

K221131 Tasso+Aug 12, 2022
116 days to decisionK221131 · Product code: **FMK** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k221131/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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