

between Data Terminal Equipment and Data Circuit-Terminating Equipment  
Proposed Solicitation Wording:

*Acquisition of General Purpose 37-Position and 9-Position Interface Between Data Terminal Equipment and Data Circuit-Terminating Equipment*

All general purpose 37-position and 9-position interfaces offered for use with data communication systems as a result of the requirements for which this is a part shall conform to the requirements established in FIPS PUB 143.

h. FIPS 146, GOSIP: Government Open Systems Interconnection Profile  
Proposed Solicitation Wording:

*Data Communications Protocols*

All computer network and communications equipment, systems, and services offered as a result of this requirement must provide the protocols specified in the standards contained in FIPS PUB 146, Government Open Systems Interconnection Profile (GOSIP).

i. FIPS 149, General Aspects of Group 4 Facsimile Apparatus  
Proposed Solicitation Wording:

*Design and Acquisition of Group 4 Facsimile Apparatus*

All Group 4 facsimile apparatus designed and offered as a result of this requirement shall comply with FIPS PUB 149.

j. FIPS 150, Facsimile Coding Schemes and Coding Control Functions for Group 4 Facsimile Apparatus  
Proposed Solicitation Wording:

*Design and Acquisition of Facsimile Coding Schemes and Coding Control Functions for Group 4 Facsimile Apparatus*

All Group 4 facsimile apparatus designed and offered as a result of this requirement shall comply with FIPS PUB 150.

k. FIPS 151, POSIX: Portable Operating Systems Interface for Computer Environments  
Proposed Solicitation Wording:

*Acquisition and Development of Portable Operating Systems*

Operating systems environments offered as a result of the requirements of which this is a part shall implement FIPS PUB 151, as well as any additional elements specified elsewhere in this requirements document [insert reference here], and shall require validation in accordance with provisions contained in FIPS PUB 151.

1. FIPS 152, Standard Generalized Markup Language (SGML)  
Proposed Solicitation Wording:

*Acquisition/Development of SGML Systems*

SGML systems offered as a result of the requirements of which this is a part shall conform to the requirements established in FIPS PUB 152 and shall implement all of the language elements of SGML, as well as any additional language features as specified elsewhere in this document [insert reference here], and shall require validation in accordance with the provisions of FIPS PUB 152.

m. FIPS 153, Programmer's Hierarchical Interactive Graphics Systems (PHIGS)  
Proposed Solicitation Wording:

*Acquisition of Programmer's Hierarchical Interactive Graphics System (PHIGS)*

Programmer's Hierarchical Interactive Graphics Systems (PHIGS) offered as a result of the requirements of which this is a part shall conform to the requirements established in FIPS PUB 153 and shall implement all of the functions contained in Part 1 thereof. PHIGS offered must also implement all of the programming language binds contained in FIPS PUB 153, Part 2, except those explicitly excluded as specified elsewhere in this document [insert reference here], as well as any additional programming language bindings specified elsewhere in this document [insert reference here].

n. FIPS 154, High Speed 25-Position Interface for Data Terminal Equipment and Data Circuit-Terminating Equipment  
Proposed Solicitation Wording:

*Acquisition of High Speed 25-Position Interface for Data*

Terminal Equipment and Data Circuit-Terminating Equipment Telecommunications systems offered as a result of this requirement which employ high speed (20,000 to 2,000,000 bits per second) interchange between DTEs and DCEs shall conform to the electrical, mechanical and procedural characteristics established for interfaces in FIPS PUB 154.

o. FIPS 155, Data Communications Systems and Services User-Oriented Performance Measurement Methods.  
Proposed Solicitation Wording:

*Data Communications Performance Testing*

Test measurements on data communications systems and services offered as a result of this requirement shall be conducted in accordance with FIPS PUB 155 to determine compliance with the user oriented performance specification contained in FIPS PUB 144.

2. Amended Specification Terminology:

a. ADP and Telecommunications Standard Index, page 69, implementation of FIPS 68-1, Minimal BASIC, change all references from FIPS 68-1 to FIPS 68-2, and change title from Minimal BASIC, to BASIC.

b. Change Solicitation Wording to read:

*Acquisition of BASIC Language Compilers*

BASIC compilers offered as a result of the requirements of which this is a part shall implement BASIC (FIPS 68-2), as well as any additional language elements as specified elsewhere in this requirements document [insert reference here].

c. Note: the current solicitation language for validation of BASIC compilers contained in the body of the Index, pages 69-70, have not been changed and remain in effect.

Dated: July 11, 1989.

Francis A McDonough,

Deputy Commissioner for Federal Information Resources Management.

[FR Doc. 89-17044 Filed 7-19-89; 8:45 am]

BILLING CODE 6820-25-M

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control**

**National Committee on Vital and Health Statistics Subcommittee on Long-Term Care Statistics; Meeting**

In accordance with the Federal Advisory Committee Act (Pub. L. 92-463), notice is hereby given that the National Committee on Vital and Health Statistics Subcommittee on Long-Term Care Statistics established pursuant to 42 U.S.C. 242k, section 306(k)(2) of the Public Health Service Act, as amended, announces the following Subcommittee meeting.

Name: National Committee on Vital and Health Statistics Subcommittee on Long-Term Care Statistics.

Time and Date: 10:30 am-5:00 pm—August 15, 1989.

Place: Hubert H. Humphrey Building, Room 337A-339A, 200 Independence Avenue, SW., Washington, DC 20201.

Status: Open.

Purpose: To be briefed on the status of mental illness screening of nursing home patients and to consider a possible information system, resulting from the HCFA Uniform Data Set for Nursing Home Dissident Assessment.

Contact Person for More Information: Substantive program information as well as

summaries of the meeting and roster of Committee members may be obtained from Gail F. Fisher, Ph.D., Executive Secretary, National Committee on Vital and Health Statistics, Room 2-12, Center Building, 3700 East West Highway, Hyattsville, Maryland 20782, telephone (301) 436-7050.

Dated: July 13, 1989.

Elvin Hilyer,

Associate Director for Policy Coordination,  
Centers for Disease Control.

[FR Doc. 89-16989 Filed 7-19-89; 8:45 am]

BILLING CODE 4160-18-M

## Food and Drug Administration

[Docket No. 89C-0203]

### Bausch & Lomb, Inc.; Filing of Color Additive Petition

AGENCY: Food and Drug Administration.

ACTION: Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that Bausch & Lomb, Inc., has filed a petition proposing that the color additive regulations be amended to provide for the safe use of 1,4-bis[4-(2-methacryloxyethyl)phenylamino] anthraquinone to color contact lenses.

#### FOR FURTHER INFORMATION CONTACT:

Sandra L. Varner, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-472-5690.

**SUPPLEMENTARY INFORMATION:** Under the Federal Food, Drug, and Cosmetic Act (sec. 706(d)(1), 74 Stat. 402-403 (21 U.S.C. 376(d)(1))), notice is given that a petition (CAP 9C0215) has been filed by Bausch & Lomb, Inc., 1400 North Goodman, Rochester, NY 14692-0450, proposing that 21 CFR Part 73 of the color additive regulations be amended to provide for the safe use of 1,4-bis[4-(2-methacryloxyethyl)phenylamino] anthraquinone to color contact lenses.

The potential environmental impact of this action is being reviewed. If the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the Federal Register in accordance with 21 CFR 25.40(c).

Dated: July 10, 1989.

Richard J. Ronk,

Deputy Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89-16951 Filed 7-19-89; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 89M-0211]

### Technomed International; Premarket Approval of the Technomed SONOLITH 2000 Lithotripter

AGENCY: Food and Drug Administration.

ACTION: Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing its approval of the application by Technomed International, Danvers, MA, for premarket approval, under the Medical Device Amendments of 1976, of the Technomed SONOLITH 2000 Lithotripter. After reviewing the recommendation of the Gastroenterology-Urology Devices Panel, FDA's Center for Devices and Radiological Health (CDRH) notified the applicant, by letter of June 23, 1989, of the approval of the application.

**DATE:** Petitions for administrative review by August 21, 1989.

**ADDRESS:** Written requests for copies of the summary of safety and effectiveness data and petitions for administrative review to the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857.

#### FOR FURTHER INFORMATION CONTACT:

Robert Gatling, Center for Devices and Radiological Health (HFZ-420), Food and Drug Administration, 1390 Piccard Dr., Rockville, MD 20850, 301-427-7750.

**SUPPLEMENTARY INFORMATION:** On June 22, 1988, Technomed International, Danvers, MA 01923, submitted to CDRH an application for premarket approval of the Technomed SONOLITH 2000 Lithotripter. The device is an extracorporeal shock wave lithotripter for use in the fragmentation of kidney stones.

On December 6, 1988, the Gastroenterology-Urology Devices Panel, an FDA advisory committee, reviewed and recommended approval of the application. On June 23, 1989, CDRH approved the application by a letter to the applicant from the Acting Director of the Office of Device Evaluation, CDRH.

A summary of the safety and effectiveness data on which CDRH based its approval is on file in the Dockets Management Branch (address above) and is available from that office upon written request. Requests should be identified with the name of the device and the docket number found in brackets in the heading of this document.

A copy of all approved labeling is available for public inspection at

CDRH—contact Robert Gatling (HFZ-420), address above.

### Opportunity for Administrative Review

Section 515(d)(3) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360e(d)(3)) authorizes any interested person to petition, under section 515(g) of the act (21 U.S.C. 360e(g)), for administrative review of CDRH's decision to approve this application. A petitioner may request either a formal hearing under Part 12 (21 CFR Part 12) of FDA's administrative practices and procedures regulations or a review of the application and CDRH's action by an independent advisory committee of experts. A petition is to be in the form of a petition for reconsideration under § 10.33(b) (21 CFR 10.33(b)). A petitioner shall identify the form of review requested (hearing or independent advisory committee) and shall submit with the petition supporting data and information showing that there is a genuine and substantial issue of material fact for resolution through administrative review. After reviewing the petition, FDA will decide whether to grant or deny the petition and will publish a notice of its decision in the Federal Register. If FDA grants the petition, the notice will state the issue to be reviewed, the form of review to be used, the persons who may participate in the review, the time and place where the review will occur, and other details.

Petitioners may, at any time on or before August 21, 1989, file with the Dockets Management Branch (address above) two copies of each petition and supporting data and information, identified with the name of the device and the docket number found in brackets in the heading of this document. Received petitions may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

This notice is issued under the Federal Food, Drug, and Cosmetic Act (secs. 515(d), 520(h), 90 Stat. 554-555, 571 (21 U.S.C. 360e(d), 360j(h))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.10) and redelegated to the Director, Center for Devices and Radiological Health (21 CFR 5.53).

Dated: July 13, 1989.

Walter E. Gundaker,

Acting Deputy Director, Center for Devices and Radiological Health.

[FR Doc. 89-16952 Filed 7-19-89; 8:45 am]

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## Health Care Financing Administration

### Public Information Collection Requirements Submitted to the Office of Management and Budget for Clearance

**AGENCY:** Health Care Financing Administration.

The Department of Health and Human Services (HHS) previously published a list of information collection packages it submitted to the Office of Management and Budget (OMB) for clearance in compliance with the Paperwork Reduction Act (Pub. L. 96-511). The Health Care Financing Administration (HCFA), a component of HHS, now publishes its own notices as the information collection requirements are submitted to OMB. The HCFA has submitted the following requirements to OMB since the last HCFA list was published.

1. *Type of Request:* Revision; *Title of Information Collection:* Hospital Survey Report Form; *Form Number:* HCFA-1537; *Frequency:* On occasion; *Respondents:* State/local governments; *Estimated Number of Responses:* 1,539; *Average Hours per Response:* 3.25; *Total Estimated Burden Hours:* 5,002.

2. *Type of Request:* Revision; *Title of Information Collection:* Medicare/Medicaid Hospital Swing-bed Survey Report Forms; *Form Number:* HCFA-1537C; *Frequency:* Annually; *Respondents:* State/local governments; *Estimated Number of Responses:* 1,500; *Average Hours per Response:* .25; *Total Estimated Burden Hours:* 375.

3. *Type of Request:* Extension; *Title of Information Collection:* Hospital Request for Certification in the Medicare/Medicaid Program; *Form Number:* HCFA-1514; *Frequency:* On occasion; *Respondents:* State/local governments; *Estimated Number of Responses:* 1,984; *Average Hours per Response:* .25; *Total Estimated Burden Hours:* 496.

4. *Type of Request:* Revision; *Title of Information Collection:* Medicare Renal Dialysis Facility Cost Report; *Form Number:* HCFA-265; *Frequency:* Annually; *Respondents:* Business/other for profit and small businesses/organizations; *Estimated Number of Responses:* 1,097; *Average Hours per Response:* 196; *Total Estimated Burden Hours:* 215,012.

5. *Type of Request:* Extension; *Title of Information Collection:* Outpatient Physical Therapy and/or Speech Pathology Services Request for Certification and Survey Report Form; *Form Number:* HCFA-1856/1893; *Frequency:* Annually; *Respondents:* State/local governments; *Estimated*

*Number of Responses:* 313; *Average Hours per Response:* 1.75; *Total Estimated Burden Hours:* 548.

6. *Type of Request:* Extension; *Title of Information Collection:*

Histocompatibility Testing Laboratories Survey Report Form; *Form Number:* HCFA-445; *Frequency:* Annually; *Respondents:* State/local governments; *Estimated Number of Responses:* 200; *Average Hours per Response:* 1; *Total Estimated Burden Hours:* 200.

7. *Type of Request:* Extension; *Title of Information Collection:* Request for Certification as a Supplier and the

Portable X-ray Survey Report Form; *Form Number:* HCFA-1880/1882; *Frequency:* Annually; *Respondents:* State/local governments; *Estimated Number of Responses:* 160; *Average Hours per Response:* 1.75; *Total Estimated Burden Hours:* 280.

*Additional Information or Comments:*

Call the Reports Clearance Officer on 301-966-2088 for copies of the clearance request packages. Written comments and recommendations for the proposed information collections should be sent directly to the following address: OMB Reports Management Branch, Attention: Allison Herron, New Executive Office Building, Room 3208, Washington, DC 20503.

Date: July 13, 1989.

Louis B. Hays,

Acting Administrator, Health Care Financing Administration.

[FR Doc. 89-16975 Filed 7-19-89; 8:45 am]

BILLING CODE 4120-03-M

### Privacy Act of 1974; Matching Program

**AGENCY:** Department of Health and Human Services (HHS), Health Care Financing Administration (HCFA).

**ACTION:** Notice of a Matching Program—Beneficiary State File (BEST) and Medicaid master files maintained by State agencies.

**SUMMARY:** The Department of Health and Human Services in providing public notice that HCFA agrees to disclose information on Medicare eligible individuals to State agencies.

The matching report set forth below is in compliance with the Computer Matching and Privacy Protection Act of 1988 (Pub. L. No. 100-503).

**DATE:** The match will begin August 21, 1989, and shall remain in effect for a period not to exceed 18 months unless three months prior to the actual expiration date, the Data Integrity Board of the Department of Health and Human Services finds that the program will be conducted without change and each

party certifies that the program has been conducted in compliance with the Matching Agreement. Under this finding, the Board may extend the Agreement for one additional year.

**ADDRESS:** The public should address comments to Richard A. DeMeo, Health Care Financing Administration, Privacy Act Officer, Office of Budget and Administration, Room G-M-1, East Low Rise Building, 6325 Security Boulevard, Baltimore, Maryland 21207. Comments will be available for inspection at this location.

**FOR FURTHER INFORMATION CONTACT:** William A. Grant, Division of Entitlement Requirements, Office of Program Operations Procedures, Bureau of Program Operations, Health Care Financing Administration, G-E-7 Meadows East Building, 6325 Security Boulevard, Baltimore, Maryland 21207, Telephone Number 301-966-6464.

**SUPPLEMENTARY INFORMATION:** The Health Insurance (HI) Master Record contains a record of each individual who is, or has been, entitled to health insurance benefits under title XVIII of the Social Security Act (the Act). The BEST file is an extract of the HI Master Record. The BEST file is an alphabetic listing, on magnetic tape, of all Medicare beneficiaries in the State by name, sex, date of birth, address, zip code, buy-in indicator and health insurance claim number (HICN). This extract is the most economical and efficient method of obtaining the claim number of a beneficiary when other identifying information is available. The BEST file has been provided to all States for the express purpose of identifying Medicaid beneficiaries to the State buy-in rolls. In most States, the activity is conducted by the State agency responsible for Medicaid.

The BEST file will enable the State agency to identify individuals eligible for inclusion in a State buy-in account. The BEST file will also enable the State agency to identify those Medicare/Medicaid dually eligible individuals for whom Medicaid has deductible and co-insurance liability. States under a buy-in agreement may enroll individuals for Medicare Part B coverage who are eligible for title XVI (Supplemental Security Income) benefits or eligible under State grants for medical assistance. The State pays the Medicare Part B premiums for these individuals eligible under title XVI.

To query the BEST file, it is necessary for the State to have the individual's name, date of birth, sex and address. With this information, the HICN can be determined. The State is then able to