FDA 510(k) Premarket Notification

K081559 GENESIS

CLEARED

Aug 28, 2008
86 days to decision

K081559 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k081559/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 3, 2008
Decision date	Aug 28, 2008
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medisiss**
Location	Sisters, OR, US
Contact	BRANDI JAMES
510(k) history	6 submissions · 6 cleared · 2003-2011