

**K885205 AQUADERM**Mar 29, 1989  
100 days to decisionK885205 · Product code: **KGX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k885205/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Tape And Bandage, Adhesive (KGX)
Date received	Dec 19, 1988
Decision date	Mar 29, 1989
Days to decision	100 days
Third-party review	No

**APPLICANT**

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Company	<b>Laivan Corp.</b>
Location	East Rockaway, NY, US
Contact	JOEL MARTZ
510(k) history	6 submissions · 2 cleared · 1988-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k885205/>; 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 7, 2026