''(3) the development of on-site access to services to address the safety, medical, mental health, and economic needs of patients either by increasing the capacity of existing health care professionals and staff to address these issues or by contracting with or hiring domestic violence advocates to provide the services, or by developing other models appropriate to the geographic and cultural needs of a site.

''(c) ELIGIBLE ENTITY.—In this section, the term 'eligible entity' shall mean a federally qualified health center as defined in section 1861(aa)(4) of the Social Security Act (42 U.S.C. 1395x(aa)(4)).

''(d) APPLICATIONS.—Each eligible entity desiring a grant under this section shall submit an application to the Secretary at such time, in such manner, and accompanied by such information as the Secretary may require.

''(e) AUTHORIZATION OF APPROPRIATIONS.—There is authorized to be appropriated to carry out this section $5,000,000 for each of the fiscal years 2004 through 2007.''.

Mr. FRIST. Mr. President, I would like to turn to several unanimous consent requests.

UNANIMOUS CONSENT AGREEMENT—H.R. 1997

Mr. FRIST. Mr. President, I ask unanimous consent that, at a time to be determined by the majority leader, in consultation with the minority leader, the Senate proceed to consideration of H.R. 1997, a bill to protect unborn victims of violence, and that the bill be considered under the following limitations: that the following amendments be the only first-degree amendments in order, with the specified time limitations: Feinstein, Motherhood Protection Act, which is at the desk, 4 hours equally divided; Murray, domestic violence prevention, which is at the desk, 2 hours equally divided; and that there be an additional 30 minutes of debate equally divided on the underlying bill.

I further ask consent that no second-degree amendments be in order to the Feinstein and Murray amendments; and that upon disposition of all amendments, the bill, as amended, if amended, be read a third time, and the Senate proceed to a vote on passage, without any intervening action or debate.

The PRESIDING OFFICER. Is there objection?

Without objection, it is so ordered.

AUTHORITY FOR COMMITTEES TO REPORT LEGISLATIVE AND EXECUTIVE MATTERS

Mr. FRIST. Mr. President, I ask unanimous consent that notwithstanding the Senate's adjournment, committees be authorized to report legislative and executive matters on Thursday, March 18, from 10 a.m. to 12 noon.

The PRESIDING OFFICER. Without objection, it is so ordered.

MEASURE READ THE FIRST TIME—S. 2207

Mr. FRIST. Mr. President, I understand that S. 2207 is at the desk, and I ask for its first reading.

The PRESIDING OFFICER. The clerk will report.

The legislative clerk read as follows:

A bill (S. 2207) to improve women's access to health care services, and the access of all individuals to emergency and trauma care services, by reducing the excessive burden the liability system places on the delivery of such services.

Mr. FRIST. I now ask for its second reading in order to place the bill on the Calendar under provisions of rule XIV and object to my own request.

The PRESIDING OFFICER. The bill will be read the second time on the next legislative day.

MEDICAL DEVICES TECHNICAL CORRECTIONS ACT

Mr. FRIST. Mr. President, I ask unanimous consent that the Chair lay before the Senate a message from the House of Representatives on (S. 1881) to amend the Federal Food, Drug, and Cosmetic Act to make technical corrections relating to the amendments by the Medical Device User Fee and Modernization Act of 2002, and for other purposes.

The PRESIDING OFFICER laid before the Senate the following message from the House of Representatives:

Resolved, That the bill from the Senate (S. 1881) entitled ''An Act to amend the Federal Food, Drug, and Cosmetic Act to make technical corrections relating to the amendments made by the Medical Device User Fee and Modernization Act of 2002, and for other purposes'', do pass with the following amendment:

Strike out all after the enacting clause and insert:

SECTION 1. SHORT TITLE.

This Act may be cited as the ''Medical Devices Technical Corrections Act''.

SEC. 2. TECHNICAL CORRECTIONS REGARDING PUBLIC LAW 107–250.

(a) TITLE I; FEES RELATING TO MEDICAL DEVICES.—Part 3 of subchapter C of chapter VII of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 379i et seq.), as added by section 102 of Public Law 107–250 (116 Stat. 1589), is amended—

(1) in section 737—

(A) in paragraph (4)(B), by striking ''and for which clinical data are generally necessary to provide a reasonable assurance of safety and effectiveness'' and inserting ''and for which substantial clinical data are necessary to provide a reasonable assurance of safety and effectiveness'';

(B) in paragraph (4)(D), by striking ''manufacturing,'';

(C) in paragraph (5)(J), by striking ''a premarket application'' and all that follows and inserting ''a premarket application or premarket report under section 515 or a premarket application under section 351 of the Public Health Service Act.''; and

(D) in paragraph (8), by striking ''The term 'affiliate' means a business entity that has a relationship with a second business entity'' and inserting ''The term 'affiliate' means a business entity that has a relationship with a second business entity (whether domestic or international)''; and

(2) in section 738—

(A) in subsection (a)(1)—

(i) in subparagraph (A)—

(I) in the matter preceding clause (i) by striking ''subsection (d),'' and inserting ''subsections (d) and (e),'';

(II) in clause (iv), by striking ''clause (i),'' and all that follows and inserting ''clause (i).''; and

(III) in clause (vii), by striking ''clause (i),'' and all that follows and inserting ''clause (i), subject to any adjustment under subsection (e)(2)(C)(ii).''; and

(ii) in subparagraph (D), in each of clauses (i) and (ii), by striking ''application'' and inserting ''application, report,'';

(B) in subsection (d)(2)(B), beginning in the second sentence, by striking ''firms. which show'' and inserting ''firms, which show'';

(C) in subsection (e)—

(i) in paragraph (1), by striking ''Where'' and inserting ''For fiscal year 2004 and each subsequent fiscal year, where''; and

(ii) in paragraph (2)—

(I) in subparagraph (B), beginning in the second sentence, by striking ''firms. which show'' and inserting ''firms, which show''; and

(II) in subparagraph (C)(i), by striking ''Where'' and inserting ''For fiscal year 2004 and each subsequent fiscal year, where'';

(D) in subsection (f), by striking ''for filing''; and

(E) in subsection (h)(2)(B)—

(i) in clause (ii), by redesignating subclauses (I) and (II) as items (aa) and (bb), respectively;

(ii) by redesignating clauses (i) and (ii) as subclauses (I) and (II), respectively;

(iii) by striking ''The Secretary'' and inserting the following:

''(i) IN GENERAL.—The Secretary''; and

(iv) by adding at the end the following:

''(ii) MORE THAN 5 PERCENT.—To the extent such costs are more than 5 percent below the specified level in subparagraph (A)(ii), fees may not be collected under this section for that fiscal year.''.

(b) TITLE II; AMENDMENTS REGARDING REGULATION OF MEDICAL DEVICES.—

(1) INSPECTIONS BY ACCREDITED PERSONS.—Section 704(g) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 374(g)), as added by section 201 of Public Law 107–250 (116 Stat. 1602), is amended—

(A) in paragraph (1), in the first sentence, by striking ''conducting inspections'' and all that follows and inserting ''conducting inspections of establishments that manufacture, prepare, propagate, compound, or process class II or class III devices, which inspections are required under section 510(h) or are inspections of such establishments required to register under section 510(i).'';

(B) in paragraph (5)(B), in the first sentence, by striking ''or poses'' and all that follows through the period and inserting ''poses a threat to public health, fails to act in a manner that is consistent with the purposes of this subsection, or where the Secretary determines that there is a financial conflict of interest in the relationship between the accredited person and the owner or operator of a device establishment that the accredited person has inspected under this subsection.'';

(C) in paragraph (6)(A)—

(i) in clause (i), by striking ''of the establishment pursuant to subsection (h) or (i) of section 510'' and inserting ''described in paragraph (1)'';

(ii) in clause (ii)—

(I) in the matter preceding subclause (I)—

(aa) by striking ''each inspection'' and inserting ''inspections''; and

(bb) by inserting ''during a 2-year period'' after ''person''; and

(II) in subclause (I), by striking ''such a person'' and inserting ''an accredited person'';

(iii) in clause (iii)—

(I) in the matter preceding subclause (I), by striking ''and the following additional conditions are met:'' and inserting ''and 1 or both of the following additional conditions are met:'';

(II) in subclause (I), by striking ''accredited'' and all that follows through the period and inserting ''(accredited under paragraph (2) and