

**K011729 SIGMA PACE EXTERNAL PACEMAKER ANALYZER
MODEL #1000**Aug 30, 2001
87 days to decisionK011729 · Product code: **DTC** · Cardiovascular
Source: <https://www.510kdatabase.net/k011729/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function (DTC)
Date received	Jun 4, 2001
Decision date	Aug 30, 2001
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Dni Nevada, Inc.
Location	Carson City, NV, US
Contact	KRISTINE BOGGS
510(k) history	1 submissions · 1 cleared · 2001-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k011729/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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