FDA 510(k) Premarket Notification

# K925317 A-DEC 2180 DUO UNIT, CUSP

**CLEARED**

Dec 9, 1993
414 days to decision

K925317 · Product code: **KLC** · Dental
Source: https://www.510kdatabase.net/k925317/

## 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...