

DATE: April 20, 2005

NOTE TO: FDA Division of Dockets Management (HFA-305) 0570 5 APR 21 A8:30

DOCKET NO.: 2004N-0456

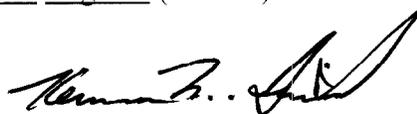
SUBJECT: Food Labeling: Serving Sizes of Products That Can Reasonable Be Consumed At One Eating Occasion; Updating of Reference Amounts Customarily Consumed; Approaches for Recommending Smaller Portion Sizes

PUB DATE: April 4, 2005

The September 30, 1993, Executive Order 12866--Regulatory Planning and Review sets forth the Administration's principles and requirements for the Federal regulatory process. Under section 6(a)(3)(E) of the Executive Order, for "significant regulatory actions," Federal agencies must make certain information available to the public after publication of the regulatory action in the Federal Register.

Pursuant to the Executive Order, the Food and Drug Administration (FDA) has attached in this docket, for the subject significant regulatory action, the following information:

- 1) A copy of the draft regulatory action as submitted to the Office of Management and Budget's (OMB) Office of Information and Regulatory Affairs (OIRA) for review, including any materials or assessments, required by the Executive Order, that accompanied the draft (TAB A);
- 2) The substantive changes between the draft submitted to OIRA for review and the regulatory action subsequently announced, including those changes that were made at the suggestion or recommendation of OIRA, and any other agency or governmental component to which this draft was submitted by OIRA for review, or FDA, if any (see mark-ups); and
- 3) A copy of the final regulatory action as sent to the Office of the Federal Register for publication or as published in the Federal Register (TAB C).



Regulations Policy and
and Management Staff (HF-26)

Attachments

044.0456

C36