Thank you for being a part of the Pacific Sickle Cell Regional Collaborative (PSCRC) Minimum Data Set! Our goal was to understand more about adults and children with sickle cell disease seen in seven Pacific Region states.

We asked adults with sickle cell disease (18 years and older) and parents of children with sickle cell disease about experiences with hydroxyurea, and about any concerns you might have with the medicine. Hydroxyurea is a medicine that can help children and adults with sickle cell disease by decreasing pain episodes, days in the hospital and episodes of acute chest syndrome.

169 adults with sickle cell disease and 238 parents of children with sickle cell disease from all seven PSCRC states filled out the surveys. Half were males and half were females. 93% were African American and 74% had the diagnosis of SS disease.

The PSCRC has expanded to 13 Western states. PSCRC providers are working together to figure out the best ways to support adults with sickle cell disease and parents of children with sickle cell disease about hydroxyurea so that you fully understand the benefits of the medicine and any risks. We know that hydroxyurea is safe when you work closely with your health care provider. Your doctor or nurse will give you instructions about getting blood work, about how often to come into clinic and about the right dose for you or your child.

Talk with your doctor or nurse if you have any questions about whether or not hydroxyurea is right for you. If you were prescribed hydroxyurea but have any concerns or barriers to taking it, let your doctor or nurse know. You can come up with a plan together with your health care provider that will lead to your best health!

For more information, visit: pacificscd.org or casicklecell.org