

(iii) a person alleging harm caused by either the silicone gel or the silicone envelope utilized in a breast implant containing silicone gel, except that—

(I) neither the exclusion provided by this clause nor any other provision of this chapter may be construed as a finding that silicone gel (or any other form of silicone) may or may not cause harm; and

(II) the existence of the exclusion under this clause may not—

(aa) be disclosed to a jury in any civil action or other proceeding; and

(bb) except as necessary to establish the applicability of this chapter, otherwise be presented in any civil action or other proceeding.

**(3) Component part**

**(A) In general**

The term “component part” means a manufactured piece of an implant.

**(B) Certain components**

Such term includes a manufactured piece of an implant that—

(i) has significant non-implant applications; and

(ii) alone, has no implant value or purpose, but when combined with other component parts and materials, constitutes an implant.

**(4) Harm**

**(A) In general**

The term “harm” means—

(i) any injury to or damage suffered by an individual;

(ii) any illness, disease, or death of that individual resulting from that injury or damage; and

(iii) any loss to that individual or any other individual resulting from that injury or damage.

**(B) Exclusion**

The term does not include any commercial loss or loss of or damage to an implant.

**(5) Implant**

The term “implant” means—

(A) a medical device that is intended by the manufacturer of the device—

(i) to be placed into a surgically or naturally formed or existing cavity of the body for a period of at least 30 days; or

(ii) to remain in contact with bodily fluids or internal human tissue through a surgically produced opening for a period of less than 30 days; and

(B) suture materials used in implant procedures.

**(6) Manufacturer**

The term “manufacturer” means any person who, with respect to an implant—

(A) is engaged in the manufacture, preparation, propagation, compounding, or processing (as defined in section 360(a)(1) of this title) of the implant; and

(B) is required—

(i) to register with the Secretary pursuant to section 360 of this title and the regulations issued under such section; and

(ii) to include the implant on a list of devices filed with the Secretary pursuant to section 360(j) of this title and the regulations issued under such section.

**(7) Medical device**

The term “medical device” means a device, as defined in section 321(h) of this title, and includes any device component of any combination product as that term is used in section 353(g) of this title.

**(8) Raw material**

The term “raw material” means a substance or product that—

(A) has a generic use; and

(B) may be used in an application other than an implant.

**(9) Secretary**

The term “Secretary” means the Secretary of Health and Human Services.

**(10) Seller**

**(A) In general**

The term “seller” means a person who, in the course of a business conducted for that purpose, sells, distributes, leases, packages, labels, or otherwise places an implant in the stream of commerce.

**(B) Exclusions**

The term does not include—

(i) a seller or lessor of real property;

(ii) a provider of professional health care services in any case in which—

(I) the sale or use of the implant is incidental to such services; and

(II) the essence of the professional health care services provided is the furnishing of judgment, skill, or services; or

(iii) any person who acts in only a financial capacity with respect to the sale of an implant.

(Pub. L. 105-230, § 3, Aug. 13, 1998, 112 Stat. 1520.)

**Statutory Notes and Related Subsidiaries**

**EFFECTIVE DATE**

Section applicable to all civil actions covered under this chapter commenced on or after Aug. 13, 1998, including any in which the harm or harmful conduct occurred before such date, see section 8 of Pub. L. 105-230, set out as a note under section 1601 of this title.

**§ 1603. General requirements; applicability; preemption**

**(a) General requirements**

**(1) In general**

In any civil action covered by this chapter, a biomaterials supplier may—

(A) raise any exclusion from liability set forth in section 1604 of this title; and

(B) make a motion for dismissal or for summary judgment as set forth in section 1605 of this title.

**(2) Procedures**

Notwithstanding any other provision of law, a Federal or State court in which an action

covered by this chapter is pending shall, in connection with a motion under section 1605 or 1606 of this title, use the procedures set forth in this chapter.

**(b) Applicability**

**(1) In general**

Except as provided in paragraph (2), this chapter applies to any civil action brought by a claimant, whether in a Federal or State court, on the basis of any legal theory, for harm allegedly caused, directly or indirectly, by an implant.

**(2) Exclusion**

A civil action brought by a purchaser of a medical device, purchased for use in providing professional health care services, for loss or damage to an implant or for commercial loss to the purchaser—

(A) shall not be considered an action that is subject to this chapter; and

(B) shall be governed by applicable commercial or contract law.

**(c) Scope of preemption**

**(1) In general**

This chapter supersedes any State law regarding recovery for harm caused by an implant and any rule of procedure applicable to a civil action to recover damages for such harm only to the extent that this chapter establishes a rule of law applicable to the recovery of such damages.

**(2) Applicability of other laws**

Any issue that arises under this chapter and that is not governed by a rule of law applicable to the recovery of damages described in paragraph (1) shall be governed by applicable Federal or State law.

**(d) Statutory construction**

Nothing in this chapter may be construed—

(1) to affect any defense available to a defendant under any other provisions of Federal or State law in an action alleging harm caused by an implant; or

(2) to create a cause of action or Federal court jurisdiction pursuant to section 1331 or 1337 of title 28 that otherwise would not exist under applicable Federal or State law.

(Pub. L. 105-230, §4, Aug. 13, 1998, 112 Stat. 1523.)

**Statutory Notes and Related Subsidiaries**

**EFFECTIVE DATE**

Section applicable to all civil actions covered under this chapter commenced on or after Aug. 13, 1998, including any in which the harm or harmful conduct occurred before such date, see section 8 of Pub. L. 105-230, set out as a note under section 1601 of this title.

**§ 1604. Liability of biomaterials suppliers**

**(a) In general**

Except as provided in section 1606 of this title, a biomaterials supplier shall not be liable for harm to a claimant caused by an implant unless such supplier is liable—

(1) as a manufacturer of the implant, as provided in subsection (b);

(2) as a seller of the implant, as provided in subsection (c); or

(3) for furnishing raw materials or component parts for the implant that failed to meet applicable contractual requirements or specifications, as provided in subsection (d).

**(b) Liability as manufacturer**

**(1) In general**

A biomaterials supplier may, to the extent required and permitted by any other applicable law, be liable for harm to a claimant caused by an implant if the biomaterials supplier is the manufacturer of the implant.

**(2) Grounds for liability**

The biomaterials supplier may be considered the manufacturer of the implant that allegedly caused harm to a claimant only if the biomaterials supplier—

(A)(i) registered or was required to register with the Secretary pursuant to section 360 of this title and the regulations issued under such section; and

(ii) included or was required to include the implant on a list of devices filed with the Secretary pursuant to section 360(j) of this title and the regulations issued under such section;

(B) is the subject of a declaration issued by the Secretary pursuant to paragraph (3) that states that the supplier, with respect to the implant that allegedly caused harm to the claimant, was required to—

(i) register with the Secretary under section 360 of this title, and the regulations issued under such section, but failed to do so; or

(ii) include the implant on a list of devices filed with the Secretary pursuant to section 360(j) of this title and the regulations issued under such section, but failed to do so; or

(C) is related by common ownership or control to a person meeting all the requirements described in subparagraph (A) or (B), if the court deciding a motion to dismiss in accordance with section 1605(c)(3)(B)(i) of this title finds, on the basis of affidavits submitted in accordance with section 1605 of this title, that it is necessary to impose liability on the biomaterials supplier as a manufacturer because the related manufacturer meeting the requirements of subparagraph (A) or (B) lacks sufficient financial resources to satisfy any judgment that the court feels it is likely to enter should the claimant prevail.

**(3) Administrative procedures**

**(A) In general**

The Secretary may issue a declaration described in paragraph (2)(B) on the motion of the Secretary or on petition by any person, after providing—

(i) notice to the affected persons; and

(ii) an opportunity for an informal hearing.

**(B) Docketing and final decision**

Immediately upon receipt of a petition filed pursuant to this paragraph, the Sec-