

**K032789 REPLACEMENT PUMP HEAD DOOR ASSEMBLY, P1,  
MODEL BX10468, REPLACEMENT PUMP HEAD DOOR  
ASSEMBLY, P2, MODEL BX10469**

Oct 9, 2003  
31 days to decision

K032789 · Product code: **FRN** · General Hospital  
Source: <https://www.510kdatabase.net/k032789/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Infusion (FRN)
Date received	Sep 8, 2003
Decision date	Oct 9, 2003
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>American I.V. Products, Inc.</b>
Location	Hanover, MD, US
Contact	GREGORY FALK
510(k) history	7 submissions · 7 cleared · 2003-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k032789/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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