

K875043 RONGEUR (NEUROSURGICAL INSTRUMENTS)Jan 22, 1988
51 days to decisionK875043 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k875043/>; Data sourced from FDA 510(k) public records (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. - Generated May 22, 2026