ingredient per 60,120 pounds of potato had residues of 0.82 ppm, 1.18 ppm, and 1.27 ppm at 0 DAT; 0.22 ppm, 0.28 ppm, and 0.41 ppm at 30 DAT; 0.10 ppm, 0.11 ppm, and 0.04 ppm at 90 DAT; and 0.03 ppm, 0.03 ppm, and 0.05

ppm at 180 DAT

Potatoes treated 3 times at day 0, 60, and 120, as described above, had 2,6-DIPN residues of 0.83 ppm, 1.28 ppm, and 1.39 ppm at 0 DAT; 0.25 ppm, 0.30 ppm, 0.37 ppm at 30 DAT; 0.80 ppm, 1.07 ppm, and 2.43 ppm at 60 DAT; 0.28 ppm, 0.42 ppm, and 0.62 ppm at 90 DAT; 1.16 ppm, 1.79 ppm, and 1.86 ppm at 120 DAT; and 0.13 ppm, 0.17 ppm, and 0.24 ppm at 180 DAT.

Magnitude of residue at the time of harvest and method used to determine the residue. A statement of why an analytical method for detecting and measuring the levels of the pesticide residue are not needed. Since the petitioner has requested a tolerance exemption, an analytical method to detect residues is not required.

D. Toxicology Profile

1. Acute toxicity. Technical 2.6-DIPN exhibits low acute toxicity. It is a toxicity category IV biopesticide. The rat oral LD₅₀ is greater than 5,000 milligram/kilogram (mg/kg), the rabbit dermal LD₅₀ is greater than 5,000 mg/kg, and the rat inhalation LC50 is greater than 2.60 milligram/Liter (mg/L) at the maximum attainable condition. In addition, 2,6-DIPN is not a skin sensitizer in guinea pigs, shows no dermal irritation at 72 hours in rabbits, and shows minimal ocular irritation in rabbits. The end use formulation is the same as the technical formulation; it contains no intentionally added inert ingredients.

Genotoxicity. Short-term assays for genotoxicity consisting of a bacterial reverse mutation assay (Ames test), an in vivo/in vitro unscheduled DNA synthesis in rat primary hepatocyte cultures at 2 time points, and an in vivo mouse micronucleus assay have been conducted for 2,6-DIPN. These studies show a lack of genotoxicity for 2,6-

DIPN.

3. Other tests. No additional mammalian toxicology testing has been conducted. Platte requested a waiver from the requirement to submit further mammalian toxicology studies on the basis of the favorable toxicological profile for 2,6-DIPN, the low residues observed in treated potatoes, the specific plant growth regulator mode of action, and the confined nature of the proposed use. No data were found in the literature that would indicate 2,6-DIPN has any adverse effect on mammals. No incidents of hypersensitivity or any

other adverse effects have been observed
G. Endocrine Effects in individuals handling the material over the past 6 years.

E. Aggregate Exposure

In examining aggregate exposure, section 408 of the FFDCA directs EPA to consider available information about exposures from the pesticide residue in food and all other non-occupational exposures, including drinking water from groundwater or surface water and exposure through pesticide use in gardens, lawns, or buildings (residential and other indoor uses).

- 1. Dietary exposure from food and drinking water. Any dietary exposure resulting from applications made under an experimental use permit (EUP) would be through potato consumption and animal products in which animals are fed potato feed stocks. Residues in treated potatoes have been shown to be low. Residues would be expected to continue to decline after potatoes are removed from storage and before consumption. Cooking and/or processing would be expected to further lower the residue level in consumed potatoes or potato products. Since 2,6-DIPN would only be used in commercial storage warehouses, there is little if any potential for drinking water exposure. There are no other established U.S. tolerances or exemptions from tolerances for 2.6-DIPN food or feed crops in the United States. The Agency has classified 2,6-DIPN as a biochemical pesticide.
- 2. Non-dietary exposure. The EUP would only cover use for direct application to potatoes when stored in commercial warehouses. There are currently no other registered uses of 2,6-DIPN. Non-dietary exposure to 2,6-DIPN via lawn care, topical treatments, etc., will not occur. Thus, the potential for non-occupational exposure to the general population is virtually nonexistent.

F. Cumulative Exposure

EPA also is required to consider the potential for cumulative effects of 2,6-DIPN and other substances that have a common mechanism of toxicity. Consideration of a common mode of toxicity is not appropriate, given that there is no indication of mammalian toxicity of 2,6-DIPN and no information that indicates toxic effects, if any, would be cumulative with any other compounds. Since, 2,6-DIPN does not exhibit a toxic mode of action in the target plant, it is appropriate to consider only the potential risks of 2,6-DIPN in this exposure assessment.

Platte has no information to suggest that 2,6-DIPN will adversely affect the immune or endocrine systems. The Agency is not requiring information on endocrine effects of this biochemical pesticide at this time.

H. Safety Determinations

1. U.S. population in general and infants and children. Since there are no anticipated residues in drinking water or from other non-occupational sources, and no reliable information exists on cumulative effects due to a common mechanism of toxicity, the aggregate exposure to 2,6-DIPN is adequately represented by the dietary route. The lack of toxicity of 2,6-DIPN has been demonstrated by the results of acute toxicity testing in mammals in which 2,6-DIPN caused no adverse effects when dosed orally, dermally, and via inhalation at the limit dose for each study. Anticipated residues in consumed potatoes are low. Moreover, 2,6-DIPN exhibits close similarity to other plant-based, naturally occurring methyl and isopropyl naphthalenes. Thus, the dietary exposure to 2,6-DIPN should pose negligible risks to human health. Based on the lack of toxicity and low exposure, there is a reasonable certainty that no harm to infants, children, or adults will result from aggregate exposure to 2,6-DIPN residues. Exempting 2,6-DIPN from the requirement of a tolerance should pose no significant risk to humans or the environment.

I. Analytical Method

An analytical method for residues is not applicable, as this proposes an exemption from the requirement of a tolerance.

J. Existing Tolerances

No codex maximum residue levels are established for residues of 2,6-DIPN in or on any food or feed crop.

[FR Doc. 98-31248 Filed 11-24-98; 8:45 am] BILLING CODE 6560-50-F

ENVIRONMENTAL PROTECTION AGENCY

[OPP-50848; FRL-6043-4]

Experimental Use Permit; Notice of Receipt of Application

AGENCY: Environmental Protection

Agency (EPA). **ACTION:** Notice.

SUMMARY: This notice announces receipt of an application [34704-EUP-RG] from

Platte Chemical Company requesting an experimental use permit (EUP) for the biochemical pesticide 2,6—diisopropylnapthalene (2,6—DIPN). The Agency has determined that the application may be of regional and national significance. Therefore, in accordance with 40 CFR 172.11(a), the Agency is soliciting comments on this application.

DATES: Written comments must be received on or before December 28, 1998.

ADDRESSES: By mail, submit written comments to: Public Information and Records Integrity Branch, Information Resources and Services Division (7502C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person, deliver comments to: Rm. 119, CM #2, 1921 Jefferson Davis Hwy., Arlington, VA.

Comments and data may also be submitted electronically to: opp-docket@epa.gov. Follow the instructions under Unit II. of this document. No Confidential Business Information (CBI) should be submitted through e-mail.

Information submitted as a comment concerning this document may be claimed confidential by marking any part or all of that information as CBI. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2. A copy of the comment that does not contain CBI must be submitted for inclusion in the public record. Information not marked confidential will be included in the public docket by EPA without prior notice. The public docket is available for public inspection in Rm. 119 at the Virginia address given above, from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays.

FOR FURTHER INFORMATION CONTACT: By mail: Rita Kumar, PM 90, Biopesticides and Pollution Prevention Division (7511C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. Office location, telephone number, and e-mail address: Rm. 902W5, CM #2, 1921 Jefferson Davis Hwy., Arlington, VA, Telephone: 703–308–8291, e-mail: kumar.rita@epa.gov.

SUPPLEMENTARY INFORMATION:

I. Background

Following the review of the Platte Chemical Company's application and any comments received in response to this notice, EPA will decide whether to issue or deny the EUP request for this EUP program, and if issued, the conditions under which it is to be conducted. Any issuance of an EUP will be announced in the **Federal Register**.

The proposed program would allow the use of 1,500 pounds of the plant growth regulator 2,6-DIPN on approximately 90 million pounds of potatoes in nine closed storage facilities (representing the harvest of approximately 3,160 acres). Platte's program would evaluate the control of potato spouting. The program would be authorized only in the States of Idaho, Maine, Minnesota, North Dakota, Oregon, Washington, and Wisconsin. This EUP is accompanied by a pesticide petition for an exemption from the requirement of a tolerance for residues of 2,6-DIPN when used to inhibit sprouting in potato held in storage. This pesticide petition is being issued elsewhere in this issue of the Federal Register.

II. Public Record and Electronic Submissions

The official record for this notice, as well as the public version, has been established for this notice under docket control number "OPP-50848" (including comments and data submitted electronically as described below). A public version of this record, including printed, paper versions of electronic comments, which does not include any information claimed as CBI, is available for inspection from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The official record is located at the Virginia address in "ADDRESSES" at the beginning of this document.

Electronic comments can be sent directly to EPA at: opp-docket@epa.gov

Electronic comments must be submitted as an ASCII file avoiding the use of special characters and any form of encryption. Comment and data will also be accepted on disks in Wordperfect 5.1/6.1 or ASCII file format. All comments and data in electronic form must be identified by the docket control number "OPP–50848." Electronic comments on this notice may be filed online at many Federal Depository Libraries.

Dated: November 4, 1998.

Janet L. Andersen,

Director, Biopesticides and Pollution Prevention Division, Office of Pesticide Programs.

[FR Doc. 98–31249 Filed 11–24–98; 8:45 am] BILLING CODE 6560–50–F

ENVIRONMENTAL PROTECTION AGENCY

[FRL-6193-2]

Notice of Proposed De Minimis Administrative Order on Consent Pursuant to Section 122(g) of the Comprehensive Environmental Response, Compensation and Liability Act (CERCLA), Osage Metals Superfund Site, Kansas City, Kansas, Docket No. VII–98–F–0023

AGENCY: Environmental Protection Agency.

ACTION: Notice of Proposed De Minimis Administrative Order on Consent, Osage Metals Superfund Site, Kansas City, Kansas.

SUMMARY: Notice is hereby given that a proposed administrative order on consent regarding the Osage Metals Superfund Site, was signed by the United States Environmental Protection Agency (EPA) on September 30, 1998 and approved by the United States Department of Justice (DOJ) on October 30, 1998.

DATES: EPA will receive comments on or before December 28, 1998 relating to the proposed agreement and covenant not to sue

ADDRESSES: Comments should be addressed to Audrey Asher, Senior Assistant Regional Counsel, United States Environmental Protection Agency, Region VII, 726 Minnesota Avenue, Kansas City KS 66101 and should refer to the Osage Metals Superfund Site Administrative Order on Consent, EPA Docket No. VII-98-F-0023.

The proposed agreement may be examined or obtained in person or by mail at the office of the United States Environmental Protection Agency, Region VII, 726 Minnesota Avenue, Kansas City, KS 66101 (913–551–7255).

SUPPLEMENTARY INFORMATION: The proposed agreement concerns the 1.7acre Osage Metals Superfund Site ("Site"), located at 120 Osage Avenue in Kansas City, Kansas. The Site was the location of metals salvage and reclamation facilities between 1948 and 1993. Samples taken at the Site in 1994 found polychlorinated biphenyls ("PCBs") in surface soils at levels as high as 334 mg/kg, and lead contamination in levels as high as 56,600 mg/kg. The EPA approved a removal action at the Site on February 13, 1995, and began cleanup in March of 1995. EPA completed its work in October 1995. No further response action is anticipated.