FDA 510(k) Premarket Notification

K782113 CAP, TAMPER PROOF PLASTIC

CLEARED

Jun 1, 1979
164 days to decision

K782113 · Product code: **KPE** · General Hospital
Source: 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