

**K861742 ELECTRO-SURGICAL PENCILS**Jul 23, 1986  
78 days to decisionK861742 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k861742/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 6, 1986
Decision date	Jul 23, 1986
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Stryker Corp.</b>
Location	Mchenry, IL, US
Contact	HARMON H WOODWORTH
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...

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