

K242968 Gifflife® Single-Step Medium PlusNov 22, 2024
57 days to decisionK242968 · Product code: **MQL** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k242968/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Sep 26, 2024
Decision date	Nov 22, 2024
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Gimbo Medical Technology Shenzhen Co., Ltd.
Location	Shenzhen, CN
Contact	Leo Guo
510(k) history	6 submissions · 6 cleared · 2024-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242968/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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