PAPERWORK REDUCTION ACT SUBMISSION WORKSHEET Part I: Information Collection Request

This template is intended for staff without an ICRAS account. Please fill out and submit to the appropriate Operating Division to enter into ICRAS. The form mirrors the screens available in the ICRAS 4 system. To request an account to log into ICRAS.

Instructions for filling out the form are available at <u>www.paperworkreduction.gov</u>.

Γ

1. Agency/Subagency originating request: Centers for Medicare & Medicaid Services		
2. Title: Electronic Health Record Demonstration		
 3. Type of information collection <i>(check one)</i> (See instructions) New collection (Request for a new OMB Control Number) Revision without change of a currently approved collection Revision of a currently approved collection Reinstatement without change of a previously approved collection Reinstatement with change of a previously approved collection Reinstatement with change of a previously approved collection Reinstatement with change of a previously approved collection Reinstatement or nonsubstantive change to a currently approved collection (formerly 83C) Existing collection in use without and OMB Control Number 	4. OCN: <u>0938-0965</u>	
 5. Type of review requested (check one) a. <u>X</u> Regular b. <u>Emergency</u> - Approval requested by: <u>//</u> c. <u>Delegated</u> 	 6. Requested expiration date (check one) a. X. Three years from approval date b. Six Months from approval date (Maximum for Emergency reviews) 	
<i>If Emergency, please attach justification.(</i> 4000 characters maximum)	c Other Specify:/ (mm/yy) or Number of Months from Approval Date:	
 7. Abstract (4000 characters maximum, attach additional sheets as necessary) The Centers for Medicare & Medicaid Services (CMS) requests clearance for the application utilized to identify and enroll practices into the Electronic Health Record demonstration. This demonstration is a high-priority Administration of this initiative. The purpose of this demonstration project is to reward the delivery of high-quality care supported by the adoption and use of electronic health records in small to medium-sized primary care physician practices. While this initiative is separate and distinct from the Medicare Care Management Performance (MCMP) Demonstration, it expands upon the foundation created by the MCMP Demonstration, which was mandated by Section 649 of the Medicare Prescription Drug, Improvement and Modernization Act of 2003. The electronic health record demonstration will be operated under section 402 demonstration waiver authority. The information to be obtained as part of the application form is necessary to document basic information for physician practices that intend to participate in this demonstration initiative. This information will be used to establish that the practices meet basic eligibility requirements for participation in this initiative. The application form requests basic physician office information (e.g., number of physicians in the practice, specialties, organizational structure, Medicare Provider Identification Number, tax identification number, contact information, etc. Practices that apply will be expected to sign a data sharing consent form that will accompany the form. It is expected that up to 2,400 application forms will be submitted by physician practices in 12 states for subsequent randomization to treatment and control groups. 		
8. Authorizing Statute(s)		
Public Law:		

US Code:			
Title	Section		Name
Executive Order: Number			Name
			hante
Statute:			
Title			Subtitle
	9. Associated Rulemaking Information Stage of Rulemaking (check one) Federal Register Citation RIN: Page number a. Proposed Rule Volume Page number		
	b Interim Final		
For a Proposed Rule, OMB will not consider a			
For a Final Rule, please put the ICR reference For ICRs associated with Interim Final or Fin			sed rule stage in Box 4.
10. Federal Register Notices & Comments			
Federal Register Citation			
	Page number		Publication Date//
30-day Notice: Volume	Page number		Publication Date//
Did the Agency receive public comments on	this ICP2 Yes No		
Unless submitted as an Emergency or Assoc		OMB will not conside	er an ICR complete until the 30-day notice
has been published. 12866, please attach a draft of the Federal I	Register document.		
11. Annual Cost to Federal Gov:		14. Agency contact:	
\$ approximately 312,000		Name: <u>Debb</u> Phone: <u>410-</u>	<u>786-6625</u>
(one-time only cost)		E-mail: <u>Debbi</u>	e.Vanhoven@cms.hhs.gov
12. Does this ICR contain surveys, censuses	, or employ statistical	confirms	
methods?			
Yes (Attach Part B of Supporting Si 13. Is the Supporting Statement intended to		{	
Assessment required by the E-Government			
YesX_No		l	

PAPERWORK REDUCTION ACT SUBMISSION WORKSHEET Part I: Information Collection Request (continued) Information Collection Budget (ICB)

If a change in burden is due to a Program Change Due to New Statute, identify the Citations for New Statutory Requirements:

Public Law:

Congress Number	Sequence Number	Section	Name

US Code:

Title	Section	Name

Executive Order:

Number	Name

Statute:

Title	Subtitle

If Program Change is due to Agency Discretion, please categorize the reduction. Burden reduction from (select one):

- a. ___ Cutting Redundancy
- b. ___ Using Information Technology
- c. ___ Changing Regulations
- d. ___ Changing Forms
- e. X Miscellaneous Actions

If Program Change is due to Agency Discretion, please categorize the increase in burden. Burden increase caused by (select one):

- a. ___ Changing Regulations
- b. <u>X</u> Miscellaneous Actions

Explain the reasons for any program changes or adjustments reported; that is, provide a short statement how the reduction in burden was achieved or why the increase in burden occurred. (If you need more space, please provide a short summary here and elaborate in the Supporting Statement.)

This demonstration is a high-priority Administration initiative conducted under section 402 demonstration waiver authority. Only physician practices that respond and express interest in participating will complete the application. The burden associated with the proposed collection of information is completely voluntary; however, it should be noted that physician practices that voluntarily respond may ultimately be eligible to earn substantial financial rewards as part of their subsequent participation in this demonstration initiative.