

**K250688 Cedar™ Endovenous Radiofrequency Catheter (ERA-C70-US)**

Oct 7, 2025  
215 days to decision

K250688 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k250688/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 6, 2025
Decision date	Oct 7, 2025
Days to decision	215 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Cedar™ Endovenous Radiofrequency Catheter (ERA-C30-US); Endovenous Radiofrequency Generator (ERA-G5-US)

**APPLICANT**

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Company	<b>Acotec Scientific Co., Ltd.</b>
Location	Beijing, CN
Contact	Anna Chan
510(k) history	1 submissions · 1 cleared · 2025-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Fabrica Consulting, LLC</b>
Contact	Sara Toyloy

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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