

**K801421 TUBE HOLDER**Jul 28, 1980  
41 days to decisionK801421 · Product code: **LDQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k801421/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jun 17, 1980
Decision date	Jul 28, 1980
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Devon Industries, Inc.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.devonmedicalproducts.com">https://www.devonmedicalproducts.com</a>
510(k) history	47 submissions · 46 cleared · 1976-1996

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

Device record: <https://www.510kdatabase.net/k801421/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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