FDA 510(k) Premarket Notification

K801802 ELECTROMAGNETIC BLOOD & ARTERIAL GAUGE

CLEARED

Jan 2, 1981
157 days to decision

K801802 · Product code: **DPW** · Cardiovascular
Source: 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