FDA 510(k) Premarket Notification

K990372 MODULAR INSTRUMENT SYSTEM FOR MINIMAL INVASIVE SURGERY

CLEARED

Oct 14, 1999
248 days to decision

K990372 · Product code: **HET** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k990372/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Feb 8, 1999
Decision date	Oct 14, 1999
Days to decision	248 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dr. Fritz GmbH**
Location	Baldwin Park, CA, US
Contact	JAMES CHITTY
510(k) history	1 submissions · 1 cleared · 1999-1999