the preliminary results are published. However, if it is not practicable to complete the review within these time periods, section 751(a)(3)(A) of the Act allows the Department to extend the time limit for the preliminary determination to a maximum of 365 days after the last day of the anniversary month.

Extension of Time Limit for Preliminary Results of Review

We determine that it is not practicable to complete the preliminary results of this administrative review within the original time limit because the Department requires additional time to analyze questionnaire responses and to evaluate surrogate value submissions.

Therefore, the Department is extending the time limit for completion of the preliminary results of the administrative review by 30 days. The preliminary results will now be due no later than April 29, 2012. As that day falls on a Sunday, the preliminary results are due no later than April 30, 2012.¹ The final results continue to be due 120 days after the publication of the preliminary results.

We are issuing and publishing this notice in accordance with sections 751(a)(3)(A) and 777(i) of the Act.


Christian Marsh,
Deputy Assistant Secretary for Antidumping and Countervailing Duty Operations.

¹ See Notice of Clarification: Application of “Next Business Day” Rule for Administrative Determination Deadlines as Amended, 70 FR 24533 (May 10, 2005).

NMFS published proposed program regulations on May 23, 2011 (76 FR 29707), and final program regulations on October 6, 2011 (76 FR 61986), to implement the reduction program. NMFS published the list of eligible voters in on March 1, 2012 (77 FR 12568). Interested persons should review these for further program details. The final regulations require NMFS to publish this notice before conducting a referendum to determine the industry’s willingness to repay a fishing capacity reduction loan to purchase the permits identified in the reduction submitted by the Southeast Revitalization Association and approved by NMFS.

As of February 24, 2012, there are 379 permits in the fishery designated as S01A by CFEC. These permanent permit holders are eligible to vote in the referendum. NMFS has updated the list and will mail referendum ballots to each. Mailed ballots will be accompanied by NMFS’ detailed voting guidance.

The referendum voting period will start March 30, 2012 and end on April 30, 2012. Any votes not received by NMFS by 5 p.m. on April 30, 2012, will not be counted.

Dated: March 26, 2012.

Lindsay Fullenkamp,
Acting Director, Office of Management and Budget, National Marine Fisheries Service.

ADDRESSES: Send comments about this notice to Paul Marx, Chief, Financial Services Division, NMFS, Attn: SE Alaska Purse Seine Salmon Buyback, 1315 East-West Highway, Silver Spring, MD 20910 (see FOR FURTHER INFORMATION CONTACT).

FOR FURTHER INFORMATION CONTACT:
Michael A. Sturtevant at (301) 427–8799, fax (301) 713–1306, or michael.a.sturtevant@noaa.gov.

SUPPLEMENTARY INFORMATION:
I. Background

The Southeast Alaska purse seine salmon fishery is a commercial fishery in Alaska state waters and adjacent Federal waters. It encompasses the commercial taking of salmon with purse seine gear, and participation is limited to fishermen designated by the Alaska Commercial Fisheries Entry Commission (CFEC).

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Lindsay Fullenkamp,
Acting Director, Office of Management and Budget, National Marine Fisheries Service.
CONSUMER PRODUCT SAFETY COMMISSION

Notice of Teleconference of the Chronic Hazard Advisory Panel on Phthalates and Phthalate Substitutes

AGENCY: Consumer Product Safety Commission.

ACTION: Notice of meeting.

SUMMARY: The Consumer Product Safety Commission (“CPSC” or “Commission”) is announcing the seventh meeting of the Chronic Hazard Advisory Panel (CHAP) on phthalates and phthalate substitutes. The Commission appointed this CHAP on April 14, 2010, to study the effects on children’s health of all phthalates and phthalate alternatives, as used in children’s toys and child care articles, pursuant to section 108 of the Consumer Product Safety Improvement Act of 2008 (CPSIA) (Pub. L. 110–314). The CHAP will discuss its progress toward completing its analysis of potential risks from phthalates and phthalate substitutes.

DATES: The teleconference will take place from 11 a.m. to 1 p.m. EDT (15:00 to 17:00 GMT) on Tuesday, April 10, 2012. Interested members of the public may listen to the CHAP’s discussion. Members of the public will not have the opportunity to ask questions, comment, or otherwise participate in the teleconference. Interested parties should contact the CPSC project manager, Michael Babich, by email (mmbabich@cpsc.gov), for call-in instructions no later than Friday, April 6, 2012.

FOR FURTHER INFORMATION CONTACT: Michael Babich, Directorate for Health Sciences, Consumer Product Safety Commission, Bethesda, MD 20814; telephone (301) 504–7253; email: mmbabich@cpsc.gov.

SUPPLEMENTARY INFORMATION: Section 108 of the CPSIA permanently prohibits the sale of any “children’s toy or child care article” containing more than 0.1 percent of each of three specified phthalates: Di-(2-ethylhexyl) phthalate (DEHP), dibutyl phthalate (DBP), and benzyl butyl phthalate (BBP). Section 108 of the CPSIA also prohibits, on an interim basis, the sale of any “children’s toy that can be placed in a child’s mouth” or “child care article” containing more than 0.1 percent of each of three additional phthalates: Dibis(2-ethylhexyl) phthalate (DINP), diisodecyl phthalate (DIDP), and di-n-octyl phthalate (DnOP).

Moreover, section 108 of the CPSIA requires the Commission to convene a CHAP “to study the effects on children’s health of all phthalates and phthalate alternatives as used in children’s toys and child care articles.” The CPSIA requires the CHAP to complete an examination of the full range of phthalates that are used in products for children and:

• Examine all of the potential health effects (including endocrine-disrupting effects) of the full range of phthalates;
• Consider the potential health effects of each of these phthalates, both in isolation, and in combination with other phthalates;
• Examine the likely levels of children’s, pregnant women’s, and others’ exposure to phthalates, based upon a reasonable estimation of normal and foreseeable use and abuse of such products;
• Consider the cumulative effect of total exposure to phthalates, both from children’s products and from other sources, such as personal care products;
• Review all relevant data, including the most recent, best-available, peer-reviewed, scientific studies of these phthalates and phthalate alternatives that employ objective data-collection practices or employ other objective methods;
• Consider the health effects of phthalates not only from ingestion, but also as a result of dermal, hand-to-mouth, or other exposure;
• Consider the level at which there is a reasonable certainty of no harm to children, pregnant women, or other susceptible individuals and their offspring, considering the best available science, and using sufficient safety factors to account for uncertainties regarding exposure and susceptibility of children, pregnant women, and other potentially susceptible individuals; and
• Consider possible similar health effects of phthalate alternatives used in children’s toys and child care articles.

The CPSIA contemplates completion of the CHAP’s examination within 18 months of the panel’s appointment on April 14, 2010. The CHAP has an additional 6 months to complete its final report to the Commission. The CHAP must review prior work on phthalates by the Commission, but it is not to be considered determinative because the CHAP’s examination must be conducted de novo.

The CHAP must make recommendations to the Commission regarding any phthalates (or combinations of phthalates), in addition to those identified in section 108 of the CPSIA, or phthalate alternatives that the panel determines should be prohibited from use in children’s toys or child care articles, or otherwise restricted. The CHAP members were selected by the Commission from scientists nominated by the National Academy of Sciences.

The CHAP met previously in April, July, and December 2010, March, July, and November 2011, and in February 2012, at the CPSC’s offices in Bethesda, MD, and by teleconference in November 2010, September 2011, December 2011, and February 2012. The CHAP heard testimony from interested parties at the July 2010, and November 2011, meetings. There will not be any opportunity for public comment during the April 2012 teleconference.


Alberta E. Mills,
Acting Secretary, Consumer Product Safety Commission.