FDA 510(k) Premarket Notification

K833541 SURGICAL INSTRUMENT

CLEARED

Nov 28, 1983
62 days to decision

K833541 · Product code: **GAD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k833541/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Sep 27, 1983
Decision date	Nov 28, 1983
Days to decision	62 days
Third-party review	No

APPLICANT

Company	**Hirata Sangyo Co. USA, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1982-1986