

K881303 TRITON MODEL 100 PULSED DOPPLER BLOOD FLOWMETERNov 29, 1988
246 days to decisionK881303 · Product code: **DPT** · Cardiovascular
Source: <https://www.510kdatabase.net/k881303/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Blood-flow, Extravascular (DPT)
Date received	Mar 28, 1988
Decision date	Nov 29, 1988
Days to decision	246 days
Third-party review	No

APPLICANT

Company	Triton Technology, Inc.
Location	San Diego, CA, US
Contact	SCOTT KEMPER
510(k) history	4 submissions · 4 cleared · 1988-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881303/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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