FDA 510(k) Premarket Notification

K811759 RETRACTING PROBES

CLEARED

Jul 20, 1981
28 days to decision

K811759 · Product code: **GAD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k811759/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Jun 22, 1981
Decision date	Jul 20, 1981
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**American V. Mueller**
Location	Mchenry, IL, US
510(k) history	31 submissions · 29 cleared · 1980-1987

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k811759/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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