

**K833541 SURGICAL INSTRUMENT**Nov 28, 1983  
62 days to decisionK833541 · Product code: **GAD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k833541/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retractor (GAD)
Date received	Sep 27, 1983
Decision date	Nov 28, 1983
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Hirata Sangyo Co. USA, Inc.</b>
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1982-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k833541/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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