

K922110 GO-CLEANMar 9, 1994
673 days to decisionK922110 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k922110/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026