FDA 510(k) Premarket Notification

K103028 EEVATM PETRI DISH

CLEARED

Aug 11, 2011
302 days to decision

K103028 · Product code: **MQK** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k103028/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Labware, Assisted Reproduction (MQK)
Date received	Oct 13, 2010
Decision date	Aug 11, 2011
Days to decision	302 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Auxogyn, Inc.**
Location	Menlo Park, CA, US
Contact	ROBERT NEWMAN
510(k) history	4 submissions · 3 cleared · 2011-2014