FDA 510(k) Premarket Notification

K781616 OMNI STIK

CLEARED

Dec 8, 1978
79 days to decision

K781616 · Product code: **CBT** · Anesthesiology
Source: https://www.510kdatabase.net/k781616/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arterial Blood Sampling Kit (CBT)
Date received	Sep 20, 1978
Decision date	Dec 8, 1978
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Unitron Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978