CADRDIOSAVE Intra-Aortic Balloon Pump

MAQUET Cardiovascular LLC, has announced U.S. Food & Drug Administration (FDA) 510(k) clearance and CE mark for its new CARDIOSAVE intra-aortic balloon pump (IABP). CARDIOSAVE is expected to be commercially available in the United States in January 2012.

The new CARDIOSAVE IABP incorporates a large state-of-the-art touchscreen display and is dramatically smaller, lighter and quieter than any pump the company has ever offered. The pump is being offered in two configurations: CARDIOSAVE Hybrid for routine in-hospital use and CARDIOSAVE Rescue for use in ambulances and aircraft.

For more information, visit ca.maquet.com [1].

Links:
[1] http://ca.maquet.com/