

**K993408 REMED MALE RIBBED CONDOM PREVENTOR,
LUCKY BOY SUMMIT**Feb 15, 2000
130 days to decisionK993408 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k993408/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Condom (HIS)
Date received	Oct 8, 1999
Decision date	Feb 15, 2000
Days to decision	130 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Remed Pharma
Location	Crofton, MD, US
Contact	E.J. Smith
510(k) history	9 submissions · 9 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k993408/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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