

K781680 MONITORING, PRESSURE KITOct 6, 1978
4 days to decisionK781680 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k781680/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Oct 2, 1978
Decision date	Oct 6, 1978
Days to decision	4 days
Third-party review	No

APPLICANT

Company	Pharmaseal Div., Baxter Healthcare Corp.
Location	Mchenry, IL, US
510(k) history	24 submissions · 24 cleared · 1976-1979

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Device record: <https://www.510kdatabase.net/k781680/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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