

**K092761 PROPLATE, MODELS 9641C, 9641F, P9641C, P9641F, 9642C, 9642F AND P9642C**

Nov 24, 2009  
76 days to decision

K092761 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k092761/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 9, 2009
Decision date	Nov 24, 2009
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Bio Protech, Inc.</b>
Location	Sparks, NV, US
Contact	JUDY BURTON
510(k) history	17 submissions · 17 cleared · 2002-2019

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

Device record: <https://www.510kdatabase.net/k092761/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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