TABLE 2—ESTIMATED ANNUAL REPORTING BURDEN 1—Continued

<table>
<thead>
<tr>
<th>Activity</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Total annual responses</th>
<th>Average burden per response (in hours)</th>
<th>Total hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of completes</td>
<td>240</td>
<td>1</td>
<td>240</td>
<td>0.42 (25 minutes)</td>
<td>101</td>
</tr>
</tbody>
</table>

Main Study

- Number to complete the screener (assumes 50% eligibility): 1,785
- Number of completes: 1,785
- Average burden per response: 0.08 (5 minutes)
- Total hours: 143

- Number of completes: 1,272
- Average burden per response: 0.42 (25 minutes)
- Total hours: 534

Total hours: 805

1 There are no capital costs or operating and maintenance costs associated with this collection of information.

References

1. Hall R.H. and P. Hanna “The Impact of Information.” Register this document publishes in the Federal Register, with web site addresses, as of the date available electronically at www.regulations.gov.

2. Baur C and C. Prue “The CDC Clear Communication Index is a Tool to Prepare and Review Health Communication.”

3. Gaeta J. and C. Prue “The CDC Clear Communication Index is a Tool to Prepare and Review Health Information.”

4. Wogalter, M.S. and W.J. Vigilante. “Effects of Label Format on Knowledge Acquisition and Perceived Readability by Younger and Older Adults.”

5. Smither, J.A.A. and C.C. Braun “Readability of Prescription Drug Labels by Older and Younger Adults.”


14. Wogalter, M.S. and W.J. Vigilante. “Effects of Label Format on Knowledge Acquisition and Perceived Readability by Younger and Older Adults.”

15. Smither, J.A.A. and C.C. Braun “Readability of Prescription Drug Labels by Older and Younger Adults.”


SUMMARY: The Food and Drug Administration is correcting a notice entitled “Novus International, Inc.: Filing of Food Additive Petition (Animal Use)” that appeared in the Federal Register of November 8, 2016 (81 FR 78528). The document announced that Novus International, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of poly(2-vinylpyridine-co-styrene) as a nutrient protectant for methionine hydroxy analog in animal food for beef cattle, dairy cattle, and replacement dairy heifers. Additionally, the petition proposes that the food additive regulations be amended to provide for the safe use of ethyl cellulose as a binder for methionine hydroxy analog to be incorporated into animal food. The document was published with the incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Lisa Granger, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 3330, Silver Spring, MD 20993–0002, 301–796–9115, lisa.granger@fda.hhs.gov.


Dated: November 22, 2016.

Leslie Kux, Associate Commissioner for Policy.