

K811234 I.V. BOTTLE (RIDGID)Jun 18, 1981
45 days to decisionK811234 · 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

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Device record: <https://www.510kdatabase.net/k811234/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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