

K221695 3M Filtek Supreme Flowable RestorativeJun 13, 2022
3 days to decisionK221695 · Product code: **EBF** · Dental
Source: <https://www.510kdatabase.net/k221695/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Material, Tooth Shade, Resin (EBF)
Date received	Jun 10, 2022
Decision date	Jun 13, 2022
Days to decision	3 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	3M Company Espe Dental Products
Location	Saint Paul, MN, US
Contact	Regina Feferman-Savvateev
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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Device record: <https://www.510kdatabase.net/k221695/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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