

**K832683 MONOJECT INJECTOMATIC**Sep 12, 1983  
34 days to decisionK832683 · Product code: **KZH** · General HospitalSource: <https://www.510kdatabase.net/k832683/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Syringe Needle (KZH)
Date received	Aug 9, 1983
Decision date	Sep 12, 1983
Days to decision	34 days
Third-party review	No

**APPLICANT**

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Company	<b>Sherwood Medical Co.</b>
Location	Mchenry, IL, US
510(k) history	191 submissions · 177 cleared · 1976-1998

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

Device record: <https://www.510kdatabase.net/k832683/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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