FDA 510(k) Premarket Notification

K926123 DUAL CHAMBER MIXING BAG

CLEARED

Sep 3, 1993
273 days to decision

K926123 · Product code: **KPE** · General Hospital
Source: https://www.510kdatabase.net/k926123/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Dec 4, 1992
Decision date	Sep 3, 1993
Days to decision	273 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Secure Medical, Inc.**
Location	Mundelein, IL, US
Contact	TED D LYJAK
510(k) history	1 submissions · 1 cleared · 1993-1993