Food and Drug Administration, HHS

subpart E of part 807 of this chapter subject to §862.9.

 $[52~{\rm FR}~16122,~{\rm May}~1,~1987,~{\rm as}~{\rm amended}~{\rm at}~65~{\rm FR}~2307,~{\rm Jan.}~14,~2000]$

§862.1470 Lipid (total) test system.

- (a) *Identification*. A lipid (total) test system is a device intended to measure total lipids (fats or fat-like substances) in serum and plasma. Lipid (total) measurements are used in the diagnosis and treatment of various diseases involving lipid metabolism and atherosclerosis.
- (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 §862.9.

 $[52\ {\rm FR}\ 16122,\ {\rm May}\ 1,\ 1987,\ {\rm as}\ {\rm amended}\ {\rm at}\ 53\ {\rm FR}\ 21449,\ {\rm June}\ 8,\ 1988;\ 66\ {\rm FR}\ 38788,\ {\rm July}\ 25,\ 2001]$

§862.1475 Lipoprotein test system.

- (a) *Identification*. A lipoprotein test system is a device intended to measure lipoprotein in serum and plasma. Lipoprotein measurements are used in the diagnosis and treatment of lipid disorders (such as diabetes mellitus), atherosclerosis, and various liver and renal diseases.
- (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 §862.9.

[52 FR 16122, May 1, 1987, as amended at 65 FR 2307, Jan. 14, 2000]

\$ 862.1485 Luteinizing hormone test system.

- (a) *Identification*. A luteinizing hormone test system is a device intended to measure luteinizing hormone in serum and urine. Luteinizing hormone measurements are used in the diagnosis and treatment of gonadal dysfunction.
- (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 §862.9.

 $[52~{\rm FR}~16122,~{\rm May}~1,~1987,~{\rm as}~{\rm amended}~{\rm at}~65~{\rm FR}~2307,~{\rm Jan.}~14,~2000]$

§862.1490 Lysozyme (muramidase) test system.

- (a) Identification. A lysozyme (muramidase) test system is a device intended to measure the activity of the bacteriolytic enzyme lysozyme (muramidase) in serum, plasma, leukocytes, and urine. Lysozyme measurements are used in the diagnosis and treatment of monocytic leukemia and kidney disease.
- (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 §862.9.

[52 FR 16122, May 1, 1987, as amended at 53 FR 21449, June 8, 1988; 66 FR 38788, July 25, 2001]

§862.1495 Magnesium test system.

- (a) Identification. A magnesium test system is a device intended to measure magnesium levels in serum and plasma. Magnesium measurements are used in the diagnosis and treatment of hypomagnesemia (abnormally low plasma levels of magnesium) and hypermagnesemia (abnormally high plasma levels of magnesium).
 - (b) Classification. Class I.

§862.1500 Malic dehydrogenase test system.

- (a) Identification. A malic dehydrogenase test system is a device that is intended to measure the activity of the enzyme malic dehydrogenase in serum and plasma. Malic dehydrogenase measurements are used in the diagnosis and treatment of muscle and liver diseases, myocardial infarctions, cancer, and blood disorders such as myelogenous (produced in the bone marrow) leukemia.
- (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 §862.9.

[52 FR 16122, May 1, 1987, as amended at 65 FR 2307, Jan. 14, 2000]

§862.1505 Mucopolysaccharides (nonquantitative) test system.

(a) *Identification*. A mucopolysaccharides (nonquantitative) test system is a device intended to