FDA 510(k) Premarket Notification

K875043 RONGEUR (NEUROSURGICAL INSTRUMENTS)

CLEARED

Jan 22, 1988
51 days to decision

K875043 · Product code: **HAE** · Neurology
Source: https://www.510kdatabase.net/k875043/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Rongeur, Manual (HAE)
Date received	Dec 2, 1987
Decision date	Jan 22, 1988
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Solway, Inc.**
Location	Hollywood, FL, US
Contact	MARTIN MUNZER
510(k) history	13 submissions · 13 cleared · 1987-1988