FDA 510(k) Premarket Notification

K940073 EXTENSION TUBE, EXTENSION TUBE LENGTH 20, & 30

CLEARED

Mar 23, 1994
76 days to decision

K940073 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k940073/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 6, 1994
Decision date	Mar 23, 1994
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Puritas Health Care, Inc.**
Location	Brookfield, CT, US
Contact	THOMAS P DIMAIO
510(k) history	24 submissions · 22 cleared · 1991-1994