FDA 510(k) Premarket Notification

K223107 MOLLI 2

CLEARED

Jan 18, 2023
110 days to decision

K223107 · Product code: **NEU** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k223107/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Radiographic, Implantable (NEU)
Date received	Sep 30, 2022
Decision date	Jan 18, 2023
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Molli Surgical, Inc.**
Location	Toronto, CA
Contact	Joseph De Croos
510(k) history	5 submissions · 5 cleared · 2021-2024