FDA 510(k) Premarket Notification

K881303 TRITON MODEL 100 PULSED DOPPLER BLOOD FLOWMETER

CLEARED

Nov 29, 1988
246 days to decision

K881303 · 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