FDA 510(k) Premarket Notification

K903392 ELECTROMEDICS BT422-120 AND BT422-40

CLEARED

Sep 6, 1990
41 days to decision

K903392 · Product code: **CAC** · Anesthesiology
Source: https://www.510kdatabase.net/k903392/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Autotransfusion (CAC)
Date received	Jul 27, 1990
Decision date	Sep 6, 1990
Days to decision	41 days
Third-party review	No

APPLICANT

Company	**Electromedics, Inc.**
Location	Mchenry, IL, US
Contact	HAROLD W BRYAN
510(k) history	41 submissions · 41 cleared · 1977-1994

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k903392/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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