SIENTRA, INC. CAPCON CARE PROGRAM
TERMS & CONDITIONS

Program Summary
The Sientra CapCon Care Program ("C³ Program") provides first-time primary breast augmentation patients implanted with Sientra Smooth or Textured Silicone Gel Breast Implants, with the limited benefit of a No-Charge Replacement Implant ("NCRI") in the event the patient is diagnosed with Baker Grade III or Baker Grade IV capsular contracture in her affected breast.

The C³ Program provides this limited benefit if the following requirements have been met:

1. Program Effective Date and Duration

a. The patient was implanted with Sientra Smooth or Textured Silicone Gel Breast Implants on or after October 1, 2014 and Baker Grade III or Baker Grade IV capsular contracture occurred within 5 years of primary augmentation surgery.

2. Patient Eligibility

a. The patient's surgery was for primary augmentation and the patient has not previously undergone any breast implant surgery; and
b. The patient has been enrolled by her primary augmentation surgeon in the Sientra Device Tracking and Limited Warranty Programs; and
c. The patient has been diagnosed by her surgeon with Baker Grade III or Baker Grade IV capsular contracture.

3. NCRI Eligibility

a. Both the original and replacement surgeries have been performed by a board-certified (or board-admissible) plastic surgeon; and
b. Both surgeries occurred within the United States (including Puerto Rico); and
c. The surgery was performed in accordance with the Sientra Silicone Gel Breast Implant's Directions for Use documentation; and
d. The surgeon has provided a completed and signed Request for an NCRI Form within the specified timeline from the original implantation in accordance with the Program Effective Date.

4. C³ Program Limitations

a. Sientra will provide an NCRI if the replacement implant is:
    (i) The same implant profile (Round, Shaped Oval Base, Shaped Round Base, Shaped Classic Base) as the original implant; and
    (ii) The same projection (Low, Moderate, Moderate Plus, Moderate High, or High) as the original implant; and
    (iii) The same gel type (High Strength Cohesive Gel, or High Strength Cohesive Plus Gel) as the original implant; and
    (iv) The same size or 1 size smaller or larger than the original implant, depending upon the availability of the requested implant size, projection and profile at the time of the NCRI request.

5. Other Limitations

a. Only a single NCRI is provided per affected breast per patient. Replacement of the contralateral breast implant or surgical spares will be billed at the list price.
b. If all program requirements have been met for both breasts and the patient diagnosis is for bilateral capsular contracture, the patient may qualify for 2 NCRI implants.
c. Revision surgery to remove and replace the original implant must occur within 180 days of shipment of the NCRI replacement implant to the surgeon performing the replacement procedure. The replacement surgery must be reported to Sientra via a completed Device Tracking Form for the replacement implant. If the Device Tracking Form is not completed and provided to Sientra within the 180 day period, the surgeon performing the replacement surgery will be charged the patient pre-approved account price for the replacement implant.
d. Sientra reserves the right to review patient medical records and operative reports and notes when determining NCRI eligibility.

6. The C^3 Program and Sientra's Limited Warranty and Lifetime Product Replacement Program

a. The C^3 Program is an added benefit to patients and is in addition to Sientra's Limited Warranty and Lifetime Product Replacement Program. C^3 Terms & Conditions and those of the Limited Warranty & Lifetime Replacement Program can be found on sientra.com.
b. The benefits of the C^3 Program and Limited Warranty & Lifetime Product Replacement Program may overlap and may allow for implant replacement in accordance with the Terms & Conditions of each Program.
c. Replacement implants carry whatever Limited Warranty or other programs are in effect at the time the replacement implant is shipped to the surgeon.
d. Patients who simultaneously experience both capsular contracture and an implant rupture may qualify for an NCRI under the C^3 Program and a replacement implant under the Lifetime Product Replacement Program; however, the patient is eligible for only one (1) NCRI via the C^3 Program per breast.

7. Refer to Product Labeling and "Directions for Use" for Additional Information

a. These Terms & Conditions do not replace the product labeling or the product's "Directions for Use" for Sientra's Silicone Gel Breast Implants. Individuals with questions about these products should read those documents or the Patient Educational Brochure available on Sientra's website, sientra.com, or consult their board-certified (or board-admissible) plastic surgeon.

8. No Cash Benefit

a. There is no cash benefit to patients in this Program; specifically, in the event that:
    (i) the NCRI has a lower list price than the patient's original implant; or
    (ii) if the patient's same style or size or requested size is not available for any reason, including supply shortage, strikes, acts of war, or acts of God, or for any other reason that Sientra is unable to supply timely (or at all) any NCRI.

9. Non Transferable

The benefit of an NCRI is provided to individual patients only and is intended exclusively for the benefit of that patient. The benefit is not transferable.

10. Program Termination

Once a patient is implanted under CapCon Care in accordance with the C^3 Program, the Program's benefits are guaranteed to the patient. However, the Program may be terminated for new enrollees at any time without notice.

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