FDA 510(k) Premarket Notification

K922110 GO-CLEAN

CLEARED

Mar 9, 1994
673 days to decision

K922110 · Product code: **FRR** · General Hospital
Source: https://www.510kdatabase.net/k922110/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chamber, Reverse Isolation, Patient Care (FRR)
Date received	May 5, 1992
Decision date	Mar 9, 1994
Days to decision	673 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Viva-Tek Assoc., Inc.**
Location	Medford, NJ, US
Contact	RONALD D RICCIUTI
510(k) history	2 submissions · 2 cleared · 1993-1994