FDA 510(k) Premarket Notification

K041435 THE STRIPPER PGD

CLEARED

Sep 29, 2004
124 days to decision

K041435 · Product code: **MQH** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k041435/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Microtools, Assisted Reproduction (pipettes) (MQH)
Date received	May 28, 2004
Decision date	Sep 29, 2004
Days to decision	124 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mid-Atlantic Diagnostics, Inc.**
Location	Marlton, NJ, US
Contact	SUSAN J BUSH
510(k) history	4 submissions · 4 cleared · 2000-2010