

K030802 CAPHOSOLNov 25, 2003
257 days to decisionK030802 · Product code: **LFD** · DentalSource: <https://www.510kdatabase.net/k030802/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Saliva, Artificial (LFD)
Date received	Mar 13, 2003
Decision date	Nov 25, 2003
Days to decision	257 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Inpharma A.S.
Location	Northborough, MA, US
Contact	BRUCE MANNING
510(k) history	2 submissions · 2 cleared · 1999-2003

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