

K893121 ROBOTRAC (TM) RETRACTOR ARMJun 1, 1989
37 days to decisionK893121 · Product code: **GAD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k893121/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Apr 25, 1989
Decision date	Jun 1, 1989
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Aesculap Instruments Corp.
Location	Mchenry, IL, US
Contact	ROBIN BUSH
510(k) history	13 submissions · 13 cleared · 1977-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893121/>; 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