FDA 510(k) Premarket Notification

K823937 SEMINAL COLLECTION DEVICE

CLEARED

Feb 24, 1983
57 days to decision

K823937 · Product code: **HIS** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k823937/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Dec 29, 1982
Decision date	Feb 24, 1983
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Health Development Corp.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1983