

**K093086 ARTHROSTEER ABLATOR & COAGULATOR,
MODELS: REF 1070 & 1080**Feb 23, 2010
145 days to decisionK093086 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k093086/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 1, 2009
Decision date	Feb 23, 2010
Days to decision	145 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Orthodynamix, LLC
Location	Jacksonville, FL, US
Contact	BRIAN S ZIMOVAN
510(k) history	1 submissions · 1 cleared · 2010-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k093086/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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