

K813133 KEYMED KEY LIGHTNov 20, 1981
15 days to decisionK813133 · Product code: **HET** · Obstetrics & GynecologySource: <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

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Device record: <https://www.510kdatabase.net/k813133/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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