

110TH CONGRESS  
2D SESSION

# H. R. 6820

To direct the Administrator of the Environmental Protection Agency to conduct a study of the presence of pharmaceuticals and personal care products in drinking water supplies in the United States, and for other purposes.

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## IN THE HOUSE OF REPRESENTATIVES

AUGUST 1, 2008

Mrs. MCCARTHY of New York (for herself, Ms. BALDWIN, and Ms. SCHWARTZ) introduced the following bill; which was referred to the Committee on Energy and Commerce

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## A BILL

To direct the Administrator of the Environmental Protection Agency to conduct a study of the presence of pharmaceuticals and personal care products in drinking water supplies in the United States, and for other purposes.

1 *Be it enacted by the Senate and House of Representa-*  
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Water Assessment and  
5 Treatment Evaluation Research Study Act of 2008” or  
6 the “WATER Study Act of 2008”.

1 **SEC. 2. PRESENCE OF PHARMACEUTICALS AND PERSONAL**  
2 **CARE PRODUCTS IN DRINKING WATER SUP-**  
3 **PLIES OF THE UNITED STATES.**

4 (a) **STUDY.**—The Administrator of the Environ-  
5 mental Protection Agency, in consultation with appro-  
6 priate government agencies (including the National Insti-  
7 tute of Environmental Health Sciences and the United  
8 States Geological Survey), shall conduct a study of the  
9 presence of pharmaceuticals and personal care products  
10 (in this section referred to as “PPCPs”) in drinking water  
11 supplies in the United States.

12 (b) **EXAMINATION OF WASTEWATER EFFLUENT AND**  
13 **RUN-OFF FROM AGRICULTURAL OPERATIONS.**—In iden-  
14 tifying sources of PPCPs under the study, the Adminis-  
15 trator shall examine wastewater effluent and run-off from  
16 agricultural operations.

17 (c) **REPORTS.**—

18 (1) **IN GENERAL.**—The Administrator shall  
19 submit to Congress an initial, interim, and final re-  
20 port on the results of the study conducted under this  
21 section.

22 (2) **INITIAL REPORT.**—The initial report shall  
23 be submitted not later than one year after the date  
24 of enactment of this Act and shall at a minimum—

25 (A) identify PPCPs that have been de-  
26 tected in the drinking water supplies of the

1 United States and the levels at which such  
2 PPCPs have been detected;

3 (B) identify the sources of PPCPs in the  
4 drinking waters of the United States; and

5 (C) identify and evaluate methods to regu-  
6 larly monitor the levels of PPCPs in the drink-  
7 ing water of the United States.

8 (3) INTERIM REPORT.—The interim report shall  
9 be submitted not later than 3 years after the date  
10 of enactment of this Act and shall at a minimum—

11 (A) identify the effects of PPCPs in drink-  
12 ing water supplies and other waters of the  
13 United States on human health and aquatic life  
14 at the levels at which PPCPs have been de-  
15 tected;

16 (B) identify methods and techniques to  
17 safely and properly dispose of unused PPCPs;

18 (C) identify and evaluate methods to re-  
19 move PPCPs from the drinking water supplies  
20 of the United States;

21 (D) identify and evaluate ways to improve  
22 the treatment of water discharged from waste-  
23 water treatment plants, including the environ-  
24 mental and economic costs of these methods.

1           (4) FINAL REPORT.—The final report shall be  
2 submitted not later than 5 years after the date of  
3 enactment of this Act and shall at a minimum—

4           (A) identify the effects of long-term expo-  
5 sure to PPCPs;

6           (B) identify levels at which the PPCPs  
7 identified in the initial report become harmful  
8 to humans;

9           (C) identify the effects on humans of inter-  
10 actions between PPCPs identified in the initial  
11 report; and

12           (D) include all information required by the  
13 initial report and interim report along with any  
14 updated information that is available on those  
15 requirements.

16           (5) INCLUSION OF AVAILABLE INFORMATION.—  
17 If the Administrator has information required for a  
18 report under paragraph (3) or (4) at the time of  
19 submission of an earlier report under this sub-  
20 section, the Administrator shall include the informa-  
21 tion in the earlier report.

22           (d) PHARMACEUTICALS AND PERSONAL CARE PROD-  
23 UCTS DEFINED.—In this section, the terms “pharma-  
24 ceuticals and personal care products” and “PPCPs” mean  
25 products used by individuals for personal health or cos-

1. medic reasons, including soaps and shampoos, or used by
2. agribusiness to enhance growth or health of livestock.

