FDA 510(k) Premarket Notification

K040625 ATAD ARD CATHETER

CLEARED

Jan 18, 2005
315 days to decision

K040625 · Product code: **HDY** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k040625/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Cervical, Hygroscopic-laminaria (HDY)
Date received	Mar 9, 2004
Decision date	Jan 18, 2005
Days to decision	315 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Atad Developments , Ltd.**
Location	Haifa, IL
Contact	JACK ATAD
510(k) history	1 submissions · 1 cleared · 2005-2005

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k040625/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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