FDA 510(k) Premarket Notification

K812005 CAPILLARY PERFUSION MONITOR #LD5000

CLEARED

Feb 4, 1982
205 days to decision

K812005 · Product code: **DPW** · Cardiovascular
Source: https://www.510kdatabase.net/k812005/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Jul 14, 1981
Decision date	Feb 4, 1982
Days to decision	205 days
Third-party review	No

APPLICANT

Company	**Nuclear Pacific, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k812005/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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