FDA 510(k) Premarket Notification

K920989 SURGICAL INSTRUMENT KIT, DISPOSABLE

CLEARED

Oct 19, 1992
231 days to decision

K920989 · Product code: **KDD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k920989/

SUBMISSION DETAILS

Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	Kit, Surgical Instrument, Disposable (KDD)
Date received	Mar 2, 1992
Decision date	Oct 19, 1992
Days to decision	231 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