

K820290 SELF RETAINING RETRACTOR FOR HAND SURGMar 5, 1982
30 days to decisionK820290 · Product code: **GAD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820290/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Feb 3, 1982
Decision date	Mar 5, 1982
Days to decision	30 days
Third-party review	No

APPLICANT

Company	The Anspach Effort, Inc.
Location	Mchenry, IL, US
510(k) history	60 submissions · 60 cleared · 1980-2022

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820290/> 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 June 28, 2026