

K202833 Sequel External Fixation DeviceJan 19, 2021
116 days to decisionK202833 · Product code: **KTT** · Orthopedic
Source: <https://www.510kdatabase.net/k202833/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Sep 25, 2020
Decision date	Jan 19, 2021
Days to decision	116 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Sequel Medical, Inc.
Location	North Miami, FL, US
Contact	Valentina Kamysheva
510(k) history	1 submissions · 1 cleared · 2021-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k202833/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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