FDA 510(k) Premarket Notification

K890828 PERIFLUX PF3

CLEARED

Jun 20, 1989
119 days to decision

K890828 · Product code: **DPW** · Cardiovascular
Source: https://www.510kdatabase.net/k890828/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Feb 21, 1989
Decision date	Jun 20, 1989
Days to decision	119 days
Third-party review	No

APPLICANT

Company	**Perimed, Inc.**
Location	Piscataway, NJ, US
Contact	OYSTEIN LIND
510(k) history	6 submissions · 6 cleared · 1989-1999