FDA 510(k) Premarket Notification

K982472 GENICON TROCAR

CLEARED

Feb 4, 1999
204 days to decision

K982472 · Product code: **GEA** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k982472/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Jul 15, 1998
Decision date	Feb 4, 1999
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Genicon, LC**
Location	Orlando, FL, US
Contact	GARY W HABERLAND
510(k) history	6 submissions · 6 cleared · 1999-2003