

§ 2.951

Laboratories.” Accreditation bodies must be approved by the FCC’s Office of Engineering and Technology, as indicated in § 0.241 of this chapter, to perform such accreditation based on ISO/IEC 58, “Calibration and Testing Laboratory Accreditation Systems—General Requirements for Operation and Recognition.” The frequency for revalidation of the test site and the information required to be filed or retained by the testing party shall comply with the requirements established by the accrediting organization.

(1) In addition to meeting the above requirements, the accreditations of laboratories located outside of the United States or its possessions will be acceptable only under one of the following conditions:

(i) If there is a mutual recognition agreement between that country and the United States and that laboratory is covered by the agreement;

(ii) If there is an agreement between accrediting bodies that permits similar accreditation of U.S. facilities to perform testing for products marketed in that country; or

(iii) If the country already accepts the accreditation of U.S. laboratories.

(2) Organizations outside of the United States that seek to become accreditors may seek agreements with approved United States accrediting bodies to mutually recognize the accreditation of laboratories. The Commission will review such agreements and will consult with the Office of the United States Trade Representative and other Executive Branch agencies before accepting them for purposes of the DoC procedure in order to ensure that the respective foreign countries accept United States accreditations and do not impose additional barriers upon United States companies. Accrediting bodies located outside of the United States will only be permitted to accredit laboratories within their own country for DoC testing.

(3) To facilitate use of the DoC procedure, the FCC will accept a laboratory that submits documentation to OET’s Equipment Authorization Division stating that it has filed an application for accreditation with an approved laboratory accreditation body and provides evidence that it meets all aspects

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of ISO/IEC Guide 25. Such labs will be provisionally accepted by the FCC for a period of one year (until August 19, 1997) or until the application for accreditation has been acted upon, whichever is sooner. A laboratory that is denied accreditation by an approved accreditation body will lose its provisional acceptance. However, any DoCs that were issued will remain valid.

[54 FR 17712, Apr. 25, 1989, as amended at 57 FR 24990, June 12, 1992; 58 FR 37430, July 12, 1993; 58 FR 44893, Aug. 25, 1993; 61 FR 31046, June 19, 1996; 62 FR 41880, Aug. 4, 1997; 63 FR 36599, July 7, 1998; 65 FR 58466, Sept. 29, 2000]

VERIFICATION

AUTHORITY: Sections 2.951 through 2.957 are issued under secs. 4, 303, 307, 48 Stat., as amended, 1066, 1082, 1083; 47 U.S.C. 154, 303, 307.

SOURCE: Sections 2.951 through 2.957 appear at 46 FR 23249, Apr. 24, 1981, unless otherwise noted.

§ 2.951 Cross reference.

The provisions of § 2.901, *et seq.*, shall apply to equipment subject to verification.

§ 2.952 Limitation on verification.

(a) Verification signifies that the manufacturer or importer has determined that the equipment has been shown to be capable of compliance with the applicable technical standards if no unauthorized change is made in the equipment and if the equipment is properly maintained and operated. Compliance with these standards shall not be construed to be a finding by the manufacturer or importer with respect to matters not encompassed by the Commission’s rules.

(b) Verification of the equipment by the manufacturer or importer is effective until a termination date is otherwise established by the Commission.

(c) No person shall, in any advertising matter, brochure, etc., use or make reference to a verification in a deceptive or misleading manner or convey the impression that such verification reflects more than a determination by the manufacturer or importer that the device or product has been shown to be capable of compliance