

**K855145 MODEL:TA-ROSE RESONANT OVERSHOOT  
ELIMINATOR**Feb 26, 1986  
64 days to decisionK855145 · Product code: **DRS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k855145/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Dec 24, 1985
Decision date	Feb 26, 1986
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Gould, Inc.</b>
Location	Mchenry, IL, US
Contact	GORDON PH.D.
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/k855145/>; 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