

Date : September 13, 2022

Travel Compliance Letter for SEFAM Medical Devices

To whom it may concern,

SEFAM respiratory assistive devices (CPAP And BPAP machines) are medical devices prescribed for use on or by the order of a physician and are approved for sale in over 60 countries.

The devices do not contain parts or materials known to cause or interfere with the safe operation of a commercial aircraft. The U.S. Department of Transportation's (DOT) final rule, "Non-discrimination on the Basis of Disability in Air Travel" (73 FR 27614, which updates 14 CFR Part 382), effective May 13, 2009, provides important new requirements for accommodating passengers with respiratory assistive devices.

In particular, respiratory assistive devices may be used on board an aircraft without further testing by the carrier, provided they have been tested for electromagnetic compatibility (EMC) in accordance with the current version of RTCA/DO-160, Section 21, Category M.

SEFAM has successfully completed testing for the respiratory assistive devices described in this letter. The devices listed below comply with RTCA/DO-160G, Section 21, Category M, and are considered compliant with the regulations.

SEFAM Medical Device	Commercial reference
Sefam S.Box	M-116412
Sefam S.Box C	M-116411

Not all products are available in each country/region. Please check with your regional representative for specific country availability.



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