on respondents; and was absorbed in the OMB burden previously approved.

NHCS Questions:

1. In the past year, did your hospital experience shortages of coronavirus disease (COVID–19) tests for any patients with presumptive positive COVID–19 infection?

2. In the past year, did your hospital create areas outside the hospital entrance to screen patients for coronavirus disease (COVID–19) infection?

3. In the past year, did your hospital need to turn away or refer elsewhere any patients with confirmed or presumptive positive coronavirus disease (COVID–19) infection?

4. In the past year, did any of the following clinical care providers in your hospital test positive for coronavirus disease (COVID–19) infection?

a. Physicians
b. Physician assistants
c. Nurse practitioners
d. Certified nurse-midwives
e. Registered nurses/licensed practical nurses
f. Other clinical care providers

5. For calendar year 2020, how many inpatient/ED visits at your hospital were related to CONFIRMED coronavirus disease (COVID–19) infections, by quarter or by year? Fill in the grid below.

6. For calendar year 2020, how many inpatient/ED visits at your hospital were Confirmed COVID–19 visits and how many were Presumptive Positive COVID–19 visits by quarter or by year?

Dated: December 14, 2020.

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2020–27820 Filed 12–17–20; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2020–0124]

Advisory Committee on Immunization Practices (ACIP); Correction

Notice is hereby given of a change in the meeting of the *Advisory Committee on Immunization Practices (ACIP);* December 18, 2020, 12:00 p.m.—6:00 p.m., EST; and December 20, 2020, 12:00 p.m.—6:00 p.m., EST (times subject to change, see the ACIP website for any updates: *http://www.cdc.gov/vaccines/acip/index.html*), which was published in the **Federal Register** on December 11, 2020, Volume 85, Number 239, page 80108.

The meeting dates and times should read as follows:

DATES: The meeting will be held on December 19—20, 2020 from 11 a.m. to 4:30 p.m., EST (times subject to change, see the ACIP website for any updates: *http://www.cdc.gov/vaccines/acip/index.html*).

Written comments must be received on or before December 21, 2020.

The meeting is open to the public.

FOR FURTHER INFORMATION CONTACT: Stephanie Thomas, ACIP Committee Management Specialist, Centers for Disease Control and Prevention, National Center for Immunization and Respiratory Diseases, 1600 Clifton Road, NE, MS–H24–8, Atlanta, GA 30329–4027; Telephone: 404–639–8367; Email: *ACIP@cdc.gov.*

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2020–28090 Filed 12–16–20; 4:15 pm]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–10346]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 (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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by January 19, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain* . Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

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 website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

2. Call the Reports Clearance Office at (410) 786–1326.

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

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. 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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension without change of a currently approved collection; *Title of Information Collection:* Appeals of Quality Bonus Payment Determinations;