

**K060852 MOONCUP**Dec 26, 2006  
273 days to decisionK060852 · Product code: **HHE** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k060852/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cup, Menstrual (HHE)
Date received	Mar 28, 2006
Decision date	Dec 26, 2006
Days to decision	273 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mooncup Limited</b>
Location	Brighton, GB
Contact	SU HARDY
510(k) history	1 submissions · 1 cleared · 2006-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060852/> 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