



Priority Research Questions—Adolescent Patients in San Francisco

1. How can patients better keep track of care management including symptoms, appointments, and other things important to report?
2. How can we make the search for a diagnosis less burdensome on the patient?
3. What will happen after I come off my parent's insurance? How will I cover medical costs?
4. What are long term effects of drugs and other treatments for chronic conditions?
5. What do you do if you can't go to school full-time? How can doctors, parents, and teachers work together?
6. How can things be made easier for parents-missing work, transportation, and researching things?
7. How to promote self-care/independence for patients so parents aren't doing everything-be more involved in their own care?
8. How can symptom data be better shared and tracked and viewed in real time using technology?
9. How can we help young people with chronic illness connect with each other?