

**K933950 LADY&apos;S CARE MAXI PADS**Apr 13, 1994  
244 days to decisionK933950 · Product code: **HHD** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k933950/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pad, Menstrual, Unscented (HHD)
Date received	Aug 12, 1993
Decision date	Apr 13, 1994
Days to decision	244 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lady&amp;apos;S Care, Inc.</b>
Location	Southampton, PA, US
Contact	LEE BORDEAUX
510(k) history	2 submissions · 2 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933950/> 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 June 29, 2026