

**K801908 SI-14**Nov 12, 1980  
93 days to decisionK801908 · Product code: **KDC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k801908/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Aug 11, 1980
Decision date	Nov 12, 1980
Days to decision	93 days
Third-party review	No

**APPLICANT**

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Company	<b>Kmi Medical, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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

Device record: <https://www.510kdatabase.net/k801908/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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