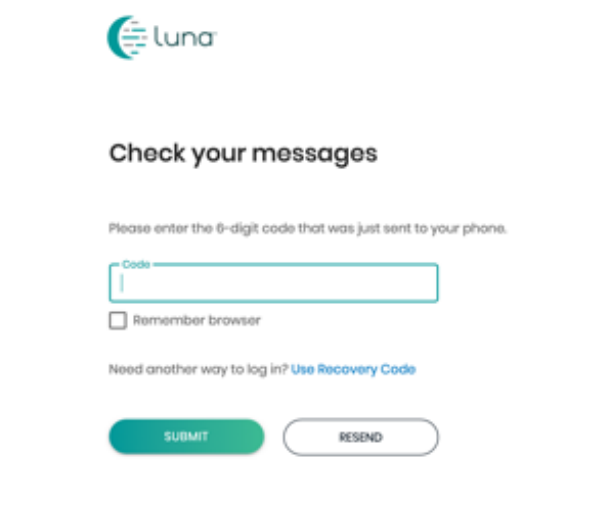
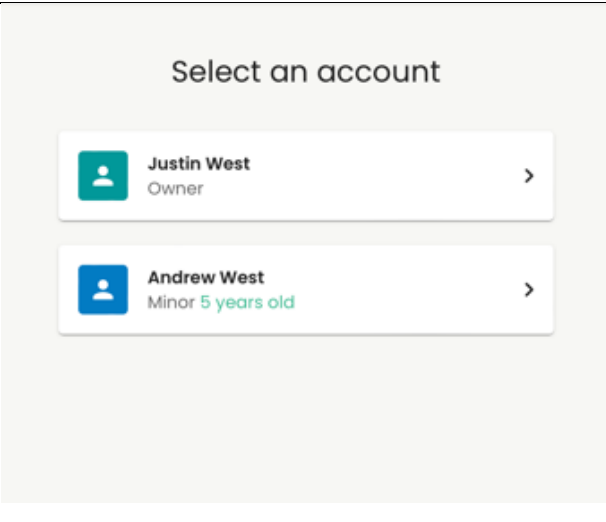


Join the KCNT1 Longitudinal Registry Study

Follow these steps to join the new longitudinal study.

<p>If you have not already, go to LunaDNA.com and log in.</p>	<p>https://id.lunadna.com/referrer/kcnt1-longitudinal?studyName=kcnt1longtermregstudy</p>
<p>Login to your account by entering the email and password for your account. You will then receive a 6-digit code to the cell phone or Google Voice number associated with your account. Enter that to proceed.</p>	
<p>Select your minor account (in blue). If you do not have a minor account, see our account set-up guide for existing users. <i>It is very important to take surveys about your child's experience from within their account.</i></p>	

You should land on your child's or ward's dashboard. Confirm this at the top of the page where it should say: "Acting on behalf of [Name of Child or Ward]."

Scroll down a little to the "My Communities" section.

You will see a box for the "KCNT1 Epilepsy Foundation". Click the box to go to the Foundation's Community Page.

On the Foundation's community page, scroll down to the "Studies" section.

You should see a box for the KCNT1 Longitudinal Registry Study. Click the box to go to the study page.

The screenshot shows the Luna dashboard interface. At the top, the user is identified as "Acting on behalf of Andrew West". The dashboard features three main sections: "DNA Files" (with an "UPLOAD A DNA FILE" button), "Health Accounts" (with a "VIEW HEALTH ACCOUNT INFO" button), and "You Tell Us" (with a "GO TO STUDY" button). Below these is a notification for "33 unread messages". The "MY COMMUNITIES" section is visible, with a purple arrow pointing to the "KCNT1 Epilepsy Foundation" card. Below this, the "MESSAGE CENTER" shows "No Messages". The "STUDIES" section contains two cards: "KCNT1 Children's Study" (marked "QUALIFIED") and "KCNT1 Longitudinal Registry Study" (marked "LAUNCHED"). A purple arrow points to the "KCNT1 Longitudinal Registry Study" card. At the bottom right, there is a "LEAVE COMMUNITY" button.

Follow these instructions to join the study and take surveys.

If you have not already, make sure you are in your minor account. Then, from within the study, click the “Join Study” button. Then click through to the “Prerequisites” screen using the link at the top.

If you have already joined the study you can skip this step.

The prerequisite is a little different. We can replace this screenshot when it is live.

On the prerequisites screen, answer the question confirming whether your child or ward has a diagnosis with a KCNT1 variant. Next, navigate to the “Data Requests” screen to begin the surveys.

If you have already joined the study you can skip this step.

The screenshot shows the Luna dashboard for the KCNT1 Children's Study. At the top, there are navigation links for DASHBOARD, MY DATA, and ALL STUDIES. The main header displays 'ALL STUDIES > KCNT1 CHILDREN'S STUDY' with status indicators for 'STUDY ENROLLMENT OPEN' and 'DATA COLLECTION'. A summary bar shows 157 STUDY PARTICIPANTS, with 128 WITH ANY SURVEYS, 0 WITH ANY DNA FILES, and 0 WITH CONNECTED HEALTH RECORDS. A progress bar below this shows four steps: OVERVIEW (active), CONSENTS, PREREQUISITES, and DATA REQUESTS. A central message reads: 'First things first – you'll need to join the study before you can proceed.' Below this, the 'KCNT1 CHILDREN'S STUDY OVERVIEW' section shows the status as 'Launched' and a description of the study. A 'JOIN STUDY' button is visible. To the right, an 'INSIGHTS' section shows 'STUDY INSIGHTS UNAVAILABLE' with a small bar chart icon.

The screenshot shows the Luna dashboard for the KCNT1 Children's Study, specifically the Prerequisites screen. The summary bar now shows 164 STUDY PARTICIPANTS, with 128 WITH ANY SURVEYS, 0 WITH ANY DNA FILES, and 0 WITH CONNECTED HEALTH RECORDS. The progress bar shows four steps: OVERVIEW, CONSENTS, PREREQUISITES (active), and DATA REQUESTS. A central message reads: 'You need to complete all of the study's prerequisites listed here before you can move on.' Below this, the 'PREREQUISITES' section shows a question: 'Diagnosis of Genetic Epilepsy Yes/no question' with a 'VIEW' button.

This page shows what **surveys** or “Data Requests” are available to complete.

Click **Begin** on the KCNT1 Demographics and Genetics survey. (Available in German, Portuguese, Italian, Spanish, Swedish, French and English. You can choose your language before beginning the survey.)

Once you have completed the KCNT1 Demographics and Genetics survey, please complete any other remaining data requests. Not all surveys are available in all languages. We will let you know as we add new surveys to the study!

The screenshot displays the Luna dashboard for the KCNT1 Children's Study. At the top, there are navigation links for DASHBOARD, MY DATA, ALL STUDIES, and MENU. Below this, the user is identified as acting on behalf of Andrew West. The dashboard shows 150 study participants, with 112 having any surveys, 0 having any DNA files, and 0 having connected health records. A progress bar indicates the completion status of four steps: OVERVIEW (100%), CONSENTS (No consents), PREREQUISITES (1/2 complete), and DATA REQUESTS (1/2 complete). A message prompts the user to complete 2 data requests. Two survey request cards are shown: 'Genetic Epilepsy in Minors' (Survey complete) and 'KCNT1 Top Tasks and Seizure Classification' (New Survey). The 'Begin' button on the second card is circled in purple.