the online instructions for submitting comments. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Additional instructions on commenting and visiting the docket, along with more information about dockets generally, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Charles Smith, Director, Registration Division (7505M), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–1030; email address: *RDFRNotices@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

You may be potentially affected by this action if you are an agricultural producer, food manufacturer, or pesticide manufacturer. The following list of North American Industrial Classification System (NAICS) codes is not intended to be exhaustive, but rather provides a guide to help readers determine whether this document applies to them. Potentially affected entities may include:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

B. What should I consider as I prepare my comments for EPA?

1. *Submitting CBI.* Do not submit CBI to EPA through email or *https://www.regulations.gov.* If you wish to include CBI in your comment, please follow the applicable instructions at *https://www.epa.gov/dockets/commenting-epa-dockets#rules* and clearly mark the information that you claim to be CBI. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

2. *Tips for preparing your comments.* When preparing and submitting your comments, see guidance provided at *https://www.epa.gov/dockets/commenting-epa-dockets.*

II. Registration Application

EPA has received the following application to register new uses for a new pesticide product containing a currently registered active ingredient. Pursuant to FIFRA section 3(c)(4) (7 U.S.C. 136a(c)(4)), EPA is hereby providing notice of receipt of this application and an opportunity to comment on the information provided below as well as the current proposed labeling associated with this application. Notice of receipt of this application does not imply a decision by the Agency on this application.

File Symbol: 7969–LNT. *Docket ID number:* EPA–HQ–OPP–2024–0154. *Applicant:* BASF Corporation, 26 Davis Drive, Research Triangle Park, North Carolina 27709–3528. *Active ingredient:* Dicamba. *Product type:* Herbicide. *Proposed use:* Dicamba-tolerant cotton and dicamba-tolerant soybeans. *Contact:* RD. This proposed new use has been coded as an R170, additional food use, which carries a PRIA 5 statutory review time of 17 months from the date that the action gets in-processed. Because EPA expects a large stakeholder interest in this application, EPA also included in the docket the BASF's current proposed labeling associated with the application.

(Authority: 7 U.S.C. 136 *et seq.*)

Dated: May 29, 2024.

Charles Smith,

Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 2024–12109 Filed 6–3–24; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–11970–01–R8]

Clean Air Act Operating Permit Program; Order on Petition for Objection to State Operating Permit for DCP Operating Company, LP: Platteville Natural Gas Processing Plant

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of final order on petition.

SUMMARY: The Environmental Protection Agency (EPA) Administrator signed an order dated April 2, 2024, granting a petition dated September 19, 2023, from the Center for Biological Diversity. The petition requested that the EPA object to the Clean Air Act (CAA) operating permits issued by the Colorado Department of Public Health and Environment (CDPHE) to DCP Operating Company, LP for its Platteville Natural Gas Processing Plant, located in Weld County, Colorado.

FOR FURTHER INFORMATION CONTACT: Donald Law, EPA Region 8, telephone number: (303) 312–7015, email address: *law.donald@epa.gov*. The final order and petition are available electronically at: *https://www.epa.gov/title-v-operating-permits/title-v-petition-database*.

SUPPLEMENTARY INFORMATION: The EPA received a petition from the Center for Biological Diversity dated September 19, 2023, requesting that the EPA object to the issuance of operating permit no. 02OPWE252 issued by CDPHE to DCP Operating Company, LP in Weld County, Colorado. On April 2, 2024, the EPA Administrator issued an order granting in part and denying in part the petition. The order itself explains the basis for the EPA's decision.

Sections 307(b) and 505(b)(2) of the CAA provide that a petitioner may request judicial review of those portions of an order that deny issues in a petition. Any petition for review shall be filed in the United States Court of Appeals for the appropriate circuit no later than August 5, 2024.

KC Becker,

Regional Administrator, Region 8.

[FR Doc. 2024–12217 Filed 6–3–24; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–24–24FY; Docket No. CDC–2024–0046]

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 proposed information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Lung Function Screening of Construction Workers with Exposure to Dusts and Chemicals. The goal of the proposed study is to determine if small airway dysfunction or early-stage disease can consistently be identified in high-risk workers with normal spirometry.

DATES: CDC must receive written comments on or before August 5, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0046 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.

Proposed Project

Lung Function Screening in Construction Workers Exposed to Dusts and Chemicals—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Construction workers are routinely exposed to a number of inhaled toxins known to contribute to the development of chronic respiratory disease. A 2010 industry study suggested that over 50% of construction workers reported occupational exposure to vapors, gases, dusts, and fumes (VGDF) at least twice a week and that nearly 18% of ever-employed construction industry workers have abnormal lung function. In fact, construction workers are approximately 1.64 times more likely to have airway obstruction than other working groups within the U.S. trades.

An emerging lung function test procedure called impulse oscillometry (IOS) is more sensitive than spirometry, the standard test for monitoring worker pulmonary function. IOS can identify small airway functional status with the potential to identify those with often reversible abnormalities within the small airways (<2mm diameter). Thus, the goal of the proposed study is to determine if small airway dysfunction or early-stage disease can consistently be identified by IOS in high-risk workers with normal spirometry.

Two construction worker groups will be enrolled: (1) those at risk for respirable silica exposure (*i.e.,* blockmasons or bricklayers); and (2) workers with occupational exposure to welding fumes and metals (*i.e.,* welders). NIOSH researchers will collect questionnaire information pertaining to respiratory symptomology, smoking history, job-type tenure, worksite mitigation strategies, and personal protective equipment use. Height, weight, blood pressure and lung function testing measures will also be performed. The amount of personal identifiable information collected will be limited, but it is necessary to collect each worker's age, month and year of birth, sex assigned at birth, race/ethnicity, and smoking history to evaluate lung function test results. Additional medical data will be collected to reduce the risk of adverse effects or transmission of infectious disease to subjects performing lung function.

CDC requests a two-year OMB approval for an estimate 83 annual burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Eligible Study Participants	Informed Consent	75	1	20/60	25
Study Participants	Height, Weight, and Demographics	75	1	10/60	13
Study Participants	Study Questionnaire	75	1	10/60	13
Spirometry Lung Function Test Results.	Spirometry Test Report	75	1	15/60	19
Impulse Oscillometry Test Results	Oscillometry Test Report	75	1	10/60	13
Total					83

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2024–12232 Filed 6–3–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–24–0900; Docket No. CDC–2024–0045]

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 proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Contact Investigation Outcome Reporting Forms. The project includes the contact investigation outcome reporting forms used to obtain data from State, local, and territorial public health professionals or conveyance operators and medical professionals on their contact tracing efforts to better assess the risk to individuals who may have been exposed to a confirmed case of a communicable disease of public health concern while traveling to or within the United States.

DATES: CDC must receive written comments on or before August 5, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0045 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.

Proposed Project

Contact Investigation Outcome Reporting Forms (OMB Control No. 0920–0900, Exp. 8/31/2024)—Revision—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This proposed information collection project includes the contact investigation outcome reporting information collection tools used by CDC to better assess the risk to interstate and arriving international travelers who may have been exposed to a confirmed case of a communicable disease of public health concern while traveling. Different forms are tailored for different diseases of public health concern that are tracked by the Division of Global Migration Health (DGMH/CDC). The information will be used to assist and collaborate with State, local, and territorial health departments, conveyance operators, air, maritime, and land port of entry partners, and international public health authorities to identify potential exposures and to determine the risk of infection, and whether future public health interventions are needed.

Methods used to collect the information are basic surveys of respondents that record information about the exposed traveler's location and activities on air or maritime conveyances or land border crossing, other potential exposures, signs/symptoms that may have occurred after their potential exposure, prior history of vaccination or disease, and other medical conditions that could influence the risk of infection or severity of illness.

Due to the COVID–19 pandemic, CDC modified how cruise ships report information on cases of influenza-like-illness. Since December 2023, CDC has been using a new surveillance system for cumulative acute respiratory illness (ARI) reporting (under OMB Control Number 0920–1335), which includes cases of influenza, COVID–19, and respiratory syncytial virus. One form (the Influenza Outbreak Enhanced Data Collection Form) is not used routinely now; however, this form may be used in limited circumstances in the future. The burden of outcome reporting forms has been adjusted based on more recent numbers, which notably excludes COVID–19 numbers because we are no longer requiring contact investigations for COVID–19 and thus these are lower than past estimates. This applies primarily to the General Contact Investigation Outcome Reporting Form. Other burden estimates have been adjusted to reflect current estimates reflecting 2021–2023 numbers.

CDC requests OMB approval for an estimated 50 annual burden hours.