FDA 510(k) Premarket Notification

K103020 ATLANTIS ABUTMENT FOR KEYSTONE IMPLANT, ATLANTIS GEMINI ABUTMENT

CLEARED

Feb 3, 2011
114 days to decision

K103020 · Product code: **NHA** · Dental
Source: https://www.510kdatabase.net/k103020/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Oct 12, 2010
Decision date	Feb 3, 2011
Days to decision	114 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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