

**K002449 BIOTEQ I.V. (INTRAVENOUS) SET**Dec 22, 2000  
135 days to decisionK002449 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k002449/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 9, 2000
Decision date	Dec 22, 2000
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bioteque Corp.</b>
Location	Levittown, PA, US
Contact	JENNIFER CHEN
510(k) history	11 submissions · 10 cleared · 1992-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002449/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 2, 2026