FDA 510(k) Premarket Notification

K060852 MOONCUP

CLEARED

Dec 26, 2006
273 days to decision

K060852 · Product code: **HHE** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k060852/

SUBMISSION DETAILS

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

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