IX. Paperwork Reduction Act

The information collection required by this permit has been submitted to OMB under the provisions of the Paperwork Reduction Act, 44 U.S.C. 3501 *et seq.,* in submission made for the NPDES permit program and assigned OMB control number 2040–0004 [(NPDES Discharge Monitoring Reports (DMRs)].

Because this permit is very similar in reporting and application requirements and in discharges which are required to be monitored as the previous Eastern Gulf of Mexico OCS general permit (GEG460000), the paperwork burdens are expected to be nearly identical. The only new requirement is entry of acute WET tests results for well treatment, completion and workover fluids discharged separately than produced wastewaters into the electronic system. When it issued the previous OCS general permit, EPA estimated it would take an affected facility three hours to prepare the request for coverage and 38 hours per year to prepare DMRs. It is estimated that the time required to prepare the request for coverage and DMRs for the reissued permit will be approximately the same.

Dated: June 2, 2023.

Denisse Diaz,

Acting Director, Water Division.

[FR Doc. 2023–12292 Filed 6–8–23; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[FR ID 147101]

Deletion of Item From June 8, 2023 Open Meeting

June 6, 2023.

The following item was adopted and released by the Commission on June 5, 2023 and deleted from the list of items scheduled for consideration at the Thursday, June 8, 2023, Open Meeting. The item was previously listed in the Commission's Sunshine Notice on Thursday, June 1, 2023.

5	Media	*Title:* Restricted Adjudicatory Matter. *Summary:* The Commission will consider a restricted adjudicatory matter.

Federal Communications Commission.

Marlene Dortch,

Secretary.

[FR Doc. 2023–12355 Filed 6–8–23; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–23–1309; Docket No. CDC–2023–0047]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of Government information, invites the general public and other Federal agencies the opportunity to comment on a continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Enterprise Laboratory Information Management System (ELIMS). This data collection is used by CDC to record specimen metadata and patient data related to test order requests submitted by external partners (SPHLs, International organizations, Federal institutions, hospitals, doctor's offices, etc.) to the CDC Infectious Diseases testing laboratories.

DATES: CDC must receive written comments on or before August 8, 2023.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2023–0047 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected;

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses; and

5. Assess information collection costs.