FDA 510(k) Premarket Notification

K241700 Tenex 2nd Generation System

CLEARED

Nov 18, 2024
158 days to decision

K241700 · Product code: **LFL** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k241700/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jun 13, 2024
Decision date	Nov 18, 2024
Days to decision	158 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Trice Medical, Inc.**
Location	King Of Prussia, PA, US
Contact	David Vancelette
510(k) history	3 submissions · 3 cleared · 2014-2024