FDA 510(k) Premarket Notification

K903329 CUI ANATOMICAL SILICONE MALAR

CLEARED

Sep 10, 1990
47 days to decision

K903329 · Product code: **LZK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k903329/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Malar (LZK)
Date received	Jul 25, 1990
Decision date	Sep 10, 1990
Days to decision	47 days
Third-party review	No

APPLICANT

Company	**Cui Corp.**
Location	Santa Barbara, CA, US
Contact	MORRIS SHERWOOD
510(k) history	7 submissions · 7 cleared · 1990-1991