

K962824 IMPLANTECH JEJ PERI-PYRIFORM IMPLANTOct 3, 1996
76 days to decisionK962824 · Product code: **FZE** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k962824/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Nose, Internal (FZE)
Date received	Jul 19, 1996
Decision date	Oct 3, 1996
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Implantech Associates, Inc.
Location	Washington, DC, US
Contact	EDWARD M BASILE
510(k) history	41 submissions · 40 cleared · 1989-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962824/> 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 20, 2026