

**K082071 ADPHARMA OCCLUDE AND OCCLUDE F DENTIN
TUBULE AGENT**Oct 2, 2008
72 days to decisionK082071 · Product code: **LBH** · Dental
Source: <https://www.510kdatabase.net/k082071/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Varnish, Cavity (LBH)
Date received	Jul 22, 2008
Decision date	Oct 2, 2008
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Adpharma, Inc.
Location	Arlington Heights, IL, US
Contact	VIVEK RAMANA
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082071/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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