

**K875123 ORCEPAD**Jun 21, 1988  
189 days to decisionK875123 · Product code: **HDD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k875123/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Obstetrical, Ac-powered (and Accessories) (HDD)
Date received	Dec 15, 1987
Decision date	Jun 21, 1988
Days to decision	189 days
Third-party review	No

**APPLICANT**

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Company	<b>Alton Dean Medical, Inc.</b>
Location	North Salt Lake City, UT, US
Contact	DIXON FORD
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k875123/>; 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 30, 2026