

K781271 MONITOR, PO2, INTRAVASCULAROct 12, 1978
80 days to decisionK781271 · Product code: **CCE** · Anesthesiology
Source: <https://www.510kdatabase.net/k781271/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Partial Pressure, Blood-phase, Indwelling (CCE)
Date received	Jul 24, 1978
Decision date	Oct 12, 1978
Days to decision	80 days
Third-party review	No

APPLICANT

Company	Roche Medical Electronics, Inc.
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1976-1979

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Device record: <https://www.510kdatabase.net/k781271/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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