Celite is a registered trademark of Celite Corporation, Santa Barbara, CA for its diatomaceous earth products.

ENDOCRINOLOGY

Hematology

Calculated—note that TCO₂ is measured on CHEM8+ cartridge and calculated on all others.

See Intended Use information at right

INTENDED USE

LACTATE

ACT® Kaolin

The ACT Kaolin Activated Clotting Time (ACT) test is an in vitro diagnostic test for the quantitative and qualitative determination of ACT in whole blood samples using the i-STAT 1 System. The test is intended for use in the early detection of bleeding and for the prevention of unnecessary blood loss during surgery or blood donation.

ACT®

The ACT® test is an in vitro diagnostic test for the quantitative measurement of anticoagulant activity in whole blood or plasma samples using the i-STAT 1 System. The test is intended for use in the early detection of bleeding and for the prevention of unnecessary blood loss during surgery or blood donation.

COAGULATION

The ACT® test is an in vitro diagnostic test for the quantitative measurement of anticoagulant activity in whole blood or plasma samples using the i-STAT 1 System. The test is intended for use in the early detection of bleeding and for the prevention of unnecessary blood loss during surgery or blood donation.

Cardiac markers

The CK-MB mass measurement is used in the diagnosis and treatment of myocardial infarction (MI).

The measurement of cardiac troponin I (cTnI) in whole blood or plasma. Measurements of cardiac troponin I may aid in the diagnosis and treatment of myocardial infarction and is used in conjunction with other clinical laboratory tests to help in the evaluation of patients with cardiac coronary artery disease and to determine the likelihood of a cardiac event.

ß-hCG

The ß-hCG test is an in vitro diagnostic test for the quantitative and qualitative determination of ß-hCG in whole blood or plasma samples using the i-STAT 1 System. The test is intended for use in the early detection of pregnancy and is for prescription use only.

This test menu is intended for use only in the United States for in vitro diagnostic use only.