

**K800191 PERITONE**Apr 16, 1980  
78 days to decisionK800191 · Product code: **HIR** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k800191/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Perineometer (HIR)
Date received	Jan 29, 1980
Decision date	Apr 16, 1980
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Laurie Ann Kaiser, Ph.D.</b>
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1980-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k800191/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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