FDA 510(k) Premarket Notification

K800309 SPECIMEN COLLECTION PAN

CLEARED

Feb 26, 1980
14 days to decision

K800309 · Product code: **FMH** · General Hospital
Source: https://www.510kdatabase.net/k800309/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Specimen, Sterile (FMH)
Date received	Feb 12, 1980
Decision date	Feb 26, 1980
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Sage Products, Inc.**
Location	Mchenry, IL, US
510(k) history	31 submissions · 30 cleared · 1978-1998