

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day–18–18AJJ; Docket No. CDC–2018–0056]

#### 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 Knowledge, Attitudes, and Practices of US Large Animal Veterinarians Concerning Common Veterinary Infection Control Measures When Working with Animal Obstetric Cases. The goals of this survey are to better describe veterinarians' current knowledge of zoonotic infectious diseases that cause abortion in large animals, determine common veterinary infection control practices when working up obstetric cases, and identify common barriers to personal protective equipment use.

**DATES:** CDC must receive written comments on or before September 18, 2018.

**ADDRESSES:** You may submit comments, identified by Docket No. CDC–2018–0056 by any of the following methods:

- *Federal eRulemaking Portal:* [Regulations.gov](http://www.Regulations.gov). Follow the instructions for submitting comments.

- *Mail:* Jeffrey Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

*Instructions:* All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to [Regulations.gov](http://www.Regulations.gov).

*Please note: Submit all comments through the Federal eRulemaking portal ([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 Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 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; and
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.
5. Assess information collection costs.

#### Proposed Project

Knowledge, Attitudes, and Practices of U.S. Large Animal Veterinarians Concerning Common Veterinary Infection Control Measures When Working with Animal Obstetric Cases—New ICR—National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

Veterinarians are particularly at risk of contracting zoonotic infectious diseases due to their close proximity to animals, especially during times of injury or illness. Some veterinarians may be unaware of recommended personal protection measures or opt not to participate in measures that would decrease their risk of contracting a zoonotic disease (Wright *et al.* 2008). In 1977, a survey conducted of 1182 veterinarians showed that approximately 43% of the respondents had contracted an infectious zoonotic disease (Schnurrenberger & Martin 1977). Today, this elevated zoonotic disease risk persists; the seroprevalence of Q fever in U.S. veterinarians is 22% (Whitney, Massung, *et al.* 2009) and the seroprevalence of leptospirosis is 2.5% (Whitney, Ailes, *et al.* 2009). Within the veterinary profession, large animal practitioners might have an increased risk of occupational exposure to infectious zoonotic diseases for many reasons, including decreased biosecurity measures available in the field and the limited space available on a mobile practice for PPE.

The goals of this study are to establish veterinarians' knowledge of zoonotic infectious disease, identify veterinarians' attitudes towards zoonotic infectious disease and personal risk, and determine practices to decrease personal risk of infection. By identifying knowledge gaps in personal protective equipment (PPE) use, transmission risk factors, and disease identification/diagnosis, we aim to determine the best methods for education of veterinarians on relevant abortion-associated zoonotic infectious diseases.

The purpose of this study is to better describe veterinarians' current knowledge of zoonotic diseases that cause abortion in large animals, determine common veterinary infection control practices when working up obstetric cases, and identify common barriers to PPE use. In order to develop effective messaging strategies, a deeper understanding of the attitudes and barriers to PPE use is needed. The information collected will be used to improve and enhance zoonotic disease education and PPE guidance targeted to veterinarians. The estimated annual burden hours are 125. There is no cost to respondents other than their time.

## 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) |
|---------------------|-----------|-----------------------|------------------------------------|--|-------------------------|
| Veterinarian .....  | .....     | 500                   | 1                                  | 15/60                                  | 125                     |
| Total .....         | .....     | .....                 | .....                              | .....                                  | 125                     |

**Jeffrey M. Zirger,**

Acting 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.

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier CMS-10675]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, HHS.

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments must be received by September 18, 2018.

**ADDRESSES:** When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and

recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number \_\_\_\_\_, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to [Paperwork@cms.hhs.gov](mailto:Paperwork@cms.hhs.gov).

3. Call the Reports Clearance Office at (410) 786-1326.

**FOR FURTHER INFORMATION CONTACT:** William Parham at (410) 786-4669.

#### SUPPLEMENTARY INFORMATION:

##### Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

#### **CMS-10675 Evaluation of the CMS Quality Improvement Organizations: Medication Safety and Adverse Drug Event Prevention**

Under the 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. The term "collection of information" is

defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

#### Information Collection

1. *Type of Information Collection Request:* New Collection of Information Request; *Title of Information Collection:* Evaluation of the CMS Quality Improvement Organizations: Medication Safety and Adverse Drug Event Prevention; *Use:* The purpose of this Information Collection Request (ICR) is to collect data to inform the program evaluation of the Centers for Medicare & Medicaid Services (CMS) Quality Improvement Organizations (QIO) current contract known as the 11th Scope of Work (SOW). The current ICR focuses on evaluating one component of the quality improvement activities of the Quality Innovation Network Quality Improvement Organizations (QIN-QIOs) and is part of a larger evaluation of the overall impact of the QIO program. This ICR aims to assess the QIN-QIO Task which focuses on Medication Safety and Adverse Drug Event Prevention. For this evaluation, we are using a mixed-methods design to compare quality improvement activities of pharmacists, physicians, and nursing home administrators or directors of nursing at nursing homes participating in the QIN-QIO program (participating) with those not participating in the QIN-QIO program (non-participating).

As mandated by Sections 1152-1154 of the Social Security Act, CMS directs the QIO program, which is one of the largest federal programs dedicated to improving health quality for Medicare beneficiaries. QIOs are groups of health quality experts, clinicians, and