

K880734 SCENTED OR SCENTED DEODORIZED MENSTRUAL PADSApr 25, 1988
61 days to decisionK880734 · Product code: **HHI** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k880734/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Abortion, Vacuum (HHI)
Date received	Feb 24, 1988
Decision date	Apr 25, 1988
Days to decision	61 days
Third-party review	No

APPLICANT

Company	Personal Products Co.
Location	Mchenry, IL, US
Contact	A. J HUETTEMAN
510(k) history	45 submissions · 45 cleared · 1976-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880734/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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