

**K963307 GDC PATIENT RETURN ELECTTRODE MODEL 45021**Dec 17, 1996  
117 days to decisionK963307 · Product code: **HCG** · Neurology  
Source: <https://www.510kdatabase.net/k963307/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Neurovascular Embolization (HCG)
Date received	Aug 22, 1996
Decision date	Dec 17, 1996
Days to decision	117 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Target Therapeutics</b>
Location	Los Angeles, CA, US
Contact	LARAIN PANGELINA
510(k) history	70 submissions · 70 cleared · 1985-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k963307/> 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 June 30, 2026