

**K801759 EFFRON PAROTID RETRACTOR**Aug 20, 1980  
23 days to decisionK801759 · Product code: **GAD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k801759/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Jul 28, 1980
Decision date	Aug 20, 1980
Days to decision	23 days
Third-party review	No

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k801759/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026