FDA 510(k) Premarket Notification

K870016 ARTERIAL BLOOD GASSING MONITORING SYSTEM (ABG)

CLEARED

May 28, 1987
146 days to decision

K870016 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k870016/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jan 2, 1987
Decision date	May 28, 1987
Days to decision	146 days
Third-party review	No

APPLICANT

Company	**Migada, Ltd.**
Location	Israel, IL
Contact	DAVID THALER
510(k) history	15 submissions · 15 cleared · 1985-1994