

the range of issues raised and alternative concepts identified in the scoping period, the draft EIS likely will assess potential impacts of several alternative fire management plan strategies. Issues could include, but may not be limited to: hazards to firefighters, the public and property; effects on native plant and animal species; impacts to rare and endangered species; adverse impacts to cultural resources; risk of fire spread across the wildland-urban interface; increased use of mechanical and manual vegetation management techniques; air and water quality impacts; and effects on park visitors and neighbors.

**Supplementary Information and Public Scoping:** As described above, the NPS is undertaking a conservation planning and environmental impact analysis effort to identify issues and concerns which should be addressed in the future fire management program, to evaluate alternative concepts for fire management, and to obtain information regarding potential impacts and appropriate mitigation strategies which should be addressed in the EIS process.

As an early step in this undertaking, public scoping meetings will be conducted in the summer and early fall of 2003 in both San Mateo and Marin counties. The San Mateo County meeting will be held in conjunction with the regularly scheduled bi-monthly GGNRA public meeting on September 16, 2003. The meeting will be held at the Pacifica City Council Chambers, 2212 Beach Boulevard, Pacifica, California. An open-house to provide information and solicit input from the public will be conducted from 6–7 p.m. The EIS FMP will also be agendaized during the bi-monthly meeting (which begins at 7 p.m.). The Marin County meeting will be held September 24, 2003 from 7–9 p.m. at the U.S. Army Corps of Engineers San Francisco Bay Model Visitor's Center, 2100 Bridgeway, Sausalito, CA 94965. Information on these and all future public meetings, and status of the EIS process, will be regularly posted on the park's Web site at <http://www.nps.gov/goga/fire>; information will also be released through direct mailings and local and regional media. For those unable to attend either of these meetings, a scoping document will be available upon request. The main topics include: background information on the fire management program; a review of relevant policy and law affecting the fire management program; an assessment of current fire management needs; and the identification of issues and potential alternatives related to future fire management in the park.

All interested individuals, organizations, and agencies are encouraged to provide comments, suggestions, and relevant information. All written comments regarding the current and future fire management program must be postmarked or transmitted not later than 60 days following the date of publication of this notice in the **Federal Register**. Immediately upon determining this date it will be posted on the park's Web site. To request the scoping materials or to provide comments, please contact: Superintendent, Golden Gate National Recreation Area, Fort Mason, Building 201, San Francisco, CA 94123, Attn: Fire Management Plan. E-mail comments or requests for scoping materials and information on meeting locations may be transmitted electronically to [GOGA\\_Public\\_Affairs@nps.gov](mailto:GOGA_Public_Affairs@nps.gov) (the subject line should indicate "EIS Scoping for FMP").

Our practice is to make comments, including respondent names and home addresses, available for public review during regular business hours. If individuals submitting comments request that their name and/or address be withheld from public disclosure, it will be honored to the extent allowable by law. Such requests must be stated prominently at the beginning of the comments. There also may be circumstances wherein the NPS may withhold a respondent's identity as allowable by law. As always: NPS will make available to public inspection all submissions from organizations or businesses, and from individuals identifying themselves as representatives or officials of organizations or businesses; and, anonymous comments may not be considered.

**Decision:** Availability of the draft EIS and fire management plan for review and written comment will be announced in the **Federal Register**, as well as via direct mailing and announcements in local and regional media. At this time it's expected these documents will be available for public review in August 2004. Availability of the final EIS will be similarly announced (which at this time is expected to occur not sooner than January 2005). As a delegated EIS, the official responsible for the NEPA decision is the Regional Director, Pacific West Region, National Park Service. Subsequently, the official responsible for implementation would be the Superintendent, Golden Gate National Recreation Area.

Dated: July 25, 2003.

**Arthur E. Eck,**

*Deputy Regional Director, Pacific West Region.*

[FR Doc. 03–19962 Filed 8–5–03; 8:45 am]

BILLING CODE 4312–FN–P

## INTERNATIONAL TRADE COMMISSION

[USITC SE–03–026]

### Sunshine Act Meeting

**AGENCY HOLDING THE MEETING:** United States International Trade Commission.

**TIME AND DATE:** August 13, 2003, at 11 a.m.

**PLACE:** Room 101, 500 E Street SW., Washington, DC 20436, Telephone: (202) 205–2000.

**STATUS:** Open to the public.

#### MATTERS TO BE CONSIDERED:

1. *Agenda for future meetings:* None.
2. Minutes.
3. Ratification List.
4. Inv. No. 731–TA–1047

(Preliminary) (Ironing Tables and Certain Parts Thereof from China)—briefing and vote. (The Commission is currently scheduled to transmit its determination to the Secretary of Commerce on or before August 14, 2003; Commissioners' opinions are currently scheduled to be transmitted to the Secretary of Commerce on or before August 21, 2003.)

5. *Outstanding action jackets:* None.

In accordance with Commission policy, subject matter listed above, not disposed of at the scheduled meeting, may be carried over to the agenda of the following meeting.

By order of the Commission.

Issued: August 4, 2003.

**Marilyn R. Abbott,**

*Secretary to the Commission.*

[FR Doc. 03–20203 Filed 8–4–03; 2:27 pm]

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## DEPARTMENT OF JUSTICE

### Antitrust Division

#### United States v. Mountain Health Care, P.A. Civil Action No. 1:02CV288–T (W.D.N.C.); Response to Public Comments

Notice is hereby given pursuant to the Antitrust Procedures and Penalties Act, 15 U.S.C. 16(b)–(h), that Public Comments and the Response of the United States have been filed with the United States District Court for the Western District of North Carolina in

*United States v. Mountain Health Care, P.A.* Civil Action No. 1:02CV288-T (W.D.N.C., filed December 13, 2002). On December 13, 2002, the United States filed a Complaint alleging that defendant, Mountain Health Care, P.A. ("MCH") and its physician owners and members, restrained competition in the sale of physician services to managed health care purchasers, in violation of section 1 of the Sherman act, 15 U.S.C. 1. The proposed Final Judgment, filed at the same time as the Complaint, requires MHC to dissolve.

Public comment was invited within the statutory 60-day comment period. Such Comments, and the Responses thereto, are hereby published in the Federal Register and have been filed with the Court. Copies of the Complaint, Stipulation, proposed Final Judgment, Competitive Impact Statement, Public Comments and the Response of the United States are available for inspection in Room 4000 of the Antitrust Division, Department of Justice, 1401 H Street, NW., Washington, DC 20530 (telephone: 202-307-0001) and at the Office of the Clerk of the United States District Court for the Western District of North Carolina, Room 212, 401 West Trade Street, Charlotte, North Carolina.

Copies of any of these materials may be obtained upon request and payment of a copying fee.

**Constance K. Robinson,**

*Director of Operations, Antitrust Division.*

[FR Doc. 03-19967 Filed 8-5-03; 8:45 am]

BILLING CODE 4410-11-M

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[DEA # 237R]

#### Controlled Substances: Proposed Revised Aggregate Production Quotas for 2003

**AGENCY:** Drug Enforcement Administration (DEA), Justice.

**ACTION:** Notice of proposed revised 2003 aggregate production quotas.

**SUMMARY:** This notice proposes revised 2003 aggregate production quotas for controlled substances in Schedules I and II of the Controlled Substances Act (CSA).

**DATES:** Comments or objections must be received on or before August 27, 2003.

**ADDRESSES:** Send comments or objections to the Acting Administrator, Drug Enforcement Administration, Washington, DC 20537, Attn.: DEA Federal Register Representative (CCR).

**FOR FURTHER INFORMATION CONTACT:** Frank L. Sapienza, Chief, Drug and Chemical Evaluation Section, Drug Enforcement Administration, Washington, DC 20537, Telephone: (202) 307-7183.

**SUPPLEMENTARY INFORMATION:** Section 306 of the CSA (21 U.S.C. 826) requires that the Attorney General establish aggregate production quotas for each basic class of controlled substance listed in Schedules I and II. This responsibility has been delegated to the Administrator of the DEA by § 0.100 of Title 28 of the Code of Federal Regulations.

On December 19, 2002, DEA published a notice of established initial 2003 aggregate production quotas for certain controlled substances in Schedules I and II (67 FR 77809). This notice stipulated that the DEA would adjust the quotas in early 2003 as provided for in part 1303 of Title 21 of the Code of Federal Regulations.

The proposed revised 2003 aggregate production quotas represent those quantities of controlled substances in Schedules I and II that may be produced in the United States in 2003 to provide adequate supplies of each substance for: The estimated medical, scientific, research and industrial needs of the United States; lawful export requirements; and the establishment and maintenance of reserve stocks. These quotas do not include imports of controlled substances for use in industrial processes.

The proposed revisions are based on a review of 2002 year-end inventories, 2002 disposition data submitted by quota applicants, estimates of the medical needs of the United States, product development, and other information available to the DEA.

Therefore, under the authority vested in the Attorney General by section 306 of the CSA of 1970 (21 U.S.C. 826), delegated to the Administrator of the DEA by § 0.100 of Title 28 of the Code of Federal Regulations, the Acting Administrator hereby proposes the following revised 2003 aggregate production quotas for the following controlled substances, expressed in grams of anhydrous acid or base:

| Basic class                                        | Previously established initial 2003 quotas | Proposed revised 2003 quotas |
|----------------------------------------------------|--------------------------------------------|------------------------------|
| <b>Schedule I</b>                                  |                                            |                              |
| 2,5-Dimethoxyamphetamine .....                     | 9,501,000                                  | 9,501,000                    |
| 2,5-Dimethoxy-4-ethylamphetamine (DOET) .....      | 2                                          | 2                            |
| 3-Methylfentanyl .....                             | 4                                          | 4                            |
| 3-Methylthiofentanyl .....                         | 2                                          | 2                            |
| 3,4-Methylenedioxymphetamine (MDA) .....           | 15                                         | 15                           |
| 3,4-Methylenedioxy-N-ethylamphetamine (MDEA) ..... | 10                                         | 10                           |
| 3,4-Methylenedioxymethamphetamine (MDMA) .....     | 19                                         | 19                           |
| 3,4,5-Trimethoxyamphetamine .....                  | 2                                          | 2                            |
| 4-Bromo-2,5-Dimethoxyamphetamine (DOB) .....       | 2                                          | 2                            |
| 4-Bromo-2,5-Dimethoxyphenethylamine (2-CB) .....   | 2                                          | 2                            |
| 4-Methoxyamphetamine .....                         |                                            |                              |
| 4-Methylaminorex .....                             | 7                                          | 7                            |
| 4-Methyl-2,5-Dimethoxyamphetamine (DOM) .....      | 2                                          | 2                            |
| 5-Methoxy-3,4-Methylenedioxymphetamine .....       | 2                                          | 2                            |
| Acetyl-alpha-methylfentanyl .....                  | 2                                          | 2                            |
| Acetyldihydrocodeine .....                         | 2                                          | 2                            |
| Acetylmethadol .....                               | 2                                          | 3                            |
| Allylprodine .....                                 | 2                                          | 2                            |
| Alphacetylmethadol .....                           | 7                                          | 7                            |
| Alpha-ethyltryptamine .....                        | 2                                          | 2                            |
| Alphameprodine .....                               | 2                                          | 2                            |
| Alphamethadol .....                                | 2                                          | 2                            |