FDA 510(k) Premarket Notification

K813133 KEYMED KEY LIGHT

CLEARED

Nov 20, 1981
15 days to decision

K813133 · Product code: **HET** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k813133/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Nov 5, 1981
Decision date	Nov 20, 1981
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Keymed, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1981-2010