FDA 510(k) Premarket Notification

K230382 3 Series NeoLux

CLEARED

Mar 24, 2023
39 days to decision

K230382 · Product code: **FTC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k230382/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Light, Ultraviolet, Dermatological (FTC)
Date received	Feb 13, 2023
Decision date	Mar 24, 2023
Days to decision	39 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Daavlin Distributing Co.**
Location	Bryan, OH, US
Contact	Michele Thiel
510(k) history	15 submissions · 15 cleared · 2002-2023