

**K992032 ACRY ISLAND BORDER DRESSING, MODELS 2031,  
2032, 2033**Aug 19, 1999  
64 days to decisionK992032 · Product code: **KMF** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k992032/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Jun 16, 1999
Decision date	Aug 19, 1999
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Acrymed, Inc.</b>
Location	Lake Oswego, OR, US
Contact	BRUCE GIBBINS
510(k) history	14 submissions · 12 cleared · 1994-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992032/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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