FDA 510(k) Premarket Notification

K221131 Tasso+

CLEARED

Aug 12, 2022
116 days to decision

K221131 · Product code: **FMK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k221131/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Apr 18, 2022
Decision date	Aug 12, 2022
Days to decision	116 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Tasso, Inc.**
Location	Seattle, WA, US
Contact	Trish Kan Brown
510(k) history	1 submissions · 1 cleared · 2022-2022