FDA 510(k) Premarket Notification

K955748 DOYLE EXTRACTOR

CLEARED

Mar 14, 1996
90 days to decision

K955748 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k955748/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Dec 15, 1995
Decision date	Mar 14, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Safetech Intl., Inc.**
Location	North Attleboro, MA, US
Contact	DIANE MINEAR
510(k) history	1 submissions · 1 cleared · 1996-1996