FDA 510(k) Premarket Notification

K962824 IMPLANTECH JEJ PERI-PYRIFORM IMPLANT

CLEARED

Oct 3, 1996
76 days to decision

K962824 · Product code: **FZE** · General & Plastic Surgery
Source: 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