FDA 510(k) Premarket Notification

K811234 I.V. BOTTLE (RIDGID)

CLEARED

Jun 18, 1981
45 days to decision

K811234 · Product code: **KPE** · General Hospital
Source: https://www.510kdatabase.net/k811234/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	May 4, 1981
Decision date	Jun 18, 1981
Days to decision	45 days
Third-party review	No

APPLICANT

Company	**Medi-Plast Intl., Inc.**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1981-1996