

**K905108 CONTORNO(R), MODELS E AND ST**Feb 4, 1991  
83 days to decisionK905108 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k905108/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 13, 1990
Decision date	Feb 4, 1991
Days to decision	83 days
Third-party review	No

**APPLICANT**

---

Company	<b>Palex Intl. SA</b>
Location	Spain, ES
Contact	MARTIN A JORDAN
510(k) history	5 submissions · 5 cleared · 1990-1991

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905108/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026