

K810157 MOI-STIRFeb 10, 1981
20 days to decisionK810157 · Product code: **LFD** · DentalSource: <https://www.510kdatabase.net/k810157/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Saliva, Artificial (LFD)
Date received	Jan 21, 1981
Decision date	Feb 10, 1981
Days to decision	20 days
Third-party review	No

APPLICANT

Company	Kingswood Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1985

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Device record: <https://www.510kdatabase.net/k810157/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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