

along with increasing visitor education and appreciation of natural and cultural resources of the park unit. This alternative will provide a comprehensive Riverways-wide approach to resource and visitor use management. Specific management zones detailing acceptable resource conditions, visitor experience and use levels, and appropriate activities and development will be applied to Riverways' lands consistent with this concept.

A mix of private and guided traditional recreational activities like boating, floating, and horseback riding will occur under this alternative. Additional trails and a small learning center at a rehabilitated Powder Mill will be developed to better orient and inform visitors. Natural resources will be restored to more natural conditions, while maintaining greater opportunities for visitor access. Most of the Big Spring Wilderness Study Area will be recommended for wilderness designation.

The selected action and three other alternatives were analyzed in the draft and final GMP/EIS. The full range of foreseeable environmental consequences was assessed. Among the alternatives the NPS considered, the selected action best achieves a high standard of natural and cultural resource protection with improved opportunities for visitors in the park. The NPS selected alternative B as its preferred alternative following an evaluation of the effectiveness of each alternative in meeting the stated objectives of the general management plan, and the environmental benefits and adverse impacts for each alternative. This alternative provides the best combination of strategies to protect the park unit's unique natural and cultural resources and visitor experience, while improving the park unit's operational effectiveness and sustainability. It also provides other advantages to the Riverways, regional communities, partners, and stakeholders.

In addition, selection of the preferred alternative, as reflected by the analysis contained in the final GMP/EIS, will not result in the impairment of park resources and will allow the National Park Service to conserve National Riverways' resources and provide for their enjoyment by visitors.

Dated: January 22, 2015.

**Patricia S. Trap,**

*Acting Regional Director, Midwest Region.*

**Editor's note:** This document was received for publication by the Office of the **Federal Register** on June 18, 2015.

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## DEPARTMENT OF THE INTERIOR

### National Park Service

**[NPS-MWR-CUVA-17695; PPMWROW2/PMP00UP05.YP0000]**

#### **Notice of Availability of a Record of Decision for the White-Tailed Deer Management Plan, Environmental Impact Statement, Cuyahoga Valley National Park, Ohio**

**AGENCY:** National Park Service, Interior.

**ACTION:** Notice of Availability.

**SUMMARY:** The National Park Service (NPS) announces the availability of the Record of Decision (ROD) for the White-Tailed Deer Management Plan/Environmental Impact Statement (plan/EIS), Cuyahoga Valley National Park (Park), Ohio.

**ADDRESSES:** Copies of the ROD are available by request by writing to the Chief of the Resource Management Division, Lisa Petit, Cuyahoga Valley National Park, 15610 Vaughn Road, Brecksville, Ohio 44141, telephone (440) 546-5970. The document is also available on the internet at the NPS Planning, Environment, and Public Comment Web site at: <http://www.parkplanning.nps.gov/>.

**FOR FURTHER INFORMATION CONTACT:** Chief of the Resource Management Division, Lisa Petit at the address or telephone number above.

**SUPPLEMENTARY INFORMATION:** We have issued a ROD for the final plan/EIS; on February 13, 2015, the Acting Regional Director for the Midwest Region approved the ROD. The plan/EIS analyzed four alternatives, including a no-action alternative. The full range of foreseeable environmental consequences was assessed, and appropriate mitigating measures were identified.

The NPS decision is to implement alternative D (hereinafter referred to as the "selected action"), which was identified as the NPS preferred alternative in the final plan/EIS. The selected action will utilize an adaptive management strategy. Under the selected action, the NPS will continue current park deer management actions including: Research, monitoring, and

data management; protection of restoration plantings; education and coordination; and enforcement of the existing wildlife feeding ban. In addition, the NPS will incorporate a combination of lethal and nonlethal actions to address high deer density. Lethal actions (including sharpshooting, with very limited capture/euthanasia if necessary) will be taken initially to reduce deer densities quickly. It is anticipated that in years one through four, 335 deer will be removed by sharpshooting (in addition to small numbers of deer removed by capture and euthanasia) to meet the deer density goal. Population maintenance will follow the initial reduction, and could be conducted by nonsurgical reproductive control methods, if an acceptable agent is available, or by sharpshooting. Both maintenance methods are included in the selected action to maintain maximum flexibility for future management. All actions will be carried out by NPS personnel or authorized agents.

The Record of Decision includes a summary of the purpose and need for action, identifies the selected action and all alternatives considered by the NPS, a listing of measures to minimize environmental harm, details about the monitoring program that will inform implementation of the selected action, the basis for the decision, and a description of the environmentally preferable alternative.

Dated: February 13, 2015.

**Patricia S. Trap,**

*Acting Regional Director, Midwest Region.*

**Editor's Note:** This document was received for publication by the Office of Federal Register on June 18, 2015.

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## INTERNATIONAL TRADE COMMISSION

### **Notice of Receipt of Complaint; Solicitation of Comments Relating to the Public Interest**

**AGENCY:** U.S. International Trade Commission.

**ACTION:** Notice.

**SUMMARY:** Notice is hereby given that the U.S. International Trade Commission has received a complaint entitled *Certain Resealable Packages with Slider Devices, DN 3072*; the Commission is soliciting comments on any public interest issues raised by the complaint or complainant's filing under section 210.8(b) of the Commission's

Rules of Practice and Procedure (19 CFR 210.8(b)).

**FOR FURTHER INFORMATION CONTACT:** Lisa R. Barton, Secretary to the Commission, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436, telephone (202) 205-2000. The public version of the complaint can be accessed on the Commission's Electronic Document Information System (EDIS) at EDIS,<sup>1</sup> and will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436, telephone (202) 205-2000.

General information concerning the Commission may also be obtained by accessing its Internet server at United States International Trade Commission (USITC) at USITC.<sup>2</sup> The public record for this investigation may be viewed on the Commission's Electronic Document Information System (EDIS) at EDIS.<sup>3</sup> Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205-1810.

**SUPPLEMENTARY INFORMATION:** The Commission has received a complaint and a submission pursuant to section 210.8(b) of the Commission's Rules of Practice and Procedure filed on behalf of Reynolds Presto Products Inc. on June 17, 2015. The complaint alleges violations of section 337 of the Tariff Act of 1930 (19 U.S.C. 1337) in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain resealable packages with slider devices. The complaint names as respondents Interplast Group, Ltd. of Livingston, NJ and Minigrip, LLC of Alpharetta, GA. The complainant requests that the Commission issue a permanent general exclusion order, cease and desist orders, and a bond upon respondents' alleged infringing articles during the 60-day Presidential review period pursuant to 19 U.S.C. § 1337(j).

Proposed respondents, other interested parties, and members of the public are invited to file comments, not to exceed five (5) pages in length, inclusive of attachments, on any public interest issues raised by the complaint or section 210.8(b) filing. Comments should address whether issuance of the

relief specifically requested by the complainant in this investigation would affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

- (i) Explain how the articles potentially subject to the requested remedial orders are used in the United States;
- (ii) identify any public health, safety, or welfare concerns in the United States relating to the requested remedial orders;
- (iii) identify like or directly competitive articles that complainant, its licensees, or third parties make in the United States which could replace the subject articles if they were to be excluded;
- (iv) indicate whether complainant, complainant's licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to the requested exclusion order and/or a cease and desist order within a commercially reasonable time; and
- (v) explain how the requested remedial orders would impact United States consumers.

Written submissions must be filed no later than by close of business, eight calendar days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above and submit 8 true paper copies to the Office of the Secretary by noon the next day pursuant to section 210.4(f) of the Commission's Rules of Practice and Procedure (19 CFR 210.4(f)). Submissions should refer to the docket number ("Docket No. 3072") in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, Electronic Filing Procedures.<sup>4</sup>) Persons with questions regarding filing should contact the Secretary (202-205-2000).

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be

directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.<sup>5</sup>

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of sections 201.10 and 210.8(c) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.8(c)).

Issued: June 18, 2015.

By order of the Commission.

**Lisa R. Barton,**

*Secretary to the Commission.*

[FR Doc. 2015-15368 Filed 6-22-15; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-392]

#### Importer of Controlled Substances Application: Midas Pharmaceuticals, Inc.

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic class, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before July 23, 2015. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 23, 2015.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODXL, 8701 Morrisette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301,

<sup>1</sup> Electronic Document Information System (EDIS): <http://edis.usitc.gov>.

<sup>2</sup> United States International Trade Commission (USITC): <http://edis.usitc.gov>.

<sup>3</sup> Electronic Document Information System (EDIS): <http://edis.usitc.gov>.

<sup>4</sup> Handbook for Electronic Filing Procedures: [http://www.usitc.gov/secretary/fed\\_reg\\_notices/rules/handbook\\_on\\_electronic\\_filing.pdf](http://www.usitc.gov/secretary/fed_reg_notices/rules/handbook_on_electronic_filing.pdf).

<sup>5</sup> Electronic Document Information System (EDIS): <http://edis.usitc.gov>.