

## § 5.405

## 47 CFR Ch. I (10–1–14 Edition)

### § 5.405 Yearly report.

Medical testing licensees must file a yearly report detailing the activity that has been performed under the license. This report is to be filed electronically to the Commission's program experimental registration Web site and must, at a minimum, include:

- (a) A list of each test performed and the testing period; and
- (b) A Description of each test, including equipment tested; and
- (c) The results of the test including any interference incidents and their resolution.

**EFFECTIVE DATE NOTE:** At 78 FR 25162, Apr. 29, 2013, §§ 5.405 was added. This section contains information collection and record-keeping requirements and will not become effective until approval has been given by the Office of Management and Budget.

### § 5.406 Responsible party, “stop-buzzer,” and notification requirements, and additional requirements related to safety of the public.

(a) Medical testing licensees must identify a single point of contact responsible for all experiments conducted under the license and must also identify a “stop buzzer” point of contact for all experiments, consistent with subpart E, §§ 5.307 and 5.308.

(b) Medical testing licensees must meet the notification and safety of the public requirements of subpart E, §§ 5.309 and 5.311.

**EFFECTIVE DATE NOTE:** At 78 FR 25162, Apr. 29, 2013, §§ 5.406 was added. This section contains information collection and record-keeping requirements and will not become effective until approval has been given by the Office of Management and Budget.

### § 5.407 Exemption from station identification requirement.

Medical testing experimental licensees are exempt from complying with the station identification requirements of § 5.115.

## Subpart G—Compliance Testing Experimental Radio Licenses

### § 5.501 Applicable rules.

In addition to the rules in this subpart, compliance testing experimental applicants and licensees must follow the rules in subparts B and C of this

part. In case of any conflict between the rules set forth in this subpart and the rules set forth in subparts B and C of this part, the rules in this subpart shall govern.

### § 5.502 Eligibility.

Compliance testing experimental radio licenses may be granted to those testing laboratories recognized by the FCC as being competent to perform measurements of equipment for equipment authorization.

### § 5.503 Scope of testing activities.

The authority of a compliance testing experimental license is limited to only those testing activities necessary for device certification (including antenna calibration, test site validation, proficiency testing, and testing in an Open Area Test Site); *i.e.*, compliance testing experimental licensees are not authorized to conduct immunity testing.

### § 5.504 Responsible party.

Compliance testing licensees must identify a single point of contact responsible for all experiments conducted under the license, including ensuring compliance with all applicable FCC rules:

(a) The responsible individual will serve as the initial point of contact for all matters involving interference resolution and must have the authority to discontinue any and all experiments being conducted under the license, if necessary.

(b) The name of the responsible individual, along with contact information, such as a phone number and email address at which he or she can be reached at any time of the day, must be identified on the license application, and this information will be listed on the license. Licensees are required to keep this information current.

**EFFECTIVE DATE NOTE:** At 78 FR 25162, Apr. 29, 2013, §§ 5.504 was added. This section contains information collection and record-keeping requirements and will not become effective until approval has been given by the Office of Management and Budget.