FDA 510(k) Premarket Notification

K082071 ADPHARMA OCCLUDE AND OCCLUDE F DENTIN TUBULE AGENT

CLEARED

Oct 2, 2008
72 days to decision

K082071 · 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