

**K231202 QuickRayPRO**Jun 22, 2023  
56 days to decisionK231202 · Product code: **MUH** · Radiology  
Source: <https://www.510kdatabase.net/k231202/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Apr 27, 2023
Decision date	Jun 22, 2023
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Denterprise International, Inc.</b>
Location	Ormond Beach, FL, US
Contact	Claude Berthoin
510(k) history	4 submissions · 4 cleared · 2012-2023

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k231202/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026