

Clinical Commissioning Policy

Breast Reduction

Category 2 Intervention - Only routinely commissioned when specific criteria are met –

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Purpose	This document is part of a suite of policies that the Integrated Care Board (ICB) uses to drive its commissioning of healthcare. Each policy in that suite is a separate public document in its own right but will be applied with reference to other policies in that suite.
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Document control:		
Date:	Version Number:	Section and Description of Change
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1. Introduction

- 1.1 This policy relates to the commissioning of interventions which optimise clinical effectiveness and represent value for money.
- 1.2 This document is part of a suite of policies which the Integrated Care Board (ICB) uses to drive its commissioning of healthcare. Each policy is a separate public document in its own right but should be considered alongside all the other policies in the suite as well as the core principles outlined in Appendix 1.
- 1.3 At the time of publication, the evidence presented per procedure/treatment was the most current available.
- 1.4 This policy is based on NHS England's Evidence-Based Interventions (EBI) recommendations see link to programme below - accurate at the point of publication <https://www.aomrc.org.uk/ebi/clinicians/breast-reduction/>.

2. Purpose

- 2.1 This policy aims to ensure a common set of criteria for treatments and procedures across the region. This is intended to reduce variation of access to NHS services in different areas and allow fair and equitable treatment for all patients.

3. Summary of intervention

- 3.1 Breast reduction surgery is a procedure used to treat women with breast hyperplasia (enlargement), where breasts are large enough to cause problems like shoulder girdle dysfunction, intertrigo and adverse effects to quality of life.

4. Policy statement

- 4.1 Breast reduction is not routinely commissioned unless all of the following criteria are met:

- 4.1.1 The woman has received a full package of supportive care from their GP such as advice on weight loss and managing pain

AND

- 4.1.2 In cases of thoracic/ shoulder girdle discomfort, a physiotherapy assessment has been provided

AND

- 4.1.3 Breast size results in functional symptoms that require other treatments/interventions (e.g. intractable candidal intertrigo; thoracic backache/kyphosis where a professionally fitted bra has not helped with backache, soft tissue indentations at site of bra straps)

AND

- 4.1.4 Breast reduction planned to be 500gms or more per breast or at least 4 cup sizes

AND

4.1.5 Body mass index (BMI) is <27 kg/m² and stable for at least twelve months

AND

4.1.6 Woman must be provided with written information to allow her to balance the risks and benefits of breast surgery

AND

4.1.7 Women should be informed that smoking increases complications following breast reduction surgery and should be advised to stop smoking

AND

4.1.8 Women should be informed that breast surgery for hypermastia can cause permanent loss of lactation.

5. Exclusions

5.1 This policy does not apply to mammoplasty associated with breast cancer treatment.

6. Rationale

6.1 One systematic review and three non-randomized studies regarding breast reduction surgery for hypermastia were identified and showed that surgery is beneficial in patients with specific symptoms. Physical and psychological improvements, such as reduced pain, increased quality of life and less anxiety and depression were found for women with hypermastia following breast reduction surgery.

6.2 Breast reduction surgery for hypermastia can cause permanent loss of lactation function of breasts, as well as decreased areolar sensation, bleeding, bruising, and scarring and often alternative approaches (e.g. weight loss or a professionally fitted bra) work just as well as surgery to reduce symptoms. For women who are severely affected by complications of hypermastia and for whom alternative approaches have not helped, surgery can be offered. The aim of surgery is not cosmetic, it is to reduce symptoms (e.g. back ache).

7. Underpinning evidence

7.1 An investigation into the relationship between breast size, bra size and mechanical back British School of Osteopathy (2010). Pages 13 & 14

7.2 Royal College of Surgeons of England (2014) Commissioning Guide: Breast Reduction

7.3 Greenbaum, a. R., Heslop, T., Morris, J., & Dunn, K. W. (2003). An investigation of the suitability of bra fit in women referred for reduction British Journal of Plastic Surgery, 56(3), 230–236.

7.4 Wood, K., Cameron, M., & Fitzgerald, K. (2008). Breast size, bra fit and thoracic pain in young women: a correlational study. Chiropractic & Osteopathy, 16(1), 1-7.

7.5 Singh KA, Losken A. Additional benefits of reduction mammoplasty: a systematic review of the literature. Plast Reconstr Surg. 2012 Mar;129(3):562-70. PubMed: PM22090252

- 7.6 Strong B, Hall-Findlay EJ. How Does Volume of Resection Relate to Symptom Relief for Reduction Mammoplasty Patients? *Ann Plast Surg.* 2014 Apr 10. PubMed: PM24727444
- 7.7 Valtonen JP, Setala LP, Mustonen PK, Blom M. Can the efficacy of reduction mammoplasty be predicted? The applicability and predictive value of breast-related symptoms questionnaire in measuring breast-related symptoms pre- and postoperatively. *J Plast Reconstr Aesthet Surg.* 2014 May;67(5):676-81. PubMed: PM24508223
- 7.8 Foreman KB, Dibble LE, Droge J, Carson R, Rockwell WB. The impact of breast reduction surgery on low-back compressive forces and function in individuals with macromastia. *Plast Reconstr Surg.* 2009 Nov;124(5):1393-9. PubMed: PM20009823
- 7.9 Shah R, Al-Ajam Y, Stott D, Kang N. Obesity in mammoplasty: a study of complications following breast reduction. *J Plast Reconstr Aesthet Surg.* 2011 Apr;64(4):508-14. doi: 10.1016/j.bjps.2010.07.001. Epub 2010 Aug 3. PubMed PMID: 20682461.
- 7.10 Oo M, Wang Z, Sakakibara T, Kasai Y. Relationship Between Brassiere Cup Size and Shoulder-Neck Pain in Women. *The Open Orthopaedics Journal.* 2012;6:140-142. doi:10.2174/1874325001206010140.
- 7.11 NHS information. <https://www.nhs.uk/conditions/breast-reduction-on-the-nhs/>
- 7.12 Chen CL(1), Shore AD, Johns R, Clark JM, Manahan M, Makary MA The impact of obesity on breast surgery complications. *Plast Reconstr Surg.* 2011 Nov;128(5):395e-402e DOI: 10.1097/PRS.0b013e3182284c05

8. Force

- 8.1 This policy remains in force until it is superseded by a revised policy or by mandatory NICE guidance or other national directive relating to this intervention, or to alternative treatments for the same condition.

9. Coding

SQL code

```
WHEN left(der.Spell_Dominant_Procedure,4) IN ('B311')
AND apcs.der_diagnosis_all not like '%Z853%'
AND APCS.Admission_Method not like ('2%')
THEN 'E_breast_red'
```

Global cancer exclusion

```
APC
WHERE 1=1
-- Cancer Diagnosis Exclusion
AND (apcs.der_diagnosis_all not like '%C[0-9][0-9]%'
AND apcs.der_diagnosis_all not like '%D0%'
AND apcs.der_diagnosis_all not like '%D3[789]%'
AND apcs.der_diagnosis_all not like '%D4[012345678]%'
OR apcs.der_diagnosis_all IS NULL)
```

10. Monitoring And Review

- 10.1 This policy may be subject to continued monitoring using a mix of the following approaches:
- Prior approval process
 - Post activity monitoring through routine data
 - Post activity monitoring through case note audits
- 10.2 This policy will be kept under regular review, to ensure that it reflects developments in the evidence base regarding effectiveness and value.

11. Quality and Equality Analysis

- 11.1 Quality and Equality Impact Analyses have been undertaken for this policy at the time of its review.

Appendix - Core Objectives and Principles

Objectives

The main objective for having healthcare commissioning policies is to ensure that:

- Patients receive appropriate health treatments
- Treatments with no or a very limited evidence base are not used; and
- Treatments with minimal health gain are restricted.

Principles

This policy aims to ensure a common set of criteria for treatments and procedures across the region. This is intended to reduce variation of access to NHS services in different areas and allow fair and equitable treatment for all patients.

Commissioning decisions by ICB Commissioners are made in accordance with the commissioning principles set out as follows:

- Commissioners require clear evidence of clinical effectiveness before NHS resources are invested in the treatment.
- Commissioners require clear evidence of cost effectiveness before NHS resources are invested in the treatment.
- Commissioners will consider the extent to which the individual or patient group will gain a benefit from the treatment.
- Commissioners will balance the needs of an individual patient against the benefit which could be gained by alternative investment possibilities to meet the needs of the community.
- Commissioners will consider all relevant national standards and consider all proper and authoritative guidance.
- Where a treatment is approved Commissioners will respect patient choice as to where a treatment is delivered, in accordance with the 'NHS Choice' framework.
- Commissioning decisions will give 'due regard' to promote equality and uphold human rights. Decision making will follow robust procedures to ensure that decisions are fair and are made within legislative frameworks.

Core Eligibility Criteria

There are a number of circumstances where a patient may meet a 'core eligibility criterion' which means they are eligible to be referred for the procedures and treatments listed, regardless of whether they meet the criteria; or the procedure or treatment is not routinely commissioned.

These core clinical eligibility criteria are as follows:

- Any patient who needs 'urgent' treatment will always be treated.
- All NICE Technology Appraisals Guidance (TAG), for patients that meet all the eligible criteria listed in a NICE TAG will receive treatment.
- In cancer care (including but not limited to skin, head and neck, breast and sarcoma) any lesion that has features suspicious of malignancy, must be referred to an appropriate specialist for urgent assessment under the 2-week rule.
- NOTE: Funding for all solid and haematological cancers are now the responsibility of NHS England.
- Reconstructive surgery post cancer or trauma including burns.
- Congenital deformities: Operations on congenital anomalies of the face and skull are usually routinely commissioned by the NHS. Some conditions are considered highly specialised and are commissioned in the UK through the National Specialised Commissioning Advisory Group (NSCAG). As the incidence of some cranio-facial congenital anomalies is small and the treatment complex, specialised teams, working in designated centres and subject to national audit, should carry out such procedures.
- Tissue degenerative conditions requiring reconstruction and/or restoring function e.g. leg ulcers, dehisced surgical wounds, necrotising fasciitis.
- For patients wishing to undergo Gender reassignment, this is the responsibility of NHS England and patients should be referred to a Gender Identity Clinic (GIC) as outlined in the Interim NHS England Gender Dysphoria Protocol and Guideline 2013/14.

Cosmetic Surgery

Cosmetic surgery is often carried out to change a person's appearance to achieve what a person perceives to be a more desirable look.

Cosmetic surgery/treatments are regarded as procedures of low clinical priority and therefore not routinely commissioned by the ICB Commissioner.

A summary of Cosmetic Surgery is provided by NHS Choices. Weblink: <http://www.nhs.uk/conditions/Cosmetic-surgery/Pages/Introduction.aspx> and <http://www.nhs.uk/Conditions/Cosmetic-surgery/Pages/Procedures.aspx>

Diagnostic Procedures

Diagnostic procedures to be performed with the sole purpose of determining whether or not a restricted procedure is feasible should not be carried out unless the eligibility criteria are met, or approval has been given by the ICB or GP (as set out in the approval process of the patients responsible ICB) or as agreed by the IFR Panel as a clinically exceptional case.

Where a General Practitioner/Optometrlist/Dentist requests only an opinion the patient should not be placed on a waiting list or treated, but the opinion given and the patient returned to the care of the General Practitioner/Optometrlist/Dentist, in order for them to make a decision on future treatment.

Clinical Trials

The ICB will not fund continuation of treatment commenced as part of a clinical trial. This is in line with the Medicines for Human Use (Clinical Trials) Regulations 2004 and the Declaration of Helsinki which stipulates that the responsibility for ensuring a clear exit strategy from a trial, and that those benefiting from treatment will have ongoing access to it, lies with those conducting the trial. This responsibility lies with the trial initiators indefinitely.

Clinical Exceptionality

If any patients are excluded from this policy, for whatever reason, the clinician has the option to make an application for clinical exceptionality. However, the clinician must make a robust case to the Panel to confirm their patient is distinct from all the other patients who might be excluded from the designated policy.

The ICB will consider clinical exceptions to this policy in accordance with the Individual Funding Request (IFR) Governance Framework consisting of: IFR Decision Making Policy; and IFR Management Policy.