- (3) promotion of scientific programs within the Centers related to the major disease area of focus of the Institute;
- (4) development of programs and enhancement of strategies to recruit, train, and provide continuing education opportunities for the personnel of the Centers with expertise related to the major disease area of focus of the Institute:
- (5) enhancement of the interactions of the Centers with patients, sponsors, and the external biomedical community regarding the major disease area of focus of the Institute; and
- (6) facilitation of the collaborative relationships of the Centers with other agencies within the Department of Health and Human Services regarding the major disease area of focus of the Institute.

(b) Public process

The Secretary shall provide a period for public comment during the time that each Institute is being implemented.

(c) Timing

The Secretary shall establish at least one Institute under subsection (a) before the date that is 1 year after December 13, 2016.

(d) Termination of Institutes

The Secretary may terminate any Institute established pursuant to this section if the Secretary determines such Institute is no longer benefitting the public health. Not less than 60 days prior to so terminating an Institute, the Secretary shall provide public notice, including the rationale for such termination.

(June 25, 1938, ch. 675, §1014, as added Pub. L. 114-255, div. A, title III, §3073(a), Dec. 13, 2016, 130

§ 399h. Grants for studying continuous drug manufacturing

(a) In general

The Secretary of Health and Human Services may award grants to institutions of higher education and nonprofit organizations for the purpose of studying and recommending improvements to the process of continuous manufacturing of drugs and biological products and similar innovative monitoring and control techniques.

(b) Definitions

In this section—

- (1) the term "drug" has the meaning given such term in section 321 of this title;
- (2) the term "biological product" has the meaning given such term in section 262(i) of title 42; and
- (3) the term "institution of higher education" has the meaning given such term in section 1001(a) of title 20.

(Pub. L. 114-255, div. A, title III, §3016, Dec. 13, 2016, 130 Stat. 1095.)

CODIFICATION

Section was enacted as part of the 21st Century Cures Act, and not as part of the Federal Food, Drug, and Cosmetic Act which comprises this chapter.

CHAPTER 10-POULTRY AND POULTRY PRODUCTS INSPECTION

Sec. 451. Congressional statement of findings. 452.

Congressional declaration of policy. Definitions.

453.

Federal and State cooperation in development and administration of State poultry product inspection programs.

455. Inspection in official establishments.

456. Operation of premises, facilities and equipment.

457. Labeling and container standards.

458. Prohibited acts.

Compliance by all establishments. 459.

Miscellaneous activities subject to regula-460. tion.

461. Offenses and punishment.

Reporting of violations; notice; opportunity 462. to present views.

463. Rules and regulations.

464 Exemptions.

465. Limitations upon entry of poultry products and other materials into official establishments.

466 Imports.

467. Inspection services.

467a Administrative detention; duration; pending judicial proceedings; notification of government authorities; release; removal of official marks.

467b. Seizure and condemnation.

Federal court jurisdiction of enforcement and 467c. injunction proceedings and other kinds of cases; limitations; United States as plaintiff; subpenas.

467d. Administration and enforcement; applicability of penalty provisions; conduct of inquiries; power and jurisdiction of courts.

Non-Federal jurisdiction of federally regulated matters; prohibition of additional or different requirements for establishments with inspection services and as to marking, labeling, packaging, and ingredients; recordkeeping and related requirements; concurrent jurisdiction over distribution for human food purposes of adulterated or misbranded and imported articles: other matters.

467f. Federal Food, Drug, and Cosmetic Act appli-

cations.

468. Cost of inspection: overtime.

469 Authorization of appropriations.

470. Omitted.

Safe Meat and Poultry Inspection Panel. 471

472 Interstate shipment of poultry inspected by Federal and State agencies for certain small establishments.

§ 451. Congressional statement of findings

Poultry and poultry products are an important source of the Nation's total supply of food. They are consumed throughout the Nation and the major portion thereof moves in interstate or foreign commerce. It is essential in the public interest that the health and welfare of consumers be protected by assuring that poultry products distributed to them are wholesome, not adulterated, and properly marked, labeled, and packaged. Unwholesome, adulterated, or misbranded poultry products impair the effective regulation of poultry products in interstate or foreign commerce, are injurious to the public welfare, destroy markets for wholesome, not adulterated, and properly labeled and packaged poultry products, and result in sundry losses to