FDA 510(k) Premarket Notification

K021419 AART MALAR IMPLANT

CLEARED

Jul 2, 2002
60 days to decision

K021419 · Product code: **LZK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k021419/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Malar (LZK)
Date received	May 3, 2002
Decision date	Jul 2, 2002
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Aesthetic and Reconstructive Technologies, Inc.**
Location	Paso Robles, CA, US
Contact	CATHERINE RIPLE
510(k) history	10 submissions · 10 cleared · 2002-2003