

**K844244 DERATA MEDI-JECTOR PRO**Jul 5, 1985  
246 days to decisionK844244 · Product code: **KZE** · General Hospital  
Source: <https://www.510kdatabase.net/k844244/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector, Fluid, Non-electrically Powered (KZE)
Date received	Nov 1, 1984
Decision date	Jul 5, 1985
Days to decision	246 days
Third-party review	No

**APPLICANT**

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Company	<b>Derata Corp.</b>
Location	Mchenry, IL, US
Contact	KENNETH W DUNLAP
510(k) history	6 submissions · 6 cleared · 1977-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k844244/>; 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 July 8, 2026