FDA 510(k) Premarket Notification

K895152 BINDER SUBMALAR FACIAL IMPLANT II

CLEARED

Oct 27, 1989
78 days to decision

K895152 · Product code: **LZK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k895152/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Malar (LZK)
Date received	Aug 10, 1989
Decision date	Oct 27, 1989
Days to decision	78 days
Third-party review	No

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.net
Device record: https://www.510kdatabase.net/k895152/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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