

K230789 Knee+Apr 21, 2023
30 days to decisionK230789 · Product code: **SBF** · Orthopedic
Source: <https://www.510kdatabase.net/k230789/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthopedic Augmented Reality (SBF)
Date received	Mar 22, 2023
Decision date	Apr 21, 2023
Days to decision	30 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Pixee Medical
Location	Besançon, FR
Contact	Lucie Pécheur
510(k) history	6 submissions · 6 cleared · 2021-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230789/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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