

**K802330 HI-TEMP LOOPTIP SURGICAL CAUTERY**Nov 12, 1980  
50 days to decisionK802330 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k802330/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 23, 1980
Decision date	Nov 12, 1980
Days to decision	50 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Products Development, Inc.</b>
Location	Mchenry, IL, US
510(k) history	13 submissions · 12 cleared · 1980-1981

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

Device record: <https://www.510kdatabase.net/k802330/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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