

K203502 MEDO-ThyroidApr 23, 2021
144 days to decisionK203502 · Product code: **QIH** · RadiologySource: <https://www.510kdatabase.net/k203502/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated Radiological Image Processing Software (QIH)
Date received	Nov 30, 2020
Decision date	Apr 23, 2021
Days to decision	144 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Medo DX Pte. , Ltd.
Location	Edmonton, CA
Contact	Dornoosh Zonoobi
510(k) history	1 submissions · 1 cleared · 2021-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k203502/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026