FDA 510(k) Premarket Notification

K022223 AART SILICONE SHEETING

CLEARED

Aug 8, 2002
30 days to decision

K022223 · Product code: **MIB** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k022223/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Elastomer, Silicone Block (MIB)
Date received	Jul 9, 2002
Decision date	Aug 8, 2002
Days to decision	30 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