FDA 510(k) Premarket Notification

K771961 DUAL INCISION TUBAL OCCLUS. BAND APPLI.

CLEARED

Oct 25, 1977
8 days to decision

K771961 · Product code: **HET** · Obstetrics & Gynecology
Source: 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