

**K051652 ATLANTIS ABUTMENT, ATLANTIS GEMINI  
ABUTMENT, ATLANTIS GEMINI+ ABUTMENT**Jul 22, 2005  
31 days to decisionK051652 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k051652/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Jun 21, 2005
Decision date	Jul 22, 2005
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051652/> 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 20, 2026