

**K896434 AUTO SUTURE SURGIPOINT CANNULA DIAMETER REDUCER\***Nov 16, 1989  
7 days to decisionK896434 · Product code: **GEA** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k896434/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Nov 9, 1989
Decision date	Nov 16, 1989
Days to decision	7 days
Third-party review	No

**APPLICANT**

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Company	<b>United States Surgical, A Division of Tyco Healthc</b>
Location	Mchenry, IL, US
Contact	CURTIS RAYMOND
510(k) history	218 submissions · 200 cleared · 1977-2007

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896434/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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