road mobile sources, and any other category of sources that the Administrator may identify; and

"(ii) reductions in such emissions will improve air quality in the petitioning State's nonattainment area or areas at least as cost-effectively as reductions in emissions from each other principal category of sources of sulfur dioxide or nitrogen oxides to the maximum extent that a methodology is reasonably available to make such a determination

In making the determination under clause (ii), the Administrator shall use the best available peer-reviewed models and methodology that consider the proximity of the source or sources to the petitioning State or political subdivision and incorporate other sources characteristics.

- "(C) The Administrator shall develop an appropriate peer reviewed methodology for making determinations under subparagraph (B) by December 31, 2006.
- "(D) The Administrator shall not make any findings with respect to an affected unit under this section prior to December 1, 2011. For any petition submitted prior to January 1, 2010, the Administrator shall make a finding or deny the petition by the December 31, 2011.
- "(E) The Administrator, by rulemaking, shall extend the compliance and implementation deadlines in subsection (c) to the extent necessary to assure that no affected unit shall be subject to any such deadline prior to January I, 2014."
- (b) TITLE III.—Section 307(d)(1)(G) of title III of the Clean Air Act is amended to read as follows:
- "(G) the promulgation or revision of any regulation under title IV,".
- (c) NoISE POLLUTION.—Title N of the Clean Air Act (relating to noise pollution) (42 U.S.C. 7641 et seq.) is redesignated as title VII and amended by renumbering sections 401 through 403 as sections 701 through 703, respectively and conforming all cross-references thereto accordingly.
- (d) Section 406.—Title IV of the Clean Air Act Amendments of 1990 (relating to acid deposition control) is amended by repealing section 406 (industrial Sulfur dioxide emissions).
- (e) MONITORING.—Section 821 (a) of title VIII of the Clean Air Act Amendments of 1990 (miscellaneous provisions) is amended to read as follows:
- "(a) MONITORING.—The Administrator shall promulgate regulations within 18 months after November 15, 1990, to require that all affected sources subject to subpart 1 of part B of title IV of the Clean Air Act as of December 31, 2009, shall also monitor carbon dioxide emissions according to the same timetable as in section 405(b). The required monitoring may be no more stringent than that required by any two of the four most populous countries for units comparable to the affected units in the United States. The regulations shall require that such data be reported to the Administrator. The provisions of section 405(e) of title IV of the Clean Air Act shall apply for purposes of this section in the same manner and to the same extent as such provision applies to the monitoring and data referred to in section 405. The Administrator shall implement this subsection under 40 CFR Part 75 (2002), amended as appropriate by the Administrator."

SUBMITTED RESOLUTIONS

CONCURRENT RESOLU-**SENATE** TION 80-URGING ΙΔΡΔΝ TO HONOR COMMITMENTS ITS UNDER THE 1986 MARKET-ORI-SECTOR-SELECTIVE ENTED (MOSS) AGREEMENT ON MED-ICAL EQUIPMENT AND PHARMA-CEUTICALS, AND FOR OTHER **PURPOSES**

Mr. COLEMAN (for himself and Mr. BAYH) submitted the following concurrent resolution; which was referred to the Committee on Foreign Relations:

S. CON. RES. 80

Whereas the revolution in medical technology has improved our ability to respond to emerging threats and prevent, identify, treat, and cure a broad range of diseases and disabilities, and has the proven potential to bring even more valuable advances in the future;

Whereas medical technology has driven dramatic productivity gains for the benefit of patients, providers, employers, and our economy:

Whereas investment from the United States medical technology industry produces the majority of the \$175,000,000,000 global business in development of medical devices, diagnostic products, and medical information systems, allowing patients to lead longer, healthier, and more productive lives;

Whereas the United States medical technology industry supports almost 1,000,000 Americans in high-value jobs located in every State, and the industry is a net contributor to the United States balance of trade, with a trade surplus of \$3,300,000,000;

Whereas Japan is one of the most important trading partners of the United States;

Whereas United States products account for roughly ½ of the global market, but garner only a ⅓ share of Japan's market;

Whereas Japan has made little progress in implementing its commitments to cut product review times, improve their reimbursement system, and consult bilaterally on policy changes under the Market-Oriented Sector-Selective (MOSS) Agreement on Medical Equipment and Pharmaceuticals, signed on January 9, 1986, between the United States and Japan;

Whereas, although regulatory reviews in Japan remain among the lengthiest in the world and Japan needs to accelerate patient access to safe and beneficial medical technologies, proposals currently under consideration in Japan would, in many cases, actually increase regulatory burdens on manufacturers and delay access without enhancing patient safety;

Whereas the general cost of doing business in Japan is among the highest in the world and is driven significantly higher by certain factors in the medical technology sector, and inefficiencies in Japanese distribution networks and hospital payment systems and unique regulatory burdens drive up the cost of bringing innovations to Japanese consumers and impede patient access to life-saving and life-enhancing medical technologies;

Whereas artificial government price caps such as the foreign average price policy adopted by the Government of Japan in 2002 restrict patient access and fail to recognize the value of innovation;

Whereas less than $\frac{1}{10}$ of 1 percent of the tens of thousands of medical technologies introduced in Japan in the last 10 years received new product pricing;

Whereas the Government of Japan has adopted artificial price caps that are targeted toward technologies predominately marketed by United States companies and is considering altering pricing rules to enable further cuts to these products; and

Whereas these discriminatory pricing policies will allow the Japanese government to take advantage of United States research and development: Now, therefore, be it

Resolved by the Senate (the House of Representatives concurring), That Congress—

(1) urges Japan to honor its commitments under the Market-Oriented Sector-Selective (MOSS) Agreement on Medical Equipment and Pharmaceuticals, signed on January 9, 1986, between the United States and Japan (hereafter in this resolution referred to as the "MOSS Agreement"), by—

(A) reducing regulatory barriers to the approval and adoption of new medical tech-

nologies; and

- (B) establishing reasonable agency performance goals for premarket approvals and an appropriate, risk-based postmarket system consistent with globally accepted practices:
- (2) urges Japan to honor its commitments under the MOSS Agreement to improve the reimbursement environment for medical technologies by actively promoting pricing policies that encourage innovation for the benefit of Japanese patients and the Japanese economy; and
- (3) urges $\tilde{\mbox{Japan}}$ to honor its commitments under the MOSS Agreement by—
- (A) implementing fair and open processes and rules that do not disproportionately harm United States medical technology products; and
- (B) providing opportunities for consultation with trading partners.

AMENDMENTS SUBMITTED & PROPOSED

SA 2143. Mr. VOINOVICH submitted an amendment intended to be proposed by him to the bill S. 150, to make permanent the moratorium on taxes on Internet access and multiple and discriminatory taxes on electronic commerce imposed by the Internet Tax Freedom Act; which was ordered to lie on the table.

SA 2144. Mr. CORNYN submitted an amendment intended to be proposed by him to the bill H.R. 2799, making appropriations for the Departments of Commerce, Justice, and State, the Judiciary, and related agencies for the fiscal year ending September 30, 2004, and for other purposes; which was ordered to lie on the table.

SA 2145. Mr. BAUCUS submitted an amendment intended to be proposed by him to the bill H.R. 2799, supra; which was ordered to lie on the table.

SA 2146. Mr. KYL submitted an amendment intended to be proposed by him to the bill H.R. 2799, supra; which was ordered to lie on the table.

SA 2147. Mr. CRAIG (for himself and Mr. HAGEL) submitted an amendment intended to be proposed by him to the bill H.R. 2799, supra; which was ordered to lie on the table.

TEXT OF AMENDMENTS

SA 2143. Mr. VOINOVICH submitted an amendment intended to be proposed by him to the bill S. 150, to make permanent the moratorium on taxes on Internet access and multiple and discriminatory taxes on electronic commerce imposed by the Internet Tax Freedom Act; which was ordered to lie on the table; as follows:

At the appropriate place, insert the following: