(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

3. Section 862.2320 is amended by revising paragraph (b) to read as follows:

§ 862.2320 Beta or gamma counter for clinical use.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

4. Section 862.2485 is amended by revising paragraph (b) to read as follows:

§ 862.2485 Electrophoresis apparatus for clinical use.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

5. Section 862.2720 is amended by revising paragraph (b) to read as follows:

§ 862.2720 Plasma oncometer for clinical use.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

6. Section 862.2800 is amended by revising paragraph (b) to read as follows:

§ 862.2800 Refractometer for clinical use.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

7. Section 862.2920 is amended by revising paragraph (b) to read as follows:

§ 862.2920 Plasma viscometer for clinical use.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

PART 872—DENTAL DEVICES

8. The authority citation for 21 CFR part 872 continues to read as follows:

Authority: Secs. 501, 510, 513, 515, 520, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 351, 360, 360c, 360e, 360j, 371).

9. Section 872.3740 is amended by revising paragraph (b) to read as follows:

§ 872.3740 Retentive and splinting pin.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

10. Section 872.3810 is amended by revising paragraph (b) to read as follows:

§ 872.3810 Root canal post.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

11. Section 872.6100 is amended by revising paragraph (b) to read as follows:

§ 872.6100 Anesthetic warmer.

* * * * *

(b) *Classification*. Class I. The device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter.

Dated: July 18, 1995.

William B. Schultz,

Deputy Commissioner for Policy.

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