

K250617 Apex FlexApr 29, 2025
60 days to decisionK250617 · Product code: **EBI** · DentalSource: <https://www.510kdatabase.net/k250617/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Feb 28, 2025
Decision date	Apr 29, 2025
Days to decision	60 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Sprintray, Inc.
Location	Los Angeles, CA, US
Contact	Sara Moghtadernejad
510(k) history	9 submissions · 9 cleared · 2021-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k250617/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026