FDA 510(k) Premarket Notification

K923176 SURGICAL INSTRUMENT KIT, DISPOSABLE

CLEARED

Jun 17, 1993
370 days to decision

K923176 · Product code: **KDD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k923176/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Jun 12, 1992
Decision date	Jun 17, 1993
Days to decision	370 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ulti-Med Intl., Inc.**
Location	Mchenry, IL, US
Contact	DAVID INSCO
510(k) history	21 submissions · 18 cleared · 1983-2011