

**K083711 MODIFICATION TO: HALO COAGULATION
CATHETER, MODELS 32041-18, 32041-20, 32041-22, 32041-25,
32041-28, 32041-31, 32041-34**

Feb 2, 2009
49 days to decision

K083711 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k083711/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 15, 2008
Decision date	Feb 2, 2009
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Barrx Medical, Incorporated
Location	Washington, Dc, DC, US
Contact	VIORICA FILIMON
510(k) history	13 submissions · 13 cleared · 2005-2010

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k083711/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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