

**K792702 VASPAR SEALANT**Jan 24, 1980  
28 days to decisionK792702 · Product code: **KEQ** · Pathology  
Source: <https://www.510kdatabase.net/k792702/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Mounting, Water Soluble (KEQ)
Date received	Dec 27, 1979
Decision date	Jan 24, 1980
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Meridian Diagnostics, Inc.</b>
Location	Walker, MI, US
510(k) history	92 submissions · 92 cleared · 1980-1999

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

Device record: <https://www.510kdatabase.net/k792702/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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