Wolfram Syndrome Global Patient Registry Procedure for Registering, Consenting and Completing Surveys

1. Go to the Wolfram Syndrome Global Patient Registry **website** at: https://wsglobalregistry.iamrare.org.

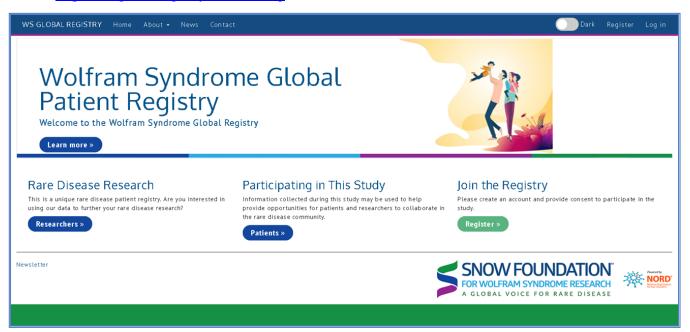


Figure 1. Landing Page

- 2. Click on the green **Register** button.
- 3. Complete the age attestation.



Figure 2. Age attestation

- 4. Read the **Terms and Conditions** document.
- 5. Fill in the requested information.
- 6. **Opt in or out** of reasons to be contacted by Study Staff.
- 7. Agree to the Terms and Conditions.
- 8. Click Create Account button.

Register You must be a lega	al adult (at least 18 years of age	e, or the age of majority in your state, province, or country) to register.
↑ Home / Register		
Please fill out the form	until all 6 symbols turn into a 2 sym	mhol
		years of age, or the age of majority in your state, province, or
country)?	- , .	,
Yes 🖸		
First Name *		0
Middle Name		
Last Name *		•
Country of Residence	v] •
Username (Email		10
Address) *		J
Confirm Username *		0
	A password must be at least 8 characters long: - contain 1 uppercase letter - contain 1 lowercase letter - contain 1 digit - contain 1 digit - contain 1 special character - not contain text from top 1000 commonly used passwords	
Password *		•
Confirm Password *		•
Additional Con	tact Information (Optiona	al)
Home Phone		
Work Phone		
Mobile Phone		
Contact Prefere	ences	
Yes, I woul	d like to be contacted about	
	☐ Reminders to update my survey	responses
	☐ Clinical trials I may be eligible f	for †
	☐ Potentially donating biospecime	en and/or DNA for future research studies ††
	☐ Facebook Group	
Terms and Condition	ns	
• Viev	v Terms and Conditions 👢 Download as PDF	
☑ I ha	ve read and agree to the terms and conditions.	•
Cres		

Figure 3. Creating an account – Blank Registration Form

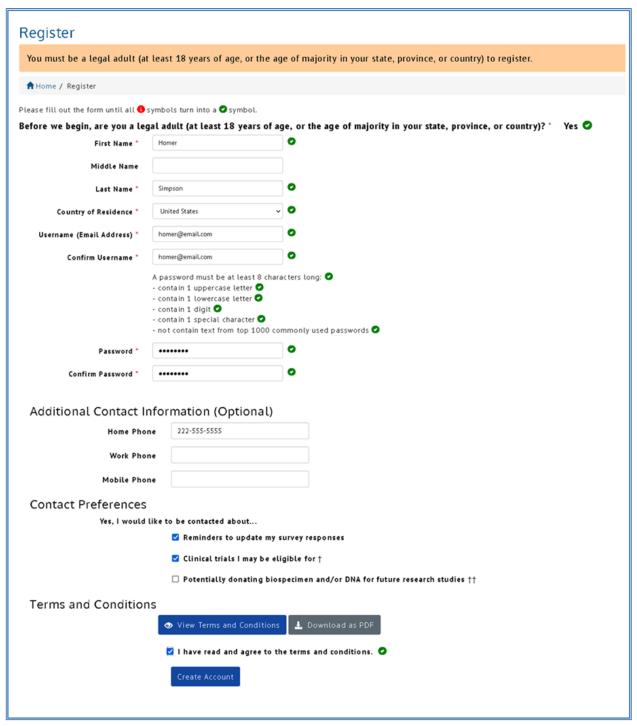


Figure 4. Creating an account – Completed Registration Form

- 9. A confirmation email will be sent to the email address that was provided during registration.
 - a. Confirm registration by copying the **Confirmation Token** from your email.
 - b. You can also confirm the email address by clicking the link in the email.

10. Verify Registration

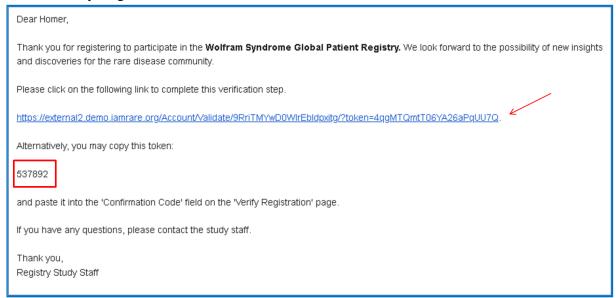


Figure 5. Registration confirmation email

Note: If the confirmation email has not been received after a few minutes, click **Resend confirmation email** at the bottom of the page. Be sure to check spam folders for the email confirmation.



Figure 6. Entering the confirmation code to complete registration

- 11. Click on **Participant Enrollment**.
- 12. Select the appropriate Option. You can add yourself, or someone else. Note: the Study Participant is the person who has the condition or diagnosis.

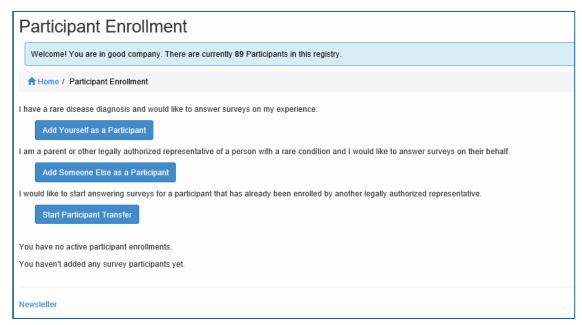


Figure 7. Adding a Participant

Alternatively, you can begin the process of transferring a Participant from one party to another. Speak to your study administrator if you have questions.

13. To enroll a Participant, fill in the fields as they apply to the **Study Participant**.

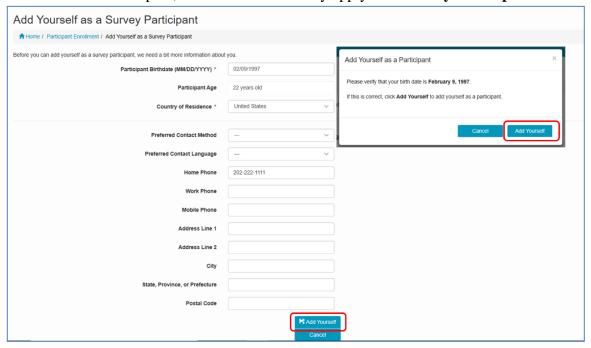


Figure 8. Adding Participants – Adding yourself as a Participant

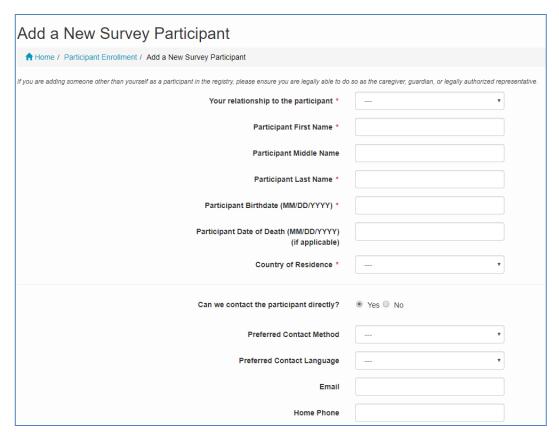


Figure 9. Adding participants - Adding another person as a Participant

14. Grant Consent



Figure 10. Accessing the Consent

Note: You must agree to all terms outlined in the Consent Form.

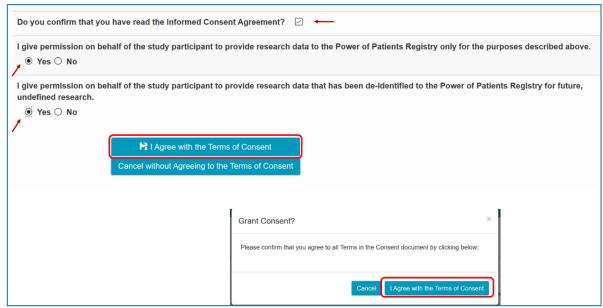


Figure 11. Granting Consent

15. Access Surveys by clicking on **Take Surveys**.



Figure 12. Participant Dashboard

- 16. Complete the **surveys** applicable to the **Study Participant**. Click the **Take** button next to the **Survey Title** to open the survey.
 - a. NOTE: The Reporter is listed in the top right hand corner of the page and the Study Participant is listed on the left side of the page. If someone is answering surveys for themself, both the Reporter and Study Participant fields will reflect the same name.

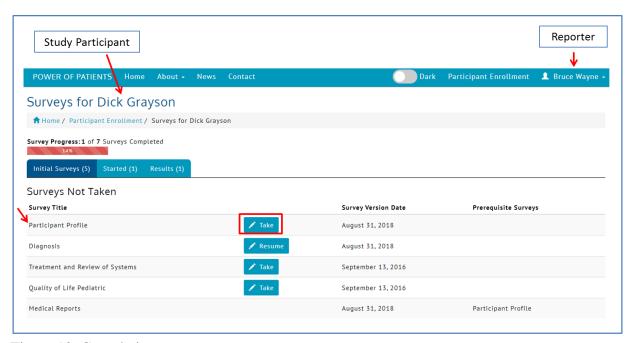


Figure 13. Completing surveys

17. Complete the survey. Questions marked with a red * require a response.

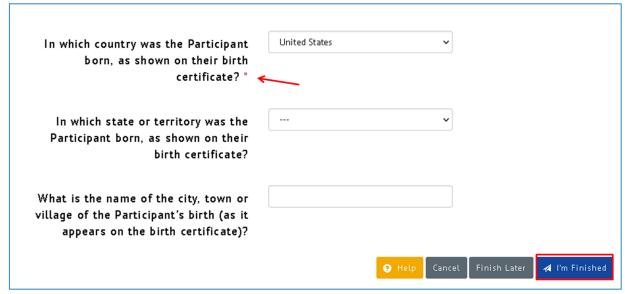


Figure 14. Completing questions and submitting surveys

18. When a survey has been completed, click on the **I'm Finished** button. If not completed, surveys can be saved as drafts by clicking the **Finish Later** button. Click the **Cancel** button to leave the survey and return later. Note that if you click Cancel, no changes will be saved.

19. Using the Calendar Feature

The registry platform supports date entry via a calendar feature.

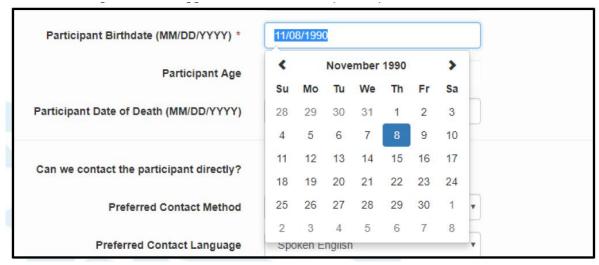


Figure 15. Date Field Displaying Calendar Widget

To use the calendar feature, click on the date field.

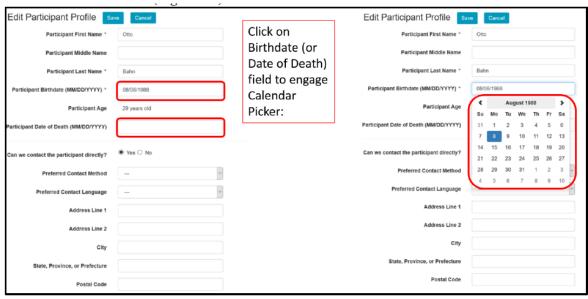


Figure 16. Using the Calendar Feature

To easily scroll through the calendar options, click on the header field. Clicking on the month will populate all the months. Clicking on the year will produce more years, etc. Use the left and right arrows to navigate to the desired time point.

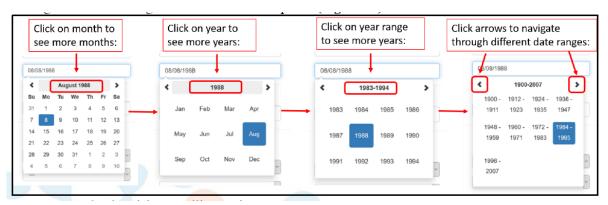


Figure 17. Calendar scrolling options

Note that dates can also be typed directly into the calendar field.

20. Updating and Retaking Surveys

Some surveys will be available to edit after submission and others will be set to a regular retake, or *longitudinal*, schedule. This supports the study of the disease over time. You may opt in for email reminders when retakes are due.

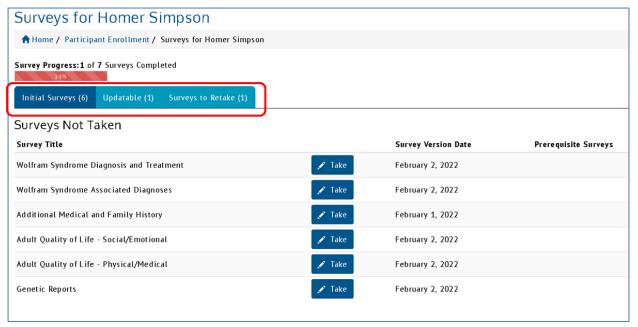


Figure 18. Participant Survey List

To update a survey, click on the **Updatable** tab and click the **Edit** button next to the survey. The survey will populate with the previous answers. Simply edit any information that has changed.

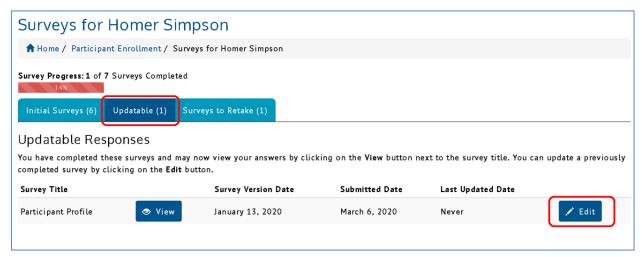


Figure 19. Accessing Updatable Surveys

To retake a survey, click on the **Surveys to Retake** tab and click the **Retake** button next to the survey. Longitudinal surveys will be blank each time, to capture fresh unbiased information.



Figure 20. Retaking Longitudinal Surveys

Here is a list of updatable and longitudinal surveys for this registry:*

Updatable	Longitudinal
Participant Profile	Adult Quality of Life - Social/Emotional
Wolfram Syndrome Diagnosis and Treatment	Adult Quality of Life - Physical/Medical
Wolfram Syndrome Associated Diagnoses	Pediatric Quality of Life - Social/Emotional
Additional Medical and Family History	Pediatric Quality of Life - Developmental/Physical
Genetic Reports	

^{*}Note: This list may change over time.