

## Rare Patient Network Partnership Levels

Partnership Types	Community Partner	Strategic Partner
<b>Who qualifies</b>	Every organization serving patients who meet the eligibility criteria: patient have at least one seizure before age of 18 and/or has developmental delays.	<ul style="list-style-type: none"> <li>→ The organization is an established Community Partner               <ul style="list-style-type: none"> <li>• Referred Ciitizen to other PAGs/Advocates</li> <li>• Added logo and Rare Patient Network description on their website</li> <li>• Actively recruited patients - social media, 1:1, webinar etc. Copies of recruitment efforts will be required</li> </ul> </li> <li>→ The organization has successfully recruited 25% of the group's US patient population</li> <li>→ The organization is able to cover the cost of data extraction and delivery or has a study sponsor</li> <li>→ The organization has identified a Principal Investigator (PI) for their data analysis and is able to submit an IRB approved protocol with Category 8 exemption.</li> </ul>
<b>Requirements</b>	<ul style="list-style-type: none"> <li>→ Allow Invitae to add your organization name and logo to the RPN landing page</li> <li>→ Promote the RPN and recruit patients using your unique link via social media, newsletter, website or other channels</li> </ul>	<ul style="list-style-type: none"> <li>→ Allow Invitae to add your organization name and logo to the RPN landing page</li> <li>→ Promote the RPN and recruit patients using your unique link via social media, newsletter, website or other channels</li> <li>→ Sign a partnership agreement</li> <li>→ Introduce us to your pharma partners</li> </ul>
<b>Benefits</b>	<ul style="list-style-type: none"> <li>→ Receive \$100 for each patient referred through the organization's unique link when patient-consented data is shared with biopharma to advance scientific research</li> <li>→ Receive the following reports quarterly:               <ul style="list-style-type: none"> <li>• Number of patients referred and their status</li> <li>• Number of patients enrolled in the RPN broken down by gene</li> </ul> </li> <li>→ Your patients will get access to the medical records we collect on their behalf. Extracted clinical data will be available if they are participating in an active research study</li> <li>→ Stepping stone to become a Strategic Partner at no risk</li> </ul>	<ul style="list-style-type: none"> <li>→ Obtain best in class, research-ready natural history datasets that is extracted from patient medical records, normalized and organized</li> <li>→ Natural history dataset will be delivered on a confirmed date and can then be shared externally to the organization, academic researchers and/or pharma partners that have submitted the Category 8 IRB protocol and signed the research agreement</li> <li>→ Specific gene(s) added to onboarding drop-down list on the Ciitizen platform</li> <li>→ Invitae will contact registered patients for additional consents (advocacy or other partners)</li> <li>→ Access to recruitment dashboard - includes consented patients' contact information and account status</li> <li>→ Access to the Invitae Explorer data platform: Ciitizen data and Genetic Testing data</li> <li>→ Advocacy information added to the Positive Report Guide sent to patients newly diagnosed through Invitae</li> <li>→ Invitae will send an email to Invitae tested patients including your organization's welcome letter</li> <li>→ Option to schedule a community specific webinar with Invitae</li> <li>→ Receive ten percent (10%) of Net Sales for each patient referred through the organization's unique link when patient-consented data is shared with biopharma to advance scientific research</li> <li>→ \$1,500 annual conference sponsorship</li> </ul>