

K242111 VIVACEFeb 6, 2025
202 days to decisionK242111 · Product code: **HGX** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k242111/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Breast, Powered (HGX)
Date received	Jul 19, 2024
Decision date	Feb 6, 2025
Days to decision	202 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Unimom.Co
Location	Osan-Si, KR
Contact	Yeong-Bin Chon
510(k) history	4 submissions · 4 cleared · 2018-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242111/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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