Food and Drug Administration, HHS

§ 864.7490 Sulfhemoglobin assay.
(a) Identification. A sulfhemoglobin assay is a device consisting of the reagents, calibrators, controls, and instrumentation used to determine the sulfhemoglobin (a compound of sulfur and hemoglobin) content of human blood as an aid in the diagnosis of sulfhemoglobinemia (presence of sulfhemoglobin in the blood due to drug administration or exposure to a poison). This measurement may be made using methods such as spectrophotometry, colorimetry, spectrophotometry, or gasometry.
(b) Classification. Class II (performance standards).
[45 FR 60621, Sept. 12, 1980]

§ 864.7500 Whole blood hemoglobin assays.
(a) Identification. A whole blood hemoglobin assay is a device consisting of reagents, calibrators, controls, or photometric or spectrophotometric instrumentation used to measure the hemoglobin content of whole blood for the detection of anemia. This generic device category does not include automated hemoglobin systems.
(b) Classification. Class II (performance standards).
[45 FR 60622, Sept. 12, 1980]

§ 864.7525 Heparin assay.
(a) Identification. A heparin assay is a device used to determine the level of the anticoagulant heparin in the patient’s circulation. These assays are quantitative clotting time procedures using the effect of heparin on activated coagulation factor X (Stuart factor) or procedures based on the neutralization of heparin by protamine sulfate (a protein that neutralizes heparin).
(b) Classification. Class II (performance standards).
[45 FR 60622, Sept. 12, 1980]

§ 864.7660 Leukocyte alkaline phosphatase test.
(a) Identification. A leukocyte alkaline phosphatase test is a device used to identify the enzyme leukocyte alkaline phosphatase in neutrophilic granulocytes (granular leukocytes stainable by neutral dyes). The cytochemical identification of alkaline phosphatase depends on the formation of blue granules in cells containing alkaline phosphatase. The results of this test are used to differentiate chronic granulocytic leukemia (a malignant disease characterized by excessive overgrowth of granulocytes in the bone marrow) and reactions that resemble true leukemia, such as those occurring in severe infections and polycythemia (increased total red cell mass).
(b) Classification. Class I (general controls). This device is exempt from the premarket notification procedures in subpart E of part 807 of this chapter subject to the limitations in § 864.9.

§ 864.7695 Platelet factor 4 radioimmunoassay.
(a) Identification. A platelet factor 4 radioimmunoassay is a device used to measure the level of platelet factor 4, a protein released during platelet activation by radioimmunoassay. This device measures platelet activation, which may indicate a coagulation disorder, such as myocardial infarction or coronary artery disease.
(b) Classification. Class II (performance standards).