Food and Drug Administration, HHS

§878.4100 Organ bag.
(a) Identification. An organ bag is a device that is a flexible plastic bag intended to be used as a temporary receptacle for an organ during surgical procedures to prevent moisture loss.
(b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §878.9.

§878.4160 Surgical camera and accessories.
(a) Identification. A surgical camera and accessories is a device intended to be used to record operative procedures.
(b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter, subject to the limitations in §878.9.

§878.4200 Introduction/drainage catheter and accessories.
(a) Identification. An introduction/drainage catheter is a device that is a flexible single or multilumen tube intended to be used to introduce nondrug fluids into body cavities other than blood vessels, drain fluids from body cavities, or evaluate certain physiologic conditions. Examples include irrigation and drainage catheters, pediatric catheters, peritoneal catheters (including dialysis), and other general surgical catheters. An introduction/drainage catheter accessory is intended to aid in the manipulation of or insertion of the device into the body. Examples of accessories include adaptors, connectors, and catheter needles.
(b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §878.9.
[53 FR 23872, June 24, 1988, as amended at 65 FR 2318, Jan. 14, 2000]

§878.4300 Implantable clip.
(a) Identification. An implantable clip is a clip-like device intended to connect internal tissues to aid healing. It is not absorbable.
(b) Classification. Class II.

§878.4320 Removable skin clip.
(a) Identification. A removable skin clip is a clip-like device intended to connect skin tissues temporarily to aid healing. It is not absorbable.
(b) Classification. Class I (general controls). The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to §878.9.
[53 FR 23872, June 24, 1988, as amended at 65 FR 2318, Jan. 14, 2000]

§878.4340 Contact cooling system for aesthetic use.
(a) Identification. A contact cooling system for aesthetic use is a device that is a combination of a cooling pad associated with a vacuum or mechanical massager intended for the disruption of adipocyte cells intended for non-invasive aesthetic use.
(b) Classification. Class II (special controls). The special controls for this device is FDA’s “Guidance for Industry and FDA Staff; Class II Special Controls Guidance Document: Contact Cooling System for Aesthetic Use.” See §878.1(e) for the availability of this guidance document.
[76 FR 6553, Feb. 7, 2011]

§878.4350 Cryosurgical unit and accessories.
(a) Identification—(1) Cryosurgical unit with a liquid nitrogen cooled cryoprobe and accessories. A cryosurgical unit with a liquid nitrogen cooled cryoprobe and accessories is a device intended to destroy tissue during surgical procedures by applying extreme cold.
(2) Cryosurgical unit with a nitrous oxide cooled cryoprobe and accessories. A cryosurgical unit with a nitrous oxide cooled cryoprobe and accessories is a device intended to destroy tissue during surgical procedures, including urological applications, by applying extreme cold.
(3) Cryosurgical unit with a carbon dioxide cooled cryoprobe or a carbon dioxide