

K072483 ATLANTIS ABUTMENT IN ZIRCONIA FOR 3I CERTAIN MICROAMINI IMPLANTS, GEMINI ABUTMENT IN ZIRCONIA FOR ASTRA IMPLANTS

Nov 16, 2007
73 days to decision

K072483 · Product code: **NHA** · Dental
Source: <https://www.510kdatabase.net/k072483/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Sep 4, 2007
Decision date	Nov 16, 2007
Days to decision	73 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k072483/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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