SONA INSTRUCTIONS FOR PRINCIPAL INVESTIGATORS AND RESEARCHERS

As you know, we are now using Sona for the scheduling and management of our subject pool. As a PI or researcher, you can set up your studies in the system, schedule the sessions (timeslots) when students may participate, and grant or deny credit after the session. All of this is handled through a simple web-based interface that you can access at any time, from any popular web browser, at this web address:

http://iupresearchparticipation.sona-systems.com

With this system, you will no longer request participants from the subject pool. Instead, you will post your study and timeslots on the website. Students will visit the website and sign up for appointments.

Each study will have both a principal investigator and a researcher. The principal investigator will always be the faculty member who is overseeing the project. Each faculty member will have a principal investigator account and receive a user ID and password to access the system. The researcher is the person who is actually running the study. Whether the students working with you should have their own researcher account is up to your discretion. It is up to faculty members to request researcher accounts for their students (to do so, email Anson.Long@iup.edu). Note that, at least for now, 290/291 students will not be permitted to have researcher accounts. Faculty members who plan to run their own studies, or who plan to post studies and timeslots and grant/deny credit on behalf of student research assistants, will select their own name as both the PI and the Researcher for their study, as both the PI and researcher fields will need to be filled in for every study.

When posting a new study on the website, you will be able to provide a title and a brief description. These may be descriptive, at your discretion. However, please make sure that they do not contain any coercive or manipulative language.

You will also have the option to specify who the study should be visible to (perhaps only males or only females, depending on your study design; note that you must have IRB approval for any inclusion/exclusion criteria).

You will also be asked to provide the IRB number and expiration date when posting a new study.

Once you have posted your study, you can post timeslots, and start watching participants fill them!

After the session is over, you should update credit right away. You will grant credit to those who participated, and indicate “Unexcused No Show” for anyone who failed to keep their appointment. (Upon accumulating two no-shows, a participant’s account will be Limited, so that they can sign up only for Read and Reviews).

Note that participants may cancel their appointment on Sona up to 24 hours beforehand. If something comes up after that, they may email you so that you can cancel their appointment for them.

The system is pretty easy and intuitive to use. However, the following pages provide additional detail about the many features of Sona. I think you’ll be impressed with all it can do.

If you have any questions about the system, please email Anson.Long@iup.edu.

DETAILED INSTRUCTIONS FOR PIs AND RESEARCHERS

Introduction

The Experiment Management System is used for the scheduling and management of research participants and the studies they participate in. Participants, researchers, principal investigators, and instructors all use the system for their respective purposes. As a researcher, you can set up your studies in the system, schedule the sessions (timeslots) when participants may participate, and grant or revoke credit after the session. All of this is handled through a simple web-based interface that you can access at any time, from any popular web browser.

System Basics

In the system, you create studies. Each study may have a number of timeslots, which are the times when you plan to run the study. Participants sign up for the timeslots by viewing a list of studies and available timeslots. You grant or revoke credit to participants after the session occurs.

Principal Investigator Special Note

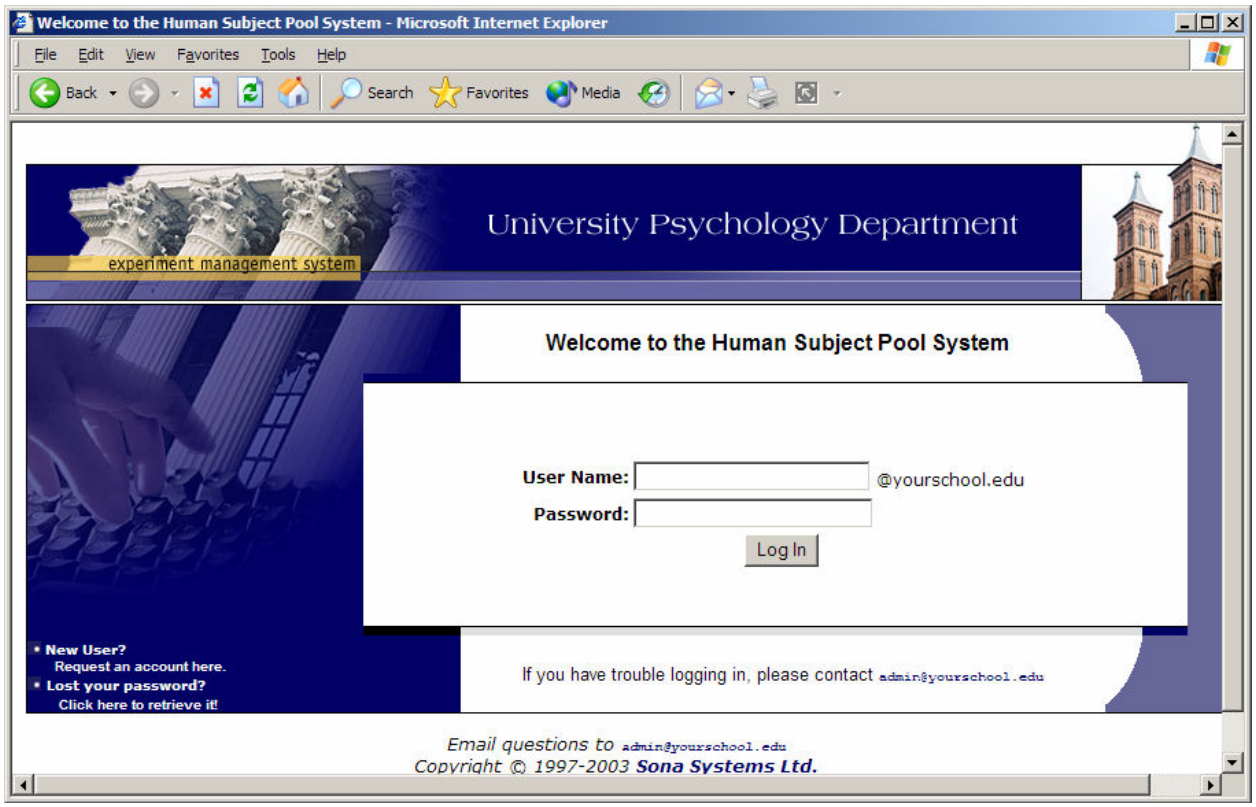
This documentation applies to both researchers and principal investigators. A P.I. can perform all the same functions on a study as a researcher. This allows a P.I. to operate in an oversight role and monitor the progress of their studies, and step in on behalf of the researcher when necessary. Because the privileges are the same, throughout this documentation, the term "researcher" can be used interchangeably with "principal investigator" except where otherwise noted. All studies must have a P.I. specified.

Getting Started

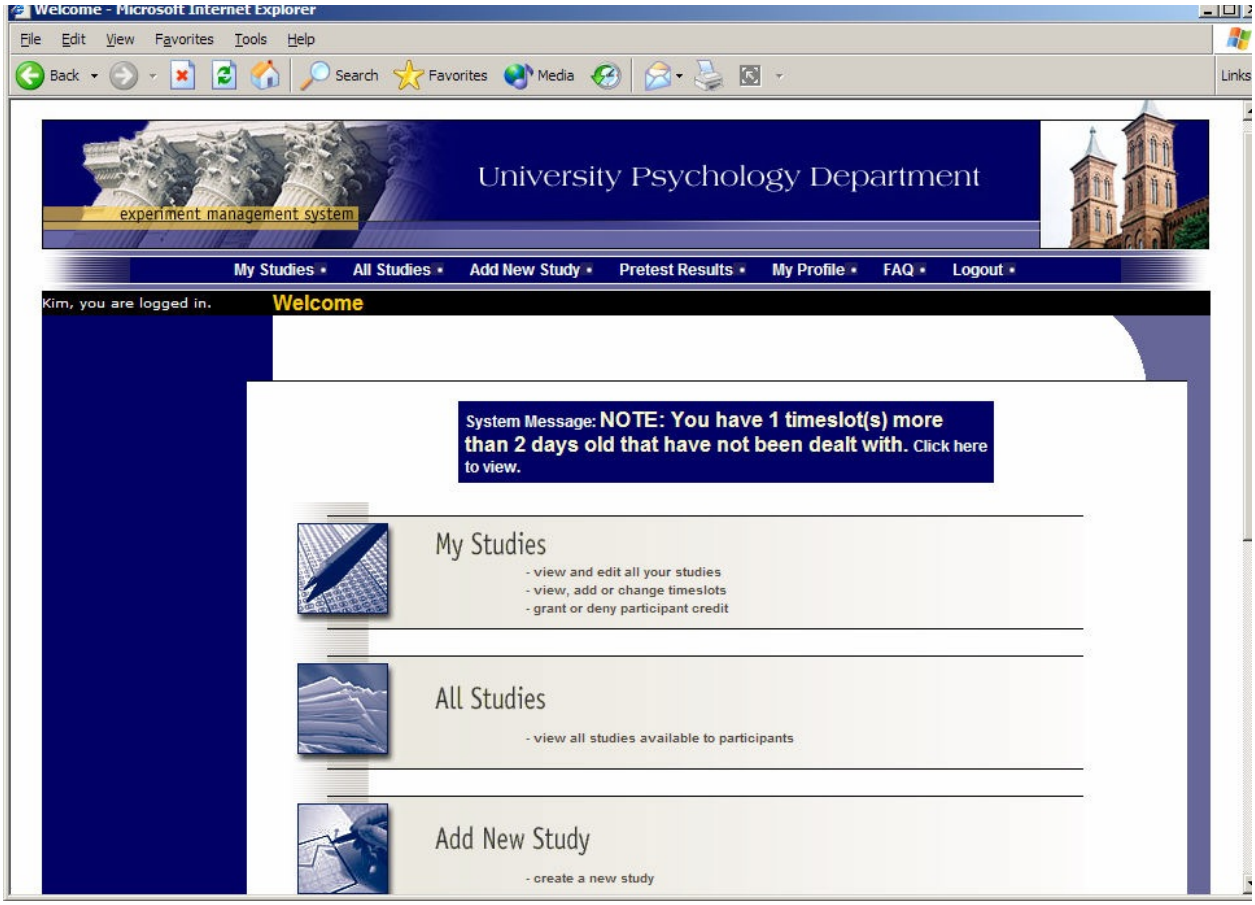
The system works best if you use any popular web browser that is less than 2 years old, like Internet Explorer, Firefox, and Safari. It will work with other web browsers, and with older versions of popular web browsers, however the layout may not be as clean. No functionality will be lost by using an older web browser. Ask your system administrator if you need help with installing or using a web browser. This documentation assumes you have a basic knowledge of how to use the web. On this system, it is not necessary to use the Back button. You can always use the toolbar on the top to navigate to anywhere on the site.

Logging In

Your administrator will provide you with a username and password to log in to the site, as well as the URL (web address).



Once you login, you will be asked to review and acknowledge your organization's human subject policy. If required by the administrator, you will need to acknowledge this once every 6 months. You will see the Main Menu after you acknowledge the policy.



Your login (also known as a session) will expire after a certain period of inactivity, usually 20 minutes. This is done for security purposes. If this happens, you can always log in again. When you are done using the system, it is better to explicitly log out, to prevent any problems that may arise if someone uses your computer before the session expires.

Retrieving a Lost Password

If you have forgotten or do not have your password, and the feature is enabled on the system, then you may choose to have your password emailed to you. You will see an option on the main login page if this feature is enabled. Your password will be emailed after you submit the form, and should arrive in your email box momentarily. If you provided an alternate email address (see the Email Address Options section of this documentation), it will be sent there. Otherwise, it will be sent to your main email address, which is derived from your user ID. If you requested that the system email a password to you, and it has not arrived after 30 minutes, then check in your email program's junk mail folder in case the email was delivered there.

Logging Out

When you are done using the system, choose Logout from the top toolbar to log out. You are now logged out. It is always a good security practice to close all your web browser windows as well, especially if you are using a computer that is shared by others.

Changing Your Password and Other Information

If you would like to change your password or other information about yourself, choose My Profile from the top toolbar. If you would like to change your password, type your new password (twice, for confirmation) in the provided boxes. If you would *not* like to change your password, simply leave these boxes empty.

If you change your password, please be sure to select a password you do not use on any other systems or websites. This is good computing practice, and especially important as in some cases, your password may be sent over email.

It is recommended you provide your phone number and office location, as most human subject committees require that this information be made available to research participants. If you are a researcher, this contact information will be displayed to participants when they view information about the study. If you are a principal investigator, this contact information will be displayed if a participant explicitly chooses to view it (since the researcher is the primary point of contact for a study).

Researchers may also choose to receive a daily reminder (by email) with information about all of their study sessions scheduled for the following day.

Email Address Options

There are certain events in the system which will cause an email notification to be sent to you. Most often, these are notifications that a participant has signed up or cancelled their sign-up for your studies, but there are a few other cases where it may be used as well. The email address is also displayed to the participant when they view information about the study, in case they need to contact you with questions.

You have two choices for your email address. When you update your personal information, you will see a box where you may provide an alternate email address. If you provide such an address (this could be a Hotmail account, for instance), this is the address where any notifications will be sent, and this is also the address that will be displayed to other users (including participants in your studies).

If you do not supply an alternate email address, the system will derive your email address from your username. Typically, it will add your organization's Internet domain to the end of your user ID to form the address, so if your user ID is "jsmith" and your organization's Internet domain is "yourschool.edu" then it would derive your email address as "jsmith@yourschool.edu".

In situations where the system is configured so you may enter an email address on this page, you will be asked to enter it twice when changing the address, to ensure it is typed correctly.

Working with Studies

Most of your time on the system will be spent, not surprisingly, using the study-related features of the system. Be sure to read this section closely, in its entirety, as there are special features and situations you should be aware of.

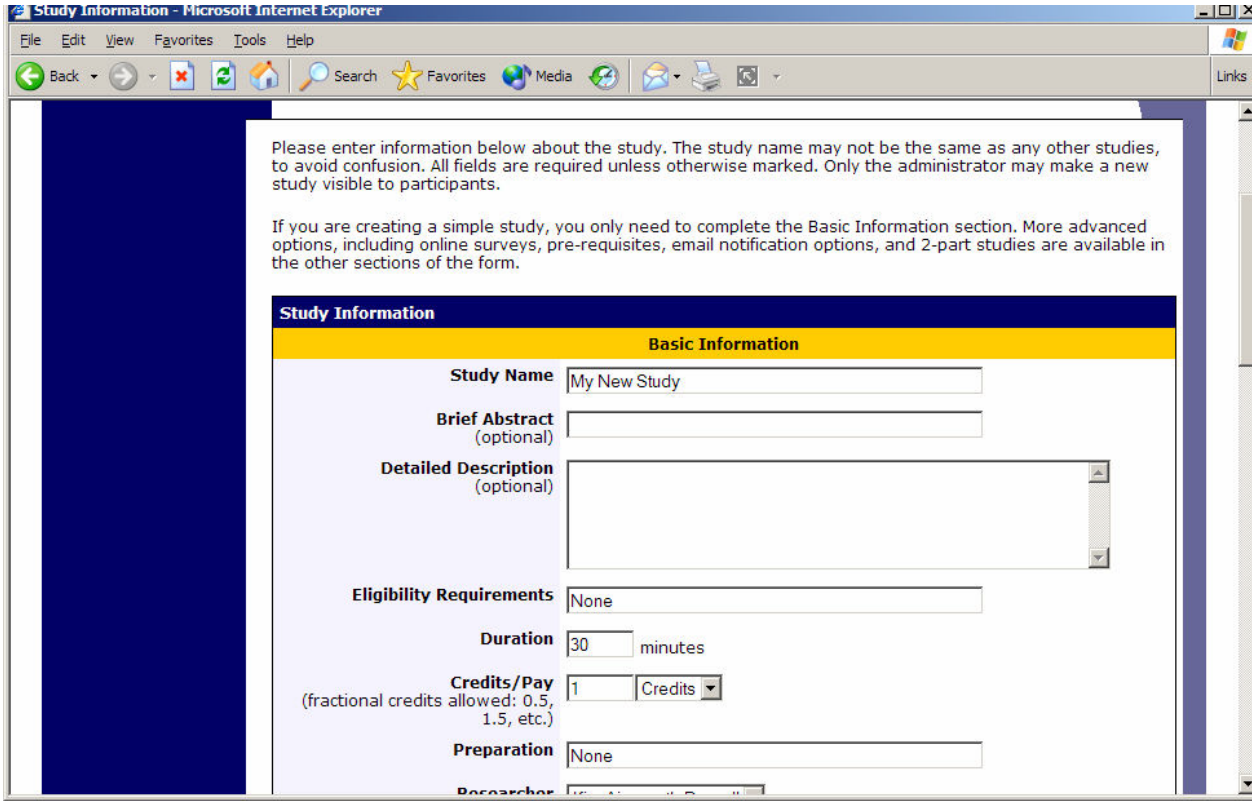
Most studies will involve participants meeting in person with a researcher for just one session and in exchange for credit toward their PSYC 101 research requirement.

However, there are some other situations that may arise:

- Online Studies: See separate documentation.
- Studies for Pay: You may have a situation where participants are compensated for their participation in the study. They may or may not also receive credit for the study. If the study is not for credit, you may set it up as a pay-only study and specify the compensation amount. If participants are compensated *and* they receive credit, you should set it up as a credit study and indicate additional compensation in the study's information section. Regardless of the type of study, after a participant participates in a study (including studies that are for pay only), you should still go into the system and indicate their participation by noting their participation or no-show when viewing their sign-up. This allows the system to properly enforce certain restrictions on the participant and their studies, like pre-requisite and disqualifier study restrictions.
- Two-Part Studies: You may create a two-part study in the system. Often, these are studies involving memory research, where the participant must return a specified number of days after the first session. When creating a study, you may specify the day range for the second part of the study (e.g. 7 to 10 days after the first part). Participants are required to sign up for both sessions at the same time, to reduce the chance they will forget to sign up for the second part. Each part of a two-part study may have a different credit value and duration, but each part must be the same type – either both parts are for credit or both parts are for compensation. Online studies may not be two-part studies. You may specify that the second part of the study must be scheduled to take place at the exact same time as the first part (on a different date), or at any time on the dates that are the specified number of days after the first part. You should ensure there are enough available timeslots for both parts of the study, or participants will be prevented from signing up for either part. Participants may cancel either part of their sign-up if necessary. If they cancel the first part, the second part is automatically cancelled as well. If they cancel only the second part and the first part has already occurred, and they would like to participate in the second part later, you will need to manually sign them up for the second part (if you are allowed to do so), or ask the administrator to handle this. If you grant a no-show for the first part of a two-part study, the second part of that participant's sign-up will *not* be cancelled automatically, but you will be reminded of the situation in case you would like to cancel the second part. The cancellation is not automatic as there are some situations where automatic cancellation is not desirable.

Adding a Study

Some researchers choose to set up their studies in the system before they have received the proper approvals (usually from their IRB) to run the study. This is supported in the system. You can set up a study but specify that is it not visible to participants (this is the approved setting). That way, as soon as your approval is received, you can simply make the study visible and everything else is already prepared. You can also post a study and make it visible immediately, if you already have IRB approval.



To add a study, choose the Add New Study option from the top toolbar. You will need to pick from four possible types of studies. Please choose this carefully as you are not able to change this later. After you choose the study type, you'll see a form asking for more information. You will need to fill out a number of fields, which are explained below. All fields must be filled out unless otherwise noted.

Study Name: A short name for the study. This is how the study is identified throughout the system. Studies will show in a random order to participants, so there is no advantage in choosing a study name that might put it at the top of an alphabetical list.
Study names must be unique, and you will be prevented from adding a study if there is already another study in the system with the same name. A study name may be up to 100 characters in length.

Brief Abstract: This is a short one or two line description of the study. This short description will be displayed to participants when they view the entire list of studies, so

you may want to list the most pertinent details here. Studies configured for payment usually have the compensation information included here, particularly if the payment varies based on certain outcomes. This field may be optional, and can be up to 255 characters in length.

Detailed Description: This can be a rather lengthy description about the study, and it will show if a participant clicks on the study to get more information, before they sign up. You may include basic HTML in this area, but please be sure you know what you are doing (ask your IT department for help if you are unsure). If you would like to add a carriage-return (paragraph break), simply type in "<p>" (without the quotes).This field may be optional. The maximum length of this field is 15,000 characters.

Eligibility Requirements: If there are any restrictions on who may participate (for instance, only those who are left-handed), list them here. Otherwise, leave the field as-is. If you list any restrictions, these will be displayed on the list of studies, when participants view a list of all available studies. Note the system does not enforce these restrictions, but it is expected a participant will only sign up for a study in which they are qualified, since they would otherwise fail to receive credit. In most cases, you will leave this field as-is and set prescreen participation restrictions, which you can do after you add the study. This field may be up to 245 characters in length.

Pre-Requisites: If there are studies a participant must participate in before participating in your study, choose them here. You may select multiple studies, and on most systems, you hold down the Ctrl key and click the desired studies. You may specify that participants must have participated in *all* of the studies you specify, or *at least one* of the studies specified. The system will handle enforcement of the pre-requisites in a strict or lenient fashion depending on how your system is configured. In strict enforcement mode, the participant must have *received credit* for (participated in) the pre-requisite studies. In lenient enforcement mode, the participant must only be *scheduled* to participate in the pre-requisite studies (it is assumed they will go on to complete the pre-requisite studies). You can ask your administrator how this is configured, if it is of concern. If your system is in lenient enforcement mode, and a participant cancels a necessary pre-requisite for you study (they are warned of this situation), and you have configured your study so that the researcher will receive notifications of cancellations or sign-ups, then the researcher will receive notification of the pre-requisite problem and can contact the participant if necessary. If there is a long list of studies for this setting, an Enlarge List button will appear. You can click this to make the list of studies larger and thus easier to click on. Participants will not see which studies you have specified as pre-requisites when they go to view your study.

Disqualifiers: If there are any studies a participant must *not* have participated in, please select them here. You may select multiple studies. The system will handle enforcements of the restriction, during the sign-up process. If a study has some other study listed as a disqualifier, and a participant signs up for this study, then they will be prevented from signing up for the disqualifier study. If there is a long list of studies for this setting, an Enlarge List button will appear. You can click this to make the list of studies larger and thus

easier to click on. Participants will not see which studies you have specified as disqualifiers when they go to view your study.

Course Restrictions: If you would only like participants enrolled in certain courses to participate in your study, select the eligible courses here. Participants who are not in at least one of the courses you selected will not see the study when they go to view the list of available studies. You may choose No Restrictions if you would like to make the study available to participants in all courses. If there is a long list of courses for this setting, an Enlarge List button will appear. You can click this to make the list of courses larger and thus easier to click on. There is a limit to how many courses can be listed as course restrictions for a study, and the limit is somewhere between 60 and 80 courses. The limit is variable depending on a few factors, and the system will simply not save the course restrictions for any courses which would take it over the limit.

Duration: The amount of time, in minutes, that each study session will take. If you are setting up a 2-part study, then this setting applies to the first part of the study. For online studies, this should be an estimate of how long participants can expect the study to take, so that they can plan accordingly.

Preparation: Enter any advanced preparation a participant must do here (e.g. "do not eat 2 hours before session"). If there are no preparation requirements, leave this field as-is.

Invitation Code: If you would like to have a special sign-up password for this study, enter it here. This is known as an invitation code, and applies just for this study. Participants must know the invitation code to sign up for this study. This is often used in cases where the researcher wants to personally select participants, so the researcher only provides the invitation code to the desired participants. Invitation codes are not case sensitive. If you do not need an invitation code, leave this field blank.

Is this a web-based study?: If this is a web-based (online) study, choose the type of online study it is. If you have set up the study on another website, you should note the study is administered outside the system. If you want to set up an online survey study to be administered by the system, select the appropriate option.

Should survey participants be identified only by a random, unique ID code?: This only applies to web-based studies administered by the system. If set to Yes, participants are only identified by a unique system-assigned ID code, to protect their privacy. Participants are also notified of this when they start the survey. Once enabled, this setting cannot be changed after participants have taken the survey, as a matter of privacy protection.

Study URL: The URL (web address, usually starting with http://) for your study. This is only required for web-based studies administered outside the system. If you are setting up a web-based study outside the system, and would like the system to pass a unique identifier in the URL so you may easily identify participants, add the text %SURVEY_CODE% in the URL where you would like the identifier to be placed. This

is discussed in further detail in the Web-Based (Online) Studies section of this documentation.

Credits/Pay: Enter the number of credits or compensation for the study. A value of 0 is acceptable, and may be desired in cases where the study is part of a set of studies, where only the final study is credit-earning. Please see the Studies for Pay section of this documentation for more information on how to fill out this field in the case of for-pay studies. If the study has a credit value, the credit value specified must be evenly divisible by the credit increment specified. For example, if the increment is 0.5, then the study can have credit values like 1 and 1.5, but not 0.75. If you are setting up a 2-part study, this is the value for the first part of the study. After a study has sign-ups, you may not change the credit value of the study. However, the administrator can change the credit value, in certain situations. A study may not be changed between a study for credit and for payment, after it has been created.

Is this a 2-part study?: Select Yes or No if this is a 2-part study. You can only decide this when creating a study (not when editing it), and this setting may not be changed after the study is created. See "Two-Part Studies" for more information.

Credits/Pay, Part 2: Enter the number of credits or compensation for part 2 of the study, if this is a two-part study (the value is ignored otherwise). A value of 0 is acceptable, and may be desired in cases where the study is part of a set of studies, where only the final study is credit-earning. Please see the Studies for Pay section of this documentation for more information on how to fill out this field in the case of for-pay studies. If the study has a credit value, the credit value specified must be evenly divisible by the credit increment specified. For example, if the increment is 0.5, then the study can have credit values like 1 and 1.5, but not 0.75.

Part 2 Duration: The amount of time, in minutes, that part 2 of the study will take.

Part 2 Scheduling Range: Specify the number of days (as a range) after part 1 is scheduled, that part 2 should be scheduled. This setting only applies to two-part studies. The range may be the same value (e.g. "between 7 and 7 days") if desired, but must be a whole number. See "Two-Part Studies" for more information.

Part 2 Scheduling Leniency: In some cases, you may want to ensure that the participant schedules the second part of the study to take place at exactly the same time (on a different date) as the first part. If so, choose Yes for this option. If there is some flexibility so they can sign up for any time within the Part 2 Scheduling range, choose No for this option.

Researcher(s): Select the researcher for this study. Most likely, this is you, and your name will automatically be selected. If you are a researcher, then you may not change who the researcher is (the P.I. for the study, as well as the administrator, can change the researcher). Depending on how your system is configured, you may be able to specify multiple researchers for a study. If you specify multiple researchers, each researcher has full control over the study. The pulldown box lists only users who are researchers.

Principal Investigator: Select the Principal Investigator for this study. The person you select will have full access to the study. If you see this option, then you must select a P.I. The pulldown box lists only users who are principal investigators.

IRB Approval Code: Enter the IRB approval code here. This field is displayed to the administrator to help them keep track of studies. This field may be required depending on how your system is configured.

IRB Approval Expiration Date: The date when IRB approval expires. This field may not appear if your system is not configured for it. If it does appear, you must provide a valid expiration date. The system will prevent you from adding new timeslots to take place after this date, and your study will become inactive (not approved and thus not visible to participants) after this date. You may not make a study active if the IRB approval has expired. Only the administrator can change the IRB approval expiration date, once it has been entered, which is why it defaults to blank to force you to choose a date. You may specify a date up to 5 years in the future.

Approved?: Select Yes if this study should show up on the list of studies which participants may sign up for. Ensure you have received the necessary approvals to run the study before choosing Yes. A study must be Approved and Active to show up on the list of studies which participants may sign up for. If you select No, the study will not be visible to participants. As a researcher, you can always make an approved study invisible to participants (by making it not approved).

Active Study?: Select Yes if this study is in progress. You must select Yes and the study must be Approved if you want the study to show up to participants so they can sign up for it. If a study is Not Approved but *is* Active, then it does not show up (to participants) on the listing of studies, but it is accessible through other links if the participant has participated in it before and they are viewing their participation history (in case the participant has follow-up questions about the study). It will also show up on the study information page (for an individual study) when it is listed as a prerequisite or disqualifier for a study. The reason to select No is if the study is being kept for historical purposes, but should not show up to participants on the list of studies they may sign up for. Often, this is done so the system can enforce prerequisites, where the inactive study is a prerequisite for an active study.

Should the Researcher receive an email notification when a participant signs up or cancels?: If set to Yes, the researcher for this study will receive an email notification whenever a participant signs up, or cancels their signup, for this study. The email notification will be sent to an email address based on the information the researcher has provided. See the Email Address Options section of this documentation for more information on how the email address is determined. Emails will contain the first 50 characters of the study name as part of the subject line, to make it easy to sort the emails with an email program that supports filtering based on subject line. If set to Yes, researchers will also receive a notification if the system is in lenient prerequisite

enforcement mode and a participant cancels a study that was a prerequisite for the current study. Read the section on Pre-Requisites in this table for more information about this situation. Emails are sent to all researchers specified for the study, unless a specific researcher is assigned to the timeslot that the email notification is being sent about. See Timeslots Linked to Specific Researchers for more information.

Researchers at Timeslot-Level: If set to Yes, it will be possible (but not required) to assign a specific researcher (from the list of researchers for the study) to a timeslot. If set to No, then it is assumed that all researchers (assigned to the study) are responsible for all timeslots. See Timeslots Linked to a Specific Researcher for more information.

Automatic Credit Granting: If set to Yes, timeslots that are more than a specified number of hours old and still in the Awaiting Action state will be changed to a credit grant. The check for timeslots in this situation is made only once per day. If an automatic credit grant is done, you may still change it later if necessary. For online external web studies, the credit grant will take place the specified number of hours after the timeslot (participation deadline) has occurred, so this feature is generally not useful in this situation. This option will not appear for online survey studies (within the system) because credit granting generally occurs automatically, immediately after the participant completes the survey.

Can a participant sign up for this study more than once?: If you would like to allow participants to sign up (and receive credit) for your study more than once (at different times), choose Yes. Otherwise, choose No. If No is chosen, participants may only sign up for the study more than once if they previously failed to show up for the study (a no-show).

Private Comments: This is an optional area where you may enter any comments or notes about the study, which are only visible to the researchers for this study, and not to participants. The maximum length of this field is 3,000 characters.

Research Alternative?: If set to Yes, then this study is considered a research alternative study. Some participants, for various reasons (typically for accruing too many unexcused no-shows, or being unable to consent to participate in studies), may be restricted such that they can only sign up for research alternative studies. Only an administrator may change this value (the default is No).

Participant Sign-Up Deadline: Enter the deadline before the study is to occur that the participant may sign up, in whole hours.

Once you have filled out the appropriate information, save it and the system will be updated immediately with the information. Your next step is likely to add timeslots (sessions). See the Working with Timeslots section of this documentation for more information.

If you need to update this study, see the following Updating a Study section of this documentation. If you would like to add participation restrictions based on prescreen

responses, you can do so when you update the study (see Prescreen Participation Restrictions).

Updating a Study

You may update any of your studies at any time. To do so, choose My Studies from the top toolbar, and you will see a list of your studies. Click on the desired study, and choose the Change Study Information link.

You will see a form remarkably similar to the one you used to add the study. A few options may no longer be changeable depending on the status of the study (e.g., if participants have already signed up for it). The fields shown are all the same as when you added the study. See the Adding a Study section of this documentation for an explanation of those fields.

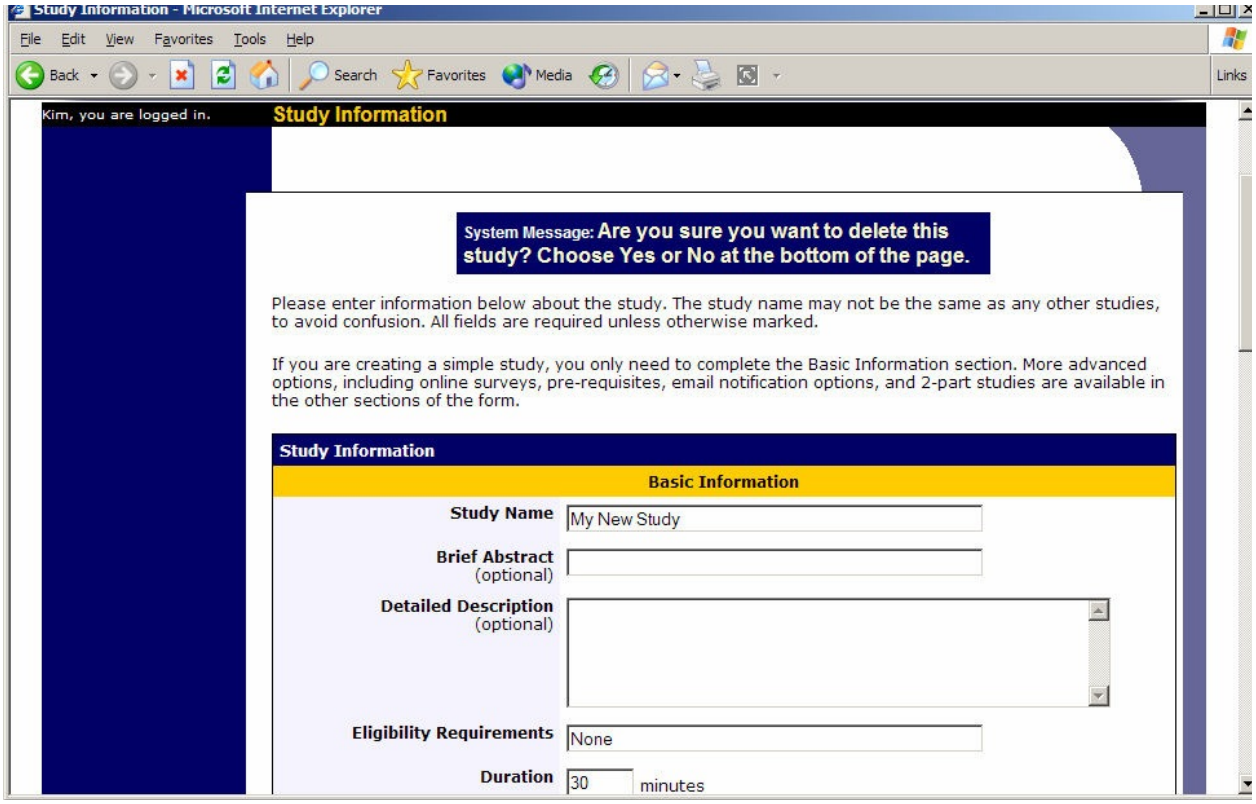
The changes you make will be will be take effect immediately after they are saved.

If you need to change the credit value for a study, and there is no option to do so, this means the study already has at least one participant signed up for it. You cannot change the credit value when a study is in this situation because there is no easy way to handle past credits for the same study (e.g. should old credit grants for the same study be adjusted to reflect the new credit value, or kept the same?). If the study is nearing the end of its run, and variable credit granting is enabled, then the easiest solution is to grant the new credit value to participants who sign up in the future. If you prefer that the credit value is changed for the entire study, contact the administrator, who can make the change for you under certain conditions (which depend on the credit status of existing sign-ups for the study).

Deleting a Study

You may delete a study only if participants have not signed up for it. If you need to delete a study which already has sign-ups, you should make it Inactive instead, if you do not want it to be visible to participants. You may not delete a study which has sign-ups, so the option will not be presented.

If you want to delete a study that has sign-ups, please contact the administrator. The administrator can delete a study with sign-ups, but only if the sign-ups are all without credit values (this usually occurs when study participation history from a previous semester was retained, but credits were zeroed out). If the study has sign-ups where the sign-ups have (non-zero) credit values linked to them, then the administrator cannot delete the study until all those credit grants are changed to a 0 value (or the participants for the sign-ups are deleted). The reason for this restriction is to ensure that the credit count for participants where they have earned credits is accurate, which means that the studies which contributed to their credit earnings must be kept intact.



To delete a study, choose My Studies from top toolbar, click on the desired study, then choose the Delete Study option. You will see a confirmation page. Choose Yes (at the bottom of the page) to delete the study.

Once a study is deleted, it cannot be restored, so use this feature very carefully. If you delete an online survey study, the survey and all data collected will also be deleted.

Timeslot Usage Summary

The timeslot usage summary is available when viewing your study. This gives some basic information about timeslot utilization in the past and in the future, as well as some basic no-show information. It also gives information on timeslots for the study by location (if the study is not an online survey study or external web study), and by researcher (if the study is configured to allow researchers to be assigned to specific timeslots).

For credit studies, the system also provides a summary of how many credits were granted.

Bulk Mail Summary

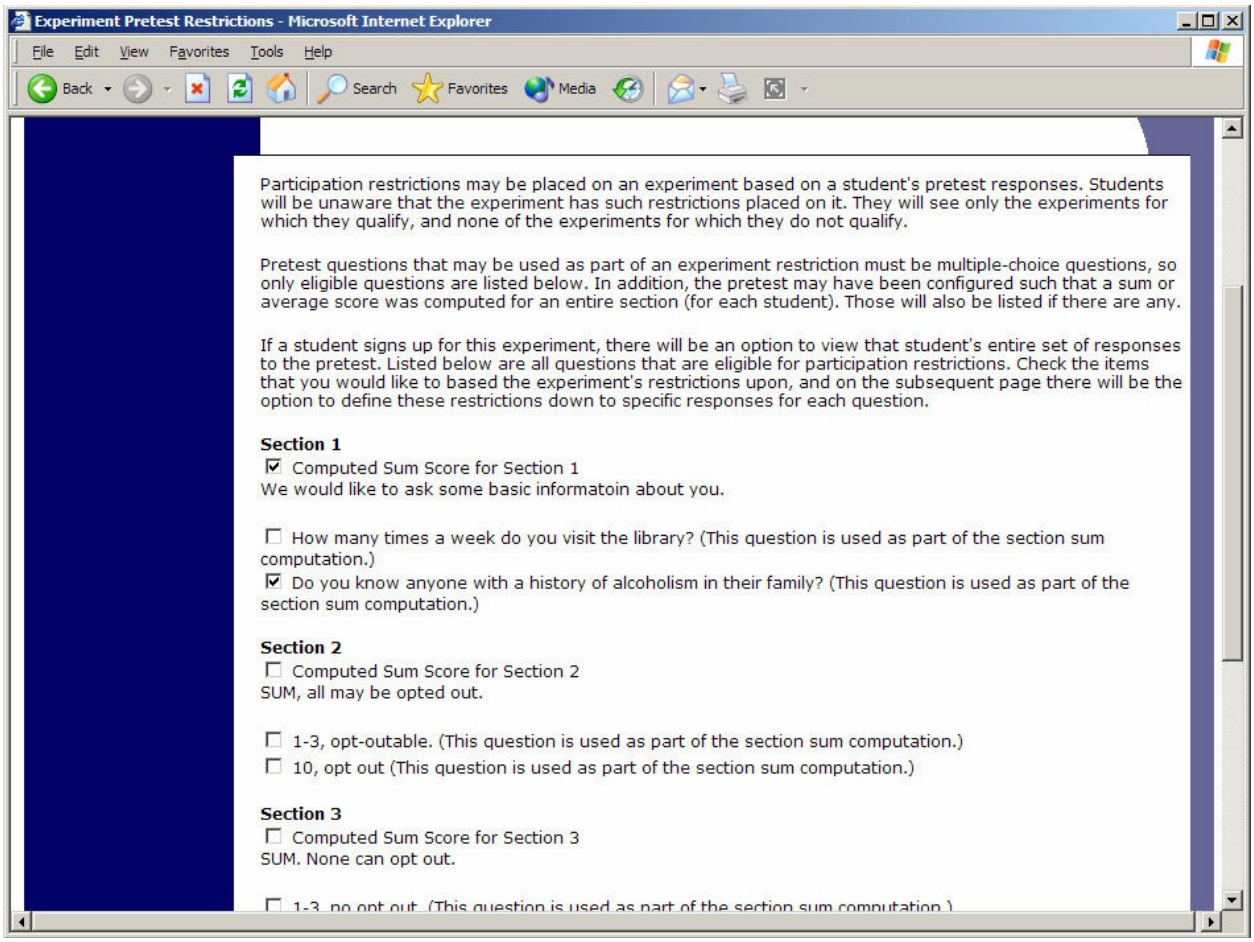
The system tracks whenever any type of bulk email is sent (by a user) related to the study. This includes inviting participants based on the study's prescreen participation restriction analysis, or contacting those who have already signed up for the study. This information is kept for 6 months, and it is tracked to ensure that all users follow generally accepted

Internet practices for responsible use of email. The administrator also has access to this information.

Prescreen Participation Restrictions

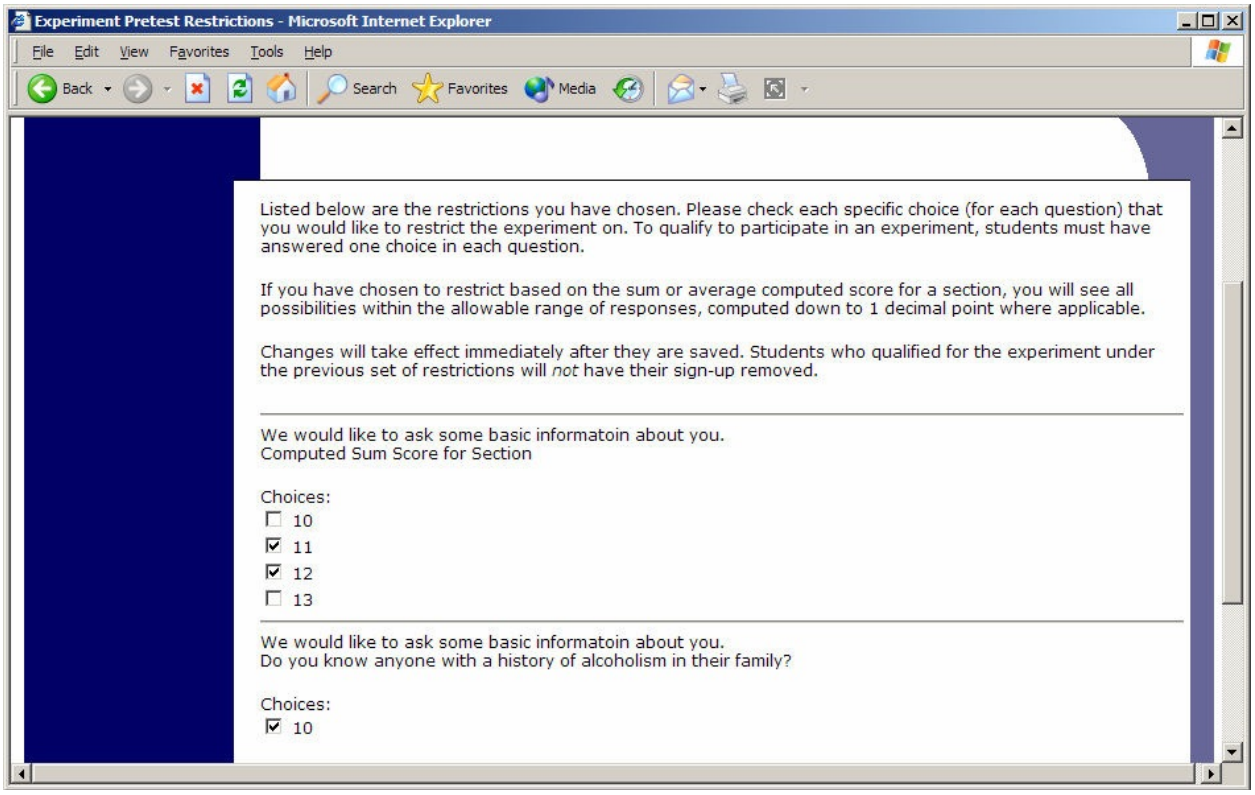
If enabled on your system, the system might contain an online prescreen that participants may (or must, depending on your system configuration) complete. You may place participation restrictions on your study based on prescreen responses. Participants are unaware that such restrictions are placed on the study. These restrictions are never listed to them. If they do not qualify to participate in a study because they do not meet the prescreen participation restrictions, then the study will simply not be listed to them. This is important to note – participants never know why a study was or was not listed to them, because they are unaware of the prescreen restrictions.

You may restrict a study on any question or questions on the prescreen that allowed for a multiple-choice answer where only one choice could be selected. You may also restrict a study based on a computed section sum or average score for a participant, if the prescreen was set up in such a manner. You may restrict to one choice or many choices for any question. If you restrict on multiple questions, it is the same as a logical "AND." For example, if you setup the prescreen restrictions so that participants must have answered "Yes" to a "Do you wear glasses?" question and "Blue" or "Grey" to "What color are your eyes?", then they must meet *both* requirements to participate. In other words, only participants who wear glasses and have either blue or grey eyes are eligible. There is no support for a logical "OR" restriction across multiple questions. The restrictions are inclusive, which means that if you select a choice as a restriction, then participants must have answered at least one of the choices selected for each question that is part of the restriction in order to see and participate in the study, as opposed to exclusive where checking the choice as a restriction would exclude them from participation.



To set participation restrictions, view (do not choose edit) your study and choose View/Modify Restrictions. You will see a list of eligible questions which you may use for your restrictions. If the study already has some restrictions, those will be checked, and you will see how many participants currently meet the restrictions. Choose the questions you would like to restrict upon (and keep the existing checked restrictions checked, unless you want to remove that restriction), and click on the Set Restrictions button. On the subsequent page, you can select each value that is acceptable for each question you have chosen. Once you have selected all the acceptable values, save your changes and they will take effect immediately. It is important to note that if you change the restrictions, it will *not* remove the study sign-ups for participants who qualified under the previous set of restrictions. For this reason, you should probably decide on your restrictions before making the study available to participants.

If you have restriction requirements where you would like to restrict participation to a percentage of the population (for instance, the responses that were chosen by the top 25% of people), but you are not sure which responses meet this requirement, you can use the prescreen response analysis feature to determine the valid responses. See Prescreen Response Analysis for more information. You may also use Analyzing Prescreen Responses to get an idea of how many participants are potential candidates for participation in your study, based on a specified set of restrictions.



Inviting Qualified Participants to a Study

While viewing the list of prescreen restrictions currently set for a study, and the number of participants who meet those restrictions, you may see the option to Invite Qualified Participants. Using this option, you may craft an email to be sent to all qualified participants. You may choose to exclude those who have already signed up for or participated in any studies you specify. The system will automatically exclude all participants who have participated or are signed up for the current study (no-shows are not excluded though, since they may sign up again). If the study is not a research alternative study, the system will also automatically exclude participants with Limited accounts, as they are ineligible to participate in studies not marked as a research alternative study.

The system will pre-fill the email text with useful information like the name of the study and how many timeslots are currently open. You cannot include attachments in the email, so if you have a document you would like to include, you should post it on a university webserver and provide a link to the document in the email you send.

If you have set participation restrictions for the study based on course enrollment, those restrictions will be taken into consideration (i.e. abided by) when determining which participants receive the email.

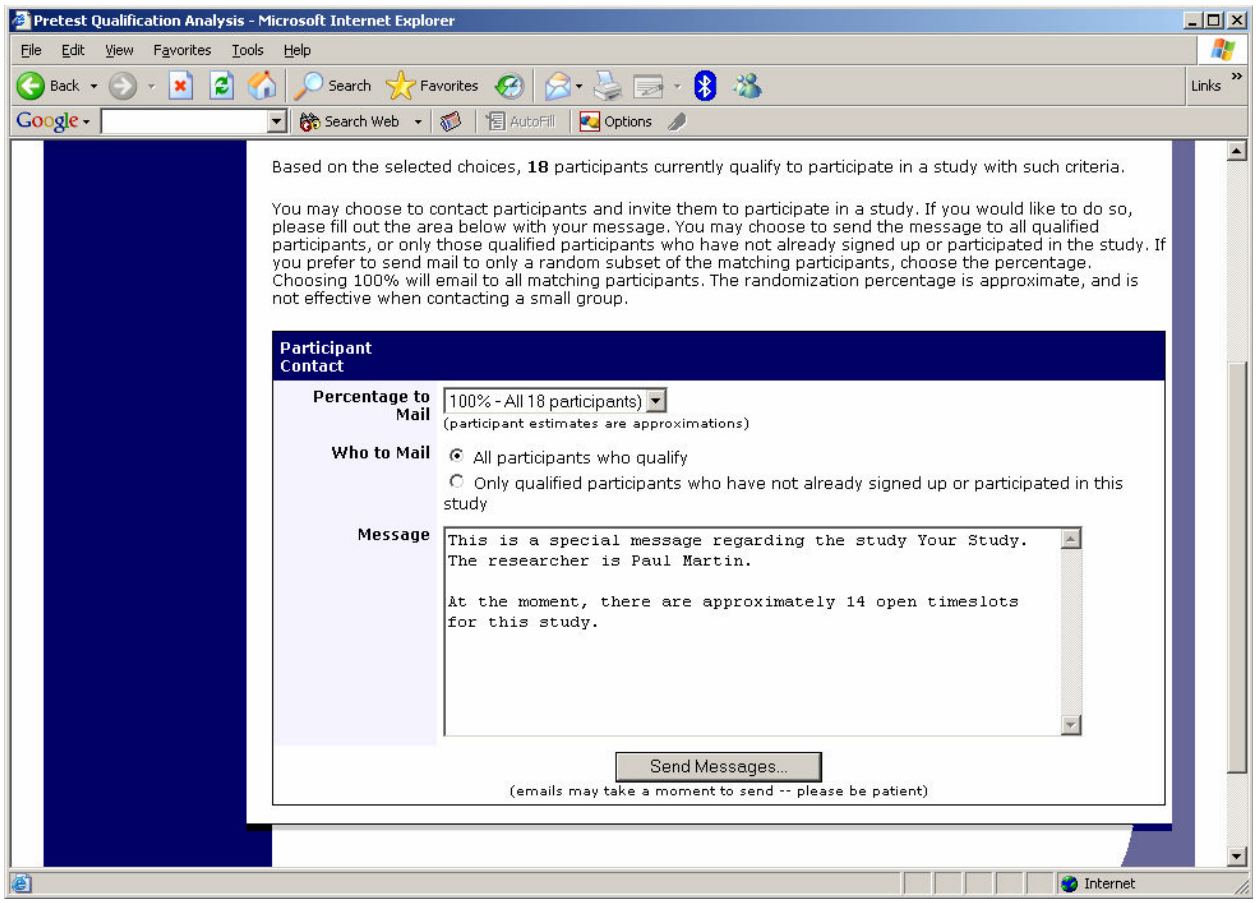
There is also an option to choose a random percentage from the overall list of matching participants to email. It is important to understand that the system does not keep track of

which random percentage of the group of matching participants is sent to each time, so that if you send to a random 30% now, and a random 30% an hour later, it could very well be the case that many participants receive the email on both those occasions.

The From (sender) address on the email will be the administrator email address, which is done to prevent the email from being blocked by junk email filters. The “Reply To” address of the email will be that of the user who is actually sending the email, so when a user chooses to reply to the email, the reply will be sent to that (the reply to) address.

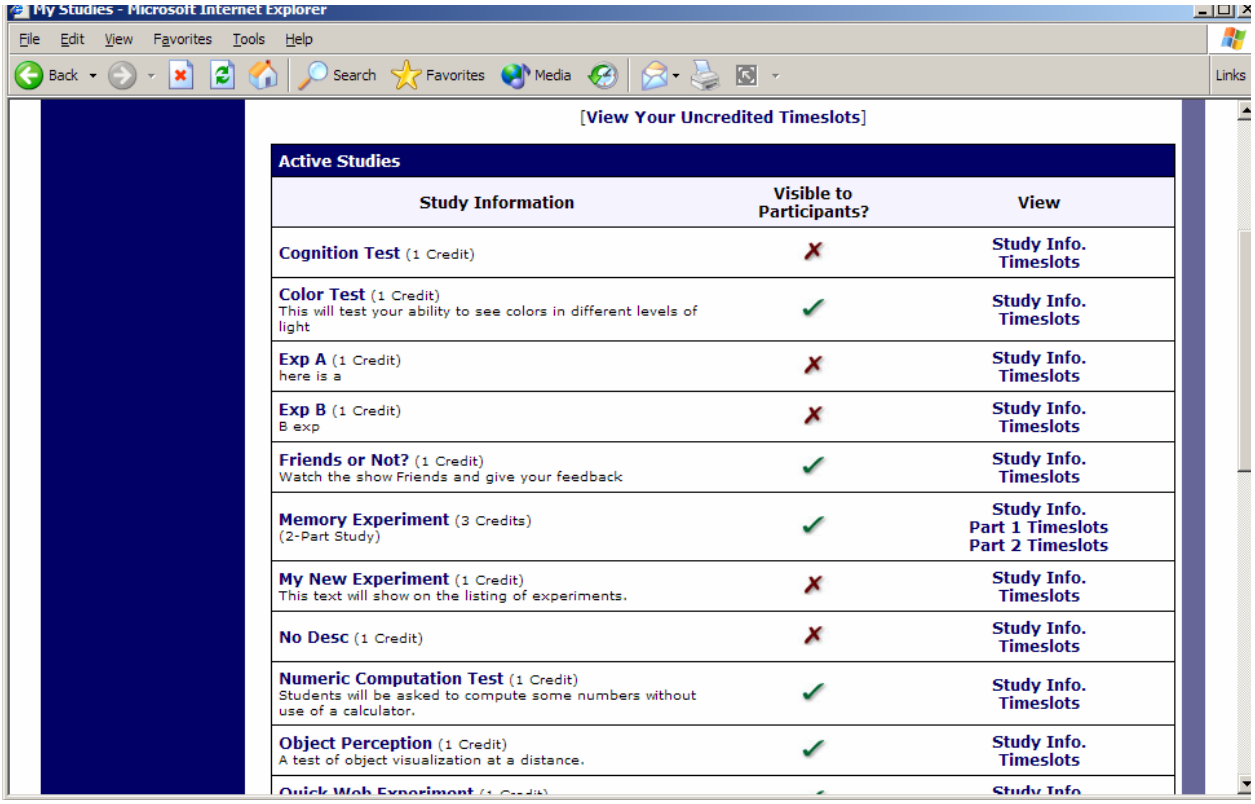
Finally, there is an option to specify a delay in sending the email, based on the number of hours from when the emailing option is used. This is useful if you want to target a certain time of day (e.g., during the evenings) when the email will be sent. The emails are generated at the time you use the emailing facility, but are stored on the server until the specified sending time. They cannot be removed from the queue once this emailing facility is used.

In some cases, the administrator may have imposed a limit on how many participants may be emailed. This is often done to prevent abuse of the system, such as cases where researchers invite too large a number of participants and this is not in accordance with generally accepted Internet principles for sending email. If there is such a limit, the system will look at the number of participants the researcher plans to email, and if that number is greater than the limit, it will block the sending entirely, as opposed to sending only enough emails up to the limit. To get around this limitation, the researcher can further restrict who they plan to send to (perhaps choosing a smaller random percentage of users, or more closely defined prescreen participation restrictions), or ask the administrator to send the email for them. The administrator is not subject to such limitations. Regardless, any use of this bulk email facility will be logged, and that information will be kept for approximately 6 months. The administrator can easily pull up a report of how many emails a specific researcher has sent, so is good to be careful about not abusing this feature. In addition, Sona Systems reserves the right to temporarily remove the right to login from a researcher if there are verifiable reports of abuse of this feature. Typically before doing so, the administrator will be notified by Sona Systems as it is preferred to have the administrator deal with such problems.



Viewing Your Studies

To view your studies (and not the studies of others), choose the My Studies option on the top toolbar. The system will list all your studies in alphabetical order by study name, grouped by studies that are active, then inactive studies.



Participant Study View

If you would like to see how your study appears when participants view it, find your study and choose the Participant Study View option. This will show exactly how the study appears to participants, with the exception that when a participant views a study, next to each pre-requisite and disqualifier study (for a study) is listed a status indicator about whether they have met that requirement. In Participant Study View, the prerequisite and disqualifier studies are listed, but there is no status indicator next to each study in the list.

If for some reason you think your study is not visible to participants, it may be due to various restrictions you have set on the study, like prescreen participation restrictions, such that few (or none) of the participants in the pool qualify. You can ask the administrator to use the Check Study Configuration tool (available to them when they view your study) to provide advice on why your study may or may not be visible to participants. Administrators there also have an option to type in a specific participant to see if that participant would qualify for your study.

Viewing Other Studies

To view all studies that are visible to participants, choose the All Studies option from the top toolbar. You will see a list first of all Active studies. These studies will show up to participants on the list of available studies. The next group of studies (if there are any) is Inactive studies. These will *not* show up on the list of available studies (to participants), but participants can access information about these individual studies on links from the page

with their progress (if they participated in the study) or if another study has the Inactive study listed as a pre-requisite or disqualifier.

Working with Timeslots (Sessions)

Timeslots (also referred to as Sessions) are the available times when a participant may participate in the study. If you are setting up timeslots for a web-based study, please read the section in this documentation on Web-Based (Online) Studies for some special information.

Timeslots allow you to specify a date, time, location, maximum number of participants, and researcher for a session.

Timeslots Linked to Specific Researchers

You will have an option to link timeslots to a specific researcher. This is done primarily for organization purposes, and has no effect on who can view and modify the study, or any timeslots for that study. This feature is useful when there are a number of researchers running a study, and researchers are responsible for running specific timeslots. If a timeslot has a specific researchers linked to it, then only that researcher will be listed as the contact point when a participant receives any emails related to their participation in that timeslot. Finally, only the researcher connected to that timeslot receives related notification emails, such as participant cancellation notification, and reminder emails (assuming such emails are enabled).

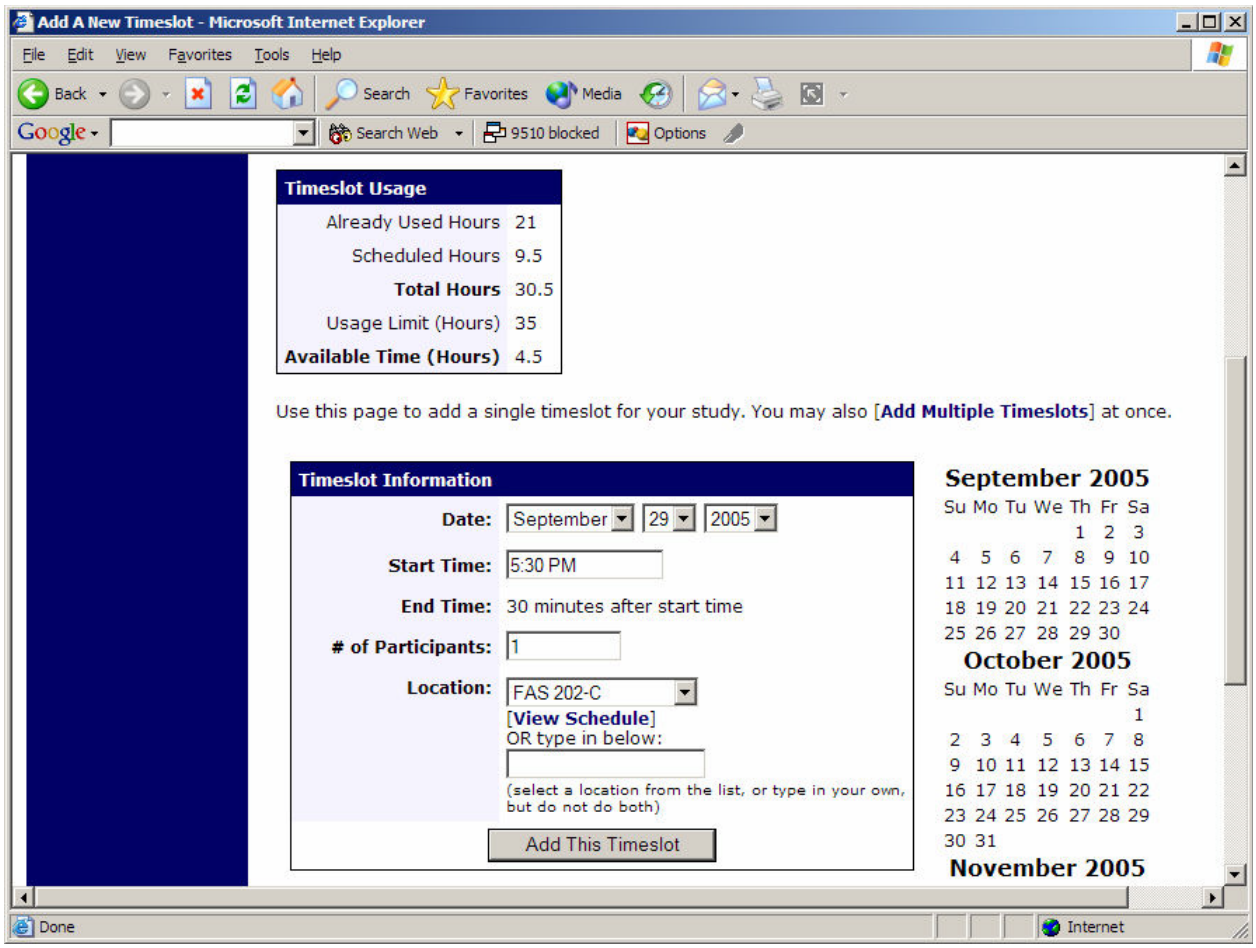
It is also possible to have some timeslots where a specific researcher is linked to them, and others where all researchers (who are assigned to the study) are responsible for the timeslot. It is not possible to link more than one, but not all of the researchers (for the study), to a specific timeslot. The options are to either link one researcher to the timeslot, or all of them.

If a researcher is removed from a study, then any timeslots that were linked to them for that study will be changed so all researchers (for the study) are now responsible for those timeslots.

To use this feature, the system must be configured to allow multiple researchers per study. Then, the study itself must be configured to allow researchers to be linked to specific timeslots. Finally, the study must have more than one researcher connected to it.

Creating Timeslots

To add a timeslot for a study, you must first choose the study that you would like to add a timeslot for. To view your studies, choose the My Studies option on the top toolbar. Click on the desired study, and choose the Timeslots choice. You will see a list of any existing timeslots, and the Add A Timeslot option at the bottom of the page. Click on Add A Timeslot.



The following table lists the information you may enter about a timeslot, along with an explanation. All fields are required.

Date: The date for the timeslot.

Start Time: The time for the timeslot. A sample time will be provided. If you want to change the time, please use the same format as the time you see presented. Note in particular how "a.m." and "p.m." are handled (if such a format is enabled on your system).

End Time: The time when the timeslot will end. This is computed automatically based on the duration you entered when you set up the study.

# of Participants: The number of participants for this timeslot. This limit is *not* visible to participants. They will only see whether the timeslot is full or not. The maximum number is 999.

Location: The physical location where the study will take place, for this timeslot. It will be automatically filled with the location of the previous timeslot, when available, to ease in data entry. The location field does not apply for web-based studies.

Researcher: The researcher assigned to this specific timeslot. The list will contain a list of all researchers for the study. Choose ALL if all researchers (for the study) should be assigned to this timeslot. See Timeslots Linked to Specific Researchers for more information.

To ease data entry, the system will automatically fill in the date, time, and location based on the ending time of the last timeslot for this study. If applicable, your current timeslot usage will be listed, and you will be prevented from adding a timeslot that would exceed your timeslot usage time limit. A convenient calendar is provided next to the form, and you can click on any date and that date will be transferred to the form.

If you add a timeslot such that there is another timeslot (for any study) that occurs in the same time, at the same location, you will receive a warning (but the addition will be allowed). If you add a timeslot that will take place outside of normal hours (for example, at 1:00am), the system will provide a warning but will allow it to be scheduled. You may not schedule a timeslot to occur after the IRB expiration date for your study, if Strict IRB mode is enabled by the administrator. The system allows adding timeslots to a study that is not available to participants (not active or not approved), but it will give a warning because participants are not able to sign up for the timeslot.

If you are running a web-based (online study), you should create a single timeslot with the participation deadline equal to the last day you would like to run the study. For number of participants, specify the maximum number of participants who may participate. If you are running a web-based study and you plan to collect data from more than 999 participants (999 is the maximum allowed in one timeslot), then once that timeslot is close to filling up, create a second timeslot as a slightly different time and/or date as the first timeslot.

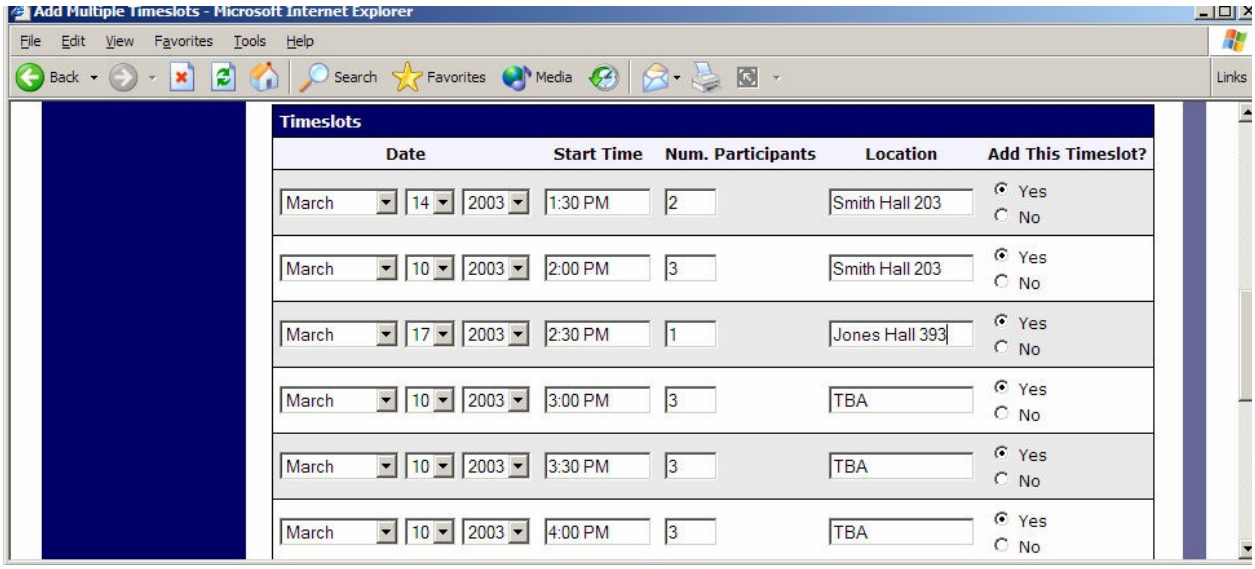
Creating Multiple Timeslots

If you would like to add multiple timeslots at once, choose the Add Multiple Timeslots link. You may choose to add a specified number of timeslots, or copy the timeslots from another week to a specified week. If you choose to copy, the system will copy the time, location, and number of participants for the specified week to the desired week, for each day of that week (starting with Monday).

If you choose to create a specified number of timeslots, you can choose the number of timeslots you would like to add, the start time and date, and the amount of time between each timeslot (to allow for breaks). You also may specify that timeslots that would occur outside normal business hours be shifted to the next business day, and specify when business hours occur. The system considers Monday-Friday to be business days. On the subsequent page, you may change any of it to deal with special cases.

Timeslots that you attempt to add, that either have errors or would result in exceeding the timeslot time usage limit, will not be added. This feature is not available for web-based (online) studies, as web-based studies rarely have more than one timeslot.
If you do not want to add a specific timeslot that is listed, choose No in the Add

This Timeslot? Column.



Modifying and Deleting Timeslots

To modify or delete a timeslot for a study, you must first choose the study that you would like to deal with. To view your studies, choose the My Studies link from the top toolbar. Choose the Timeslots option in the timeslots column for the desired study. You will see a list of all recent timeslots. Recent timeslots in the past with no participants signed up will not be displayed. To work with timeslots more than a few days old and to see all timeslots, you will see a link to view all timeslots for the study. Select the timeslot you would like to deal with, and click the Modify button.

If the timeslot has no participants signed up for it, you will see a Delete button. You may not delete a timeslot that has participants signed up for it. If you would like to delete the timeslot, click the Delete button, and you will see a confirmation page. Choose Delete again to delete the timeslot.

If you would like to modify the timeslot, modify the desired information and click the Update button just below the timeslot information. It should be noted that participants will *not* be notified (by email) of any changes you make to the timeslot, so you should contact them if information needs to be passed on to them (a link is provided on the same page to do so). If you change the date or time of the timeslot, you will be warned that this was changed in case the change was unintended. You may not update the size of the timeslot (number of participants) to a value lower than the current number of participants signed up for the timeslot. Generally, researchers only update timeslots with sign-ups to update the location, if it was not available when the timeslot was originally created.

Timeslot Change Tracking

The system automatically tracks certain changes that occur with a timeslot, including any time key information about the timeslot (date, time, etc.) is changed, as well as any time a manual sign-up or cancellation is performed (i.e., not a sign-up or cancellation done by the participant). This information is tracked for the last 3 months of changes for each timeslot.

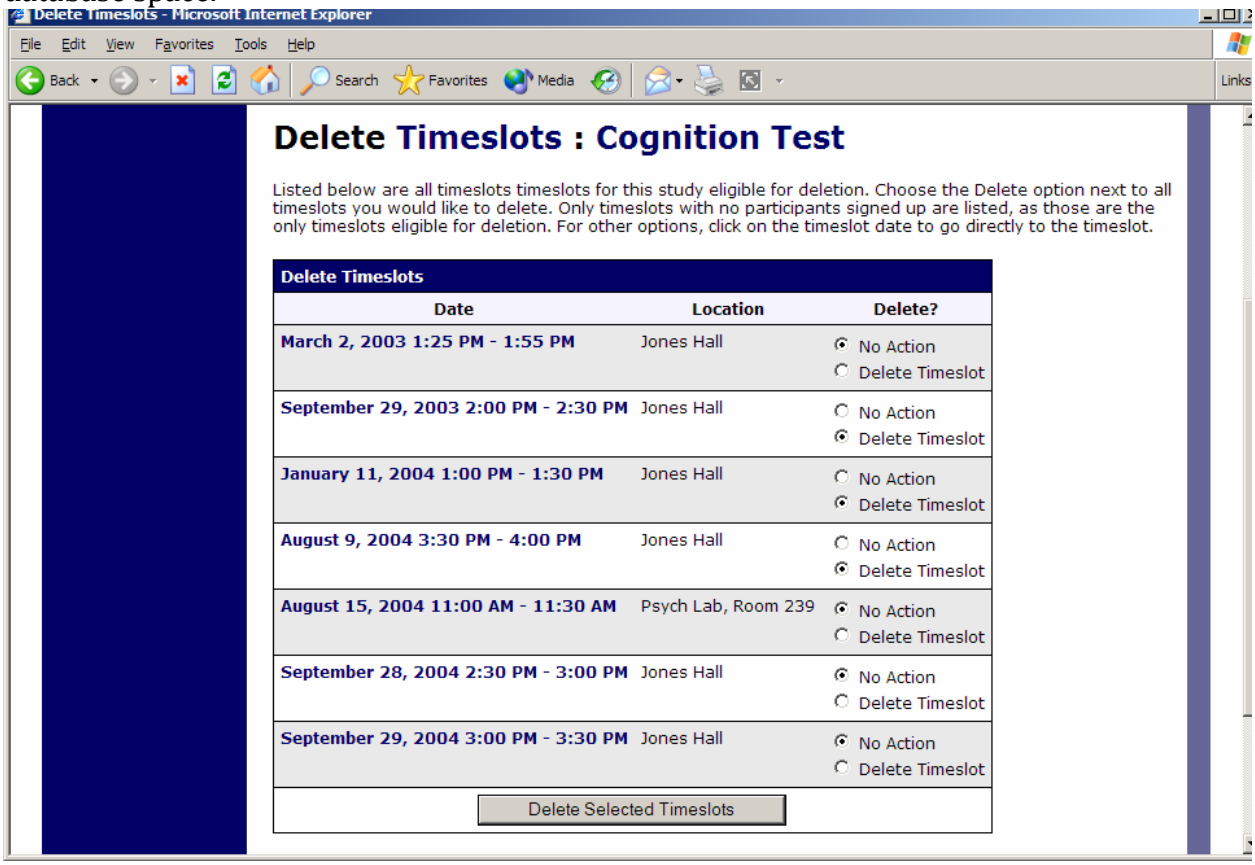
To view this information, choose the View Timeslot Modification Log when viewing a timeslot, and you will see this information.

Deleting Multiple Timeslots

If you would like to delete multiple timeslots at once, you may do that as well. Such a feature is only available for timeslots which have no participants signed up. To do so, select the desired experiment and choose Timeslots. At the bottom of the Timeslots page, you will see a Delete Multiple Timeslots option. The option may not appear in certain cases where such an option is not available because of a lack of available timeslots to delete.

After going to that page, you will see a list of timeslots eligible for deletion. Choose the timeslots you would like to delete, and choose Delete Selected Timeslots to proceed. If you would like to delete all empty timeslots, there is a Select All option at the bottom of this page that will automatically select all timeslots listed on the page for deletion. Click the Reset button to revert the effect of choosing the Select All option.

The system routinely deletes all empty timeslots more than 3 months old to preserve database space.



Manual Sign-Up

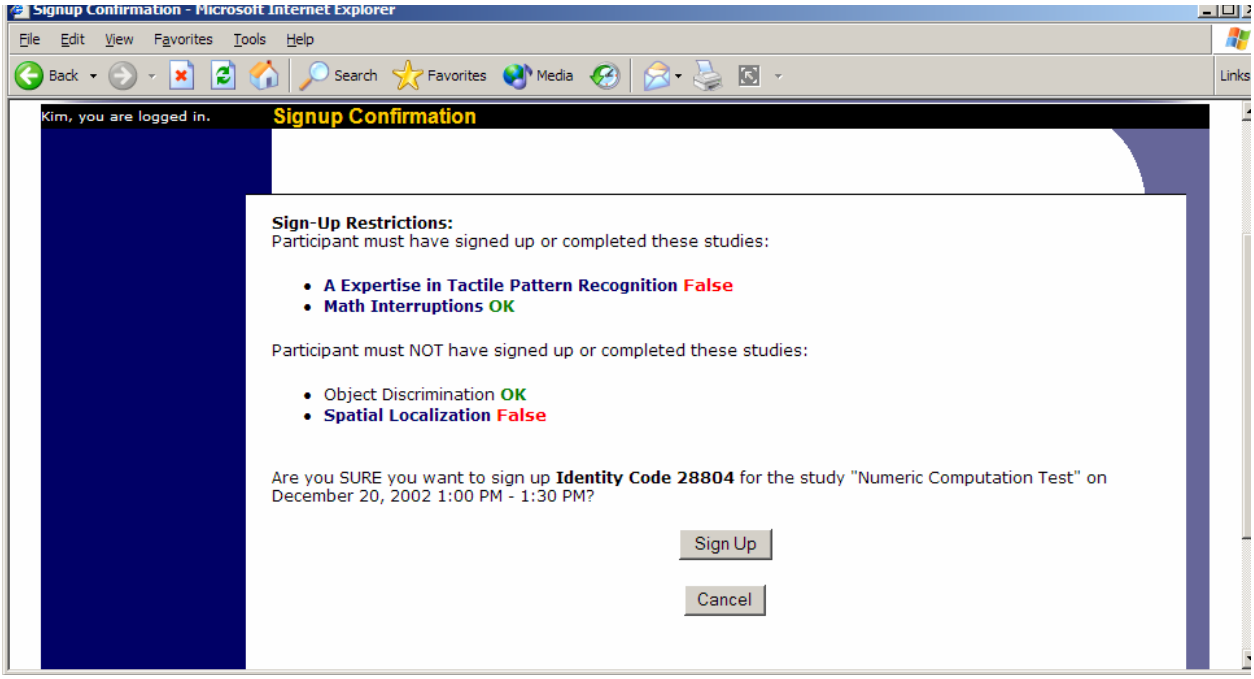
If enabled on your system, you may manually sign up participants for your study. There are a number of situations where this is desirable. If the participant happens to show up for a timeslot they were not signed up for, and you elect to let them participate, you can sign them up on the spot for the timeslot. The participant in many cases cannot sign up on their own in this situation, because the sign-up deadline has passed. You may also sign up a participant for a study that has already occurred, if necessary.

Also, a manual sign-up overrides any restrictions you have placed on the study (e.g. prerequisites), though you will be warned if you are overriding any restrictions. You may not sign up a participant for the same timeslot that they are already signed up for. You are allowed to sign them up for a study even if they are already signed up for a different timeslot for that same study, though you will receive a warning in this case. You may not sign up a participant for a study if it would cause them to exceed their maximum credit limit. If it is necessary to do so, please ask the administrator to do this, as they are allowed to do a manual sign-up even when it will violate maximum credit earning limits.
You also may not sign up a participant whose account is Limited, if your study is not a research alternative study, as those participants are ineligible for your study (the administrator can still do this).

If the system is configured as such, the participant will receive a confirmation email when you sign them up for a study. In that case, you are also given the option to enter comments to be included in this email that may better explain to the participant why they were signed up. If you are signing up a participant for a timeslot more than one year old, a confirmation will *not* be sent despite the system configuration. This is to make it easier when transitioning from an existing system, as you may sign up old participants for the purposes of preventing them from signing up for the same study again in Sona.

You may only sign up participants for your own study.

To sign up a participant for a timeslot, you must first find the desired study and timeslot. To view your studies, choose the My Studies option from the top toolbar. Click on Timeslots for the desired study, then select the timeslot you would like to deal with, and click the Modify button. At the bottom of the page, you will see a Manual Sign-Up option, if it is enabled. Type in the participant's User ID (you may have to ask them for this) and click Sign Up. You may also have the choice to enter their last name and choose from a list of participants. In all cases, after submitting the form, you will see a confirmation page that also lists any restrictions on the study. Choose Sign Up to complete the sign-up.



If you are doing a manual sign-up for a two-part study, you must do a manual sign-up for each part separately. The system will overlook the scheduling range restrictions as well.

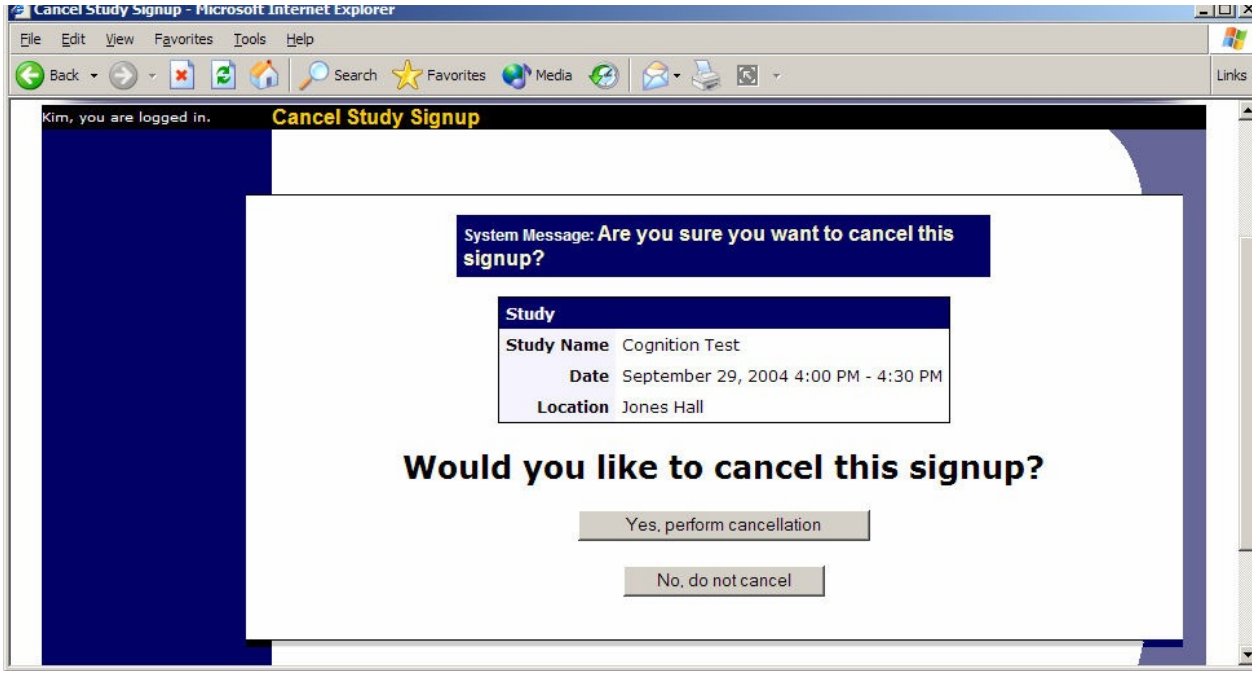
You cannot use the manual sign-up feature for online survey studies, because the sign-up for the study is integrated with the administration of the survey.

Manual Cancellation

If enabled on your system, you may have the opportunity to cancel a participant's signup.

You may only cancel sign-ups that are in a No Action Taken state. To cancel a signup, find the desired timeslot and participant, and click Cancel next to their name. The participant will be sent an email about the cancellation (and who performed it), along with a confirmation code, and their sign-up will be immediately cancelled. The administrator may also receive a copy of this cancellation email, depending on how the system is configured.

You may cancel all participants for the same timeslot at one time, when applicable. The option will appear below the list of signups, in cases where there are two or more participants signed up for the timeslot who are eligible for cancellation (No Action Taken state).



Viewing the Participant List

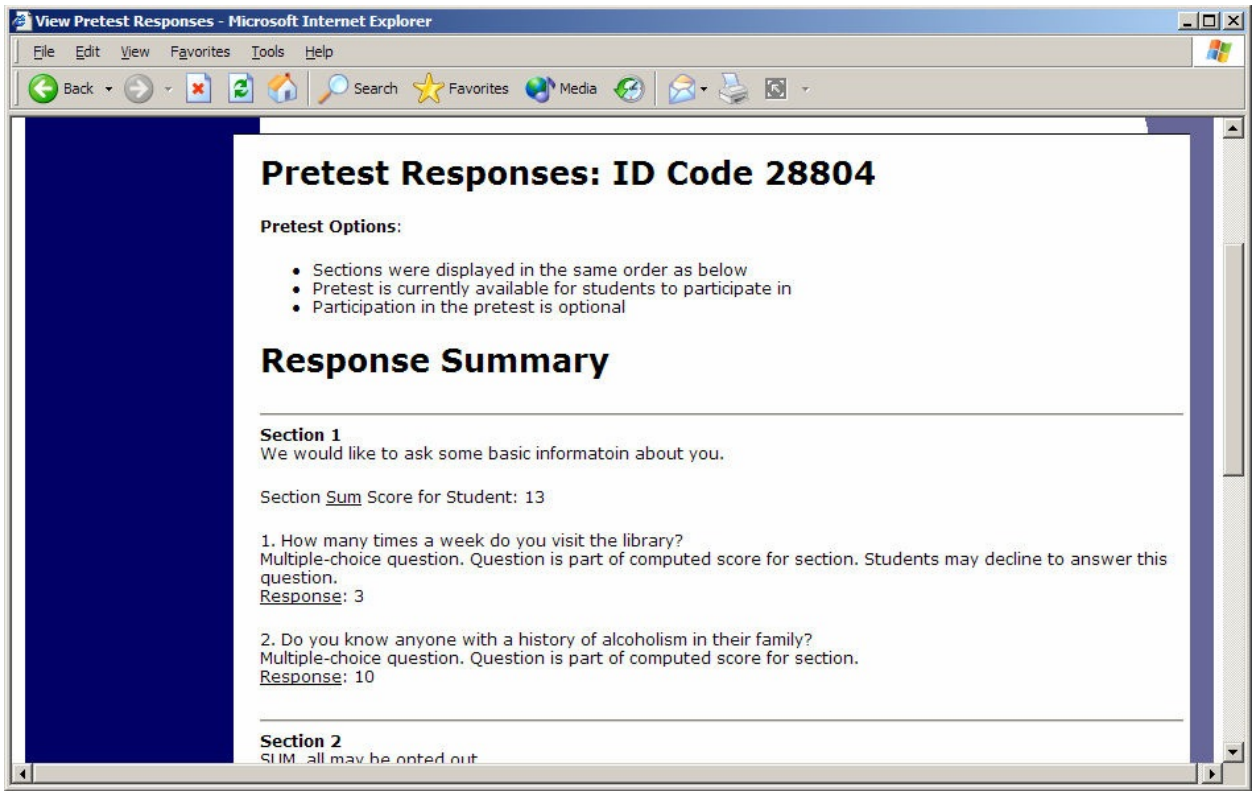
To view the list of participants who have signed up for your study, you must first select the study and timeslot you wish to see. To view your studies, choose the My Studies option from the top toolbar. Click on the Edit link in the timeslots column for the desired study, then select the timeslot you would like to see, and click the Modify button. The list of participants, along with their email addresses, will be listed.

Viewing Prescreen Responses

If online prescreens are enabled on your system, and you are also allowed to view an individual participant's prescreen responses, then you will see a Prescreen link next to each participant's name when you view the information for a timeslot. Click on that link to view the participant's prescreen responses.

If you would like to download the prescreen data for all participants in your study, choose the Download Prescreen Responses option after clicking on your study. That will allow you to download all the data at once, in CSV (comma-separated) format, for further analysis. The download will not contain data for participants marked as a no-show.

If you would like to analyze responses in aggregate (across all participants in the system), see Analyzing Prescreen Responses in this documentation.



Granting or Revoking Credit

At the completion of a session, you should promptly deal with the participants, in the system, to ensure proper credit grants. The reason for the prompt handling of this situation is in the event your study is a pre-requisite for another study, and a few other situations. You do not want to hold up other studies that are waiting on your response to the study you just ran.

To grant or revoke credit for a timeslot, you must first find the desired study and timeslot. To view your studies, choose the My Studies option from the top toolbar. Click on the Edit link in the timeslots column for the desired study, then select the timeslot you would like to see, and click the Modify button. You will see a list of participants, identified by name. If the participant properly participated in the study, click the Credit Granted button next to their name (this text may appear as Participated if the study is set up for payment).

If the participant did not appear for the timeslot, you may choose to mark their no-show as excused or unexcused. Depending on how your system is configured, an unexcused no-show may result in a penalty being assessed for the participant (the system will compute this automatically), or their privileges to use the system may be restricted. You should ask your administrator for guidelines about when to grant an excused no-show or an unexcused no-show. Generally, excused no-shows are granted for extenuating circumstances, like if the participant was involved in a car accident on their way to the appointment. An unexcused no-show is generally used when the participant did not show

up and had no reasonable excuse. For most schools, the majority of no-shows are unexcused and are due to carelessness on the part of participants.

If desired, enter any comments about the session in the Comments section (generally, this is used to indicate the reason for denying credit). Participants will see anything you enter in the Comments section for their sign-up, and these comments will be included in the email sent to participants when a credit grant/revocation occurs, if notification emails are enabled on your system.

Click on the Update Sign-Ups button at the bottom of the list of sign-ups to save your changes. Credit will be granted or a penalty assessed as necessary. The participant(s) will be emailed about this if the system is configured in such a manner.

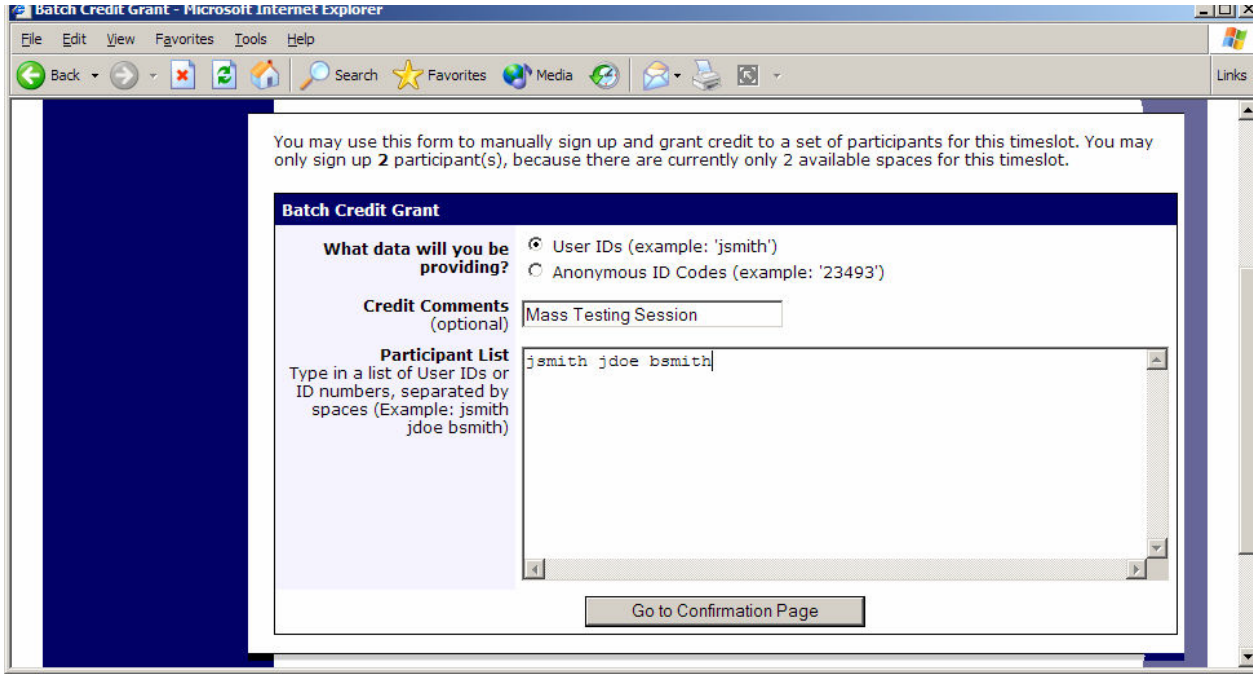
It is not recommended to leave any sign-up for a timeslot that has occurred in the "No Action Taken" stage. This is a credit "limbo" and the system will warn you upon your next login about the offending timeslot that has not been dealt with properly. Note that if Manual Cancellation is enabled and you would like to cancel a participant's sign-up, the sign-up must be in No Action Taken state.

If you need to do a simple credit grant across many timeslots, see the Uncredited Timeslots section which offers such a feature.

Batch Credit Granting

In some cases, you may wish to automatically sign up and immediately credit a group of participants. This is often useful if you administered a study on an ad-hoc basis, and you want to credit participants after the fact.

To do so, go to the appropriate timeslot (you may want to create a timeslot specifically for this purpose), and click on Modify Timeslot. In the Manual Sign-Up section (if enabled), you will see a Batch Credit Grant link. Click that and you can provide the list of User IDs of users you would like to sign up and credit. Users will be signed up and credited immediately. This feature overrides any sign-up restrictions on the study, just as a normal manual sign-up does.

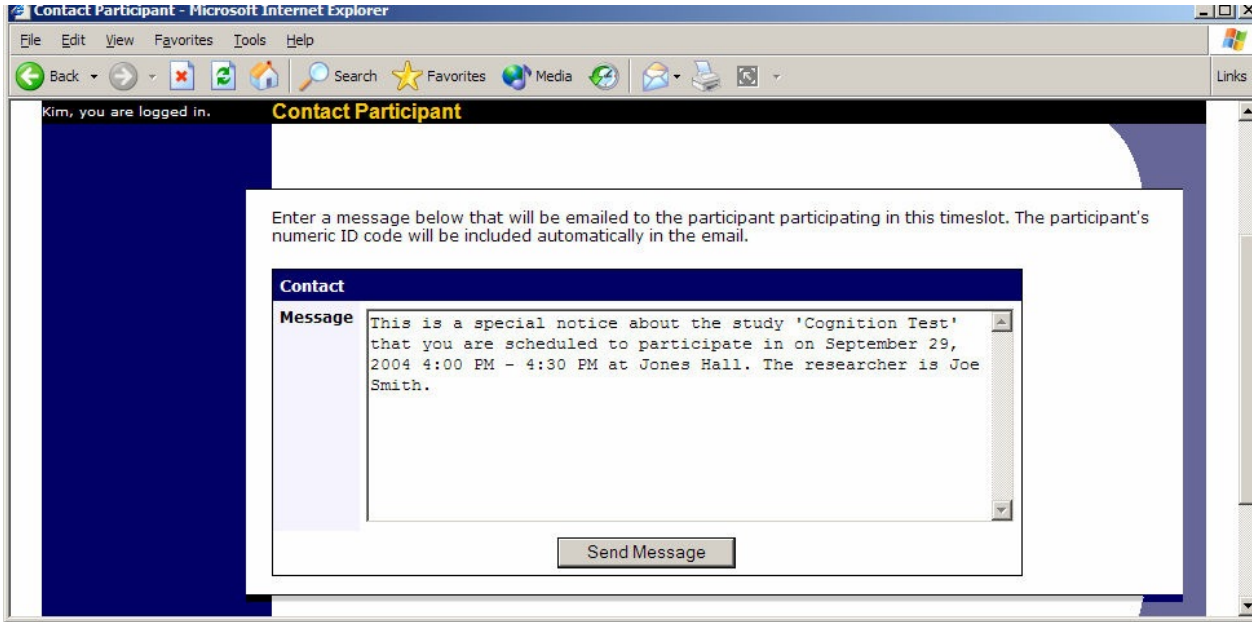


If you are signing up a participant for a timeslot more than one year old, a sign-up confirmation will *not* be sent despite the system configuration. This is to make it easier when transitioning from an existing system, as you may sign up old participants for the purposes of preventing them from signing up for the same study again in Sona.

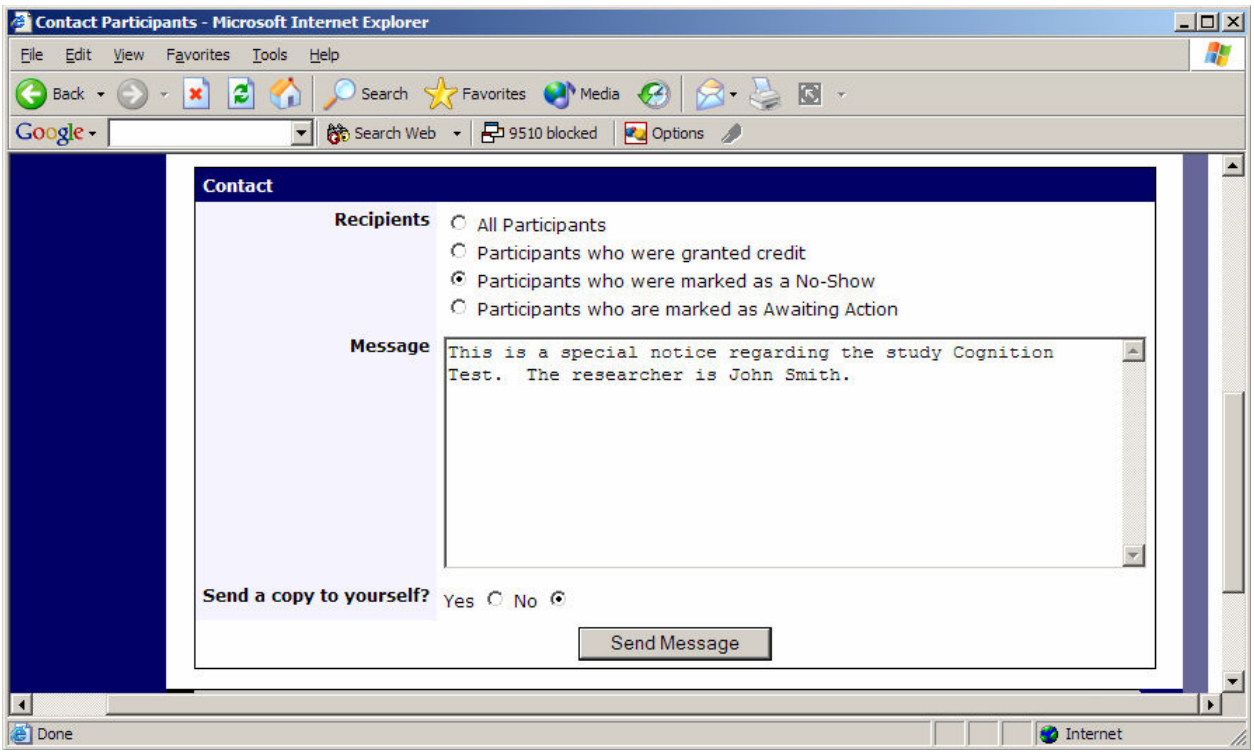
Emailing Participants

If you wish to contact participants in a particular timeslot for any reason, you may click on the Contact link that will appear next to each participant's name (or ID code) to contact an individual participant. To email the group of participants for a particular timeslot, click the Contact All Participants choice at the bottom of the Modify Timeslot page for that timeslot. You will be taken to a page where you can fill out a message that the system will send to the selected participants. The message is auto-filled with some basic information about the study, so participants are aware of which study you are referring to. You may remove this information if desired. You may choose to receive a copy of the email that you send.

Depending on how your system is configured, participants may already be receiving a reminder about upcoming studies the day before they are scheduled to participate. Ask your administrator for more information.



In some cases, you may find it useful to contact all participants for the study, across all timeslots. This feature may be particularly useful if you are sending debriefing information when a study has concluded. To do so, go to My Studies, click Study Info. next to the desired study, and choose the Contact Participants option. You will then be able to select which group of participants to send to, and a message to send. Messages will be sent in groups of 300 (or less, depending on how your system is configured) to avoid overloading email servers. You cannot include attachments in the email, so if you have a document you would like to include, you should post it on a university webserver and provide a link to the document in the email you send.



The From (sender) address on the email will be the administrator email address, which is done to prevent the email from being blocked by junk email filters. The "Reply To" address of the email will be that of the user who is actually sending the email, so when a user chooses to reply to the email, the reply will be sent to that (the reply to) address.

There is also the option to restrict the emails so they only go to participants who signed up for timeslots in a specified date range. The date range is based on the date of the timeslot, not when the participant signed up for, completed, or received credit for the study.

Finally, there is an option to specify a delay in sending the email, based on the number of hours from when the emailing option is used. This is useful if you want to target a certain time of day (e.g., during the evenings) when the email will be sent. The emails are generated at the time you use the emailing facility, but are stored on the server until the specified sending time. They cannot be removed from the queue once this emailing facility is used.

In most cases, summary information about the email you sent, and in particular to how many recipients it was sent to, will be logged and made available to the administrator. This is done to ensure there is no abuse of the email facility in the system, in compliance with generally accepted Internet practices for sending emails.

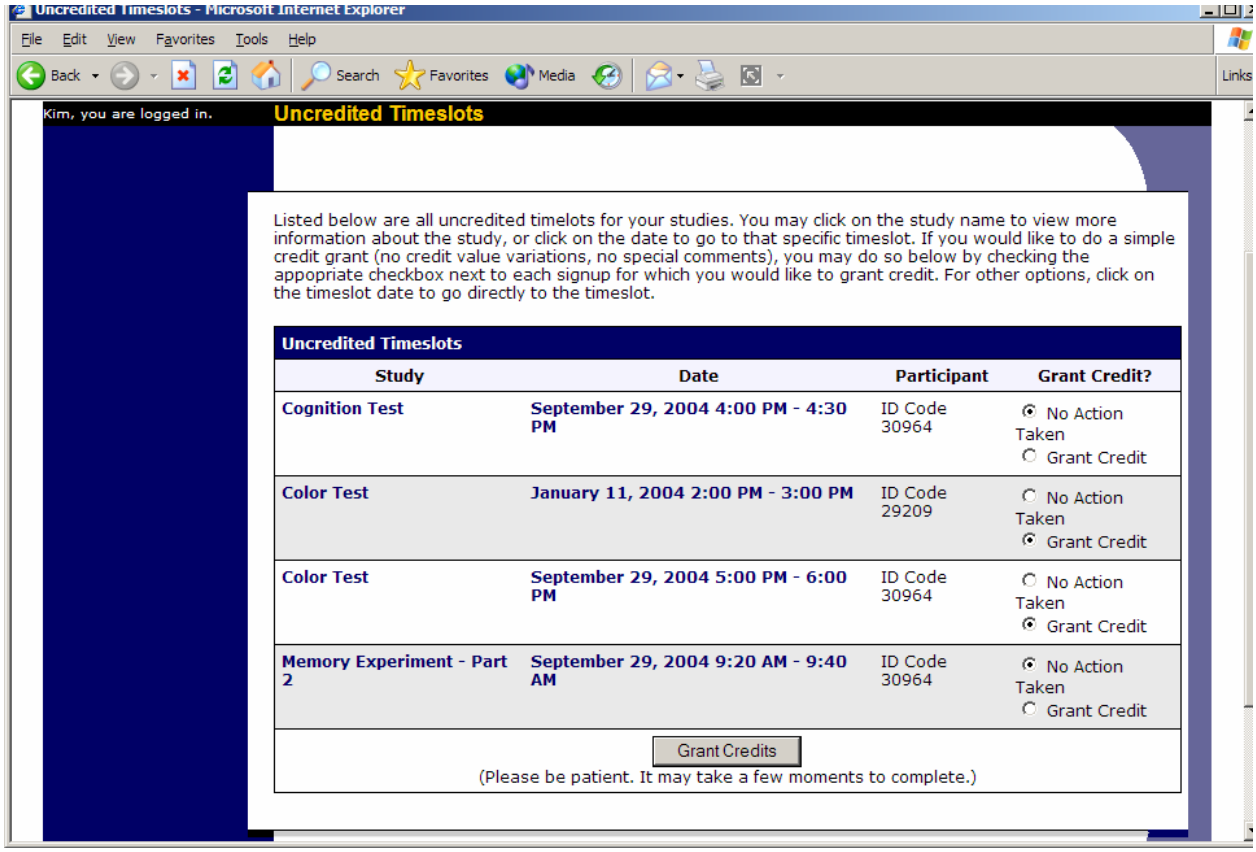
Viewing Uncredited Timeslots

When you login to the system, you will receive a warning if you have any timeslots that are more than 2 days old and haven't been dealt with. You may view a list of all timeslots that have not been dealt with by choosing the View Uncredited Timeslots option from the My Studies page. The default view will show in-person studies with timeslots in the past, as well as all uncredited timeslots for online studies. Timeslots for online studies, including those in the future, are always considered in need of a response. See the Web-Based (Online) Studies section of this documentation for more information.

If you would like to do a simple credit grant (standard credit grant, no comments), you may do so directly from this page. Select the desired sign-ups/timeslots, and then choose Grant Credits. The action may take a short time to complete, so please be patient while the credit grants are processed.

If you need to do something more complex, like mark a no-show, add comments, or perform a special credit grant with a non-standard credit amount, you can easily click on the timeslot's date and time, and go directly to that timeslot.

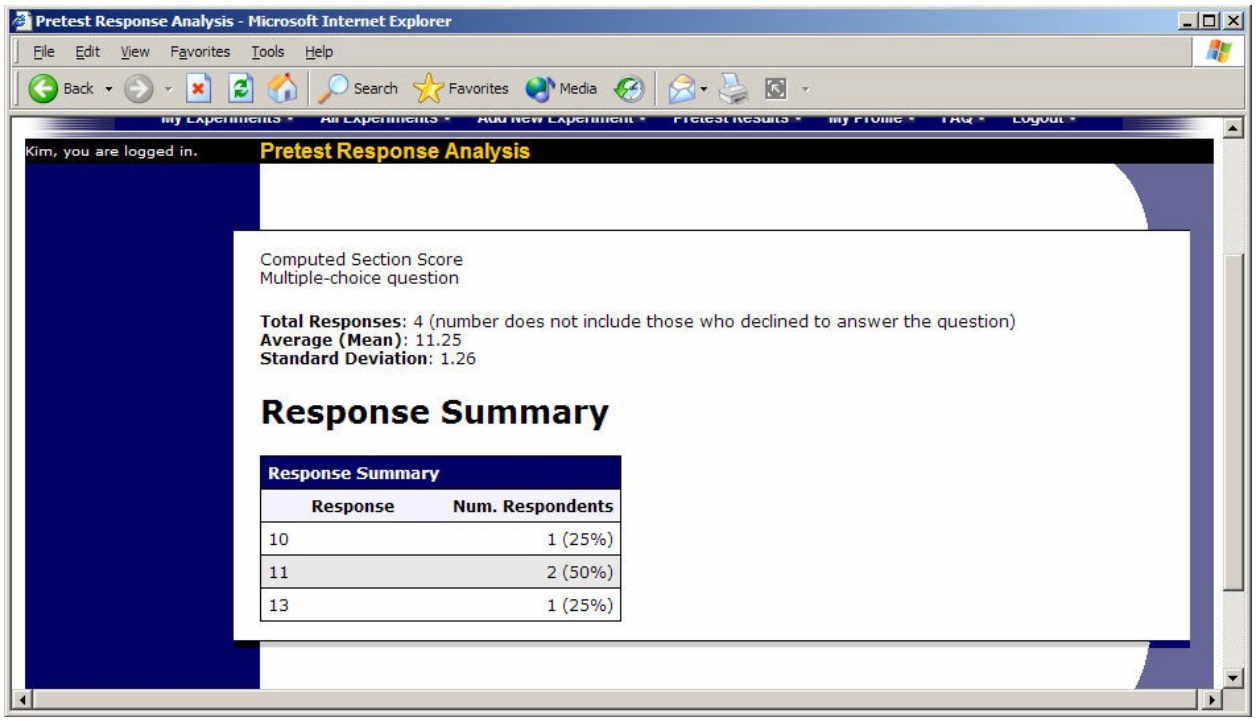
In cases where a study has timeslots linked to specific researchers, you will see the warning only for timeslots that are specifically linked to you, or to everyone in the study (i.e., not timeslots linked to someone else in the study). However, when you view uncredited timeslots, you will see all uncredited timeslots for your studies, even if someone else is linked to one of the timeslots for your study. This is done to make it easier to give your fellow researchers (for your studies) assistance in dealing with uncredited timeslots.



Analyzing Prescreen Responses

If online prescreening is enabled on your system, then you might also have the opportunity to analyze prescreen responses in aggregate or as raw data. Choose the Prescreen Results option from the top menu bar. You can then select which question you would like to analyze, and whether you would like to see summary data or raw data (in CSV format) for the selected question. The raw data will identify each participant only by a unique ID code, not by their name, for privacy reasons. If for some reason you need the participants' real names, ask the Administrator to run the same analysis, as they can also pull the real names with their report. This gives you access to all prescreen data across all participants in the system.

If you would like to analyze the prescreen data for just those who participated in your study, select the Download Prescreen Responses after clicking on your study. See Viewing Prescreen Responses in this documentation for further information.



Prescreen Qualification Analysis

If you would like to get an idea of how many participants meet a set of requirements (for help in setting prescreen restrictions on your study), use the Prescreen Qualification Analysis link from the Prescreen Responses page. Using this feature, you can select multiple questions (only questions that qualify for study participation restrictions are listed), and then the desired responses for those questions, and you will see how many participants meet that criteria.

If enabled, you may also contact participants and invite them to participate in any of your studies. See the Invite Qualified Participants to a Study section in this documentation for more information on how this works. The functionality is the same as the functionality described in that section, though a few options may be unavailable when not inviting directly from a study, because those options do not apply. Be sure to include information about how to sign up for the study in your communication to them, as a direct link to the study is not provided in the email.

Frequently Asked Questions (FAQ)

Why do I have to acknowledge the Human Subjects Policy?

Certain regulations and research guidelines either require or recommend it. You only need to do it once every 6 months, so it should not be too intrusive.

What is the best way to set up a study where participants receive monetary compensation?

You may set it up as a pay study only (indication a compensation amount), or a credit study if it is for both pay and credits. See the Studies for Pay section of this documentation for more information.

I want to set up a study so that participants can choose to receive credit or payment. How do I set this up?

Set it up as a study for credit, and note in the study description that participants may opt to receive payment instead, and they should notify the researcher of this when they come to their appointment. If the participant at that time chooses to receive monetary compensation, the researcher should grant credit, but mark the credit as 0 credits (Variable Credit Granting must be enabled in System Settings by the administrator), and then note in the comments for the timeslot that payment was received.

The monetary compensation a participant receives for a study depends on decisions they make during the study. How do I indicate this?

You must enter one value when setting up the study, so enter the minimum payment value (or 0) or whatever you feel is appropriate, and then note in the study description the entire range of compensation that is possible.

I want a participant to participate in an upcoming session, but the system says it is too late for them to sign up. What do I do?

If enabled, you can perform a manual sign-up. See the Manual Sign-Up section of this documentation. If not enabled, your administrator can still perform a manual sign-up.

Where are email notifications to me sent?

Email notifications (e.g. sign-up notices) are sent to either an address derived from your user ID or your alternate email address. See the Email Address Options section of this documentation for more information.

How do I deal with dyads?

A dyad is a study which requires a pair of people to participate, but often the second participant is not a “real” participant, but rather a colleague of the researcher who is “colluding” with the researcher as part of the study itself.

You do not need to deal with dyads in the system itself. Participants cannot see how many people have signed up for a timeslot, nor how many spaces are available for a timeslot. So, your “fake” participant can just act like a real participant and the real participant will be unaware of this.

I have finished running my study. What should I do?

So it does not clutter the list of studies for participants, you should make the study Inactive. See the Updating a Study section of this documentation for more information.

Who has access to my studies?

All users can see the information about your studies and the available timeslots. Administrators, the principal investigator, and the researchers for the study are the only people who can see who has signed up, and modify the study.

Regulatory Compliance Guidelines

Introduction

This software complies with all major regulations governing human subject research and privacy of data stored online. The system complies with both HIPAA and Common Rule for customers in the United States. For customers in Canada, it complies with the Personal Information Protection and Electronic Documents Act as well as the Tri-Council Statement. For customers in the European Union or in countries that follow OECD rules, it complies with OECD privacy rules and the European Union Directive of Data Protection. Your organization may or may not need to comply with the relevant regulations. Your subject pool administrator can advise you on this situation.

Even if you are not required to comply, compliance is still a good idea, as protecting sensitive data is always a good thing. Compliance in the context of this system is as simple as reading the remaining paragraphs of this section (that apply to your organization) and following the guidelines contained therein. The remaining compliance issues involving software, privacy and electronic data storage are all handled automatically by the software. You should still consult with your IRB or organization to learn about additional compliance rules you must follow outside of use of this software (the handling of the data you collect during your study would be one example).

Some regulations (particularly the US HIPAA regulations) are focused primarily on health data. You may think the system does not store confidential health data (in HIPAA terms, it is called PHI -- Protected Health Information), but depending on how your organization uses the software, there may very well be confidential data in the system.
Consider the case of a study that requires that a participant come from a family that has a history of mental illness. Merely knowing who signed up for that study can be considered confidential because that type of information should not be revealed to the public. It may turn out that your studies are not of such a nature, but even more benign situations, like a study that requires that participants be regular contact lens wearers, can be construed as confidential information. Organizations typically err on the side of caution given the criminal and civil penalties for violation of these types of regulations.

Data Handling and Security Guidelines

In your role, you have access to your studies and you can see who has signed up for those studies. You may also have access to prescreen responses. Because of these privileges, you should follow these simple guidelines:

- Secure Your Account. Use a password that is difficult to guess. The most secure passwords contain a combination of letters and numbers, do not spell a real word, and are at least 8 characters long. Your university IT department can provide you with assistance on choosing a secure password.
- Secure Your Work Area. If you are logged into the system and you leave your computer, you should logout of the system or use a password lock on your computer. Ask your network administrator for help with setting up a password lock.
- Handle Paper Documents Carefully. Any printouts from the system should be kept reasonably secure. Store them in desk drawer out of the public view. Documents you decide to discard should be shredded if possible.

Human Subjects/Privacy Policy Acknowledgment

Upon your first login to the system, and every 6 months thereafter, you may be required to acknowledge your organization's policy on these matters, and this acknowledgement will be logged. Ask your subject pool administrator if you have any questions.