FDA 510(k) Premarket Notification

K182422 Lucitone Denture Base Materials

CLEARED

Dec 18, 2018
103 days to decision

K182422 · Product code: **EBI** · Dental
Source: https://www.510kdatabase.net/k182422/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Sep 6, 2018
Decision date	Dec 18, 2018
Days to decision	103 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Dentsply Sirona**
Location	York, PA, US
Contact	Karl Nittinger
Website	https://www.dentsplysirona.com
510(k) history	65 submissions · 65 cleared · 2016-2026

Dentsply Sirona is an American dental equipment manufacturer and consumables producer headquartered in York, US. The company markets products in over 120 countries and operates factories across 21 nations. Dentsply Sirona has received FDA 510(k) clearances from total submissions since 2016. Dental devices represent 78% of the company's regulatory submissions, reflecting its core focus on laboratory equipment, specialty products, and consumables including abutments, CAD/CAM blocks, and restorative materials. The company's latest clearance in 2026 demonstrates continued reg...