

K791839 H59 CRITICAL CARE MONITOROct 1, 1979
17 days to decisionK791839 · Product code: **DSK** · CardiovascularSource: <https://www.510kdatabase.net/k791839/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Blood-pressure (DSK)
Date received	Sep 14, 1979
Decision date	Oct 1, 1979
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Medtel Pty. , Ltd.
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1979-1983

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k791839/> 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 27, 2026