

**K834144 UNSCENTED MENSTRUAL PAD**Dec 29, 1983  
28 days to decisionK834144 · Product code: **HHD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k834144/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pad, Menstrual, Unscented (HHD)
Date received	Dec 1, 1983
Decision date	Dec 29, 1983
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Procter &amp; Gamble Mfg. Co.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.pg.com">https://www.pg.com</a>
510(k) history	72 submissions · 72 cleared · 1976-1988

Procter & Gamble Mfg. Co. is a consumer health and personal care manufacturer headquartered in McHenry, US. The company has a long history of developing products across multiple healthcare categories. The company received FDA 510(k) clearances from total submissions, with no denied submissions on record. Regulatory activity spans from 1976 to 1988, establishing a historical record primarily in Obstetrics & Gynecology and Gastroenterology & Urology device categories. The company is inactive in the FDA 510(k) clearance database, with no submissions recorded in the past seven...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k834144/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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