FDA 510(k) Premarket Notification

K900302 UMBILICAL CORD BLOOD SAMPLING KIT

CLEARED

Apr 10, 1990
78 days to decision

K900302 · 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