

**K913303 LAP SPONGES, MODIFIED**Aug 6, 1992  
378 days to decisionK913303 · Product code: **GDY** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k913303/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal, X-ray Detectable (GDY)
Date received	Jul 25, 1991
Decision date	Aug 6, 1992
Days to decision	378 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Carwild Corp.</b>
Location	Mchenry, IL, US
Contact	JOEL S WILDSTEIN
510(k) history	6 submissions · 6 cleared · 1979-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913303/> 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