

**K121270 BIOJECT NEEDLE-FREE INJECTION MANAGEMENT SYSTEM**Jul 19, 2012  
84 days to decisionK121270 · Product code: **KZE** · General Hospital  
Source: <https://www.510kdatabase.net/k121270/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector, Fluid, Non-electrically Powered (KZE)
Date received	Apr 26, 2012
Decision date	Jul 19, 2012
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bioject Medical Technologies, Inc.</b>
Location	Bedminster, NJ, US
Contact	CHRISTINE BREITBACH
510(k) history	3 submissions · 3 cleared · 2004-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121270/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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