

**K103020 ATLANTIS ABUTMENT FOR KEYSTONE IMPLANT,
ATLANTIS GEMINI ABUTMENT**Feb 3, 2011
114 days to decisionK103020 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103020/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 19, 2026