

K840807 THROAT MOI-STIRMay 25, 1984
91 days to decisionK840807 · Product code: **LFD** · DentalSource: <https://www.510kdatabase.net/k840807/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Saliva, Artificial (LFD)
Date received	Feb 24, 1984
Decision date	May 25, 1984
Days to decision	91 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k840807/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026