

K771961 DUAL INCISION TUBAL OCCLUS. BAND APPLI.Oct 25, 1977
8 days to decisionK771961 · Product code: **HET** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k771961/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Oct 17, 1977
Decision date	Oct 25, 1977
Days to decision	8 days
Third-party review	No

APPLICANT

Company	Kli
Location	Walker, MI, US
510(k) history	18 submissions · 18 cleared · 1977-1980

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Device record: <https://www.510kdatabase.net/k771961/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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