FDA 510(k) Premarket Notification

K062069 ATLANTIS ABUTMENT FOR 3I MICROMINI IMPLANT

CLEARED

Feb 14, 2007
208 days to decision

K062069 · Product code: **NHA** · Dental
Source: https://www.510kdatabase.net/k062069/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jul 21, 2006
Decision date	Feb 14, 2007
Days to decision	208 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Atlantis Components, Inc.**
Location	Skokie, IL, US
Contact	BETSY A BROWN
510(k) history	18 submissions · 18 cleared · 1999-2011