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to people must satisfy the requirements of subpart E of part 500 of this chapter.


§ 70.51 Advisory committee on the applicability of the anticancer clause.

All requests for and procedures governing any advisory committee on the anticancer clause shall be subject to the provisions of part 14 of this chapter, and particularly subpart H of that part.

§ 70.55 Request for scientific studies.

The Commissioner will consider requests by any interested person who desires the Food and Drug Administration to conduct scientific studies to support a petition for a regulation for a color additive. If favorably acted upon, such studies will be limited to pharmacological investigations, studies of the chemical and physical structure of the color additive, and methods of analysis of the pure color additive (including impurities) and its identification and determination in foods, drugs, or cosmetics, as the case may be. All requests for such studies shall be accompanied by the fee prescribed in §70.19.

PART 71—COLOR ADDITIVE PETITIONS

Subpart A—General Provisions

Sec. 71.1 Petitions.

71.2 Notice of filing of petition.

71.4 Samples; additional information.

71.6 Extension of time for studying petitions; substantive amendments; withdrawal of petitions without prejudice.

71.15 Confidentiality of data and information in color additive petitions.

71.18 Petition for exemption from certification.

Subpart B—Administrative Action on Petitions

71.20 Publication of regulation.

71.22 Deception as a basis for refusing to issue regulations; deceptive use of a color additive for which a regulation has issued.

71.25 Condition for certification.

71.26 Revocation of exemption from certification.

71.27 Listing and exemption from certification on the Commissioner’s initiative.

71.30 Procedure for filing objections to regulations.

71.37 Exemption of color additives for investigational use.


SOURCE: 42 FR 15639, Mar. 22, 1977, unless otherwise noted.