§ 2.1076

- (viii) A description of any modifications made to the EUT by the testing company or individual to achieve compliance with the regulations;
- (ix) All of the data required to show compliance with the appropriate regulations:
- (x) The signature of the individual responsible for testing the product along with the name and signature of an official of the responsible party, as designated in §2.909; and
- (xi) A copy of the compliance information, as described in §2.1077, required to be provided with the equipment.
- (b) If the equipment is assembled using modular components that, by themselves, are subject to authorization under a Declaration of Conformity and/or a grant of certification, and the assembled product is also subject to authorization under a Declaration of Conformity but, in accordance with the applicable regulations, does not require additional testing, the assembler shall maintain the following records in order to show the basis on which compliance with the standards was determined:
- (1) A listing of all of the components used in the assembly;
- (2) Copies of the compliance information, as described in §2.1077 for all of the modular components used in the assembly;
- (3) A listing of the FCC Identifier numbers for all of the components used in the assembly that are authorized under a grant of certification:
- (4) A listing of equipment modifications, if any, that were made during assembly; and
- (5) A copy of any instructions included with the components that were required to be followed to ensure the assembly of a compliant product, along with a statement, signed by the assembler, that these instructions were followed during assembly. This statement shall also contain the name and signature of an official of the responsible party, as designated in §2.909.
- (c) The records listed in paragraphs (a) and (b) of this section shall be retained for two years after the manufacture or assembly, as appropriate, of said equipment has been permanently discontinued, or until the conclusion of an investigation or a proceeding if the

responsible party is officially notified that an investigation or any other administrative proceeding involving the equipment has been instituted. Requests for the records described in this section and for sample units also are covered under the provisions of §2.946.

[61 FR 31047, June 19, 1996]

§ 2.1076 FCC inspection and submission of equipment for testing.

- (a) Each responsible party, upon receipt of a reasonable request, shall submit to the Commission the records required by §2.1075 or one or more sample units for measurements at the Commission's laboratory.
- (b) Shipping costs to the Commission's Laboratory and return shall be borne by the responsible party. In the event the responsible party believes that shipment of the sample to the Commission's Laboratory is impractical because of the size or weight of the equipment, or the power requirement, or for any other reason, the responsible party may submit a written explanation why such shipment is impractical and should not be required.

[61 FR 31047, June 19, 1996]

§2.1077 Compliance information.

- (a) If a product must be tested and authorized under a Declaration of Conformity, a compliance information statement shall be supplied with the product at the time of marketing or importation, containing the following information:
- (1) Identification of the product, e.g., name and model number;
- (2) A statement, similar to that contained in §15.19(a)(3) of this chapter, that the product complies with part 15 of this chapters; and
- (3) The identification, by name, address and telephone number, of the responsible party, as defined in §2.909. The responsible party for a Declaration of Conformity must be located within the United States.
- (b) If a product is assembled from modular components that, by themselves, are authorized under a Declaration of Conformity and/or a grant of certification, and the assembled product is also subject to authorization under a Declaration of Conformity but,