§ 101–25.108 Multiyear subscriptions for publications.

Subscriptions for periodicals, newspapers, and other publications for which it is known in advance that a continuing requirement exists should be for multiple years rather than for a single year where such method is advantageous for the purpose of economy or otherwise. Where various bureaus or offices in the same agency are subscribing to the same publication, consideration shall be given to consolidating these requirements, to the extent practical, on an agency-wide basis and on a multiyear basis. Payment covering issues to be delivered during the entire subscription period may be made in advance from currently available appropriations (31 U.S.C. 530a).


§ 101–25.109 Laboratory and research equipment.

(a) This section prescribes controls for use by Federal agencies in managing laboratory and research equipment in Federal laboratories. Agencies may establish such additional controls as are appropriate to increase the use of already-owned equipment instead of procuring similar equipment.

(b) The term Federal laboratory, as used in this section, means any laboratory or laboratory facility in any Government-owned or -leased building which is equipped and/or used for scientific research, testing, or analysis, except clinical laboratories operating in direct support of Federal health care programs. To the extent practicable, agencies should observe the provisions of this section with regard to commercial laboratories and laboratory facilities which operate under contract with the Government and use Government-furnished equipment.

[43 FR 29004, July 5, 1978]

§ 101–25.109–1 Identification of idle equipment.

(a) The provisions of this §101–25.109–1 apply to all Federal laboratories regardless of size.

(b) Inspection tours of Federal laboratories shall be conducted on a scheduled basis, annually, if feasible, but no less than every 2 years, for the purpose of identifying idle and unneeded laboratory and research equipment. Following each tour, a report of findings shall be prepared by the inspection team and, as determined by the agency head or his designee,

(a) The provisions of this § 101–25.109–2 apply to Federal laboratories which occupy an area of 10,000 square feet or more and employ 25 or more technical or scientific personnel.

(b) Equipment pools shall be established in Federal laboratories so that laboratory and research equipment can be shared or allocated on a temporary basis to laboratory activities and individuals whose average use does not warrant the assignment of the equipment on a permanent basis. In determining the number and location of equipment pools, consideration shall be given to economy of operation, mobility of equipment, accessibility to users, frequency of use of the equipment, and impact on research programs. Pooling operations should begin expeditiously, within 120 days, if feasible, following decisions regarding the number and location of pools. If it is determined that an equipment pool would not be practical or economical or for any other reason is not needed at a particular laboratory, a written report supporting that determination shall be submitted to the agency head or his designee. Federal laboratories which do not meet the size and staffing criteria in § 101–25.109–2(a) should also establish equipment pools whenever feasible; however, these facilities need not submit written reports regarding determinations not to establish pools.

(c) Laboratory inspection teams shall be comprised of senior program management, property management, and scientific personnel who are familiar with the plans and programs of the laboratory(ies) and who have a knowledge of laboratory and research equipment utilization. As determined by the agency head or his designee, members of an inspection team shall be appointed by either the head of the laboratory or a higher agency official having laboratories management responsibility.

(d) The agency head or his designee shall ensure compliance by responsible personnel with the requirements of this § 101–25.109–1 and shall require that periodic independent reviews of walk-through procedures employed in Federal laboratories under his control be conducted to determine their effectiveness and to effect modifications as appropriate.

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