FDA 510(k) Premarket Notification

K792710 PARA-PAK CLEAN VIAL

CLEARED

Jan 28, 1980
32 days to decision

K792710 · Product code: **FMH** · General Hospital
Source: https://www.510kdatabase.net/k792710/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Specimen, Sterile (FMH)
Date received	Dec 27, 1979
Decision date	Jan 28, 1980
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Meridian Diagnostics, Inc.**
Location	Walker, MI, US
510(k) history	92 submissions · 92 cleared · 1980-1999