

K851702 MORTARA INSTRUMENT MODEL EL-1Jul 29, 1985
96 days to decisionK851702 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k851702/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 24, 1985
Decision date	Jul 29, 1985
Days to decision	96 days
Third-party review	No

APPLICANT

Company	Mortara Instrument, Inc.
Location	Walker, MI, US
Contact	TIMOTHY C MICKELSON
510(k) history	51 submissions · 51 cleared · 1983-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k851702/>; 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 June 19, 2026