FDA 510(k) Premarket Notification

K992071 TRANSVAGINAL ULTRASOUND PROBE HOLDER DEVICE

CLEARED

Aug 27, 1999
70 days to decision

K992071 · Product code: **HDC** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k992071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tenaculum, Uterine (HDC)
Date received	Jun 18, 1999
Decision date	Aug 27, 1999
Days to decision	70 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ron-Tech Medical , Ltd.**
Location	Petach-Tikva, IL
Contact	AHAVA STEIN
510(k) history	2 submissions · 2 cleared · 1999-2004