

K850870 MEDIFETTE ERGOMETERJun 12, 1985
103 days to decision

K850870 · Physical Medicine

Source: <https://www.510kdatabase.net/k850870/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Mar 1, 1985
Decision date	Jun 12, 1985
Days to decision	103 days
Third-party review	No

APPLICANT

Company	Ketronic, Inc.
Location	Netherlands, US
Contact	VANDILL
510(k) history	10 submissions · 9 cleared · 1984-1985

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k850870/> 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 21, 2026