FDA 510(k) Premarket Notification

K831912 ROUND-BEEHIVE- DISSECTORS 3/8

CLEARED

Oct 31, 1983
139 days to decision

K831912 · Product code: **GDI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k831912/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dissector, Surgical, General & Plastic Surgery (GDI)
Date received	Jun 14, 1983
Decision date	Oct 31, 1983
Days to decision	139 days
Third-party review	No

APPLICANT

Company	**Ormed Mfg., Inc.**
Location	Mchenry, IL, US
510(k) history	23 submissions · 23 cleared · 1983-1994