



June 19, 2008

Clay G. Small
Senior Vice President
Legal Affairs

The Honorable John D. Dingell
Chairman
House Committee on Energy and Commerce
2125 Rayburn House Office Building
Washington, D.C. 20515

The Honorable Bart Stupak
Chairman
Subcommittee on Oversight and Investigations
House Committee on Energy and Commerce
2125 Rayburn House Office Building
Washington, D.C. 20515

Re: PepsiCo, Inc. Response to May 8, 2008 Inquiry

Dear Chairman Dingell and Chairman Stupak:

I am responding to your May 8, 2008 letter to Ms. Indra K. Nooyi requesting certain information regarding the safety of food processed and sold by PepsiCo, Inc. (PepsiCo). In this letter we provide responses to questions one through six on behalf of our international business units, including Puerto Rico and U.S. territories, for food products intended for consumption in the United States and its territories. As the Committee staff have requested, Ms. Nooyi will be submitting an additional letter next week covering all of PepsiCo's responses.

PepsiCo has numerous international business units throughout the world. Although the majority of the markets for the PepsiCo international business units are external to the United States, a limited number of business units produce and distribute products for consumption in the United States and its territories. PepsiCo's international business units adhere to PepsiCo's global standards for food safety and quality and meet local and international standards, where applicable.

In the preparation of this response, we followed the same procedures and process explained in our earlier submissions, which we do not repeat again in our responses to each question. As with our previous submissions, we are attaching to this letter confidential information, which we have identified as "confidential." For your convenience, we first repeat each question and then provide our response.

1. A list of all food recalls and food safety alerts issued by your company. For each recall or safety alert, please provide the date of the recall or alert, the product and brand affected, and the reason for the recall or alert. If the food was affected by microbial or chemical contamination, please identify the contaminant.

Attachment 1 contains a table identifying all recalls initiated by PepsiCo's international business units for products intended for U.S. consumption since January 1, 2000. For each recall we have identified the date of the recall, the brand and product affected, the reason for the recall, and whether FDA was notified. We have also identified the FDA class of recall.

2. For each brand or kind of product, please list all instances when internal microbiological testing was found to be positive for the presence of E. coli, Salmonella, Cyclospora cayetanensis, Cryptosporidium, hepatitis A, Clostridium botulinum, or Listeria in excess of the highest limit acceptable to the Food and Drug Administration (FDA) or any State regulatory authority.

We have no additional information to report in response to this question.

3. For each brand or kind of product, please list the instances when internal testing was found to be positive for the presence of a chemical contaminant at levels in excess of the highest limit acceptable to FDA or any State regulatory authority.

Attachment 3 contains the information that is responsive to this question.

4. For products imported into the United States for handling or processing by any facility operated by your firm, please list the instances when internal or outside laboratory testing was positive for the presence of either a chemical or microbiological contaminant in excess of FDA or State regulatory limits.

Other than the information provided in our domestic response on June 5th or in response to question 3, above, we have no additional information to report in response to this question.

5. For each of the above items, please specify whether FDA was notified, and if not, why not.

The information in the attachments to this response and our earlier responses identifies whether we notified FDA.

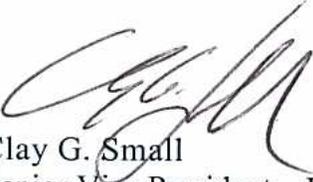
6. Please supply a list of all instances where FDA or any State regulatory authority was denied entrance to any facility, foreign or domestic, or denied access to any records regarding microbiological or chemical testing

performed on products processed at the facility. This request encompasses denials of initial requests for entry or any such testing record regarding of whether the plant or its records were to be made available for inspection at a later date.

Attachment 6 contains the information that is responsive this question.

We do not view this submission as waiving any rights, privileges, or immunities. If we can answer any questions regarding the information provided in this letter, please let us know.

Respectfully submitted,



Clay G. Small
Senior Vice President – Legal Affairs

cc: The Honorable Joe Barton, Ranking Member
House Committee on Energy and Commerce

The Honorable John Shimkus, Ranking Member
Subcommittee on Oversight and Investigations