

**K782080 VENEER SYSTEM, MOSTIQUE LAMINATE**Jan 26, 1979  
44 days to decisionK782080 · Product code: **KLE** · DentalSource: <https://www.510kdatabase.net/k782080/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Dec 13, 1978
Decision date	Jan 26, 1979
Days to decision	44 days
Third-party review	No

**APPLICANT**

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Company	<b>Caulk Co., Div. Dentsply</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1981

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

Device record: <https://www.510kdatabase.net/k782080/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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