FDA 510(k) Premarket Notification

K923321 EDGE SYRINGES AND NEEDLES

CLEARED

Sep 28, 1993
448 days to decision

K923321 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k923321/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jul 7, 1992
Decision date	Sep 28, 1993
Days to decision	448 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Trico Intl., Inc.**
Location	New York, NY, US
Contact	ESTHER LAI
510(k) history	2 submissions · 2 cleared · 1992-1993