

One Hundred Eleventh Congress
of the
United States of America

AT THE FIRST SESSION

*Begun and held at the City of Washington on Tuesday,
the sixth day of January, two thousand and nine*

An Act

Making omnibus appropriations for the fiscal year ending September 30, 2009,
and for other purposes.

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

SECTION 1. SHORT TITLE.

This Act may be cited as the “Omnibus Appropriations Act,
2009”.

SEC. 2. TABLE OF CONTENTS.

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SEC. 3. REFERENCES.

Except as expressly provided otherwise, any reference to “this Act” contained in any division of this Act shall be treated as referring only to the provisions of that division.

SEC. 4. EXPLANATORY STATEMENT.

The explanatory statement regarding this Act printed in the House of Representatives section of the Congressional Record on or about February 23, 2009 by the Chairman of the Committee on Appropriations of the House shall have the same effect with respect to the allocation of funds and implementation of this Act as if it were a joint explanatory statement of a committee of conference.

SEC. 5. STATEMENT OF APPROPRIATIONS.

The following sums in this Act are appropriated, out of any money in the Treasury not otherwise appropriated, for the fiscal year ending September 30, 2009.

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\$4,985,000 shall be transferred to and merged with the appropriation for “Foreign Agricultural Service, Salaries and Expenses”, and of which \$348,000 shall be transferred to and merged with the appropriation for “Farm Service Agency, Salaries and Expenses”.

MC GOVERN-DOLE INTERNATIONAL FOOD FOR EDUCATION AND CHILD
NUTRITION PROGRAM GRANTS

For necessary expenses to carry out the provisions of section 3107 of the Farm Security and Rural Investment Act of 2002 (7 U.S.C. 1736o-1), \$100,000,000, to remain available until expended: *Provided*, That the Commodity Credit Corporation is authorized to provide the services, facilities, and authorities for the purpose of implementing such section, subject to reimbursement from amounts provided herein.

TITLE VI

RELATED AGENCY AND FOOD AND DRUG ADMINISTRATION

DEPARTMENT OF HEALTH AND HUMAN SERVICES

FOOD AND DRUG ADMINISTRATION

SALARIES AND EXPENSES

For necessary expenses of the Food and Drug Administration, including hire and purchase of passenger motor vehicles; for payment of space rental and related costs pursuant to Public Law 92-313 for programs and activities of the Food and Drug Administration which are included in this Act; for rental of special purpose space in the District of Columbia or elsewhere; for miscellaneous and emergency expenses of enforcement activities, authorized and approved by the Secretary and to be accounted for solely on the Secretary's certificate, not to exceed \$25,000; and notwithstanding section 521 of Public Law 107-188; \$2,622,267,000, of which \$7,641,000 shall be for the purposes, and in the amounts, specified in the final paragraph under “Food and Drug Administration, Salaries and Expenses” in the explanatory statement described in section 4 (in the matter preceding division A of this consolidated Act): *Provided*, That of the amount provided under this heading, \$510,665,000 shall be derived from prescription drug user fees authorized by 21 U.S.C. 379h shall be credited to this account and remain available until expended, and shall not include any fees pursuant to 21 U.S.C. 379h(a)(2) and (a)(3) assessed for fiscal year 2010 but collected in fiscal year 2009; \$52,547,000 shall be derived from medical device user fees authorized by 21 U.S.C. 379j, and shall be credited to this account and remain available until expended; \$15,260,000 shall be derived from animal drug user fees authorized by 21 U.S.C. 379j, and shall be credited to this account and remain available until expended; and \$4,831,000 shall be derived from animal generic drug user fees authorized by 21 U.S.C. 379f, and shall be credited to this account and shall remain available until expended: *Provided further*, That fees derived from prescription drug, medical device, animal drug, and animal generic drug assessments for fiscal year 2009 received during fiscal year 2009, including any such fees assessed prior to fiscal year 2009 but credited for fiscal year 2009, shall be subject to the

fiscal year 2009 limitations: *Provided further*, That none of these funds shall be used to develop, establish, or operate any program of user fees authorized by 31 U.S.C. 9701: *Provided further*, That of the total amount appropriated: (1) \$648,722,000 shall be for the Center for Food Safety and Applied Nutrition and related field activities in the Office of Regulatory Affairs; (2) \$777,437,000 shall be for the Center for Drug Evaluation and Research and related field activities in the Office of Regulatory Affairs, of which no less than \$41,358,000 shall be available for the Office of Generic Drugs; (3) \$271,490,000 shall be for the Center for Biologics Evaluation and Research and for related field activities in the Office of Regulatory Affairs; (4) \$134,344,000 shall be for the Center for Veterinary Medicine and for related field activities in the Office of Regulatory Affairs; (5) \$310,547,000 shall be for the Center for Devices and Radiological Health and for related field activities in the Office of Regulatory Affairs; (6) \$52,511,000 shall be for the National Center for Toxicological Research; (7) not to exceed \$111,758,000 shall be for Rent and Related activities, of which \$41,281,000 is for White Oak Consolidation, other than the amounts paid to the General Services Administration for rent; (8) not to exceed \$155,425,000 shall be for payments to the General Services Administration for rent; and (9) \$160,033,000 shall be for other activities, including the Office of the Commissioner; the Office of Scientific and Medical Programs; the Office of Policy, Planning and Preparedness; the Office of International and Special Programs; the Office of Operations; and central services for these offices: *Provided further*, That none of the funds made available under this heading shall be used to transfer funds under section 770(n) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379dd): *Provided further*, That funds may be transferred from one specified activity to another with the prior approval of the Committees on Appropriations of both Houses of Congress.

In addition, mammography user fees authorized by 42 U.S.C. 263b, export certification user fees authorized by 21 U.S.C. 381, and priority review user fees authorized by 21 U.S.C. 360n may be credited to this account, to remain available until expended.

BUILDINGS AND FACILITIES

For plans, construction, repair, improvement, extension, alteration, and purchase of fixed equipment or facilities of or used by the Food and Drug Administration, where not otherwise provided, \$12,433,000, to remain available until expended.

INDEPENDENT AGENCY

FARM CREDIT ADMINISTRATION

LIMITATION ON ADMINISTRATIVE EXPENSES

Not to exceed \$49,000,000 (from assessments collected from farm credit institutions, including the Federal Agricultural Mortgage Corporation) shall be obligated during the current fiscal year for administrative expenses as authorized under 12 U.S.C. 2249: *Provided*, That this limitation shall not apply to expenses associated with receiverships.

School Lunch Act as amended by section 4304 of Public Law 110–246 in excess of \$16,000,000 until October 1, 2009: *Provided further*, of the unobligated balances under section 32 of the Act of August 24, 1935, \$293,530,000 are hereby rescinded.

SEC. 724. Notwithstanding any other provision of law, the Secretary of Agriculture is authorized to make funding and other assistance available through the emergency watershed protection program under section 403 of the Agricultural Credit Act of 1978 (16 U.S.C. 2203) to repair and prevent damage to non-Federal land in watersheds that have been impaired by fires initiated by the Federal Government and shall waive cost sharing requirements for the funding and assistance.

SEC. 725. There is hereby appropriated \$3,497,000, to remain available until expended, for a grant to the National Center for Natural Products Research for construction or renovation to carry out the research objectives of the natural products research grant issued by the Food and Drug Administration.

SEC. 726. There is hereby appropriated \$469,000, to remain available until expended, for the planning and design of construction of an agriculture pest facility in the State of Hawaii.

SEC. 727. None of the funds made available in this Act may be used to establish or implement a rule allowing poultry products to be imported into the United States from the People's Republic of China.

SEC. 728. There is hereby appropriated \$794,000 to the Farm Service Agency to carry out a pilot program to demonstrate the use of new technologies that increase the rate of growth of re-forested hardwood trees on private non-industrial forests lands, enrolling lands on the coast of the Gulf of Mexico that were damaged by Hurricane Katrina in 2005.

SEC. 729. None of the funds made available to the Department of Agriculture in this Act may be used to implement the risk-based inspection program in the 30 prototype locations announced on February 22, 2007, by the Under Secretary for Food Safety, or at any other locations, until the USDA Office of Inspector General has provided its findings to the Food Safety and Inspection Service and the Committees on Appropriations of the House of Representatives and the Senate on the data used in support of the development and design of the risk-based inspection program and FSIS has addressed and resolved issues identified by OIG.

SEC. 730. Notwithstanding any other provision of law, and until receipt of the decennial Census in the year 2010, the Secretary of Agriculture shall consider—

(1) the City of Palmview, Texas; the City of Pharr, Texas; the City of Hidalgo, Texas; the City of Alton, Texas; the City of La Joya, Texas; the City of Penitas, Texas; the City of Schertz, Texas; the City of Converse, Texas; the City of Cibolo, Texas; and the Township of Bern, Pennsylvania (including individuals and entities with projects within the cities), eligible for loans and grants funded through the Rural Business Program account;

(2) the County of Nueces, Texas (including individuals and entities with projects within the county), eligible under the

[House Appropriations Committee Print]

Omnibus Appropriations Act, 2009
(H.R. 1105; Public Law 111-8)

DIVISION A—AGRICULTURE, RURAL DEVELOPMENT, FOOD AND DRUG ADMINISTRATION, AND RELATED AGENCIES APPROPRIATIONS ACT, 2009

TITLE VI

RELATED AGENCY AND FOOD AND DRUG ADMINISTRATION

DEPARTMENT OF HEALTH AND HUMAN SERVICES

FOOD AND DRUG ADMINISTRATION

SALARIES AND EXPENSES

The bill provides total appropriations, including Prescription Drug User Fee Act, Medical Device User Fee and Modernization Act, Animal Drug User Fee Act and Animal Generic Drug User Fee collections, of \$2,622,267,000 for the salaries and expenses of the Food and Drug Administration. The bill provides a direct appropriation of \$2,038,964,000.

FDA is directed to provide all reports and studies requested in this statement to the Committees on Appropriations of the House of Representatives and the Senate (hereafter referred to as “the Committees”) in both an electronic and hard copy format within 60 days after the enactment of this Act, unless another date is specified for a particular report.

Food and Drug Administration, Salaries and Expenses

[In thousands of dollars]

Program	Budget authority
Foods	648,722
Center for Food Safety and Applied Nutrition	210,486
Field Activities	438,236
Human Drugs	413,482
Center for Drug Evaluation and Research	302,386
Field Activities	111,096
Biologics	183,451
Center for Biologics Evaluation and Research	148,134
Field Activities	35,317
Animal Drugs & Feeds	116,471
Center for Veterinary Medicine	73,035
Field Activities	43,436
Device & Radiological Products	280,587
Center for Devices and Radiological Health	209,061
Field Activities	71,526
National Center for Toxicological Research	52,511
Other Activities	120,560
White Oak Consolidation	38,536
Other Rent & Rent-Related	50,293
GSA Rent	134,351
Total Salaries & Expenses	2,038,964

The Food and Drug Administration has received significant increases in appropriated funds over the past several years. Specifically, the agency received an increase of \$145,093,000, or nine percent, in the fiscal year 2008 appropriations act. An additional \$150,000,000 in supplemental funding was provided to the agency during fiscal year 2008. This bill provides an increase of \$324,672,000, or 19 percent, above the fiscal year 2008 funding level.

It is expected that these substantial funding increases will lead to significant improvements in food and medical product safety. FDA should also build on these improvements by taking broader approaches to addressing safety issues. In the foods area, for example, FDA might focus on the identification of the most significant food safety hazards, prioritized by risk and the ability to reduce such risks, and develop a plan with findings, resources and tools to address those specific risks. Further, FDA could do a compliance audit for a whole category of foods with significant safety issues, providing the agency with a much more complete understanding of their risks and what needs to be done to address them. This approach could have helped FDA in the investigation into the Salmonella outbreak during the summer of 2008, which originally focused on tomatoes before additional evidence suggested that peppers may have been the source of the contamination. This audit approach also could be taken in other product areas, especially with regard to foreign drug facilities.

To ensure that FDA efficiently applies this funding increase to its most pressing needs, FDA is directed to provide an expenditure report to the Committees no later than 15 days after the end of each fiscal year quarter following the date of enactment of this Act. This report shall include specific information for:

- The number of new hires and their estimated costs;
- The number of inspections and their estimated costs; and
- Information technology acquisition and development spending.

All cost estimates and spending in the quarterly reports must be shown on a center/field basis.

To provide a basis of comparison for the new activities in these reports, FDA is directed to include information on base funding, FTEs, inspections, and any other applicable base activity levels for each activity that has received increased funding. Finally, the reports must include up-to-date dollar obligation data for each enhanced activity.

The bill provides an increase of \$54,531,000 for cost of living adjustments instead of \$25,000,000 as requested in the budget. The amount provided reflects the full estimated cost associated with maintaining FDA's current staff levels in light of the hiring surge undertaken by the agency in fiscal year 2008. The bill provides an increase of \$3,739,000 for rental payments to the General Services Administration.

The bill includes an increase of \$141,526,000 for activities related to food safety. An increase of more than \$55,000,000 for food safety was provided in fiscal year 2008, and FDA received an additional \$72,000,000 in supplemental funds during fiscal year 2008 for food safety related activities. It is expected that this funding will result in increased safety measures for both domestic and imported food from production to consumption.

Specifically, these funding increases will, at a minimum, provide FDA with the capability to:

- Significantly increase the number of domestic and foreign, risk-based, food production and/or processing facility inspections (and other high-risk products), significantly increase the number of import food field exams, and achieve greater laboratory capacity to

support increases in risk-based inspections and sampling of domestic and imported foods;

- Develop and deploy risk-based screening technologies that will allow FDA to target high-risk products at the border;

- Identify, develop and deploy new screening tools and methods to identify pathogens and other contaminants, including more rapid screening tools to be used by field investigators and analysts;

- Establish an early warning surveillance and notification system to identify adulteration of the pet food supply and outbreaks of illnesses associated with pet food;

- Enhance FDA's national food emergency system, allowing FDA to develop a risk communication strategy that would result in more rapid responses and reductions in the risk of consumer contamination when food contamination occurs;

- Expand its presence in foreign countries. FDA recently opened its first foreign office in China. With this increase, FDA will be able to establish FDA offices in other foreign locations, as appropriate. These offices are intended to enhance the ability of FDA inspectors to enter foreign food facilities and to gain a greater understanding of manufacturing processes overseas, resulting in more rapid identification of and response to any potential food safety issues. The FDA also will be able to assess the ability of foreign government systems to manage food safety risks;

- Enhance FDA's traceability capabilities for more rapid and precise product tracking, which will allow FDA to contain contaminated product and to provide consumers more specific information when problems occur;

- Improve risk communications to the public during food-related events and make it easier to receive adverse events reports, including creating a reportable food registry;

- Identify additional data and information needed to increase understanding of food protection risk and vulnerabilities by improving the quality of foodborne illness attribution data; and

- Support partnerships with state and local partners through information technology, training and data sharing.

The bill provides an increase of \$1,000,000 for the Office of Cosmetics and Colors.

The bill provides an increase of \$114,211,000 for medical product safety. An increase of more than \$21,000,000 for medical product safety was provided in fiscal year 2008, and FDA received an additional \$58,000,000 in supplemental funds during fiscal year 2008 for medical product safety. As noted above, it is expected that this funding will result in safer drugs, devices, and biologic products for consumers. Similar to a recent approach FDA has taken to address overall food safety issues, FDA is directed to prepare and provide to the Committees on Appropriations a comprehensive approach to ensuring the safety of medical products from the manufacturing of raw ingredients or components to consumer use.

Specifically, these funding increases will, at a minimum, provide FDA with the capability to:

- Significantly increase foreign and domestic medical product facility inspections, improve laboratory infrastructure and rapid analysis tools, and conduct many more laboratory analyses and several thousand import exams and samples;

- Establish a unique device identification system to track devices, facilitate recalls, and support inventory management;
- Begin to implement the safety requirements outlined in the Food and Drug Administration Amendments Act;
- Upgrade the agency’s information technology to enable data sharing and enhanced analysis of adverse events;
- Develop a regulated product information data warehouse that will enable information sharing with other regulatory agencies; and
- Integrate risk-based information into data systems that will support FDA’s ability to improve electronic screening of imports and allow the agency to proactively identify problems and risks associated with imported products.

The bill provides \$16,000,000 for the critical path initiative, including not less than \$4,000,000 for competitive contracts or grants to universities and non-profit organizations to support critical path projects. Funding for critical path activities is distributed throughout FDA’s program areas, and the Office of Critical Path Programs (OCPP) is responsible for coordinating these activities at the agency. It is understood that OCPP, working with FDA’s centers, will play a primary role in determining which critical path efforts the agency will undertake.

The bill provides an increase of \$6,620,000 for the Division of Drug Marketing, Advertising and Communication in CDER. The funding provided is to be used for the review of direct-to-consumer advertisements and is equal to the amount of funding that the budget estimated would have been raised by the fee in fiscal year 2009.

The bill provides \$2,000,000 for Demonstration Grants for Improving Pediatric Device Availability, as authorized by the Food and Drug Administration Amendments Act of 2007, in the Center for Devices and Radiological Health. Medical device products are typically developed for adults, limiting children’s access to safe and effective medical devices. This program will provide grants to non-profit pediatric medical device consortia, which will assist scientists and innovators with technical and financial resources to improve the number of medical devices available to children. The Office of Orphan Products Development will be responsible for carrying out this program.

The bill provides \$6,000,000 for the Office of Women’s Health, an increase of \$1,000,000.

There is concern about the contamination of farm-raised shrimp imports with banned antibiotics. FDA currently inspects less than two percent of imported shrimp. FDA is strongly encouraged to develop, in cooperation with state testing programs, a program for increasing the inspection of imported shrimp for banned antibiotics.

FDA is encouraged to conduct workshops and engage in other forms of communication with federal agencies, organizations involved in blood collection and others, to ensure that those organizations and the public understand the latest scientific information available on blood safety issues.

There are poor survival rates and a lack of new therapies associated with many pediatric cancers, including high-risk neuroblastoma. FDA is encouraged to prioritize review of new treatments

and clinical trials for pediatric oncology patients and provide a report on these activities.

The bill provides no less than the fiscal year 2008 level in appropriated funds for activities related to the Mammography Quality Standards Act (MQSA). Appropriations for this program fund research grants and various activities to develop and enforce quality standards for mammography service. On June 26, 2008, the Committees received a report on actions being taken to implement recommendations made in the Institute of Medicine report entitled "Breast Imaging Quality Standards." The report stated that FDA held an open public meeting on September 28 and 29, 2006, and has been considering potential amendments to MQSA, which would address the IOM report, since this meeting. To date, FDA has not acted on any of these recommendations. This is an unacceptable delay. FDA is directed to report to the Committees on which amendments that FDA will propose to MQSA, if any, in response to the IOM report recommendations, and provide a timeline for these amendments.

The importance of seafood to a healthy diet is recognized, but there are concerns that FDA does not focus sufficient attention on economic integrity issues, particularly with respect to mislabeling of species, weights, country of origin, and treatment. FDA is encouraged to work with states to more aggressively combat fraud in parts of the seafood industry.

The Hawaii Department of Agriculture has proposed a state-wide standardized food safety certification system. FDA is encouraged to work with the State of Hawaii on this system and to provide funding if appropriate.

Serious concerns have been raised about illnesses and deaths from Methicillin Resistant Staphylococcus aureus (MRSA). Estimates suggest that tens of thousands of persons develop serious MRSA infections in the United States each year and thousands die. While both FDA and USDA fund research on this issue, more may need to be done. FDA is encouraged to work with USDA and CDC, through the National Antibiotic Resistance Monitoring System and/or the Antibiotic Resistance Interagency Task Force, to address the issue of the prevalence of MRSA in domestic farm animals.

The bill provides funding for the following items: \$1,650,000 for the Agricultural Products Food Safety Laboratory at New Mexico State University; \$525,000 for collaborative drug safety research at the Critical Path Institute and the University of Utah; \$1,608,000 for dietary supplements research at the National Center for Natural Products Research in Mississippi; \$2,077,000 for the National Center for Food Safety and Technology, Summit-Argo, Illinois; \$139,000 for the Interstate Shellfish Sanitation Conference (ISSC); \$174,000 for ISSC vibrio vulnificus education; \$69,000 for the Waste Management Education and Research Consortium at New Mexico State University; and \$1,399,000 for the Western Region Center of Excellence at the University of California-Davis. Funding for these items was included in the budget request.

BUILDINGS AND FACILITIES

The bill provides \$12,433,000 for FDA buildings and facilities, as requested in the amended budget. This funding shall be used to up-

grade FDA facilities and laboratories that are currently below public safety standards and incapable of performing agency requirements. In providing this funding, specific projects are not approved, as the backlog of maintenance and repairs at FDA locations is significant. FDA is directed to prioritize this funding consistent with the backlog of maintenance and repairs and improve the average facility condition index at FDA sites. Within 30 days of the date of enactment of this Act, FDA is directed to provide a plan for allocating the funding to the Committees. The plan should include the methodology used to allocate the resources; the specific maintenance or repairs that will be conducted; whether the funding allocated to the site will complete a project or is partial funding for the project; and if partial funding for a project is provided, the full cost of completing the project.

INDEPENDENT AGENCY

FARM CREDIT ADMINISTRATION

LIMITATION ON ADMINISTRATIVE EXPENSES

The bill includes a limitation of \$49,000,000 on administrative expenses of the Farm Credit Administration.

TITLE VII—GENERAL PROVISIONS

(INCLUDING RESCISSION AND TRANSFERS OF FUNDS)

Section 701.—The bill includes language making funds available for the purchase, replacement, and hire of passenger motor vehicles.

Section 702.—The bill includes language regarding appropriation items that remain available until expended.

Section 703.—The bill includes language allowing for unobligated balances to be transferred to the Working Capital Fund.

Section 704.—The bill includes language limiting the funding provided in the bill to one year, unless otherwise specified.

Section 705.—The bill includes language limiting indirect costs on cooperative agreements between the Department of Agriculture and nonprofit organizations to 10 percent.

Section 706.—The bill includes language making appropriations to the Department of Agriculture for the cost of direct and guaranteed loans available until expended to disburse obligations for certain Rural Development programs.

Section 707.—The bill includes language for funds to cover necessary expenses related to advisory committees.

Section 708.—The bill includes language prohibiting the use of funds to establish an inspection panel at the Department of Agriculture.

Section 709.—The bill includes language regarding detailed employees.

Section 710.—The bill includes language regarding the appropriations hearing process.

Section 711.—The bill includes language regarding the transfer of funds to the Office of the Chief Information Officer and information technology funding obligations.

Cooperative State Research Education and Extension Service	SRG	Wood Utilization, AK, ID, ME, MI, MN, MS, NC, OR, TN, WV	\$4,545,000	Alexander, Burr, Byrd, Cochran, Coleman, Collins, Craig, Crapo, Dole, Klobuchar, Landrieu, Levin, Murkowski, Smith, Snowe, Stabenow, Vitter, Wicker, Wyden	Duncan, Jr., John J.; Price, David E.; Wu, David; Rogers (MI), Mike; Michaud, Michael H.; Hooley, Darlene; Pickering, Charles W. Chip; Oberstar, James L.; Allen, Thomas H.; Etheridge, Bob
Cooperative State Research Education and Extension Service	SRG	Wool Research, MT, TX, WY	\$206,000		Conaway, K. Michael; Rodriguez, Ciro D.
Food and Drug Administration	Salaries and expenses	Collaborative drug safety research, Critical Path Institute and University of Utah	\$525,000	Bennett	
Food and Drug Administration	Salaries and expenses	Dietary supplements research, National Center for Natural Products Research, Oxford, Mississippi	\$1,608,000	Cochran, Wicker	
Food and Drug Administration	Salaries and expenses	Interstate Shellfish Sanitation Conference, ISSC	\$139,000	Cochran, Shelby	
Food and Drug Administration	Salaries and expenses	ISSC <i>vibrio vulnificus</i> education	\$174,000	Cochran, Shelby	Melancon, Charlie
Food and Drug Administration	Salaries and expenses	National Center for Food Safety and Technology, IL	\$2,077,000	Durbin	Jackson, Jr., Jesse L.; Lipinski, Daniel
Food and Drug Administration	Salaries and expenses	New Mexico State University Agricultural Products Food Safety Laboratory	\$1,650,000	Bingaman, Domenici	Wilson, Heather
Food and Drug Administration	Salaries and expenses	Waste Management Education and Research Consortium, New Mexico State University	\$69,000	Bingaman, Domenici	Pearce, Stevan
Food and Drug Administration	Salaries and expenses	Western Region FDA Center of Excellence, University of California Davis	\$1,399,000	Boxer, Feinstein	Lungren, Daniel E.; Thompson, Mike
General Provision		Speciality Markets, Wisconsin Department of Agriculture, Trade, and Consumer Protection	\$338,000	Kohl	Kagen, Steve; Obey, David R.
General Provision		Bill Emerson National Hunger Fellowship Program and the Mickey Leland International Hunger Fellowship Program	\$2,347,000		Emerson, Jo Ann; McGovern, James P.; Kaptur, Marcy