

challenging research activities. Oversight systems and tools are critical for the NIH to ensure participant safety, data integrity, and accountability of the use of public funds. The NIH has been engaged in a multi-year effort to examine how clinical trials are supported and the level of oversight needed. The collection of more

structured information in the PHS applications and pre-award reporting requirements as well as continued monitoring and update during the post-award reporting requirements will facilitate the NIH's oversight of clinical trials. In addition, some of the data reported in the RPPR will ultimately be accessible to investigators to update

certain sections of forms when registering or reporting their trials with *ClinicalTrials.gov*.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 519,408.

ESTIMATED ANNUALIZED BURDEN HOURS

Information collection forms	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
<b>Reporting:</b>				
PHS 416-7 .....	12,580	1	30/60	6,290
PHS 6031-1 .....	1,778	1	20/60	593
PHS 568 .....	11,180	1	5/60	932
iEdison .....	5,697	1	15/60	1,424
PHS 2271 .....	22,035	1	15/60	5,509
PHS 2590 .....	243	1	18	4,374
RPPR—Core Data .....	32,098	1	8	256,784
Biosketch (Part of RPPR) .....	2,544	1	2	5,088
Data Tables (Part of RPPR) .....	758	1	4	3,032
Trainee Diversity Report (Part of RPPR) .....	480	1	15/60	120
PHS Human Subjects and Clinical Trial Information (Part of RPPR, includes inclusion enrollment report) .....	6,420	1	4	25,680
Publication Reporting .....	97,023	3	5/60	8,085
Final RPPR—Core Data .....	18,000	1	10	180,000
Data Tables (Part of Final RPPR) .....	758	1	4	3,032
Trainee Diversity Report (Part of Final RPPR) .....	480	1	15/60	120
PHS Human Subjects and Clinical Trial Information (Part of Final RPPR, includes inclusion/enrollment) .....	3,600	1	4	14,400
PHS 3734 .....	479	1	30/60	240
Final Progress Report .....	2,000	1	1	2,000
SBIR/STTR Phase II Final Progress Report .....	1,330	1	1	1,330
<b>Reporting Burden Total</b> .....				<b>499,033</b>
<b>Recordkeeping:</b>				
SBIR/STTR Life Cycle Certification .....	1,500	1	15/60	375
<b>Grand Total</b> .....				<b>519,408</b>

Dated: February 1, 2017.

**Lawrence A. Tabak,**  
Deputy Director, National Institutes of Health.  
[FR Doc. 2017-02471 Filed 2-6-17; 8:45 am]

**BILLING CODE 4140-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Institute of Mental Health; Amended Notice of Meeting**

Notice is hereby given of a change in the meeting of the National Institute of Mental Health Special Emphasis Panel, February 03, 2017, 10:00 a.m. to February 03, 2017, 03:00 p.m., National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD, 20852 which was published in the **Federal Register** on January 18, 2017, 82FR5588.

This meeting notice is amended to change the meeting time to 1:00 p.m.–5:00 p.m. The meeting is closed to the public.

Dated: February 1, 2017.

**Melanie J. Pantoja,**  
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017-02430 Filed 2-6-17; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Submission for OMB Review; 30-Day Comment Request PHS Applications and Pre-Award Reporting Requirements (OD/OPERA)**

**AGENCY:** National Institutes of Health, HHS.

**ACTION:** Notice.

**SUMMARY:** In compliance with the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on November 2, 2016, Volume 81, No. 212, pages 76368–76370 and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment.

**DATES:** Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

**ADDRESSES:** Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden