

K123310 FINESSE 3RD GENERATION (MODELS FINESSE+ AND FINESSEII+)Jan 23, 2013
90 days to decisionK123310 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k123310/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 25, 2012
Decision date	Jan 23, 2013
Days to decision	90 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Utah Medical Products, Inc.
Location	Mchenry, IL, US
Contact	BEN SHIRLEY
510(k) history	38 submissions · 38 cleared · 1979-2015

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