

**K994079 REFLEX TREATMENT SYSTEM**Mar 1, 2000  
90 days to decisionK994079 · Product code: **HIR** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k994079/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Perineometer (HIR)
Date received	Dec 2, 1999
Decision date	Mar 1, 2000
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Deschutes Medical Products, Inc.</b>
Location	Bend, OR, US
Contact	DENISE BESTWICK
510(k) history	8 submissions · 8 cleared · 1995-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k994079/> 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 29, 2026