

K041435 THE STRIPPER PGDSep 29, 2004
124 days to decisionK041435 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k041435/> 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 June 20, 2026