FDA 510(k) Premarket Notification

K855145 MODEL:TA-ROSE RESONANT OVERSHOOT ELIMINATOR

CLEARED

Feb 26, 1986
64 days to decision

K855145 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k855145/

SUBMISSION DETAILS

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

Company	**Gould, Inc.**
Location	Mchenry, IL, US
Contact	GORDON PH.D.
510(k) history	31 submissions · 31 cleared · 1976-1991