

**Request for Approval under the “Generic Clearance for Improving Customer Experience:
OMB Circular A-11, Section 280 Implementation”
(OMB Control Number: 0938-1382)**

TITLE OF INFORMATION COLLECTION: Medicaid.gov Static Feedback (CMS-10911)

TYPE OF ACTIVITY

Select all that apply.

☐ Screener (for example, distributed before or during a usability testing session or other kind of session)

☐ Question script for interview, focus group, discussion group, etc. Scripts for usability testing sessions are, in general, exempt from PRA review subject to the caveats on page 3 of this [OMB Memorandum](#).

☒ Survey to obtain feedback immediately following a transaction - **limited to 15 questions and 5-minutes of burden maximum.**

☐ Other survey

TYPE OF SUBMISSION

☐ New collection: Select this if you do not already have an approved collection.

☒ Change to an already approved collection: Select this if you have an already approved collection but are making changes to the questions, the number of respondents, etc.

Please attach your instrument as a separate document using the Word template that pertains to your particular engagement, i.e., either a survey/screener or script. OMB has prepared special templates that must be used for submitting your instrument. Please ask your agency’s PRA officer for these templates. ***Do not paste the question list within this document.*** As a reminder, usability testing session scripts are, in general, exempt from PRA review subject to the caveats on p.3 [here](#), but screeners for those sessions must be submitted for PRA review.

PURPOSE OF COLLECTION:

What are you hoping to learn / improve? How do you plan to use what you learn to support service delivery improvement activities? Are there artifacts (user personas, journey maps, digital roadmaps, summary of customer insights to inform service improvements, performance dashboards) the data from this collection will inform?

The CMCS/ Operation Services Group/ Division of Communication and Outreach is updating the current Medicaid.gov Static Feedback survey activity OMB 0190-0008. The Medicaid.gov Feedback survey remains within OMB and A-11 requirements and is fully aligned with other CMS website surveys to reduce costs while gathering useful, quality data. We are using this juncture to update the current Medicaid.gov Feedback Survey (survey OMB 0190-0008) to be succinct, streamlined, and consistent with the data gathered by the other Medicaid.gov and IKN surveys. The

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survey contains only 5 questions to minimize user burden while focusing on program priorities and the user experience. The updated survey will ask users to voluntarily self-identify their user type, their trust in Medicaid.gov, satisfaction, and reason for visiting, while offering the option for detailed written feedback. This combination of data will provide a better understanding of each user and their journey on the site allowing us to better understand user needs and how we can improve our services.

Our approach does not use any statistical methods – no statistical techniques or models have been applied to the data. Since this is a survey, we have added Supporting Statement B to this submission.

ACTIVITY DETAILS

1. If this is a survey for a High Impact Service Provider, will the results of this survey be reported to OMB as part of quarterly reporting obligations?

- ☐ Yes
☒ No
☐ Not applicable

2. How will you collect the information? (Check all that apply)

- ☐ Social Media
☒ Website
☐ Telephone
☐ In-person
☐ Video conference (e.g., Zoom, WebEx, Teams, etc.)
☐ Mail
☐ Other, Explain

3. If this is a survey OR a screener, what platform will be used (e.g., Medallia, Microsoft Forms, Qualtrics, SurveyMonkey, Touchpoints, etc.) *Qualtrics*

4. Who will you collect the information from?

Please describe exactly how you will select the people who you will invite to take the screener or survey, or to participate in the session. Some examples:

- If you are requesting approval for a survey that will appear as a “feedback button” on a website, you can write: “100% of people who visit the website will have the opportunity to take the survey, since the invitation appears on each page of the website.”
- If you are requesting approval for a focus group to conduct discovery phase research, you can write: “of the people who provided their email address to the call center rep, 15% of them will be selected at random to be invited to take part in the focus group.”

These are just examples. The key is you must describe the exact methodology you used to determine who gets the invitation to take part in the screener/survey/session.

All users who access Medicaid.gov will have the opportunity to provide feedback. Typically, these individuals self-identify as 'consumers' of health care services or policy professionals who use the extensive data on Medicaid.gov to support their work. The feedback tab will always be available to leave feedback. No personal information will be requested or collected.

5. Please describe the activity or methodology

Describe the information collection activity – e.g., what happens when a person agrees to participate? Will facilitators or interviewers be used? What's the format of the interview/focus group?

This is a feedback tab on Medicaid.gov without any direct supervision or external involvement. If a site user would like to leave feedback they can click on the tab.

- ***How will you ask a respondent to provide information?***

(e.g., after an application is submitted online, the final screen will present the opportunity to provide feedback by presenting a link to a feedback form / an actual feedback form)

Survey questions will mostly be multiple choice however, depending on the respondents' choices, 2 free-text opportunities will be available for expanded feedback.

- ***When will the activity happen?***

Describe the time frame or number of events that will occur (e.g., "We will conduct focus groups on May 13, 14, 15 of XXXX (year)"; "We plan to conduct customer intercept interviews over the course of the summer of XXXX (year) at the field offices identified in response to #2 based on scheduling logistics concluding by Sept. 10th", or "This survey will remain on our website in alignment with the timing of the overall clearance." If you are uncertain as to how long it will take to complete your research, you can write: "Until all participants complete/are interviewed.")

This revised survey will go live September 2026 and will be available as a clickable tab to leave feedback if desired.

6. Will you be compensating participants?

It may be appropriate to compensate participants for reasonable out-of-pocket costs and to recruit those who are hard to reach, such as those with multiple jobs or are caretakers. It is not appropriate to compensate those participating in their work capacity and those who are vulnerable to coercion or undue influence such as children and people with certain mental disabilities.

☐ Yes

☒ No

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If yes, please describe: (1) the dollar amount and the type of compensation (e.g., gift card) per participant; (2) the reason you are compensating; and (3) why you believe no one in your sample is vulnerable to coercion or undue influence.

PERSONALLY IDENTIFIABLE INFORMATION

Please complete the following questions in coordination with your agency's privacy office to ensure that the information collection complies with any applicable requirements under the Privacy Act of 1974 (5 U.S.C. § 552a) and other laws and policies:

1. Could any of the questions offer the opportunity for respondents to provide personally identifiable information (PII)?¹

☐ Yes

☒ No

2. If the answer to Question 1 is "Yes," will the information collected be covered by the Privacy Act of 1974?

☐ Yes

☐ No

3. If the answer to Question 2 is "Yes," what is the status of the applicable system of records notice(s) (SORNs)?

☐ Published (include citation here) and no modification needed. Include citation:

☐ Published (include citation here), but modification needed. Include citation:

☐ No SORN exists; need to publish new SORN(s)

4. If a privacy impact assessment (PIA) is required, please provide a link to where the PIA is published on the agency's website:

BURDEN HOURS

Active

We estimate that there are two primary types of respondents, health care 'consumers' and policy professionals.

Category of Respondent	No. of Respondents	Participation Time	Burden Hours
Beneficiaries	700	1 minute	11.6
Grantees, States, Outreach	600	1 minute	10.0
Total	1,300		21.6

No Burden Change

¹ PII could include, but is not limited to, name, email address, phone number, or any other information that could be used to distinguish or trace an individual's identity, either alone or when combined with other information that is linked or linkable to a specific individual.

IF YOU SELECTED “NEW COLLECTION” IN RESPONSE TO “TYPE OF SUBMISSION” ON PAGE 1:

Please fill in each applicable cell in the table below.

IF YOU SELECTED “CHANGE TO AN ALREADY APPROVED COLLECTION” IN RESPONSE TO “TYPE OF SUBMISSION” ON PAGE 1:

First, you must figure out whether your change will affect the number of respondents per year or the participation time for the instrument.

- If the number of respondents and the participation time are both staying the same: input “1” for “No. of Respondents per year,” input 60 for “Participation Time in minutes,” and input “1” for “Total Burden per year in hrs.” This is because “1 hour” is the minimum that can be inputted into ROCIS for total burden hours.
- If the number of respondents or the participation time is changing: Contact the CX Desk Officer before completing this table.

Type of Instrument	No. of Respondents per year	Participation Time in minutes	Total Burden per year in hrs
Screeners (e.g., distributed before or during a usability testing session or other kind of session)			
Question script for focus group, interview group, etc. See discussion of usability testing on page 1.			
Survey to obtain feedback immediately following a transaction			
Other survey	1	60	1
Total	1	60	1

CERTIFICATION: Please read the certification carefully. If you incorrectly certify, the collection will be returned as improperly submitted or it will be disapproved.

I certify the following to be true:

1. The collections are voluntary;
2. The collections are low-burden for respondents (based on considerations of total burden hours or burden-hours per respondent) and are low-cost for both the respondents and the Federal Government;
3. The collections are non-controversial, meaning they do not raise issues that warrant public comment;
4. Any collection is targeted to the solicitation of opinions from respondents who have experience with the program or may have experience with the program in the near future;
5. Personally identifiable information (PII) is collected only to the extent necessary and the agency will comply with applicable legal and policy requirements to ensure its protection;
6. Information gathered is intended to be used for general service improvement and program management purposes;

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7. The agency will follow the procedures specified in any relevant OMB guidance for the required reporting to OMB of data from surveys;
8. Outside of the reporting mentioned in the bullet immediately above, if the agency intends to release journey maps, user personas, reports, or other data-related summaries stemming from this collection, the agency must include appropriate caveats around those summaries, noting that conclusions should not be generalized beyond the sample, considering the sample size and response rates. The agency must submit the data summary itself (e.g., the report) and the caveat language mentioned above to OMB before it releases them outside the agency. OMB will engage in a passback process with the agency.

Name of Person(s) who developed the screener/question script/survey: *Jayne Burleson*

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