

K812765 DATASCRIBE #DS-40Nov 16, 1981
42 days to decisionK812765 · Product code: **BXW** · Anesthesiology
Source: <https://www.510kdatabase.net/k812765/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrator, Volume, Gas (BXW)
Date received	Oct 5, 1981
Decision date	Nov 16, 1981
Days to decision	42 days
Third-party review	No

APPLICANT

Company	Timeter Instrument Corp.
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1976-1988

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Device record: <https://www.510kdatabase.net/k812765/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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