

explanations in support of the commenter's recommendations. Comments received after the time indicated under **DATES** or at locations other than the Casper Field Office will not necessarily be considered in the final rulemaking or included in the administrative record.

## 2. Public Hearing

Persons wishing to testify at the public hearing should contact the person listed under **FOR FURTHER INFORMATION CONTACT** by 4:00 p.m., m.d.t., on April 25, 1996. Any disabled individual who has need for a special accommodation to attend a public hearing should contact the individual listed under **FOR FURTHER INFORMATION CONTACT**. The location and time of the hearing will be arranged with those persons requesting the hearing. If no one requests an opportunity to testify at the public hearing, the hearing will not be held.

Filing of a written statement at the time of the hearing is requested as it will greatly assist the transcriber. Submission of written statements in advance of the hearing will allow OSM officials to prepare adequate responses and appropriate questions.

The public hearing will continue on the specified date until all persons scheduled to testify have been heard. Persons in the audience who have not been scheduled to testify, and who wish to do so, will be heard following those who have been scheduled. The hearing will end after all persons scheduled to testify and persons present in the audience who wish to testify have been heard.

## 3. Public Meeting

If only one person requests an opportunity to testify at a hearing, a public meeting, rather than a public hearing may be held. Persons wishing to meet with OSM representatives to discuss the proposed amendment may request a meeting by contacting the person listed under **FOR FURTHER INFORMATION CONTACT**. All such meetings will be open to the public and, if possible, notices of meetings will be posted at the locations listed under **ADDRESSES**. A written summary of each meeting will be made a part of the administrative record.

## IV. Procedural Determinations

### 1. Executive Order 12866

This rule is exempted from review by the Office of Management and Budget (OMB) under Executive Order 12866 (Regulatory Planning and Review).

### 2. Executive Order 12778

The Department of the Interior has conducted the reviews required by section 2 of Executive Order 12778 (Civil Justice Reform) and has determined that this rule meets the applicable standards of subsections (a) and (b) of that section. However, these standards are not applicable to the actual language of State regulatory programs and program amendments since each such program is drafted and promulgated by a specific State, not by OSM. Under sections 503 and 505 of SMCRA (30 U.S.C. 1253 and 1255) and the Federal regulations at 30 CFR 730.11, 732.15, and 732.17(h)(10), decisions on proposed State regulatory programs and program amendments submitted by the States must be based solely on a determination of whether the submittal is consistent with SMCRA and its implementing Federal regulations and whether the other requirements of 30 CFR Parts 730, 731, and 732 have been met.

### 3. National Environmental Policy Act

No environmental impact statement is required for this rule since section 702(d) of SMCRA (30 U.S.C. 1292(d)) provides that agency decisions on proposed State regulatory program provisions do not constitute major Federal actions within the meaning of section 102(2)(C) of the National Environmental Policy Act of 1969 (42 U.S.C. 4332(2)(C)).

### 4. Paperwork Reduction Act

This rule does not contain information collection requirements that require approval by OMB under the Paperwork Reduction Act (44 U.S.C. 3507 *et seq.*).

### 5. Regulatory Flexibility Act

The Department of the Interior has determined that this rule will not have a significant economic impact on a substantial number of small entities under the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*). The State submittal that is the subject of this rule is based upon counterpart Federal regulations for which an economic analysis was prepared and certification made that such regulations would not have a significant economic effect upon a substantial number of small entities. Accordingly, this rule will ensure that existing requirements previously promulgated by OSM will be implemented by the State. In making the determination as to whether this rule would have a significant economic impact, the Department relied upon the data and assumptions for the counterpart Federal regulations.

## List of Subjects in 30 CFR Part 926

Intergovernmental relations, Surface mining, Underground mining.

Dated: April 3, 1996.

Russell F. Price,

*Acting Regional Director, Western Regional Coordinating Center.*

[FR Doc. 96-8921 Filed 4-9-96; 8:45 am]

BILLING CODE 4310-05-M

## ENVIRONMENTAL PROTECTION AGENCY

### 40 CFR Part 180

[OPP-300420; FRL-5361-2]

### Potassium Citrate; Tolerance Exemption

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Proposed rule.

**SUMMARY:** This document proposes that residues of potassium citrate (CAS Reg. No. 866-84-2) be exempted from the requirement of a tolerance when used as an inert ingredient (chelating agent and pH control) in pesticide formulations applied to growing crops, raw agricultural commodities after harvest, and animals. This proposed regulation was requested by Monsanto Company and Zeneca Ag Products pursuant to the Federal Food, Drug, and Cosmetic Act (FFDCA).

**DATES:** Comments, identified by the docket control number [OPP-300420], must be received on or before May 10, 1996.

**ADDRESSES:** By mail, submit written comments to: Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person deliver comments to: Rm. 1132, Crystal Mall CM #2, 1921 Jefferson Davis Highway, Arlington, VA.

Information submitted as a comment concerning this document may be claimed confidential by marking any part or all of that information as "Confidential Business Information" (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. 1132 at the address given above,

from 8 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays.

Comments and data may also be submitted electronically by sending electronic mail (e-mail) to: opp-docket@epamail.epa.gov. Electronic comments must be submitted as an ASCII file avoiding the use of special characters and any form of encryption. Comments and data will also be accepted on disks in WordPerfect in 5.1 file format or ASCII file format. All comments and data in electronic form must be identified by the docket number, [OPP-300420]. No CBI should be submitted through e-mail. Electronic comments on this proposed rule may be filed online at many Federal Depository Libraries. Additional information on electronic submissions can be found below in this document.

**FOR FURTHER INFORMATION CONTACT:** By mail: Amelia M. Acierto, Registration Support Branch, Registration Division (7505W), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. Office location, telephone number, and e-mail address: 2800 Crystal Drive, North Tower, Arlington, VA, (703)308-8375, e-mail:

acierto.amelia@epamail.epa.gov.

**SUPPLEMENTARY INFORMATION:** Monsanto Company, 700 14th Street, NW., Washington, DC 20005, has submitted pesticide petition (PP) 6E04607 and Zeneca Ag Products, 1800 Concord Pike, Wilmington, DE 19850-5458, has submitted pesticide petition (PP) 6E04637, to EPA requesting that the Administrator, pursuant to section 408(e) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 346a(e), propose to amend 40 CFR 180.1001(c) and (e) by establishing an exemption from the requirement of a tolerance for potassium citrate when used as an inert ingredient (chelating agent and pH control) in pesticide formulations applied to growing crops, raw agricultural commodities after harvest, and animals.

Inert ingredients are all ingredients that are not active ingredients as defined in 40 CFR 153.125 and include, but are not limited to, the following types of ingredients (except when they have a pesticidal efficacy of their own): solvents such as alcohols and hydrocarbons; surfactants such as polyoxyethylene polymers and fatty acids; carriers such as clay and diatomaceous earth; thickeners such as carrageenan and modified cellulose; wetting, spreading, and dispersing agents; propellants in aerosol dispensers; microencapsulating agents;

and emulsifiers. The term "inert" is not intended to imply nontoxicity; the ingredient may or may not be chemically active.

The data submitted in the petition and other relevant material have been evaluated. As part of the EPA policy statement on inert ingredients published in the Federal Register of April 22, 1987 (52 FR 13305), the Agency set forth a list of studies which would generally be used to evaluate the risks posed by the presence of an inert ingredient in a pesticide formulation. However, where it can be determined without that data that the inert ingredient will present minimal or no risk, the Agency generally does not require some or all of the listed studies to rule on the proposed tolerance or exemption from the requirement of a tolerance for an inert ingredient. The Agency has decided that no data, in addition to that described below, for potassium citrate will need to be submitted. The rationale for this decision is described below:

1. Potassium citrate is a potassium salt of citric acid which is a naturally occurring metabolite in plant and animal tissues.

2. Potassium citrate has been affirmed as generally recognized as safe (GRAS) by the U.S. Food and Drug Administration (FDA) as a direct human food ingredient when used as Multiple Purpose GRAS Food Substance (21 CFR 184.1625). The conditions of use specify only that potassium citrate is generally recognized as safe when used in accordance with good manufacturing practice.

3. Other citrates are already exempt from the requirements of a tolerance under 40 CFR 180.1001 when used as inert ingredient in pesticide formulations applied to growing plants or to raw agricultural commodities after harvest (citric acid and calcium citrate), to growing plants only (sodium citrate) or to animals (citric acid). Furthermore, other related citrates (triethyl citrate and dibasic ammonium citrate) have been considered by the Agency as inert ingredients of minimal concern (i.e., classified as List 4B inert ingredients).

Based on the extensive FDA review of potassium citrate, resulting in this chemical being affirmed as GRAS, its structure and physico-chemical properties, and the fact that other chemically related citrates have also been approved for food use applications, the Agency does not believe that a potential for significant hazard exists when potassium citrate is used in accordance with good agricultural practices. The Agency believes that this ingredient is useful and a tolerance is not necessary to

protect the public health. Therefore, EPA proposes that the exemption from the requirement of a tolerance be established as set forth below.

Any person who has registered or submitted an application for registration of a pesticide, under the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) as amended, which contains any of the ingredients listed herein, may request within 30 days after publication of this document in the Federal Register that this proposal be referred to an Advisory Committee in accordance with section 408(e) of FFDCA.

Interested persons are invited to submit written comments on the proposed regulation. Comments must bear a notation indicating the docket control number, [OPP-300420]. All written comments filed in response to this petition will be available in the Public Response and Program Resources Branch, at the address given above from 8 a.m. to 4:30 p.m. Monday through Friday, except legal holidays.

A record has been established for this rulemaking under docket number [OPP-300420] (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 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The public record is located in Room 1132 of the Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, Crystal MallCM #2, 1921 Jefferson Davis Highway, Arlington, VA.

Electronic comments can be sent directly to EPA at:

opp-docket@epamail.epa.gov

Electronic comments must be submitted as an ASCII file avoiding the use of special characters and any form of encryption.

The official record for this rulemaking, as well as the public version, as described above will be kept in paper form. Accordingly, EPA will transfer all comments received electronically into printed, paper form as they are received and will place the paper copies in the official rulemaking record which will also include all comments submitted directly in writing. The official rulemaking record is the paper record maintained at the Virginia address in "ADDRESSES" at the beginning of this document.

The Office of Management and Budget has exempted this proposed rule from

the requirements of section 3 of Executive Order 12866.

Pursuant to the requirements of the Regulatory Flexibility Act (Pub. L. 96-354, 94 Stat. 1164, 5 U.S.C. 601-612), the Administrator has determined that regulations establishing new tolerances or raising tolerance levels or establishing exemptions from tolerance requirements do not have a significant economic impact on a substantial number of small entities. A certification statement to this effect was published in the Federal Register of May 4, 1981 (46 FR 24950).

#### List of Subjects in 40 CFR Part 180

Environmental protection, Administrative practice and procedure, Agricultural commodities, Pesticides and pests, Reporting and recordkeeping requirements.

Dated: March 29, 1996.

Peter Caulkins,

*Acting Director, Registration Division, Office of Pesticide Programs.*

Therefore, it is proposed that 40 CFR part 180 be amended as follows:

#### PART 180—[AMENDED]

1. The authority citation for part 180 continues to read as follows:

Authority: 21 U.S.C. 346a and 371.

2. Section 180.1001 is amended by adding alphabetically to the tables in paragraphs (c) and (e) the inert ingredient "Potassium citrate (CAS Reg. No. 866-84-2)" to read as follows:

#### § 180.1001 Exemptions from the requirements of a tolerance.

\* \* \* \* \*

(c) \* \* \*

Inert Ingredients	Limits	Uses
Potassium citrate (CAS Reg. No. 866-84-2) .....	* * *	* * Chelating agent, pH control
* *	* *	* *

(e) \* \* \*

Inert Ingredients	Limits	Uses
Potassium citrate (CAS Reg. No. 866-84-2) .....	* * *	* * Chelating agent, pH control
* *	* *	* *

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#### 40 CFR Part 180

[OPP-300417; FRL-5353-5]

RIN 2070-AB18

#### 1,1,1,2-Tetrafluoroethane; Tolerance Exemption

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Proposed rule.

**SUMMARY:** This document proposes that residues of 1,1,1,2-tetrafluoroethane be exempted from the requirement of a tolerance when used as an inert ingredient (aerosol propellant) in insecticide aerosol formulations intended to be applied in food handling establishments. This proposed regulation was requested by Whitmire Research Laboratories, Inc.

**DATE:** Comments, identified by the document control number [OPP-300417], must be received on or before May 10, 1996.

**ADDRESS:** By mail, submit written comments to: Public Response and

Program Resources Branch, Field Operations Division (7506C) Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person deliver comments to: Rm. 1128, Crystal Mall, Building #2, 1921 Jefferson Davis Highway, Arlington, VA 22202. Information submitted as a comment concerning this document may be claimed confidential by marking any part or all of that information as "Confidential Business Information" (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 the EPA without prior notice. The public docket is available for public inspection in Room 1128 at the address given above, from 8 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays.

Comments and data may also be submitted electronically by sending electronic mail (e-mail) to: opp-docket@epamail.epa.gov. Electronic comments must be submitted as an

ASCII file avoiding the use of special characters and any form of encryption. Comments and data will also be accepted on disks in WordPerfect 5.1 file format or ASCII file format. All comments and data in electronic form must be identified by the docket number [OPP-300417]. No CBI should be submitted through e-mail. Electronic comments on this proposed rule may be filed online at many Federal Depository Libraries. Additional information on electronic submissions can be found below in this document.

**FOR FURTHER INFORMATION CONTACT:** By mail: Amelia M. Acierto Registration Support Branch, Registration Division (7505W), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. Office location and telephone number: 2800 Crystal Drive, North Tower, Arlington, VA 22202, (703) 308-8375, e-mail: acierto.amelia@epamail.epa.gov.

**SUPPLEMENTARY INFORMATION:** Whitmire Research Laboratories, Inc. 3568 Tree Court Industrial Boulevard, Saint Louis, MO 63122-6620 submitted pesticide petition (PP) number 5E4439 to EPA requesting that the Administrator, pursuant to Section 408(e) of the Federal Food, Drug, and Cosmetic Act,