

**K772371 BAYONET SURGICAL INSTRUMENTS**Jan 6, 1978  
10 days to decisionK772371 · Product code: **HAO** · Neurology  
Source: <https://www.510kdatabase.net/k772371/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Non-powered (HAO)
Date received	Dec 27, 1977
Decision date	Jan 6, 1978
Days to decision	10 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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Device record: <https://www.510kdatabase.net/k772371/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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