

K801652 TOTAL PARENTERAL NUTRITION SET BAGAug 27, 1980
40 days to decisionK801652 · Product code: **KPE** · General Hospital
Source: <https://www.510kdatabase.net/k801652/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Jul 18, 1980
Decision date	Aug 27, 1980
Days to decision	40 days
Third-party review	No

APPLICANT

Company	Cutter Laboratories, Inc.
Location	Mchenry, IL, US
Website	https://www.bayer.com
510(k) history	39 submissions · 39 cleared · 1976-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k801652/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026