

Silicone Or Saline Breast Implant Removal

Issue Date: June 30, 1993

Authority: [32 CFR 199.4\(a\)\(1\)](#), [\(e\)\(8\)\(iv\)](#), and [\(e\)\(9\)](#)

1.0 CPT¹ PROCEDURE CODES

19328, 19330

2.0 DESCRIPTION

The removal of silicone or saline mammary implant material.

3.0 POLICY

3.1 Removal of silicone or saline breast implants is covered if the initial silicone or saline breast implantation was or would have been a covered benefit.

3.2 Signs or symptoms of complications must be present and documented. Current medical literature supports removal of silicone or saline breast implants for the following indications:

- Signs and symptoms that may signal implant rupture; and
- Capsular contracture.

3.3 If the initial silicone or saline breast implant surgery was for an indication not covered or coverable by TRICARE, implant removal may be covered only if it is necessary treatment of a complication which represents a separate medical condition. See [Section 1.1](#).

3.4 Breast Magnetic Resonance Imaging (MRI) to detect implant rupture is covered. The implantation of the breast implants must have been covered by TRICARE.

4.0 EXCLUSIONS

4.1 Removal of silicone or saline breast implants for the presence of autoimmune or connective tissue disorders.

4.2 In the case of implants not originally covered or coverable, implant damage, hardening, leakage, and autoimmune disorder do not qualify as separate medical conditions. They are

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TRICARE Policy Manual 6010.57-M, February 1, 2008

Chapter 4, Section 5.5

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considered unfortunate sequelae resulting from the initial non-covered surgery, and, therefore, are excluded.

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