FDA 510(k) Premarket Notification

K840807 THROAT MOI-STIR

CLEARED

May 25, 1984
91 days to decision

K840807 · Product code: **LFD** · Dental
Source: 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