secure means of their choice (*e.g.,* web-based application, fax or email).

As part of this revision, CDC requests approval for a number of changes and adjustments:

- CDC is discontinuing all Ebola related forms;
- CDC is discontinuing all current maritime-related forms except a condensed maritime TB contact investigation follow-up form in an Excel format;
- CDC is requesting a downward revision of the estimated number of TB contact investigation forms used annually, but an upward revision of the amount of time requested from each respondent;
- CDC is requesting addition of varicella and influenza like illness outbreak contact investigation follow up forms;
- no changes are requested of the Air or Land associated forms; however adjustments in burden are requested.

The proposed changes will result in a decrease of 673 burden hours (from 782 burden hours to 109 hours).

There is no cost to respondents other than their time to complete the form and submit the data to CDC.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Cruise Ship Physicians/Cargo Ship Managers	Clinically Active TB Contact Investigation Outcome Reporting Form—Maritime.	15	1	20/60
Cruise Ship Physicians/Cargo Ship Managers	Varicella Investigation Outcome Reporting Form.	29	1	20/60
Cruise Ship Physicians/Cargo Ship Managers	Influenza Like Illness Investigation Outcome Reporting Form.	45	1	20/60
State/Local public health staff	General Contact Investigation Outcome Reporting Form—Air.	34	1	5/60
State/Local public health staff	TB Contact Investigation Outcome Reporting Form—Air.	547	1	5/60
State/Local public health staff	Measles Contact Investigation Outcome Reporting Form—Air.	324	1	5/60
State/Local public health staff	Rubella Contact Investigation Outcome Reporting Form—Air.	27	1	5/60
State/Local public health staff	General Contact Investigation Outcome Reporting Form—Land.	15	1	5/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2018–08163 Filed 4–18–18; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: TANF Office Culture Study.

OMB No.: New Collection.

Description: The Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS) is proposing data collection activities as part of a project to identify and describe exemplars of TANF organizational culture as well as successful strategies human services offices have undertaken to improve their organizational culture. This qualitative study intends to use this information to increase understanding of how various agencies' organizational cultures influence TANF clients' experiences, service delivery, and frontline workers.

The information collection activities to be submitted in the package include:

(1) Leadership and supervisor interviews will collect information on program structure and staffing, client experiences, agency goals and performance management, organizational learning and innovation, cultural congruence across service providers, and the perception of the organizational culture change, if applicable.

(2) Frontline workers' interviews will collect information about frontline staffs' role in service delivery, client experiences, peer interaction and social institutions within the agency, agency goals, organizational learning and innovation, and the perception of the organizational culture change initiative, if applicable.

(3) The focus groups will collect information about program participants' perceptions of agency processes, their communication with agency staff, and their assessment of the agency's organizational culture.

Respondents: Individuals receiving TANF and related services, TANF directors, and managers and staff at local TANF offices.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Leadership and Supervisor Interview Guide	24	8	1	1.5	12
Frontline Staff Interview Guide	12	4	1	1	4
Focus Group Guide	54	18	1	1.5	27

Estimated Total Annual Burden Hours: 43.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, *Email: OIRA_SUBMISSION@OMB.EOP.GOV*, *Attn:* Desk Officer for the Administration for Children and Families.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–08233 Filed 4–18–18; 8:45 am]

BILLING CODE 4184–09–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–D–1098]

Metered Dose Inhaler and Dry Powder Inhaler Drug Products—Quality Considerations; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for industry entitled ''Metered Dose Inhaler (MDI) and Dry Powder Inhaler (DPI) Drug Products—Quality Considerations.'' The purpose of this guidance is to provide recommendations to industry on the development and manufacture of inhalation aerosols (also known as metered dose inhalers, or MDIs) and inhalation powders (also known as dry powder inhalers, or DPIs). Although not explicitly discussed, some of the principles and recommendations provided in this guidance may be applicable to nasal delivery products, as well. The recommendations in this guidance can apply to MDI and DPI products intended for local or systemic effect.

DATES: Submit either electronic or written comments on the draft guidance by June 18, 2018 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov*.
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2018–D–1098 for ''Metered Dose Inhaler (MDI) and Dry Powder Inhaler (DPI) Drug Products—Quality Considerations.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the draft guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10001 New Hampshire Ave. Hillandale Building, 4th Floor, Silver Spring, MD 20993–0002. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT: Richard Lostritto, Center for Drug