FDA 510(k) Premarket Notification

K060526 TIPI

CLEARED

Jun 9, 2006
102 days to decision

K060526 · Product code: **HHW** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k060526/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pessary, Vaginal (HHW)
Date received	Feb 27, 2006
Decision date	Jun 9, 2006
Days to decision	102 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Contipi , Ltd.**
Location	Mccordsville, IN, US
Contact	PAUL DRYDEN
510(k) history	2 submissions · 2 cleared · 2006-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k060526/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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