1998. Since the last approval, the Demonstration Partnership Program no longer exists. Therefore, this request covers 6 programs, rather than the 7 programs previously covered.

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Estimated number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per respondent</th>
<th>Total burden hours</th>
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<td>CFN Announcement</td>
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<td>1</td>
<td>10</td>
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<td>LIHEAP Announcement</td>
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<td>Community Economic Dev. Announcement</td>
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<td>JOLI Announcement</td>
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<td>CSBG T&amp;A Announcement</td>
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<td>Family Violence Announcement</td>
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<td>1</td>
<td>40</td>
<td>4000</td>
</tr>
</tbody>
</table>

Estimated Total Annual Burden: 20,340.

Additional Information:
Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, 370 L’Enfant Promenade, S.W., Washington, D.C. 20447; Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the Federal Register. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503; Attn: Ms. Wendy Taylor.


Bob Sargis,
Acting Reports Clearance Officer.
[FR Doc. 98–23299 Filed 8–28–98; 8:45 am]
BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Notice of Food and Drug Administration

[Docket No. 98F–0714]

Asahi Denka Kogyo K.K.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Asahi Denka Kogyo K.K. has filed a petition proposing that the food additive regulations be amended to provide for the expanded safe use of 2, 2′-methylenebis (4,6-di-tert-butylphenyl) 2-ethylhexyl phosphite as an antioxidant and/or stabilizer in linear low density polyethylene articles intended for contact with food.

The agency has determined under 21 CFR 25.32(i) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.


Laure M. Tarantino,
Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.
[FR Doc. 98–23269 Filed 8–28–98; 8:45 am]
BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Notice of Food and Drug Administration

[Docket No. 98F–0716]

Dainippon Ink and Chemicals, Inc.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Dainippon Ink and Chemicals, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the expanded safe use of a polyester-polyurethane resin-acid dianhydride adhesive in retortable pouches for use in contact with fatty food.

FOR FURTHER INFORMATION CONTACT: Mark A. Hepp, Center for Food Safety