

**K992071 TRANSVAGINAL ULTRASOUND PROBE HOLDER  
DEVICE**Aug 27, 1999  
70 days to decisionK992071 · Product code: **HDC** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k992071/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992071/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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