

K792710 PARA-PAK CLEAN VIALJan 28, 1980
32 days to decisionK792710 · 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

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Device record: <https://www.510kdatabase.net/k792710/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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