FDA 510(k) Premarket Notification

K953700 UNSCENTED CURVED MAXI PAD

CLEARED

Sep 27, 1995
50 days to decision

K953700 · Product code: **HHD** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k953700/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pad, Menstrual, Unscented (HHD)
Date received	Aug 8, 1995
Decision date	Sep 27, 1995
Days to decision	50 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Paragon Trade Brands, Inc.**
Location	Federal Way, WA, US
Contact	TONY SILWANOWICZ
510(k) history	14 submissions · 14 cleared · 1995-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k953700/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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