

(b) The final report shall be signed and dated by the study director.

(c) Corrections or additions to a final report shall be in the form of an amendment by the study director. The amendment shall clearly identify that part of the final report that is being added to or corrected and the reasons for the correction or addition, and shall be signed and dated by the person responsible. Modification of a final report to comply with the submission requirements of EPA does not constitute a correction, addition, or amendment to a final report.

(d) A copy of the final report and of any amendment to it shall be maintained by the sponsor and the test facility.

§ 160.190 Storage and retrieval of records and data.

(a) All raw data, documentation, records, protocols, specimens, and final reports generated as a result of a study shall be retained. Specimens obtained from mutagenicity tests, specimens of soil, water, and plants, and wet specimens of blood, urine, feces, and biological fluids, do not need to be retained after quality assurance verification. Correspondence and other documents relating to interpretation and evaluation of data, other than those documents contained in the final report, also shall be retained.

(b) There shall be archives for orderly storage and expedient retrieval of all raw data, documentation, protocols, specimens, and interim and final reports. Conditions of storage shall minimize deterioration of the documents or specimens in accordance with the requirements for the time period of their retention and the nature of the documents of specimens. A testing facility may contract with commercial archives to provide a repository for all material to be retained. Raw data and specimens may be retained elsewhere provided that the archives have specific reference to those other locations.

(c) An individual shall be identified as responsible for the archives.

(d) Only authorized personnel shall enter the archives.

(e) Material retained or referred to in the archives shall be indexed to permit expedient retrieval.

§ 160.195 Retention of records.

(a) Record retention requirements set forth in this section do not supersede the record retention requirements of any other regulations in this subchapter.

(b) Except as provided in paragraph (c) of this section, documentation records, raw data, and specimens pertaining to a study and required to be retained by this part shall be retained in the archive(s) for whichever of the following periods is longest:

(1) In the case of any study used to support an application for a research or marketing permit approved by EPA, the period during which the sponsor holds any research or marketing permit to which the study is pertinent.

(2) A period of at least 5 years following the date on which the results of the study are submitted to the EPA in support of an application for a research or marketing permit.

(3) In other situations (e.g., where the study does not result in the submission of the study in support of an application for a research or marketing permit), a period of at least 2 years following the date on which the study is completed, terminated, or discontinued.

(c) Wet specimens, samples of test, control, or reference substances, and specially prepared material which are relatively fragile and differ markedly in stability and quality during storage, shall be retained only as long as the quality of the preparation affords evaluation. Specimens obtained from mutagenicity tests, specimens of soil, water, and plants, and wet specimens of blood, urine, feces, and biological fluids, do not need to be retained after quality assurance verification. In no case shall retention be required for longer periods than those set forth in paragraph (b) of this section.

(d) The master schedule sheet, copies of protocols, and records of quality assurance inspections, as required by § 160.35(c) shall be maintained by the quality assurance unit as an easily accessible system of records for the period of time specified in paragraph (b) of this section.

(e) Summaries of training and experience and job descriptions required to be

maintained by §160.29(b) may be retained along with all other testing facility employment records for the length of time specified in paragraph (b) of this section.

(f) Records and reports of the maintenance and calibration and inspection of equipment, as required by §160.63 (b) and (c), shall be retained for the length of time specified in paragraph (b) of this section.

(g) If a facility conducting testing or an archive contracting facility goes out of business, all raw data, documentation, and other material specified in this section shall be transferred to the archives of the sponsor of the study. The EPA shall be notified in writing of such a transfer.

(h) Specimens, samples, or other non-documentary materials need not be retained after EPA has notified in writing the sponsor or testing facility holding the materials that retention is no longer required by EPA. Such notification normally will be furnished upon request after EPA or FDA has completed an audit of the particular study to which the materials relate and EPA has concluded that the study was conducted in accordance with this part.

(i) Records required by this part may be retained either as original records or as true copies such as photocopies, microfilm, microfiche, or other accurate reproductions of the original records.

PART 161—DATA REQUIREMENTS FOR REGISTRATION

Subpart A—General Provisions

- Sec.
- 161.20 Overview.
 - 161.25 Applicability of data requirements.
 - 161.30 Timing of the imposition of data requirements.
 - 161.32 Format of data submission.
 - 161.33 Procedures for claims of confidentiality of data.
 - 161.34 Flagging of studies for potential adverse effects.
 - 161.35 Flexibility of the data requirements.
 - 161.40 Consultation with the Agency.
 - 161.45 Waivers.
 - 161.55 Agricultural vs. non-agricultural pesticides.
 - 161.60 Minor uses.
 - 161.70 Acceptable protocols.
 - 161.75 Requirements for additional data.

- 161.80 Acceptability of data.
- 161.85 Revision of data requirements and guidelines.

Subpart B—How To Use Data Tables

- 161.100 How to determine registration data requirements.
- 161.101 Required vs. conditionally required data.
- 161.102 Distinguishing between what data are required and what substance is to be tested.
- 161.108 Relationship of Pesticide Assessment Guidelines to data requirements.

Subpart C—Product Chemistry Data Requirements

- 161.150 General.
- 161.153 Definitions.
- 161.155 Product composition.
- 161.160 Description of materials used to produce the product.
- 161.162 Description of production process.
- 161.165 Description of formulation process.
- 161.167 Discussion of formation of impurities.
- 161.170 Preliminary analysis.
- 161.175 Certified limits.
- 161.180 Enforcement analytical method.
- 161.190 Physical and chemical characteristics.

Subpart D—Data Requirement Tables

- 161.202 Purposes of the registration data requirements.
- 161.240 Residue chemistry data requirements.
- 161.290 Environmental fate data requirements.
- 161.340 Toxicology data requirements.
- 161.390 Reentry protection data requirements.
- 161.440 Spray drift data requirements.
- 161.490 Wildlife and aquatic organisms data requirements.
- 161.540 Plant protection data requirements.
- 161.590 Nontarget insect data requirements.
- 161.640 Product performance data requirements.

APPENDIX A TO PART 161—DATA REQUIREMENTS FOR REGISTRATION: USE PATTERN INDEX.

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Subpart A—General Provisions

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