

The question was taken.

Mr. BOEHNER. Mr. Speaker, on that I demand the yeas and nays.

The yeas and nays were ordered.

The SPEAKER pro tempore. Pursuant to clause 8 of rule XX and the Chair's prior announcement, further proceedings on this motion will be postponed.

GENERAL LEAVE

Mr. BOEHNER. 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 the subject of the concurrent resolution just considered.

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

There was no objection.

PERMITTING OFFICIAL PHOTOGRAPHS OF HOUSE WHILE IN SESSION

Mr. BOEHNER. Mr. Speaker, I move to suspend the rules and agree to the resolution (H. Res. 407) permitting official photographs of the House of Representatives to be taken while the House is in actual session.

The Clerk read as follows:

H. RES. 407

Resolved, That at a time designated by the Speaker of the House of Representatives, official photographs of the House may be taken while the House is in actual session. Payment for the costs associated with taking, preparing, and distributing such photographs may be made from the applicable accounts of the House of Representatives.

The SPEAKER pro tempore. Pursuant to the rule, the gentleman from Ohio (Mr. BOEHNER) and the gentleman from Maryland (Mr. HOYER) each will control 20 minutes.

The Chair recognizes the gentleman from Ohio (Mr. BOEHNER).

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Mr. BOEHNER. Mr. Speaker, I yield myself such time as I may consume.

Mr. Speaker, this resolution is very straightforward and simply authorizes the use of the Chamber for a photo while we are in session. The Speaker would set the date for such photo and payment as authorized from the applicable accounts of the House.

As Members know, in the last session of Congress there was a photo taken of all of the Members of the House, something that was rather routine in sessions past, but over a period of 3 or 4 sessions it did not occur. Several years ago when this was done the Members were very supportive of the effort, and the Committee on House Administration voted for it. The Members thereof have suggested that the House take another photograph in this session.

Mr. Speaker, I reserve the balance of my time.

Mr. HOYER. Mr. Speaker, I yield myself such time as I may consume.

Mr. Speaker, my staff behind me has suggested that Members should not forget to smile. I think it is appropriate that we take a picture of the House of Representatives and its Members on an annual basis, or at least once during every Congress. I think this is not only a substantial memento for those who have the great honor and privilege of serving here, but as well, an historical record of those who are here, and of course I rise in strong support of the resolution.

Mr. Speaker, I yield back the balance of my time.

Mr. BOEHNER. Mr. Speaker, I yield back the balance of my time.

The SPEAKER pro tempore (Mr. PETRI). The question is on the motion offered by the gentleman from Ohio (Mr. BOEHNER) that is House suspend the rules and agree to the resolution, H. Res. 407.

The question was taken; and (two-thirds having voted in favor thereof), the rules were suspended and the resolution was agreed to.

A motion to reconsider was laid on the table.

HILLORY J. FARIAS AND SAMANTHA REID DATE-RAPE DRUG PROHIBITION ACT OF 1999

Mr. UPTON. Mr. Speaker, I move to suspend the rules and concur in the Senate amendments to the bill (H.R. 2130) to amend the Controlled Substances Act to add gamma hydroxybutyric acid and ketamine to the schedules of controlled substances, to provide for a national awareness campaign, and for other purposes.

The Clerk read as follows:

Senate amendments:

Strike out all after the enacting clause and insert:

SECTION 1. SHORT TITLE.

This Act may be cited as the "Hillory J. Farias and Samantha Reid Date-Rape Drug Prohibition Act of 1999".

SEC. 2. FINDINGS.

Congress finds as follows:

(1) Gamma hydroxybutyric acid (also called G, Liquid X, Liquid Ecstasy, Grievous Bodily Harm, Georgia Home Boy, Scoop) has become a significant and growing problem in law enforcement. At least 20 States have scheduled such drug in their drug laws and law enforcement officials have been experiencing an increased presence of the drug in driving under the influence, sexual assault, and overdose cases especially at night clubs and parties.

(2) A behavioral depressant and a hypnotic, gamma hydroxybutyric acid ("GHB") is being used in conjunction with alcohol and other drugs with detrimental effects in an increasing number of cases. It is difficult to isolate the impact of such drug's ingestion since it is so typically taken with an ever-changing array of other drugs and especially alcohol which potentiates its impact.

(3) GHB takes the same path as alcohol, processes via alcohol dehydrogenase, and its symptoms at high levels of intake and as impact builds are comparable to alcohol ingestion/in-

torication. Thus, aggression and violence can be expected in some individuals who use such drug.

(4) If taken for human consumption, common industrial chemicals such as gamma butyrolactone and 1,4-butanediol are swiftly converted by the body into GHB. Illicit use of these and other GHB analogues and precursor chemicals is a significant and growing law enforcement problem.

(5) A human pharmaceutical formulation of gamma hydroxybutyric acid is being developed as a treatment for cataplexy, a serious and debilitating disease. Cataplexy, which causes sudden and total loss of muscle control, affects about 65 percent of the estimated 180,000 Americans with narcolepsy, a sleep disorder. People with cataplexy often are unable to work, drive a car, hold their children or live a normal life.

(6) Abuse of illicit GHB is an imminent hazard to public safety that requires immediate regulatory action under the Controlled Substances Act (21 U.S.C. 801 et seq.).

SEC. 3. EMERGENCY SCHEDULING OF GAMMA HYDROXYBUTYRIC ACID AND LISTING OF GAMMA BUTYROLACTONE AS LIST I CHEMICAL.

(a) EMERGENCY SCHEDULING OF GHB.—

(1) IN GENERAL.—The Congress finds that the abuse of illicit gamma hydroxybutyric acid is an imminent hazard to the public safety. Accordingly, the Attorney General, notwithstanding sections 201(a), 201(b), 201(c), and 202 of the Controlled Substances Act, shall issue, not later than 60 days after the date of the enactment of this Act, a final order that schedules such drug (together with its salts, isomers, and salts of isomers) in the same schedule under section 202(c) of the Controlled Substances Act as would apply to a scheduling of a substance by the Attorney General under section 201(h)(1) of such Act (relating to imminent hazards to the public safety), except as follows:

(A) For purposes of any requirements that relate to the physical security of registered manufacturers and registered distributors, the final order shall treat such drug, when the drug is manufactured, distributed, or possessed in accordance with an exemption under section 505(i) of the Federal Food, Drug, and Cosmetic Act (whether the exemption involved is authorized before, on, or after the date of the enactment of this Act), as being in the same schedule as that recommended by the Secretary of Health and Human Services for the drug when the drug is the subject of an authorized investigational new drug application (relating to such section 505(i)). The recommendation referred to in the preceding sentence is contained in the first paragraph of the letter transmitted on May 19, 1999, by such Secretary (acting through the Assistant Secretary for Health) to the Attorney General (acting through the Deputy Administrator of the Drug Enforcement Administration), which letter was in response to the letter transmitted by the Attorney General (acting through such Deputy Administrator) on September 16, 1997. In publishing the final order in the Federal Register, the Attorney General shall publish a copy of the letter that was transmitted by the Secretary of Health and Human Services.

(B) In the case of gamma hydroxybutyric acid that is contained in a drug product for which an application is approved under section 505 of the Federal Food, Drug, and Cosmetic Act (whether the application involved is approved before, on, or after the date of the enactment of this Act), the final order shall schedule such drug in the same schedule as that recommended by the Secretary of Health and Human Services for authorized formulations of the drug. The recommendation referred to in the preceding sentence is contained in the last sentence of the fourth paragraph of the letter referred to in subparagraph (A) with respect to May 19, 1999.