

K022223 AART SILICONE SHEETINGAug 8, 2002
30 days to decisionK022223 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k022223/> 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 July 3, 2026