

K801802 ELECTROMAGNETIC BLOOD & ARTERIAL GAUGEJan 2, 1981
157 days to decisionK801802 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k801802/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Jul 29, 1980
Decision date	Jan 2, 1981
Days to decision	157 days
Third-party review	No

APPLICANT

Company	Medical Testing System, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1981

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Device record: <https://www.510kdatabase.net/k801802/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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