

K782113 CAP, TAMPER PROOF PLASTICJun 1, 1979
164 days to decisionK782113 · Product code: **KPE** · General HospitalSource: <https://www.510kdatabase.net/k782113/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Dec 19, 1978
Decision date	Jun 1, 1979
Days to decision	164 days
Third-party review	No

APPLICANT

Company	Acacia, Inc.
Location	Mchenry, IL, US
510(k) history	17 submissions · 17 cleared · 1979-2014

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Device record: <https://www.510kdatabase.net/k782113/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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