

**K181331 JUVORA Oyster White Dental Disc, CERAMILL PEEK
by JUVORA, Oyster White**

Dec 21, 2018
214 days to decision

K181331 · Product code: **EBI** · Dental
Source: <https://www.510kdatabase.net/k181331/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	May 21, 2018
Decision date	Dec 21, 2018
Days to decision	214 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Juvora, Ltd.
Location	Thornton-Clevelys, GB
Contact	Tim Leyva
510(k) history	1 submissions · 1 cleared · 2018-2018

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k181331/> 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 June 28, 2026