

**K780797 TRANSDUCER, PRESSURE, OPTION, H-P, P23ID**Jun 13, 1978  
29 days to decisionK780797 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k780797/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	May 15, 1978
Decision date	Jun 13, 1978
Days to decision	29 days
Third-party review	No

**APPLICANT**

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Company	<b>Gould, Inc.</b>
Location	Mchenry, IL, US
Contact	none none
510(k) history	31 submissions · 31 cleared · 1976-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k780797/>; 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 May 27, 2026