

K925319 A-DEC 2182 DOCTOR'S UNITDec 9, 1993
414 days to decisionK925319 · Product code: **KLC** · Dental
Source: <https://www.510kdatabase.net/k925319/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chair, Dental, With Operative Unit (KLC)
Date received	Oct 21, 1992
Decision date	Dec 9, 1993
Days to decision	414 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	A-Dec, Inc.
Location	Newberg, OR, US
Contact	PAT RIDENOUR
Website	http://www.a-dec.com/
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...
