

obligations under the Hague Convention, which is the substance in S. 2883.

This measure has the support of the relevant House and Senate Committees and the Departments of State and Justice. If this measure is not enacted into law, NCMC may not be able to continue its operations on behalf of the Federal Government since its resources would be lost in the defense of frivolous lawsuits. Left-behind parents would suffer the prolonged loss of their children, and our Nation potentially would lose its advantage in pressing other nations to return abducted children.

I wish to extend my personal gratitude to the National Center for Missing and Exploited Children for its critical work on reuniting families, to Chairman JIM SENSENBRENNER of the House Judiciary Committee, and to Senators HATCH and LEAHY of the Senate Judiciary Committee and to Senators LUGAR and BIDEN of the Senate Foreign Relations Committee, for working tirelessly to implement this measure.

GENERAL LEAVE

Mr. CANNON. Mr. Speaker, I ask unanimous consent that all Members may have 5 legislative days within which to revise and extend their remarks and include extraneous material on S. 2883.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

ANABOLIC STEROID CONTROL ACT OF 2004

Mr. CANNON. Mr. Speaker, I ask unanimous consent to take from the Speaker's table the Senate bill (S. 2195) to amend the Controlled Substances Act to clarify the definition of anabolic steroids and to provide for research and education activities relating to steroids and steroid precursors, and ask for its immediate consideration in the House.

The Clerk read the title of the Senate bill.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

The Clerk read the Senate bill, as follows:

S. 2195

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 "Anabolic Steroid Control Act of 2004".

SEC. 2. AMENDMENTS TO THE CONTROLLED SUBSTANCES ACT.

(a) DEFINITIONS.—Section 102 of the Controlled Substances Act (21 U.S.C. 802) is amended—

(1) in paragraph (41)—

(A) by realigning the margin so as to align with paragraph (40); and

(B) by striking subparagraph (A) and inserting the following:

“(A) The term ‘anabolic steroid’ means any drug or hormonal substance, chemically and pharmacologically related to testosterone

(other than estrogens, progestins, corticosteroids, and dehydroepiandrosterone), and includes—

“(i) androstenediol—

“(I) 3 β ,17 β -dihydroxy-5 α -androstane; and

“(II) 3 α ,17 β -dihydroxy-5 α -androstane; and

“(ii) androstenedione (5 α -androstane-3,17-dione);

“(iii) androstenediol—

“(I) 1-androstenediol (3 β ,17 β -dihydroxy-5 α -androst-1-ene);

“(II) 1-androstenediol (3 α ,17 β -dihydroxy-5 α -androst-1-ene);

“(III) 4-androstenediol (3 β ,17 β -dihydroxy-androst-4-ene); and

“(IV) 5-androstenediol (3 β ,17 β -dihydroxy-androst-5-ene);

“(iv) androstenedione—

“(I) 1-androstenedione ([5 α]-androst-1-en-3,17-dione);

“(II) 4-androstenedione (androst-4-en-3,17-dione); and

“(III) 5-androstenedione (androst-5-en-3,17-dione);

“(v) bolasterone (7 α ,17 α -dimethyl-17 β -hydroxyandrost-4-en-3-one);

“(vi) boldenone (17 β -hydroxyandrost-1,4-diene-3-one);

“(vii) calusterone (7 β ,17 α -dimethyl-17 β -hydroxyandrost-4-en-3-one);

“(viii) clostebol (4-chloro-17 β -hydroxyandrost-4-en-3-one);

“(ix) dehydrochloromethyltestosterone (4-chloro-17 β -hydroxy-17 α -methyl-androst-1,4-dien-3-one);

“(x) Δ 1-dihydrotestosterone (a.k.a. ‘1-testosterone’) (17 β -hydroxy-5 α -androst-1-en-3-one);

“(xi) 4-dihydrotestosterone (17 β -hydroxy-androst-3-one);

“(xii) drostanolone (17 β -hydroxy-2 α -methyl-5 α -androst-3-one);

“(xiii) ethylestrenol (17 α -ethyl-17 β -hydroxyestr-4-ene);

“(xiv) fluoxymesterone (9-fluoro-17 α -methyl-11 β ,17 β -dihydroxyandrost-4-en-3-one);

“(xv) formebolone (2-formyl-17 α -methyl-11 α ,17 β -dihydroxyandrost-1,4-dien-3-one);

“(xvi) furazabol (17 α -methyl-17 β -hydroxyandrostano[2,3-c]-furan);

“(xvii) 13 β -ethyl-17 α -hydroxygon-4-en-3-one;

“(xviii) 4-hydroxytestosterone (4,17 β -dihydroxy-androst-4-en-3-one);

“(xix) 4-hydroxy-19-nortestosterone (4,17 β -dihydroxy-estr-4-en-3-one);

“(xx) mestanolone (17 α -methyl-17 β -hydroxy-5 α -androst-3-one);

“(xxi) mesterolone (1 α -methyl-17 β -hydroxy-[5 α]-androst-3-one);

“(xxii) methandienone (17 α -methyl-17 β -hydroxyandrost-1,4-dien-3-one);

“(xxiii) methandriol (17 α -methyl-3 β ,17 β -dihydroxyandrost-5-ene);

“(xxiv) methenolone (1-methyl-17 β -hydroxy-5 α -androst-1-en-3-one);

“(xxv) 17 α -methyl-3 β , 17 β -dihydroxy-5 α -androstane;

“(xxvi) 17 α -methyl-3 α ,17 β -dihydroxy-5 α -androstane;

“(xxvii) 17 α -methyl-3 β ,17 β -dihydroxy-androst-4-ene.

“(xxviii) 17 α -methyl-4-hydroxynandrolone (17 α -methyl-4-hydroxy-17 β -hydroxyestr-4-en-3-one);

“(xxix) methylidenolone (17 α -methyl-17 β -hydroxyestra-4,9(10)-dien-3-one);

“(xxx) methyltrienolone (17 α -methyl-17 β -hydroxyestra-4,9-11-trien-3-one);

“(xxxi) methyltestosterone (17 α -methyl-17 β -hydroxyandrost-4-en-3-one);

“(xxxii) mibolerone (7 α ,17 α -dimethyl-17 β -hydroxyestr-4-en-3-one);

“(xxxiii) 17 α -methyl- Δ 1-dihydrotestosterone (17 β -hydroxy-17 α -methyl-5 α -androst-1-en-3-one) (a.k.a. ‘17 α -methyl-1-testosterone’);

“(xxxiv) nandrolone (17 β -hydroxyestr-4-en-3-one);

“(xxxv) norandrostenediol—

“(I) 19-nor-4-androstenediol (3 β , 17 β -dihydroxyestr-4-ene);

“(II) 19-nor-4-androstenediol (3 α , 17 β -dihydroxyestr-4-ene);

“(III) 19-nor-5-androstenediol (3 β , 17 β -dihydroxyestr-5-ene); and

“(IV) 19-nor-5-androstenediol (3 α , 17 β -dihydroxyestr-5-ene);

“(xxxvi) norandrostenedione—

“(I) 19-nor-4-androstenedione (estr-4-en-3,17-dione); and

“(II) 19-nor-5-androstenedione (estr-5-en-3,17-dione);

“(xxxvii) norbolethone (13 β ,17 α -diethyl-17 β -hydroxygon-4-en-3-one);

“(xxxviii) norclostebol (4-chloro-17 β -hydroxyestr-4-en-3-one);

“(xxxix) norethandrolone (17 α -ethyl-17 β -hydroxyestr-4-en-3-one);

“(xl) normethandrolone (17 α -methyl-17 β -hydroxyestr-4-en-3-one);

“(xli) oxandrolone (17 α -methyl-17 β -hydroxy-2-oxa-[5 α]-androst-3-one);

“(xlii) oxymesterone (17 α -methyl-4,17 β -dihydroxyandrost-4-en-3-one);

“(xliii) oxymetholone (17 α -methyl-2-hydroxymethylene-17 β -hydroxy-[5 α]-androst-3-one);

“(xliv) stanozolol (17 α -methyl-17 α -hydroxy-[5 α]-androst-2-eno[3,2-c]-pyrazole);

“(xlv) stenbolone (17 β -hydroxy-2-methyl-[5 α]-androst-1-en-3-one);

“(xlii) testolactone (13-hydroxy-3-oxo-13,17-secoandrost-1,4-dien-17-oic acid lactone);

“(xlvii) testosterone (17 β -hydroxyandrost-4-en-3-one);

“(xlviii) tetrahydrogestrinone (13 β ,17 α -diethyl-17 β -hydroxygon-4,9,11-trien-3-one);

“(xlix) trenbolone (17 β -hydroxyestr-4,9,11-trien-3-one); and

“(xlx) any salt, ester, or ether of a drug or substance described in this paragraph.

The substances excluded under this subparagraph may at any time be scheduled by the Attorney General in accordance with the authority and requirements of subsections (a) through (c) of section 201; and

(2) in paragraph (44), by inserting “anabolic steroids,” after “marihuana,”.

(b) AUTHORITY AND CRITERIA FOR CLASSIFICATION.—Section 201(g) of the Controlled Substances Act (21 U.S.C. 811(g)) is amended—

(1) in paragraph (1), by striking “substance from a schedule if such substance” and inserting “drug which contains a controlled substance from the application of titles II and III of the Comprehensive Drug Abuse Prevention and Control Act (21 U.S.C. 802 et seq.) if such drug”; and

(2) in paragraph (3), by adding at the end the following:

“(C) Upon the recommendation of the Secretary of Health and Human Services, a compound, mixture, or preparation which contains any anabolic steroid, which is intended for administration to a human being or an animal, and which, because of its concentration, preparation, formulation or delivery system, does not present any significant potential for abuse.”.

(c) ANABOLIC STEROIDS CONTROL ACT.—Section 1903 of the Anabolic Steroids Control Act of 1990 (Public Law 101-647) is amended—

(1) by striking subsection (a); and

(2) by redesignating subsections (b) and (c) as subsections (a) and (b), respectively.

(d) EFFECTIVE DATE.—The amendments made by this section shall take effect 90 days after the date of enactment of this Act.

SEC. 3. SENTENCING COMMISSION GUIDELINES.

The United States Sentencing Commission shall—

(1) review the Federal sentencing guidelines with respect to offenses involving anabolic steroids;

(2) consider amending the Federal sentencing guidelines to provide for increased penalties with respect to offenses involving anabolic steroids in a manner that reflects the seriousness of such offenses and the need to deter anabolic steroid trafficking and use; and

(3) take such other action that the Commission considers necessary to carry out this section.

SEC. 4. PREVENTION AND EDUCATION PROGRAMS.

(a) IN GENERAL.—The Secretary of Health and Human Services (referred to in this Act as the “Secretary”) shall award grants to public and nonprofit private entities to enable such entities to carry out science-based education programs in elementary and secondary schools to highlight the harmful effects of anabolic steroids.

(b) ELIGIBILITY.—

(1) APPLICATION.—To be eligible for grants under subsection (a), an entity shall prepare and submit to the Secretary an application at such time, in such manner, and containing such information as the Secretary may require.

(2) PREFERENCE.—In awarding grants under subsection (a), the Secretary shall give preference to applicants that intend to use grant funds to carry out programs based on—

(A) the Athletes Training and Learning to Avoid Steroids program;

(B) The Athletes Targeting Healthy Exercise and Nutrition Alternatives program; and

(C) other programs determined to be effective by the National Institute on Drug Abuse.

(c) USE OF FUNDS.—Amounts received under a grant under subsection (a) shall be used for education programs that will directly communicate with teachers, principals, coaches, as well as elementary and secondary school children concerning the harmful effects of anabolic steroids.

(d) AUTHORIZATION OF APPROPRIATIONS.—There is authorized to be appropriated to carry out this section, \$15,000,000 for each of fiscal years 2005 through 2010.

SEC. 5. NATIONAL SURVEY ON DRUG USE AND HEALTH.

(a) IN GENERAL.—The Secretary of Health and Human Services shall ensure that the National Survey on Drug Use and Health includes questions concerning the use of anabolic steroids.

(b) AUTHORIZATION OF APPROPRIATIONS.—There is authorized to be appropriated to carry out this section, \$1,000,000 for each of fiscal years 2005 through 2010.

The Senate bill was ordered to be read a third time, was read the third time, and passed, and a motion to reconsider was laid on the table.

GENERAL LEAVE

Mr. CANNON. Mr. Speaker, I ask unanimous consent that all Members may have 5 legislative days within which to revise and extend their remarks and include extraneous material on S. 2195.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

FEDERAL REGULATORY IMPROVEMENT ACT OF 2004

Mr. CANNON. Mr. Speaker, I ask unanimous consent that the Com-

mittee on the Judiciary be discharged from further consideration of the bill (H.R. 4917) to amend title 5, United States Code, to authorize appropriations for the Administrative Conference of the United States for fiscal years 2005, 2006, and 2007, and for other purposes, and ask for its immediate consideration in the House.

The Clerk read the title of the bill.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

The Clerk read the bill, as follows:

H.R. 4917

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 “Federal Regulatory Improvement Act of 2004”.

SEC. 2. PURPOSES.

(a) PURPOSES.—Section 591 of title 5, United States Code, is amended to read as follows:

“§ 591 Purposes

“The purposes of this subchapter are—

“(1) to provide suitable arrangements through which Federal agencies, assisted by outside experts, may cooperatively study mutual problems, exchange information, and develop recommendations for action by proper authorities to the end that private rights may be fully protected and regulatory activities and other Federal responsibilities may be carried out expeditiously in the public interest;

“(2) to promote more effective public participation and efficiency in the rulemaking process;

“(3) to reduce unnecessary litigation in the regulatory process;

“(4) to improve the use of science in the regulatory process; and

“(5) to improve the effectiveness of laws applicable to the regulatory process.”.

(b) CONFORMING AMENDMENTS.—Title 5 of the United States Code is amended—

(1) in section 594 by striking “purpose” and inserting “purposes”; and

(2) in the table of sections of chapter 5 of part I by amending the item relating to section 591 to read as follows:

“591. Purposes”.

SEC. 3. AUTHORIZATION OF APPROPRIATIONS.

Section 596 of title 5, United States Code, is amended to read as follows:

“§ 596. Authorization of appropriations

“There are authorized to be appropriated to carry out this subchapter not more than \$3,000,000 for fiscal year 2005, \$3,100,000 for fiscal year 2006, and \$3,200,000 for fiscal year 2007. Of any amounts appropriated under this section, not more than \$2,500 may be made available in each fiscal year for official representation and entertainment expenses for foreign dignitaries.”.

The bill was ordered to be engrossed and read a third time, was read the third time, and passed, and a motion to reconsider was laid on the table.

GENERAL LEAVE

Mr. CANNON. Mr. Speaker, I ask unanimous consent that all Members may have 5 legislative days within which to revise and extend their remarks and include extraneous material on H.R. 4917.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

FAMILY FARMER BANKRUPTCY RELIEF ACT OF 2004

Mr. CANNON. Mr. Speaker, I ask unanimous consent that the Committee on the Judiciary be discharged from further consideration of the Senate bill (S. 2864) to extend for eighteen months the period for which chapter 12 of title 11, United States Code, is reenacted, and ask for its immediate consideration in the House.

The Clerk read the title of the Senate bill.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

The Clerk read the Senate bill, as follows:

S. 2864

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 “Family Farmer Bankruptcy Relief Act of 2004”.

SEC. 2. EIGHTEEN-MONTH EXTENSION OF PERIOD FOR WHICH CHAPTER 12 OF TITLE 11, UNITED STATES CODE, IS REENACTED.

(a) AMENDMENTS.—Section 149 of title I of division C of Public Law 105-277 (11 U.S.C. 1201 note) is amended—

(1) by striking “January 1, 2004” each place that term appears and inserting “July 1, 2005”; and

(2) in subsection (a)—

(A) by striking “June 30, 2003” and inserting “December 31, 2003”; and

(B) by striking “July 1, 2003” and inserting “January 1, 2004”.

(b) EFFECTIVE DATE.—The amendments made by subsection (a) are deemed to have taken effect on January 1, 2004.

The Senate bill was ordered to be read a third time, was read the third time, and passed, and a motion to reconsider was laid on the table.

GENERAL LEAVE

Mr. CANNON. Mr. Speaker, I ask unanimous consent that all Members may have 5 legislative days within which to revise and extend their remarks and include extraneous material on S. 2864.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from Utah?

There was no objection.

ASSISTIVE TECHNOLOGY ACT OF 2004

Mr. McKEON. Mr. Speaker, I ask unanimous consent to take from the Speaker's table the bill (H.R. 4278) to amend the Assistive Technology Act of 1998 to support programs of grants to States to address the assistive technology needs of individuals with disabilities, and for other purposes, with a