

**K063025 LINA POWERBLADE, MODEL 5005,5005-E,5005-C,5005  
-150,5000-C-150,5005-E-  
C-150,5005-420,5005-C-420,5005-E-420,5005-E-C-420,**

Dec 1, 2006  
60 days to decision

K063025 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k063025/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 2, 2006
Decision date	Dec 1, 2006
Days to decision	60 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lina Medical Aps</b>
Location	Chapel Hill, NC, US
Contact	WALT BRITTLE
510(k) history	16 submissions · 16 cleared · 2006-2024

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

Device record: <https://www.510kdatabase.net/k063025/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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