FDA 510(k) Premarket Notification

K991322 ENHANCE-S PLUS

CLEARED

Jul 26, 1999
98 days to decision

K991322 · Product code: **MQL** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k991322/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Apr 19, 1999
Decision date	Jul 26, 1999
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Conception Technology, Inc.**
Location	Irvine, CA, US
Contact	GREG HOLLAND
510(k) history	5 submissions · 5 cleared · 1998-2003