

**K832050 MORTARA INSTRUMENT #101**Dec 13, 1983  
172 days to decisionK832050 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k832050/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Jun 24, 1983
Decision date	Dec 13, 1983
Days to decision	172 days
Third-party review	No

**APPLICANT**

---

Company	<b>Mortara Instrument, Inc.</b>
Location	Walker, MI, US
510(k) history	51 submissions · 51 cleared · 1983-2019

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k832050/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 19, 2026