

**K233361 Sunny Cup and Applicator**Jun 21, 2024  
266 days to decisionK233361 · Product code: **HHE** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k233361/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cup, Menstrual (HHE)
Date received	Sep 29, 2023
Decision date	Jun 21, 2024
Days to decision	266 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Menstrual Mates</b>
Location	Lewes, DE, US
Contact	Cindy Belardo
510(k) history	1 submissions · 1 cleared · 2024-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233361/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026