FDA 510(k) Premarket Notification

K103160 NUPRO MODEL 13016901

CLEARED

Jan 6, 2011
72 days to decision

K103160 · Product code: **LBH** · Dental
Source: https://www.510kdatabase.net/k103160/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Varnish, Cavity (LBH)
Date received	Oct 26, 2010
Decision date	Jan 6, 2011
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**CAO Group, Inc.**
Location	West Jordan, UT, US
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510(k) history	31 submissions · 31 cleared · 2001-2026