

**K062069 ATLANTIS ABUTMENT FOR 3I MICROMINI IMPLANT**Feb 14, 2007  
208 days to decisionK062069 · 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	<b>Atlantis Components, Inc.</b>
Location	Skokie, IL, US
Contact	BETSY A 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/k062069/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 19, 2026