

**K021419 AART MALAR IMPLANT**Jul 2, 2002  
60 days to decisionK021419 · Product code: **LZK** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k021419/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k021419/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 3, 2026