

K900302 UMBILICAL CORD BLOOD SAMPLING KITApr 10, 1990
78 days to decisionK900302 · Product code: **CBT** · Anesthesiology
Source: <https://www.510kdatabase.net/k900302/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arterial Blood Sampling Kit (CBT)
Date received	Jan 22, 1990
Decision date	Apr 10, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	Ge Medical Systems Information Technologies
Location	Mchenry, IL, US
Contact	BRIAN R BARRY
510(k) history	136 submissions · 132 cleared · 1978-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900302/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026