

# Part II+ Project

# New Consent Project: Part 2+ Data Sharing

## Opportunity for new data sharing to support whole patient care

- Made possible by VITL's new platform and spurred by requirement for exchange of DA data
- Using opportunity to create universal policies and procedures for sharing Part 2 data, and possibly other sensitive data types, on a patient-driven opt-in basis

## Advisory Workgroup Convened

- DVHA: Emily Richards, Naomi Hahr, Mary Kate Mohlman
- VITL: Carolyn Stone, Maurine Gilbert, (consultant)
- VCP: Ken Gingras
- BiState: Heather Skeels
- Looking for representative of SUD treatment providers

# Workstreams & Stakeholder Engagement

## Part 2+ Project Workstreams

- Policy and procedure development
- Technical development and implementation
- Provider education and onboarding to new consent processes
- Patient education
- And stakeholder engagement (to inform all areas above)

## Stakeholder engagement opportunities

- Introductory meetings to explore provider / health care organization perspectives and preferences, to inform options
- Will use more structured engagement including interviews and workshops to explore specific design questions and inform development of provider and patient education strategy, materials, and consent processes
- Participants will include patients, providers, health care organizations, and advocacy groups
- Reach out to participate or refer participants: [mgilbert@vitl.net](mailto:mgilbert@vitl.net)

# Part 2+ Project Draft Timeline

## April and May

- Met with other HIEs to understand their approaches ([summarized here](#))

## June

- Convened Advisory Workgroup

## July

- Begin stakeholder engagement
- Define a set of options for how we might manage Part 2+ data

## September

- Debrief Advisory Workgroup on stakeholder engagement to date, identify an option to move forward based on stakeholder feedback

## October

- Develop draft policy and procedures
- Begin technical work to prepare for ingesting new data

## November

- Develop agreements with DAs, begin interface projects
- Begin education and outreach strategy development, with provider and patient input

## December

- Final preparations for new platform launch
- Proposal to HIE Steering Committee of updated consent policy and procedures for managing DA data

## February 2021

- Finalize configuration and testing of Part 2+ consent management, data ingestion, view filtering

## TBD in 2021

- Begin patient education about options
- VHIE begins making Part 2+ data available to providers