

**K870016 ARTERIAL BLOOD GASSING MONITORING SYSTEM
(ABG)**May 28, 1987
146 days to decisionK870016 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k870016/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026