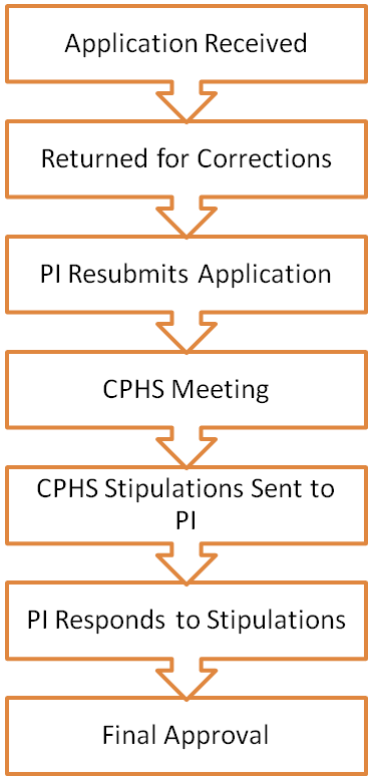
TIME TO APPROVAL—FULL BOARD ONLY



Duration 1 – Median time in days between the date the IRB office receives the application and the date the IRB office sends notification to the PI requesting changes.

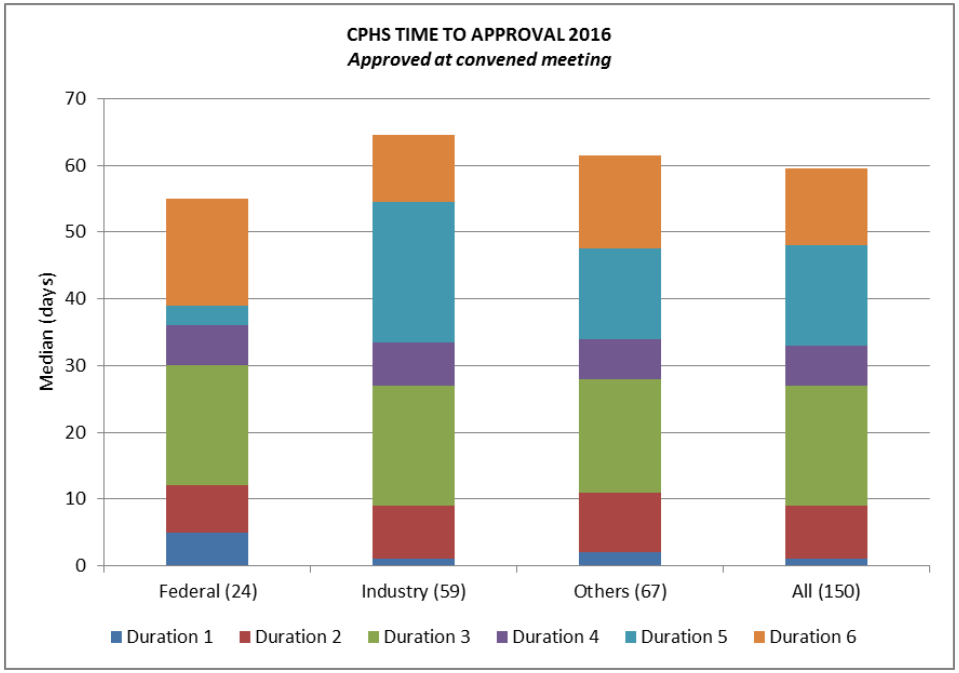
Duration 2 - Median time in days between the date the IRB office returns the application for corrections to the PI and the date the PI re-submits a corrected application.

Duration 3 - Median time in days between the date the PI re-submits the application and the date the protocol is reviewed by the fully convened IRB.

Duration 4 - Median time in days between the IRB meeting date and the date the IRB sends stipulations to the PI.

Duration 5 - Median time in days between the date the IRB sends stipulations to the PI and the date that the PI submits responses to the stipulations.

Duration 6 - Median time in days between the date that a response to the stipulations is received by the IRB office and the date final approval is granted by the IRB with no contingencies remaining.



Acknowledgements: Thanks to Chunyan Cai, PhD, Assistant Professor, Internal Medicine—CCTS for data analysis.

UTHealth
The University of Texas Health Science Center at Houston

REPORT TO FACULTY AND STAFF ON CPHS ACTIVITIES

-2016-

from

Anne Dougherty, MD
Vice President, Human Research Protection Program

Panel 1
Chair: Rebecca Lunstroth, JD
Vice Chair: Kathleen Kennedy, MD
Coordinator: Stephanie Francisco, BA

Panel 2
Chair: Ben Barnett, MD
Vice Chair: George Delclos, PhD
Sr. Coordinator: Audrey Williams, PhD

Panel 3
Chair: Charles Miller, PhD
Vice Chair: Rita Swinford, MD
Vice Chair: Cathy Thompson, RN
Coordinator: Vanessa Fuller, BS

Panel 4
Chair: Max Buja, MD
Vice Chair: Ralph Frankowski, PhD
Coordinator: Cristal Casanova, BS

IRB Support Staff
Director: Cynthia Edmonds, MLA
Sr. IRB Coordinator: Sylvia Romo, BSBM
Sr. Systems Analyst: Barbara Legate, BS
IRB Assistant: Olufemi Popoola, MPH
Email: cphs@uth.tmc.edu
Website: www.uth.edu/cphs

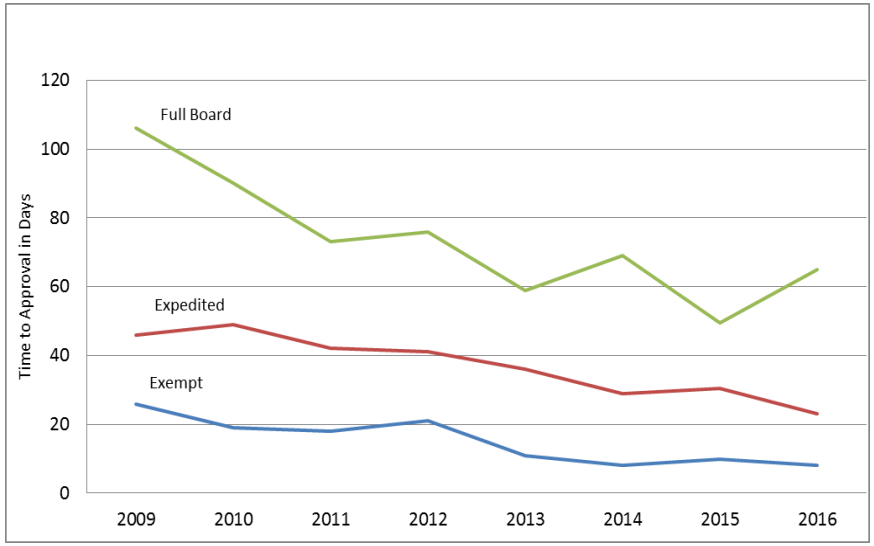
Research Compliance
Director: Sujatha Sridhar, MBBS, MCE
Sr. Compliance Specialist: Carolyn McKinney, RN
Sr. Compliance Specialist: Elizabeth Gendel, PhD
Graduate Assistant: Noopur Singh, BSE
Graduate Assistant: Chaitra Muthalgiri, MBBS
Email: clinicaltrials@uth.tmc.edu
Website: www.uth.edu/ctrc

CPHS Office
6410 Fannin Street, Suite 1100, Houston, Texas 77030
Phone: 713.500.7943

Association for the Accreditation of Human Research Protection Programs, Inc. — Full Accreditation

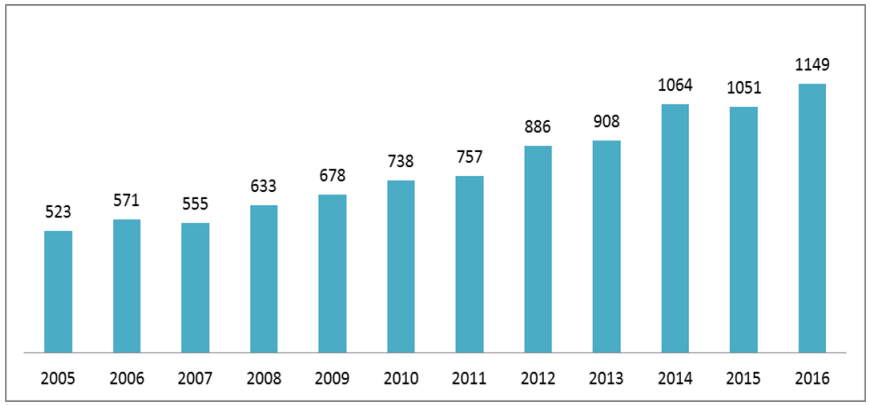
TIME TO APPROVAL

The median turnaround time (which is the time between initial submission of the protocol and final approval) has steadily decreased. This includes the time that the protocol was on the researcher's queue to address pre-screening concerns, such as missing documents and post review stipulations. requests.

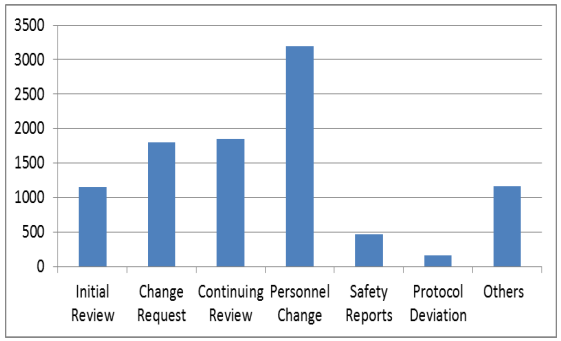


NEW APPLICATIONS TO CPHS

The number of initial applications to CPHS has been increasing since UT Houston began using iRIS. From just over 500 new applications in 2005, CPHS received 1,149 initial applications in 2016 for review and approval.



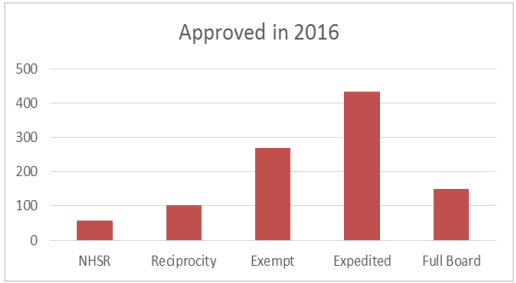
2016 SUBMISSIONS



In 2016, CPHS reviewed and processed 12,229 submissions. The safety reports include reportable adverse events as well as data safety monitoring board reports.

2016 APPROVALS

In 2016, CPHS approved 1010 new submissions of which only 150 were reviewed at a convened full board meeting. CPHS made 57 submissions not human subjects research determinations.



CPHS FACULTY SURVEY

Responses to the CPHS Faculty Survey, including free text responses, are shared with the CPHS Executive Committee each quarter. The responses are helpful in continuous quality improvement of CPHS processes.