

§ 888.1

- 888.3720 Toe joint polymer constrained prosthesis.
- 888.3730 Toe joint phalangeal (hemi-toe) polymer prosthesis.
- 888.3750 Wrist joint carpal lunate polymer prosthesis.
- 888.3760 Wrist joint carpal scaphoid polymer prosthesis.
- 888.3770 Wrist joint carpal trapezium polymer prosthesis.
- 888.3780 Wrist joint polymer constrained prosthesis.
- 888.3790 Wrist joint metal constrained cemented prosthesis.
- 888.3800 Wrist joint metal/polymer semi-constrained cemented prosthesis.
- 888.3810 Wrist joint ulnar (hemi-wrist) polymer prosthesis.

Subpart E—Surgical Devices

- 888.4150 Calipers for clinical use.
- 888.4200 Cement dispenser.
- 888.4210 Cement mixer for clinical use.
- 888.4220 Cement monomer vapor evacuator.
- 888.4230 Cement ventilation tube.
- 888.4300 Depth gauge for clinical use.
- 888.4540 Orthopedic manual surgical instrument.
- 888.4580 Sonic surgical instrument and accessories/attachments.
- 888.4600 Protractor for clinical use.
- 888.4800 Template for clinical use.
- 888.5850 Nonpowered orthopedic traction apparatus and accessories.
- 888.5890 Noninvasive traction component.
- 888.5940 Cast component.
- 888.5960 Cast removal instrument.
- 888.5980 Manual cast application and removal instrument.

AUTHORITY: 21 U.S.C. 351, 360, 360c, 360e, 360j, 371.

SOURCE: 52 FR 33702, Sept. 4, 1987, unless otherwise noted.

Subpart A—General Provisions

§ 888.1 Scope.

(a) This part sets forth the classification of orthopedic devices intended for human use that are in commercial distribution.

(b) The identification of a device in a regulation in this part is not a precise description of every device that is, or will be, subject to the regulation. A manufacturer who submits a premarket notification submission for a device under part 807 cannot show merely that the device is accurately described by the section title and identification provision of a regulation in this part, but shall state why the de-

21 CFR Ch. I (4–1–03 Edition)

vice is substantially equivalent to other devices, as required by § 807.87.

(c) To avoid duplicative listings, an orthopedic device that has two or more types of uses (e.g., used both as a diagnostic device and as a surgical device) is listed in one subpart only.

(d) References in this part to regulatory sections of the Code of Federal Regulations are to chapter I of title 21 unless otherwise noted.

(e) Guidance documents referenced in this part are available on the Internet at <http://www.fda.gov/cdrh/guidance.html>.

[52 FR 33702, Sept. 4, 1987, as amended at 68 FR 14137, Mar. 24, 2003]

§ 888.3 Effective dates of requirement for premarket approval.

A device included in this part that is classified into class III (premarket approval) shall not be commercially distributed after the date shown in the regulation classifying the device unless the manufacturer has an approval under section 515 of the act (unless an exemption has been granted under section 520(g)(2) of the act). An approval under section 515 of the act consists of FDA's issuance of an order approving an application for premarket approval (PMA) for the device or declaring completed a product development protocol (PDP) for the device.

(a) Before FDA requires that a device commercially distributed before the enactment date of the amendments, or a device that has been found substantially equivalent to such a device, has an approval under section 515 of the act, FDA must promulgate a regulation under section 515(b) of the act requiring such approval, except as provided in paragraphs (b) and (c) of this section. Such a regulation under section 515(b) of the act shall not be effective during the grace period ending on the 90th day after its promulgation or on the last day of the 30th full calendar month after the regulation that classifies the device into class III is effective, whichever is later. See section 501(f)(2)(B) of the act. Accordingly, unless an effective date of the requirement for premarket approval is shown in the regulation for a device classified into class III in this part, the device