

**AMENDMENT TO THE AMENDMENT IN THE  
NATURE OF A SUBSTITUTE TO H.R. 5320  
OFFERED BY MS. BALDWIN OF WISCONSIN**

At the end add the following new section (and revise the table of contents in section 1(b) accordingly):

1 **SEC. \_\_\_\_ . PRESENCE OF PHARMACEUTICALS AND PER-**  
2 **SONAL CARE PRODUCTS IN SOURCES OF**  
3 **DRINKING WATER.**

4 Subsection (a) of section 1442 (42 U.S.C. 300j-1)  
5 is amended by adding at the end the following:

6 “(11) PRESENCE OF PHARMACEUTICALS AND PER-  
7 SONAL CARE PRODUCTS IN SOURCES OF DRINKING  
8 WATER.—

9 “(A) STUDY.—The Administrator shall carry  
10 out a study on the presence of pharmaceuticals and  
11 personal care products in sources of drinking water,  
12 which shall—

13 “(i) identify pharmaceuticals and personal  
14 care products that have been detected in  
15 sources of drinking water and the levels at  
16 which such pharmaceuticals and personal care  
17 products have been detected;

1           “(ii) identify the sources of pharma-  
2           ceuticals and personal care products in sources  
3           of drinking water, including point sources and  
4           nonpoint sources of pharmaceutical and per-  
5           sonal care products;

6           “(iii) identify the effects of such products  
7           on humans, the environment, and the safety of  
8           drinking water; and

9           “(iv) identify methods to control, limit,  
10          treat, or prevent the presence of such products.

11          “(B) CONSULTATION.—The Administrator shall  
12          conduct the study described in subparagraph (A) in  
13          consultation with the Secretary of Health and  
14          Human Services (acting through the Commissioner  
15          of Food and Drugs), the Director of the United  
16          States Geological Survey, the heads of other appro-  
17          priate Federal agencies (including the National In-  
18          stitute of Environmental Health Sciences), and other  
19          interested stakeholders (including manufacturers of  
20          pharmaceuticals and personal care products and  
21          consumer groups and advocates).

22          “(C) REPORT.—Not later than 2 years after  
23          the date of the enactment of this paragraph, the Ad-  
24          ministrator shall submit to the Congress a report on

1 the results of the study carried out under this para-  
2 graph.

3 “(D) DEFINITIONS.—In this paragraph:

4 “(i) The term ‘personal care product’ has  
5 the meaning given the term ‘cosmetic’ in section  
6 201 of the Federal Food, Drug, and Cosmetic  
7 Act.

8 “(ii) The term ‘pharmaceutical’ has the  
9 meaning given the term ‘drug’ in section 201 of  
10 the Federal Food, Drug, and Cosmetic Act.”.



