

**K770944 INSTATEMP**Jul 27, 1977  
63 days to decisionK770944 · Product code: **KDP** · General Hospital  
Source: <https://www.510kdatabase.net/k770944/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	May 25, 1977
Decision date	Jul 27, 1977
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>H.I. Silverman, D.Sc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977

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

Device record: <https://www.510kdatabase.net/k770944/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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