FDA 510(k) Premarket Notification

K222606 V-HYADASE

CLEARED

May 26, 2023
270 days to decision

K222606 · Product code: **MQL** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k222606/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Aug 29, 2022
Decision date	May 26, 2023
Days to decision	270 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vitromed GmbH**
Location	Jena, DE
Contact	Greg Holland
510(k) history	7 submissions · 7 cleared · 2020-2025