

**K172049 CryoIQ DERM, CryoIQ PRO, CryoIQ EquiMED**Aug 30, 2017  
55 days to decisionK172049 · Product code: **GEH** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k172049/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryosurgical, Accessories (GEH)
Date received	Jul 6, 2017
Decision date	Aug 30, 2017
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cry IQ AB</b>
Location	Onsala, SE
Contact	Stefan Skafte
510(k) history	1 submissions · 1 cleared · 2017-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172049/> 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 31, 2026