

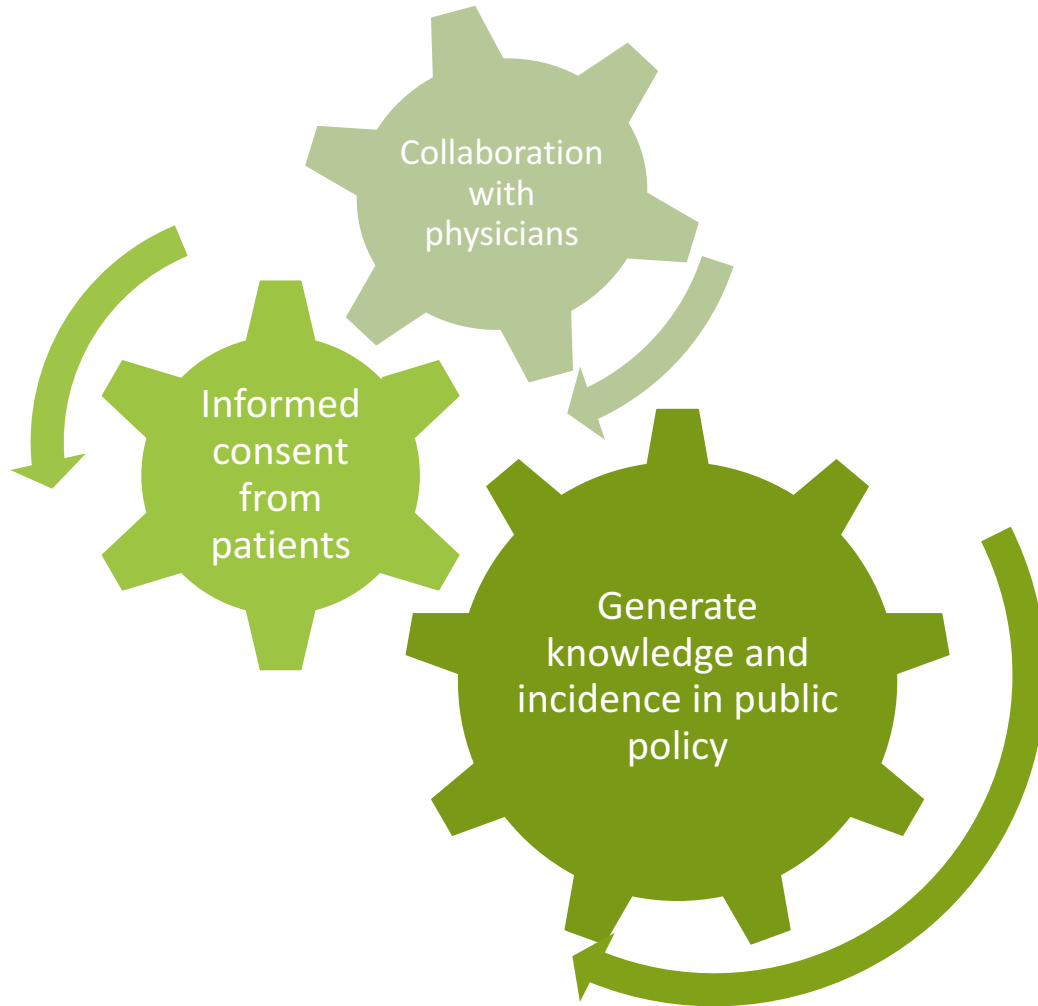


THE ROLE OF A PATIENT REGISTRY IN DRUG ACCESS AND COVERAGE IN MEXICO

WHAT DO WE DO?

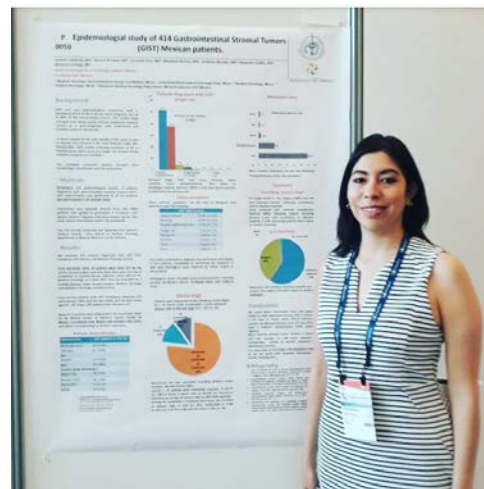
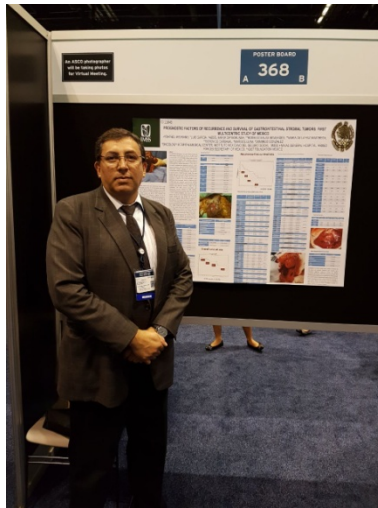
- Collect data from patients
- Follow up frequently and empower patients with information so they can take control of the disease (questions for physicians, contraindicated drugs, how to manage side effects)
- Generate data so we understand better the development of the disease in our countries
- Publish scientific information with physicians
- Use the knowledge acquired for incidence in public policy (access to treatments: Ley Ricarde Soto, Seguro Popular, etc.)

HOW DO WE DO IT?



SALUD CON DATOS

- Gathered for the first time medical and patient stakeholders and decided to start the project in Mexico, Chile and Argentina



	Patients	Papers
México	380	2
Chile	200	Collaboration with Universidad Católica de Chile
Argentina	250	In progress

GOALS

SHORT TERM

- Generate joint papers with physicians from the 3 countries and obtain information from the region
- Pharmacovigilance

LONG TERM

- Use the network to promote clinical trials