

**K020265 SERRALENE, MODEL CATALOG NO 1S**Mar 8, 2002  
42 days to decisionK020265 · Product code: **GAW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k020265/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable, Synthetic, Polypropylene (GAW)
Date received	Jan 25, 2002
Decision date	Mar 8, 2002
Days to decision	42 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Serral, S.A. DE C.V.</b>
Location	Williamsburg, VA, US
Contact	SCOTT HENDERSON
510(k) history	6 submissions · 6 cleared · 2002-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020265/> 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