

K233539 MAMMOMAT B.brilliantMar 27, 2024
145 days to decisionK233539 · Product code: **MUE** · Radiology
Source: <https://www.510kdatabase.net/k233539/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Full Field Digital, System, X-ray, Mammographic (MUE)
Date received	Nov 3, 2023
Decision date	Mar 27, 2024
Days to decision	145 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
Contact	Denise Adams
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233539/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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