"(f) The President shall conduct a thorough review of existing disaster relief legislation as it relates to emergency loans and housing loans administered by the Farmers Home Administration of the United States Department of Agriculture, and not later than January 31, 1973, he shall transmit to the Committee on Agriculture and Forestry of the Senate and the Committee on Agriculture of the House of Representatives a report containing specific legislative proposals for the comprehensive revision of such legislation in order to—

"(1) adjust the benefits and the coverage available to persons affected by disasters;

"(2) improve the execution of the program by simplifying and eliminating unnecessary administrative procedures; and

"(3) prevent the misuse of benefits made available under the program."

Disaster relief legislation, Presidential review.
Report to congressional committees.

SEC. 6. Section 231 of the Disaster Relief Act of 1970 is amended by—

(1) inserting "(a)" after "SEC. 231."; and

(2) adding at the end of such section the following new subsection:

"(b) Loans to which this section applies may also be made for the purpose of providing small business concerns with working capital, the payment of operating expenses, and any purpose for which loans may be made under section 7(a) of the Small Business Act (15 U.S.C. 636(a))."

84 Stat. 1752.
42 USC 4451.

72 Stat. 387;
81 Stat. 268.

Approved August 16, 1972.

Public Law 92-386

JOINT RESOLUTION

August 16, 1972
[S. J. Res. 254]

To authorize the printing and binding of a revised edition of Senate Procedure and providing the same shall be subject to copyright by the author.

Resolved by the Senate and House of Representatives of the United States of America in Congress assembled, That there shall be printed and bound for the use of the Senate one thousand five hundred copies of a revised edition of Senate Procedure (originally prepared by Charles L. Watkins and Floyd M. Riddick), to be prepared by Floyd M. Riddick, Parliamentarian of the United States Senate, to be printed under the supervision of the author and to be distributed to the Members of the Senate.

"Senate Procedure."
Revised edition, printing.

SEC. 2. That notwithstanding any provision of the copyright laws and regulations with respect to publications in the public domain, such revised edition of Senate Procedure shall be subject to copyright by the author thereof.

Approved August 16, 1972.

Public Law 92-387

AN ACT

August 16, 1972
[H. R. 9936]

To amend the Federal Food, Drug, and Cosmetic Act to provide for a current listing of each drug manufactured, prepared, propagated, compounded, or processed by a registrant under that Act, and for other purposes.

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,

Drug Listing Act of 1972.

SECTION 1. This Act may be cited as the "Drug Listing Act of 1972".

SEC. 2. The Federal Government which is responsible for regulating

drugs has no ready means of determining what drugs are actually being manufactured or packed by establishments registered under the Federal Food, Drug, and Cosmetic Act except by periodic inspection of such registered establishments. Knowledge of which particular drugs are being manufactured or packed by each registered establishment would substantially assist in the enforcement of Federal laws requiring that such drugs be pure, safe, effective, and properly labeled. Information on the discontinuance of a particular drug could serve to alleviate the burden of reviewing and implementing enforcement actions against drugs which, although commercially discontinued, remain active for regulatory purposes. Information on the type and number of different drugs being manufactured or packed by drug establishments could permit more effective and timely regulation by the agencies of the Federal Government responsible for regulating drugs, including identification of which drugs in interstate commerce are subject to section 505 or 507, or to other provisions of the Federal Food, Drug, and Cosmetic Act.

52 Stat. 1040.
21 USC 301.

76 Stat. 781.
59 Stat. 463.
21 USC 355, 357.

SEC. 3. Section 510 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360) is amended by adding at the end of the following new subsection:

Registration, drug list requirements.
76 Stat. 794.

"(j)(1) Every person who registers with the Secretary under subsection (b), (c), or (d) shall, at the time of registration under any such subsection, file with the Secretary a list of all drugs (by established name (as defined in section 502(e)) and by any proprietary name) which are being manufactured, prepared, propagated, compounded, or processed by him for commercial distribution and which he has not included in any list of drugs filed by him with the Secretary under this paragraph or paragraph (2) before such time of registration. Such list shall be prepared in such form and manner as the Secretary may prescribe and shall be accompanied by—

52 Stat. 1050; 76 Stat. 790.
21 USC 352.

"(A) in the case of a drug contained in such list and subject to section 505, 506, 507, or 512, a reference to the authority for the marketing of such drug and a copy of all labeling for such drug;

55 Stat. 851.
82 Stat. 343.
21 USC 356, 360b.
65 Stat. 648.
21 USC 353.

"(B) in the case of any other drug contained in such list—

"(i) which is subject to section 503(b)(1), a copy of all labeling for such drug, a representative sampling of advertisements for such drug, and, upon request made by the Secretary for good cause, a copy of all advertisements for a particular drug product, or

"(ii) which is not subject to section 503(b)(1), the label and package insert for such drug and a representative sampling of any other labeling for such drug;

"(C) in the case of any drug contained in such list which is described in subpararaph (B), a quantitative listing of its active ingredient or ingredients, except that with respect to a particular drug product the Secretary may require the submission of a quantitative listing of all ingredients if he finds that such submission is necessary to carry out the purposes of this Act; and

"(D) if the registrant filing the list has determined that a particular drug product contained in such list is not subject to section 505, 506, 507, or 512, a brief statement of the basis upon which the registrant made such determination if the Secretary requests such a statement with respect to that particular drug product.

"(2) Each person who registers with the Secretary under this section shall report to the Secretary once during the month of June of each year and once during the month of December of each year the following information: Reports.

"(A) A list of each drug introduced by the registrant for commercial distribution which has not been included in any list previously filed by him with the Secretary under this subparagraph or paragraph (1) of this subsection. A list under this subparagraph shall list a drug by its established name (as defined in section 502(e)) and by any proprietary name it may have and shall be accompanied by the other information required by paragraph (1). 52 Stat. 1050; 76 Stat. 790. 21 USC 352.

"(B) If since the date the registrant last made a report under this paragraph (or if he has not made a report under this paragraph, since the effective date of this subsection) he has discontinued the manufacture, preparation, propagation, compounding, or processing for commercial distribution of a drug included in a list filed by him under subparagraph (A) or paragraph (1); notice of such discontinuance, the date of such discontinuance, and the identity (by established name (as defined in section 502 (e)) and by any proprietary name) of such drug.

"(C) If since the date the registrant reported pursuant to subparagraph (B) a notice of discontinuance he has resumed the manufacture, preparation, propagation, compounding, or processing for commercial distribution of the drug with respect to which such notice of discontinuance was reported; notice of such resumption, the date of such resumption, the identity of such drug (by established name (as defined in section 502(e)) and by any proprietary name), and the other information required by paragraph (1), unless the registrant has previously reported such resumption to the Secretary pursuant to this subparagraph.

"(D) Any material change in any information previously submitted pursuant to this paragraph or paragraph (1).

"(3) The Secretary may also require each registrant under this section to submit a list of each drug product which (A) the registrant is manufacturing, preparing, propagating, compounding, or processing for commercial distribution, and (B) contains a particular ingredient. The Secretary may not require the submission of such a list unless he has made a finding that the submission of such a list is necessary to carry out the purposes of this Act." Drug product list.

SEC. 4. (a) Section 510(e) of such Act is amended by adding at the end thereof the following: "The Secretary may also assign a listing number to each drug or class of drugs listed under subsection (j). Any number assigned pursuant to the preceding sentence shall be the same as that assigned pursuant to the National Drug Code.". 76 Stat. 794. 21 USC 360. *Ante*, p. 560.

(b) Section 510(f) of such Act is amended by inserting before the period the following: "; except that any list submitted pursuant to paragraph (3) of subsection (j) and the information accompanying any list or notice filed under paragraph (1) or (2) of that subsection shall be exempt from such inspection unless the Secretary finds that such an exemption would be inconsistent with protection of the public health".

76 Stat. 794.
21 USC 360.

(c) The second sentence of section 510(i) of such Act is amended by inserting "shall require such establishment to provide the information required by subsection (j) and" immediately before "shall include".

Ante, p. 560.

76 Stat. 781.

(d) Clause (1) of the second sentence of section 505(e) of such Act (21 U.S.C. 355(e)) is amended by inserting "or to comply with the notice requirements of section 510(j)(2)" immediately after "subsection (j)".

76 Stat. 795.

(e) Section 301(p) of such Act (21 U.S.C. 331(p)) is amended to read as follows:

"(p) The failure to register in accordance with section 510, the failure to provide any information required by section 510(j), or the failure to provide a notice required by section 510(j)(2)."

Effective date.

SEC. 5. The amendments made by this Act shall take effect on the first day of the sixth month beginning after the date of enactment of this Act.

Approved August 16, 1972.

Public Law 92-388

August 17, 1972
[H. R. 1462]

AN ACT

To provide for the establishment of the Puukohola Heiau National Historic Site, in the State of Hawaii, and for other purposes.

Puukohola Heiau National Historic Site, Hawaii. Establishment.

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled, That in order to restore and preserve in public ownership the historically significant temple associated with Kamehameha the Great, who founded the historic Kingdom of Hawaii, and the property of John Young who fought for Kamehameha the Great during the period of his ascendancy to power, the Secretary of the Interior is authorized to acquire, by donation or purchase with donated funds, such lands and interests in lands, together with structures and improvements thereon, not to exceed one hundred acres, in the vicinity of Kawaihae, Hawaii, as generally depicted on a map entitled "Boundary Map, Proposed Puukohola Heiau National Historic Site," numbered NHS–PK 20,002, dated February 1970, which shall be on file and available for public inspection in the offices of the National Park Service, Washington, District of Columbia. The Secretary of the Interior may from time to time revise the boundaries of the proposed historic site, but the total acreage of the site shall not exceed one hundred acres.

Limitation.

SEC. 2. The Secretary of the Interior shall establish the area as the "Puukohola Heiau National Historic Site" at such time as he deems sufficient interests in lands have been acquired to constitute an administrable unit. Pending and after establishment, the Puukohola Heiau National Historic Site shall be administered, developed, preserved, and maintained in accordance with the provisions of the Act entitled "An Act to establish a National Park Service, and for other purposes", approved August 25, 1916 (39 Stat. 535), as amended and supplemented (16 U.S.C. 1 et seq.), and the Act entitled "An Act to provide for the preservation of historic American sites, buildings, objects, and antiquities of national significance, and for other purposes", approved August 21, 1935 (49 Stat. 666; 16 U.S.C. 461 et seq.).

Land acquisition.

SEC. 3. Notwithstanding the acreage limitation contained in section 1 of this Act, the Secretary of the Interior is authorized to acquire by donation, purchase, or exchange, such additional lands and interests therein outside the boundary of the site as he deems necessary to relocate portions of State and county roads which are currently within