

DUPUYTREN'S CONTRACTURE RELEASE SURGERY IN ADULTS PRIOR APPROVAL (PA) POLICY

Version:	2223.v4a
Recommendation by:	NHS Somerset ICB Clinical Commissioning Policy Forum (CCPF)
Date Ratified:	July 2022
Name of Originator/Author:	EBI Service
Approved by Responsible Committee/Individual:	NHS Somerset ICB Clinical Executive Committee (CEC)
Publication/issue date:	September 2022
Review date:	Earliest of either NICE publication or 3 years from issue
Target audience:	<p>NHS Somerset ICB:</p> <ul style="list-style-type: none"> • NHS Providers • GP Practices • Contracts Team <p>Medical Directors:</p> <ul style="list-style-type: none"> • Somerset Foundation Trust • Yeovil District Hospital NHS FT • Royal United Hospitals Bath NHS FT
Application Form	Prior Approval Form

**DUPUYTREN'S CONTRACTURE RELEASE SURGERY
IN ADULTS
PRIOR APPROVAL (PA) POLICY**

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VERSION CONTROL

Document Status:	Current policy
Version:	2223.v4a

DOCUMENT CHANGE HISTORY

Version	Date	Comments
1718.v1	April 2017	Amended to include NICE update on CCH
1718.v2	January 2018	New policy template and PALs email address
1718.V3	April 2019	IFR replaced with EBI name change. 'Regard 'to Section 14Z8 of the NHS Act 2006
1819.v3a	December 2019	Removal of CCH treatment
1920.v4	September 2022	3-year review. Amendment from SCCG to NHS Somerset ICB. New PALS email address

Equality Impact Assessment EIA	20 February 2017 1718.v1
Quality Impact Assessment QIA	20180302 v1
Sponsoring Director:	Dr A Murray
Document Reference:	2223.v4a

1 GENERAL PRINCIPLES (PRIOR APPROVAL)

- 1.1 Funding approval must be secured by primary care/secondary/community care prior to referring/treating patients for this prior approval treatment
- 1.2 Funding approval must be secured prior to a referral for an assessment/surgery. Referring patients without funding approval secured not only incurs significant costs in out-patient appointments for patients that may not qualify for surgery, but inappropriately raises the patient's expectation of treatment
- 1.3 On limited occasions, we may approve funding for an assessment only in order to confirm or obtain evidence demonstrating whether a patient meets the criteria for funding. In such cases, patients should be made aware that the assessment does not mean that they will be provided with surgery and surgery will only be provided where it can be demonstrated that the patients meet the criteria to access treatment in this policy
- 1.4 Funding approval will only be given where there is evidence that the treatment requested is effective and the patient has the potential to benefit from the proposed treatment. Where it is demonstrated that patients have previously been provided with the treatment with limited or diminishing benefit, funding approval is unlikely to be agreed
- 1.5 Receiving funding approval does not confirm that they will receive treatment or surgery for a condition as a consent discussion will need to be undertaken with a clinician prior to treatment
- 1.6 The policy does not apply to patients with suspected malignancy who should continue to be referred under 2 week wait pathway rules for assessment and testing as appropriate
- 1.7 Patients with an elevated BMI of 30 or more may experience more post-surgical complications including post-surgical wound infection so should be encouraged to lose weight further prior to seeking surgery.
<https://www.sciencedirect.com/science/article/pii/S1198743X15007193>
(Thelwall, 2015)
- 1.8 Patients who are smokers should be referred to smoking cessation services in order to reduce the risk of surgery and improve healing
- 1.9 Where prior approval funding is secured by the EBI service it will be available for a specified period of time, normally one year

2 POLICY CRITERIA PRIOR APPROVAL

- 2.1 The ICB does **NOT** commission Radiation Therapy or Collagenase Clostridium Histolyticum (CCH) for Dupuytren's contracture

- 2.2 The ICB commissions surgical intervention for Dupuytren's contracture where the following criteria have been met;
- 2.2.1 The patient has a 30 degree or greater fixed flexion deformity (contracture) at either the:
- a) Metacarpophalangeal joint **OR**
 - b) Proximal interphalangeal joint **OR**

- 2.2.2 Severe thumb contractures which interfere with function **OR**

In adults with a palpable cord and where ALL of the following apply:

- 2.3 There is evidence of moderate disease (functional problems and metacarpophalangeal joint contracture of 30° to 60° and proximal interphalangeal joint contracture of less than 30° or first web contracture) plus up to 2 affected joints
- 2.4 Patients who are not eligible for treatment under this policy, please refer to section 3 EVIDENCE BASED INTERVENTIONS APPLICATION PROCESS on how to apply for funding with evidence of clinical exceptionalty

3 EVIDENCE BASED INTERVENTIONS APPLICATION PROCESS

- 3.1 Patients who are not eligible for treatment under this policy may be considered on an individual basis where their GP or Consultant believes clinical exceptional circumstances exist that warrant deviation from the rule of this policy
- 3.2 Completion of a **Generic EBI Application Form** by a patient's GP or Consultant is required
- 3.3 Applications cannot be considered from patients personally
- 3.4 Only electronically completed EBI applications will be accepted to the EBI Service
- 3.5 It is expected that clinicians will have ensured that the patient, on behalf of who they are forwarding the application for, is appropriately informed about the existing policies prior to an application to the EBI Panel. This will reassure the Panel that the patient has a reasonable expectation of the outcome of the application and its context
- 3.6 EBI applications are reviewed and considered against clinical exceptionalty

For further information on 'clinical exceptionalty' please refer to the NHS England information using the link below page 9-13;

- <https://www.england.nhs.uk/wp-content/uploads/2017/11/comm-policy-individual-funding-requests.pdf>

Social, Emotional and Environmental factors *i.e., income, housing, environmental pollution, access to services, family, friends, ethnicity, life experiences etc.* CANNOT be considered with an application

3.7 Where appropriate photographic supporting evidence can be forwarded with the application form

3.8 An application put forward for consideration must demonstrate some unusual or unique clinical factor about the patient that suggests they are exceptional as defined below:

- Significantly different to the general population of patients with the condition in question
- Likely to gain significantly more benefit from the intervention than might be expected from the average patient with the condition

4 ACCESS TO POLICY

4.1 If you would like further copies of this policy or need it in another format, such as Braille or another language, please contact the Patient Advice and Liaison Service on Telephone number: 08000 851067

4.2 **Or write to us:** NHS Somerset ICB, Freepost RRKL-XKSC-ACSG, Yeovil, Somerset, BA22 8HR or **Email us:** somicb.pals@nhs.net

5 REFERENCES

- The following sources have been considered when drafting this policy:
- 5.1 NHS Choices. (2015, May 29th). Dupuytren's contracture - Treatment. Retrieved from NHS Choices:
<http://www.nhs.uk/Conditions/Dupuytren's-contracture/Pages/Surgery.aspx>
- 5.2 NICE. (DEC 2016). Radiation therapy for early Dupuytren's disease. Retrieved from <https://www.nice.org.uk/guidance/indevelopment/gid-ipg10022/documents>
- 5.3 NICE, 2017. Collagenase clostridium histolyticum for treating Dupuytren's contracture. : <https://www.nice.org.uk/guidance/ta459>
- 5.4 Thelwall, S. P. (2015). Impact of obesity on the risk of wound infection following surgery: results from a nationwide prospective multicentre cohort study in England.
- 5.5 Clinical microbiology and infection: the official publication of the European Society of Clinical Microbiology and Infectious Diseases, vol. 21, no. 11, p. 1008.e1.
- 5.6 Von Campe, A., Kende, K., Omaren, H., & Meuli-Simmen, C. (2012). Painful Nodules and Cords in Dupuytren Disease. The Journal of Hand Surgery, 1313-1318.
- 5.7 Cochrane review dated 2014
<https://www.ncbi.nlm.nih.gov/pubmed/24671929>
- 5.8 <https://cks.nice.org.uk/dupuytren's-disease>
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- 5.10 Krefter C, Marks M, Hensler S, Herren DB, Calcagni M. Complications after treating dupuytren's disease. A systematic literature review. Hand surgery & rehabilitation. 2017, 36: 322-9
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- 5.13 Scherman P, Jenmalm P, Dahlin LB. Three-year recurrence of Dupuytren's contracture after needle fasciotomy and collagenase injection: a two-centre randomized controlled trial. J Hand Surg Eur Vol. 2018;43(8):836-40
- 5.14 Skov ST, Bisgaard T, Sondergaard P, Lange J. Injectable Collagenase Versus Percutaneous Needle Fasciotomy for Dupuytren Contracture in Proximal Interphalangeal Joints: A Randomized Controlled Trial. J Hand Surg Am. 2017;42(5):321-8 e3
- 5.15 Stromberg J, Ibsen Sorensen A, Friden J. Percutaneous Needle Fasciotomy Versus Collagenase Treatment for Dupuytren Contracture: A Randomized Controlled Trial with a Two-Year Follow-up. J Bone Joint Surg Am. 2018;100(13):1079-86
- 5.16 van Rijssen AL, Gerbrandy FS, Ter Linden H, Klip H, Werker PM. 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, 31: 717-25
- 5.17 van Rijssen AL, ter Linden H, Werker PM. Five-year results of a randomized clinical trial on treatment in Dupuytren's disease: Percutaneous needle fasciotomy versus limited fasciectomy. Plast Reconstr Surg. 2012, 129: 469-77
- 5.18 NHS England EBI List 1
[NHS England » Evidence-Based Interventions Programme](#)
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