

**K760316 REGULATORS, LOW SUCTION**Aug 4, 1976  
9 days to decisionK760316 · Product code: **KDP** · General Hospital  
Source: <https://www.510kdatabase.net/k760316/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	Jul 26, 1976
Decision date	Aug 4, 1976
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Harris-Lake, Inc.</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1976-1977

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Device record: <https://www.510kdatabase.net/k760316/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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