

K973924 FRIOS BONESHIELDMay 7, 1998
204 days to decision

K973924 · Product code: JEY · Dental

Source: <https://www.510kdatabase.net/k973924/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Oct 15, 1997
Decision date	May 7, 1998
Days to decision	204 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Friatec AG
Location	Potomac, MD, US
Contact	ROBERT J SLOMOFF
Website	http://www.friatec.de/
510(k) history	6 submissions · 6 cleared · 1998-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k973924/> 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 2, 2026