CRISP

Consent Tool Webinar

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Agenda

- Purpose of the CRISP Consent Tool
- SUD and MH Data Workflow
- Next Steps
- Demo of the Tool
- Questions

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Purpose of the Consent Tool

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- SUD data is protected by regulation 42 CFR Part 2
- Priority of MDH and Medicaid to share SUD data through the HIE for purposes of care coordination between SUD providers and other healthcare providers
- Consent management solution = consent tool
- Goal: ease workflow burden when obtaining and disclosing information

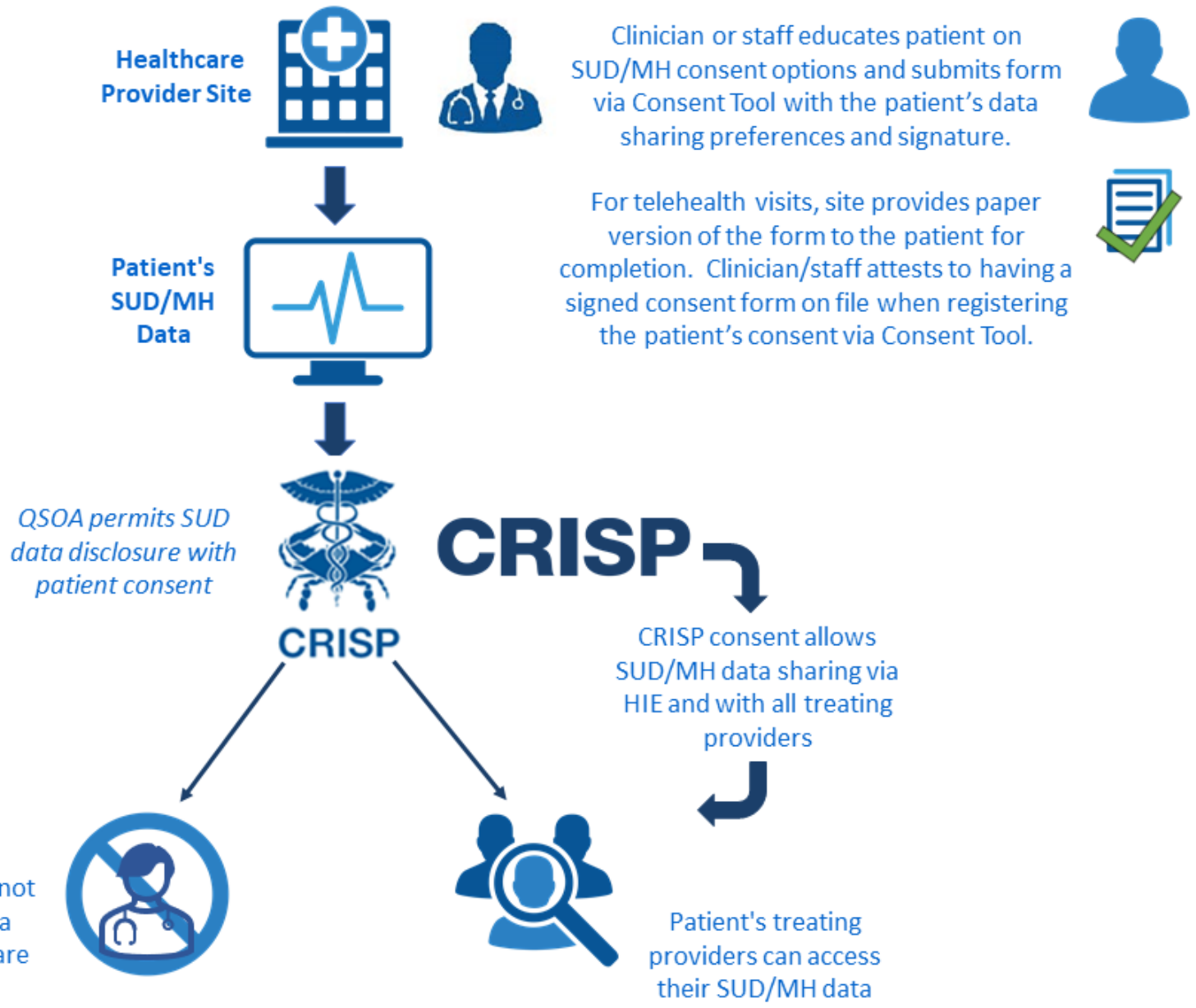
What Is My Patient Consenting To?:

- To allow their SUD or MH data to be shared with members of their patient care team who participate with CRISP
  - Option to share all SUD treatment data, or just their SUD providers contact information

CRISP Consent Form – Paper Version

- Patient must complete and sign the CRISP Consent Form prior to provider registering consent in the tool
  - Form can be found **here**
- Please keep the consent on file
- Afterwards, complete the registration via the Consent Tool. The patient's SUD and MH information will stay masked until the consent is registered via the Consent Tool online

SUD/MH Data Flow Diagram



Next Steps

- Working on getting more SUD and MH information flowing in the HIE
- Sensitive document repository is built and ready for organizations to begin sending sensitive CCDs to CRISP

Consent Tool Materials

- Training and informational materials can be found on our website
  - One Pager
  - Training Guide
  - FAQS
  - Consent Paper Form

https://www.crisphealth.org/consent-tool/

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Consent Tool Demo

Login - Unified Landing Page (crisphealth.org)

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Questions?

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