FDA 510(k) Premarket Notification

K072309 MICROLUX/BLU

CLEARED

Nov 19, 2007
94 days to decision

K072309 · Product code: **EAZ** · Dental
Source: https://www.510kdatabase.net/k072309/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Operating, Dental (EAZ)
Date received	Aug 17, 2007
Decision date	Nov 19, 2007
Days to decision	94 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Addent, Inc.**
Location	Danbury, CT, US
Contact	JOSHUA FRIEDMAN, DDS
510(k) history	7 submissions · 7 cleared · 2005-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k072309/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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