

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: [RDFFRNotices@epa.gov](mailto:RDFFRNotices@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 PRA 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](mailto: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 symptomatology, 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](http://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](http://www.regulations.gov).

*Please note:* Submit all comments  
through the Federal eRulemaking portal  
([www.regulations.gov](http://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](mailto: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.