FDA 510(k) Premarket Notification

K933950 LADY'S CARE MAXI PADS

CLEARED

Apr 13, 1994
244 days to decision

K933950 · Product code: **HHD** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k933950/

SUBMISSION DETAILS

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

Company	**Lady'S Care, Inc.**
Location	Southampton, PA, US
Contact	LEE BORDEAUX
510(k) history	2 submissions · 2 cleared · 1994-1994