

Agenda: At the workshop, FDA will hear presentations and receive comments from stakeholders likely to be affected by FDA policies or procedures regarding the review and approval of innovative medical products. Stakeholders include, but are not limited to device, drug, and biological product manufacturers; members of the academic and clinical communities; and consumer and patient advocacy groups.

Registration: Preregistration is required by July 1, 2003, and will be accepted on a first-come, first-served basis; however, notwithstanding attendance at the workshop, interested persons are encouraged to provide comments (see the *Request for Comments* section of this document). There will be no onsite registration. FDA is pleased to provide the opportunity for interested persons to listen from a remote location to the live proceedings of the public workshop. In order to ensure that a sufficient number of call-in lines are available, please register to listen to the meeting at <http://www.fda.gov/cdrh/meetings/070803.html>. Persons without Internet access may call 1-888-203-6161. The registration deadline is July 1, 2003. For technical reasons, persons wishing to make an oral presentation at the public workshop must do so in person. Those who wish to make presentations should submit written notification including: (1) The specific issue related to the topic you intend to address; (2) the names and addresses of all individuals that will participate in your presentation; (3) the approximate amount of time your presentation will require; and (4) two copies of all presentation materials to Cynthia Benson by June 27, 2003. Presentations will be limited to the topics outlined in the **SUPPLEMENTARY INFORMATION** section of this document and, depending on the number of speakers, FDA may limit the time allotted for each presentation. If you need special accommodations due to a disability, please contact Anne Marie Williams at 301-594-1283 at least 7 days in advance.

Request for Comments: Regardless of attendance at the workshop, interested persons may submit written or electronic comments to the Dockets Management Branch (see the *Addresses* section of this document). You should annotate and organize your comments to identify the specific issues to which they refer. Submit two paper copies of any mailed comments. Individuals may submit one copy. Identify comments with the docket number found in brackets in the heading of this document. The comments that FDA receives will be made available at the

Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Transcripts: Following the workshop, transcripts will be available for review at the Dockets Management Branch (see the *Addresses* section of this document).

SUPPLEMENTARY INFORMATION: FDA believes that innovative and novel medical technologies have the potential to greatly improve the public health in many different areas. By addressing and clarifying regulatory uncertainty, the agency believes that the development of these technologies will be expedited and the predictability in product development will be increased, thus allowing more of these products to reach the marketplace in a timely manner. As part of a broad effort to increase the development of novel medical technologies, FDA is seeking information on how to expedite the review and approval of innovative devices for the delivery of drugs and biologics. For this effort, these products will be broadly defined. We are including any combination of drug and device or biologic and device products in which the two components work together to have a desired effect on the patient. Some examples of the innovative products to be included in this effort are:

- Novel, specialized catheters to permit localized delivery of drugs or biologics (e.g., chemotherapeutic agents, thrombolytics, cells/biologics);
- Lasers or other energy delivery devices for delivery or enhancement of drug or biologic effectiveness (e.g., electroporetic or laser systems to enhance the transport of drugs to the target site);
- Device/drug or device/biologic combinations that permit new routes of administration for drugs (e.g., devices for inhalation of drugs formerly administered intravenously);
- Devices that activate drugs in the body (e.g., photodynamic therapy);
- Drug-eluting stents designed to prevent restenosis; and
- Orthopedic repair products containing bone morphogenic proteins or other cytokines.

The lead for review of the products to be discussed in the workshop may be in any of the FDA medical products centers (the Center for Drug Evaluation and Research, the Center for Biologics Evaluation and Research, and the Center for Devices and Radiological Health, i.e., CDER, CBER or CDRH) and the products may reach the market through several different regulatory pathways (e.g., investigational device exemption/premarket approval applications (IDE/

PMA), investigational new drug application/new drug application (IND/NDA), IND/biological license application (BLA), IDE/510(k), or a combination of these). This workshop is being held to provide a forum for the academic and clinical communities, industry, consumer and patient advocacy groups and FDA to discuss the latest scientific and clinical developments for these products as well as any regulatory concerns and challenges. In addition to increasing our understanding of the latest technological developments in this field, FDA is seeking input to specifically address the following:

1. What are the most critical challenges in developing and bringing to market a novel, innovative technology for delivery of drugs or biologics?
2. Which areas are most important for the agency to provide guidance to developers of these novel products?
3. How can the agency best collaborate with industry, academia, other government agencies, and other scientific bodies in this area of rapidly evolving technology?

The agency hopes to use the information from the workshop to guide the future development of guidance documents, memoranda of understanding, or other position papers.

Dated: May 27, 2003.

Linda S. Kahan,
Deputy Director, Center for Devices and Radiological Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 03N-0168]

Current Status of Useful Written Prescription Drug Information for Consumers: Public Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public meeting to discuss the current status of the private sector's efforts to provide useful written prescription drug information to consumers. Public Law 104-180 adopted a goal that useful written information would be distributed to 75 percent of individuals receiving new prescriptions by the year 2000. An FDA-commissioned study of