

K892517 TULUX-FILTERAug 21, 1989
132 days to decisionK892517 · Product code: **EBZ** · DentalSource: <https://www.510kdatabase.net/k892517/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Activator, Ultraviolet, For Polymerization (EBZ) |
| Date received | Apr 11, 1989 |
| Decision date | Aug 21, 1989 |
| Days to decision | 132 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Espe GmbH (Us) |
| Location | Mchenry, IL, US |
| Contact | HAROLD OPOTOW |
| 510(k) history | 59 submissions · 57 cleared · 1977-1993 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k892517/>; 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 July 3, 2026