

**K802966 I.V. ADDITIVE CAP**Jan 7, 1981  
47 days to decisionK802966 · Product code: **KPE** · General Hospital  
Source: <https://www.510kdatabase.net/k802966/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Nov 21, 1980
Decision date	Jan 7, 1981
Days to decision	47 days
Third-party review	No

**APPLICANT**

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Company	<b>Baxa Corp., Sub. of Cook Group, Inc.</b>
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1978-1991

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Device record: <https://www.510kdatabase.net/k802966/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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