

**K960506 A-DEC PERFORMER CONTROL UNIT**Mar 25, 1996  
49 days to decisionK960506 · Product code: **EIA** · DentalSource: <https://www.510kdatabase.net/k960506/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Operative Dental (EIA)
Date received	Feb 5, 1996
Decision date	Mar 25, 1996
Days to decision	49 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>A-Dec, Inc.</b>
Location	Newberg, OR, US
Contact	PATIRCK RIDENOUR
Website	<a href="http://www.a-dec.com/">http://www.a-dec.com/</a>
510(k) history	69 submissions · 69 cleared · 1989-2022

A-Dec, Inc. is a dental equipment manufacturer based in Newberg, Oregon. The company designs and produces integrated dental operatory systems, delivery equipment, handpieces, and mechanical room solutions for dental practices worldwide. A-Dec has maintained a strong FDA 510(k) regulatory record since 1989. The company has received FDA 510(k) clearances from total submissions, with no denied submissions. All cleared devices fall within the Dental category, reflecting the company's specialization in dental equipment and accessories. The latest clearance on record dates to 2...

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