FDA 510(k) Premarket Notification

K172049 CryoIQ DERM, CryoIQ PRO, CryoIQ EquiMED

CLEARED

Aug 30, 2017
55 days to decision

K172049 · Product code: **GEH** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k172049/

SUBMISSION DETAILS

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

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