

K072747 UNIDUS MAGNUM MALE LATEX CONDOMNov 15, 2007
49 days to decisionK072747 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k072747/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Condom (HIS)
Date received	Sep 27, 2007
Decision date	Nov 15, 2007
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Unidus Corp.
Location	Seoul, Korea, KR
Contact	CLAUDE L WRIGHT
510(k) history	7 submissions · 7 cleared · 1989-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072747/> 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 26, 2026