

K781616 OMNI STIKDec 8, 1978
79 days to decisionK781616 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k781616/> 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 28, 2026