

**K791983 SPATULA'S #S  
7618,7248,7240,7242,5108**Oct 4, 1979  
9 days to decisionK791983 · Product code: **HND** · Ophthalmic  
Source: <https://www.510kdatabase.net/k791983/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spatula, Ophthalmic (HND)
Date received	Sep 25, 1979
Decision date	Oct 4, 1979
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Edward Weck, Inc.</b>
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k791983/> 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 May 23, 2026