

**Request for Approval under the “Generic Clearance for the Collection of Routine Customer Feedback” (OMB#: 0925-0648 Exp., date: 05/2021)**

---

**TITLE OF INFORMATION COLLECTION:** Customer Satisfaction Survey for the Innovative Clinical Trials Resource (ICTR) Website.

**PURPOSE:** The purpose of this customer satisfaction survey is to collect feedback on the public facing Innovative Clinical Trials Resource (ICTR) Website. The National Heart, Lung, and Blood Institute (NHLBI) funded the ICTR program to: 1) provide infrastructure and expertise to support awardees of the NHLBI “Catalyzing Innovation in Late Phase Clinical Trial Design and Statistical Analysis Plans Initiative funding opportunity announcement; and 2) develop and provide an educational program on the application of non-traditional clinical trial design and analysis for the NHLBI research community. The ICTR program website houses educational material on non-traditional clinical trial designs; resources for the scientific research community; and programmatic announcements and updates. This survey will solicit feedback on the quality and usefulness of the materials and resources developed by the program.

**DESCRIPTION OF RESPONDENTS:**

We anticipate that respondents will primarily be Physician Scientists from the Heart, Lung, Blood, and Sleep scientific research community

**TYPE OF COLLECTION:** (Check one)

- |   |  |
|---|--|
| <input type="checkbox"/> Customer Comment Card/Complaint Form         | <input checked="" type="checkbox"/> Customer Satisfaction Survey |
| <input type="checkbox"/> Usability Testing (e.g., Website or Software | <input type="checkbox"/> Small Discussion Group                  |
| <input type="checkbox"/> Focus Group                                  | <input type="checkbox"/> Other: _____                            |

**CERTIFICATION:**

I certify the following to be true:

1. The collection is voluntary.
2. The collection is low-burden for respondents and low-cost for the Federal Government.
3. The collection is non-controversial and does not raise issues of concern to other federal agencies.
4. The results are not intended to be disseminated to the public.
5. Information gathered will not be used for the purpose of substantially informing influential policy decisions.
6. The 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 future.

Name: Erin E Smith

To assist review, please provide answers to the following question:

**Personally Identifiable Information:**

1. Is personally identifiable information (PII) collected?  Yes  No
2. If Yes, is the information that will be collected included in records that are subject to the Privacy Act of 1974?  Yes  No
3. If Applicable, has a System or Records Notice been published?  Yes  No

**Gifts or Payments:**

Is an incentive (e.g., money or reimbursement of expenses, token of appreciation) provided to participants? [ ] Yes [ X] No

**ESTIMATED BURDEN HOURS and COSTS**

Category of Respondent	No. of Respondents	No. of Responses per Respondent	Time per Response (in hours)	Total Burden Hours
Physician Scientists	50	1	5/60	4
<b>Totals</b>	<b>50</b>	50		4

Category of Respondent	Total Burden Hours	Hourly Wage Rate*	Total Burden Cost
Physician Scientists	4 hours	99.00	396.00
<b>Totals</b>			396.00

[https://www.bls.gov/oes/current/oes\\_nat.htm#19-0000](https://www.bls.gov/oes/current/oes_nat.htm#19-0000)

**FEDERAL COST:** The estimated annual cost to the Federal government is \$545.07

Staff	Grade/Step	Salary*	% of Effort	Fringe (if applicable)	Total Cost to Gov't
<b>Federal Oversight</b>					
Contracting Office Representative (COR)	14/5	129,869	0.002		259.70
<b>Contractor Cost</b>		57.07 (per hour)	5 hours		285.35
Travel					
Other Cost					
<b>Total</b>					545.05

\*the Salary in table above is cited from <https://www.opm.gov/policy-data-oversight/pay-leave/salaries-wages/salary-tables/pdf/2018/DCB.pdf>

**If you are conducting a focus group, survey, or plan to employ statistical methods, please provide answers to the following questions:**

**The selection of your targeted respondents**

1. Do you have a customer list or something similar that defines the universe of potential respondents and do you have a sampling plan for selecting from this universe?  
 Yes  No

If the answer is yes, please provide a description of both below (or attach the sampling plan)? If the answer is no, please provide a description of how you plan to identify your potential group of respondents and how you will select them?

We anticipate that respondents will primarily be Physician Scientists from the Heart, Lung, Blood, and Sleep scientific research community.

**Administration of the Instrument**

1. How will you collect the information? (Check all that apply)  
 Web-based or other forms of Social Media  
 Telephone  
 In-person  
 Mail  
 Other, Explain
2. Will interviewers or facilitators be used?  Yes  No