomized, controlled trial. *J Hand Surg Am.* 2013 Jul;38(7):1285-94. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/23790420?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/23790420?tool=bestpractice.bmj.com)
55. McIndoe A, Beare RL. The surgical management of Dupuytren's contracture. *Am J Surg.* 1958;95:197-203. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/13487940?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/13487940?tool=bestpractice.bmj.com)
56. Zachariae L. Extensive versus limited fasciectomy for Dupuytren's contracture. *Scand J Plast Reconstr Surg.* 1967;1(2):150-3. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/5605144?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/5605144?tool=bestpractice.bmj.com)
57. Abe Y, Rokkaku T, Kuniyoshi K, et al. Clinical results of dermofasciectomy for Dupuytren's disease in Japanese patients. *J Hand Surg Eur Vol.* 2007 Aug;32(4):407-10. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/17287058?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/17287058?tool=bestpractice.bmj.com)
58. Searle AE, Logan AM. A mid-term review of the results of dermofasciectomy for Dupuytren's disease. *Ann Chir Main Memb Super.* 1992;11(5):375-80. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/1284018?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/1284018?tool=bestpractice.bmj.com)
59. Ketchum LD, Hixson FP. Dermofasciectomy and full-thickness grafts in the treatment of Dupuytren's contracture. *J Hand Surg Am.* 1987 Sep;12:659-64. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/3309018?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/3309018?tool=bestpractice.bmj.com)
60. Kelly C, Varian J. Dermofasciectomy: a long term review. *Ann Chir Main Memb Super.* 1992;11(5):381-2. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/1284019?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/1284019?tool=bestpractice.bmj.com)
61. Weinzierl G, Flügel M, Geldmacher J. Lack of effectiveness of alternative non-surgical treatment procedures of Dupuytren contracture. *Chirurg.* 1993 Jun;64(6):492-4. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/8359061?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/8359061?tool=bestpractice.bmj.com)

62. Sakellariou VI, Brault J, Rizzo M. Ultrasound-assisted percutaneous needle fasciotomy for Dupuytren's contracture. *Orthopedics*. 2015 May;38(5):299-303. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/25970356?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/25970356?tool=bestpractice.bmj.com)

63. Nanchahal J, Ball C, Rombach I, et al. Anti-tumour necrosis factor therapy for early-stage Dupuytren's disease (RIDD): a phase 2b, randomised, double-blind, placebo-controlled trial. *Lancet Rheumatol*. 2022 Jun;4(6):E407-16. [Full text \(https://www.thelancet.com/journals/lanrhe/article/PIIS2665-9913\(22\)00093-5/fulltext\)](https://www.thelancet.com/journals/lanrhe/article/PIIS2665-9913(22)00093-5/fulltext) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/35949922?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/35949922?tool=bestpractice.bmj.com)

64. Gudmundsson KG, Arngrimsson R, Jónsson T. Eighteen years follow-up study of the clinical manifestations and progression of Dupuytren's disease. *Scand J Rheumatol*. 2001;30(1):31-4. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/11252689?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/11252689?tool=bestpractice.bmj.com)

65. Boyer MI, Gelberman RH. Complications of the operative treatment of Dupuytren's disease. *Hand Clin*. 1999 Feb;15(1):161-6. [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/10050251?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/10050251?tool=bestpractice.bmj.com)

66. Alser O, Craig RS, Lane JCE, et al. Serious complications and risk of re-operation after Dupuytren's disease surgery: a population-based cohort study of 121,488 patients in England. *Sci Rep*. 2020 Oct 5;10(1):16520. [Full text \(https://www.nature.com/articles/s41598-020-73595-y\)](https://www.nature.com/articles/s41598-020-73595-y) [Abstract \(http://www.ncbi.nlm.nih.gov/pubmed/33020582?tool=bestpractice.bmj.com\)](http://www.ncbi.nlm.nih.gov/pubmed/33020582?tool=bestpractice.bmj.com)

Images



Figure 1: Preoperative view of a small finger flexion contracture with surgical indications

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Figure 2: Preoperative view of a small finger flexion contracture with surgical indications

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Figure 3: Preoperative view of the ring finger of a patient with a flexion contracture with surgical indications, showing the incision marking, demonstrating a transverse incision overlying the distal palmar crease, and oblique Brunner incisions coursing from it proximally and distally

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Figure 4: Intraoperative view of the ring finger of a patient with a flexion contracture, with the radial digital neurovascular bundle identified and isolated coursing volar over the Dupuytren cord, which is being held up by forceps as it is excised in a proximal-to-distal direction

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Figure 5: Postoperative view of the ring finger of a patient with a flexion contracture, showing the closed wound over a Penrose drain, which is used to minimise subsequent haematoma formation

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Figure 6: One-month postoperative view of the ring finger of a patient with a flexion contracture, demonstrating full active digital extension

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Figure 7: One-month postoperative view of the ring finger of a patient with a flexion contracture, demonstrating active digital flexion

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This approach is in line with the guidance of the [International Bureau of Weights and Measures Service](#).

Figure 1 – BMJ Best Practice Numeral Style

5-digit numerals: 10,000

4-digit numerals: 1000

numerals < 1: 0.25

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// Acknowledgements:

Dr Craig M. Rodner would like to gratefully acknowledge Dr Thomas H. Trojian and Dr Daniel M. Avery, both previous contributors to this topic. THT and DMA declare that they have no competing interests.

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