

Consent Tool User Guide for HIE Portal and SSO

Updated 1/16/2025

10480 Little Patuxent Parkway, Suite 800 Columbia, MD 21044 877.952.7477 | info@crisphealth.org www.crisphealth.org



Consent Tool Overview

Purpose of the Consent Tool



- The CRISP Consent Tool was designed as a platform for providers and staff to register patient consents to share 42 CFR Part 2protected data.
- This application has since grown to cover other unique scenarios where patients may need to "opt in" to sharing additional data types via the HIE
- What is my patient consenting to with a 42 CFR Part 2 consent?
 - To allow their 42 CFR Part 2-covered provider to share information about their SUD treatment via the Health Information Exchange (HIE).
 - The HIE will then share it with other members of the patient's health care team who participate with CRISP HIEs
 - Including Maryland, DC, West Virginia, Connecticut, Alaska, Virginia, Rhode Island, and any HIE affiliates in the future.
- Find our complete list of FAQs <u>here</u>.



Registering Consents Already on File

Registering Consents Already on File



- If the consent has been captured outside of the HIE portal, a credentialed staff member may complete the registration in the Consent Tool, based on the patient's designation, before checking the "Attestation for Consent on File" box in the signature section.
- Please keep the signed copy of the consent form on file. It is required by federal law to have a patient signature to share the patient's SUD information available upon request.



Registering a Consent During In-Person Visits

Using the CRISP Consent Form for In-Person Visits



- HIE user searches for their patient in HIE Portal or through SSO in their EHR.
- After launching the tool, provider explains the consent to their patient, educating them on what data they are sharing and with whom.
- Patient designates to share "all SUD data".
- Patient (or parent/guardian) signs directly in the tool during the in-person visit.
- The provider registers their own legal attestations in the tool and then adds their name before submitting the consent.



Steps to Register a Consent via Single Sign-On (SSO) from an EHR

Launch the Consent Tool from the InContext App in your EHR



- Click on the consent tool tab on the left-hand side of your screen
- The consent tool will open in a new tab in a new window
- Follow the Portal registration instructions (on subsequent slides) to register the patient's consent the same way as you would via the HIE Portal

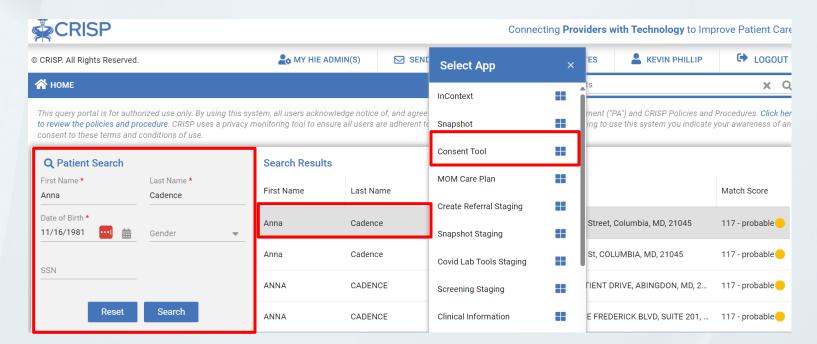


Steps to Register a Consent in the CRISP Portal

Launch the CRISP Portal and Search for a Patient



- Enter patient name and date of birth into Patient Search
- Select the patient from search results returned
- Click on the square icon next to the Consent Tool app







- If you are registering consent for a new patient (one not currently on your existing CRISP panel), a "Attest to Relationship" prompt will appear.
- Click 'Proceed' to continue.

Attest to Relationship





This patient is not currently linked to a active or existing relationship at your organization. As a reminder, CRISP prohibits access to patient records where there is no active relationship. All access to patient records are monitored. Do you wish to continue?

PROCEED

CANCEL





 After clicking "proceed", you must select a reason for searching the patient. Please select the option that applies to you.

Please select a reason ×				
Reason: New patient	Treatment	Care coordination	Quality improvement	O Public health
				SUBMIT

Select Part 2 Form





Consent Types

Provider Consent - Patient Consent to Disclose SUD and MH Treatment Information

Prevention of Harm - Block Patient Access Form

Review the Information Section with patient, using the Accounting of Disclosures and



- FAQ links as needed
 - https://www.crisphealth.org/consent-tool/#sharing-data--faq



Information about this Consent

By completing and signing this form, you will be allowing your 42 CFR Part 2 - Substance Use Disorder treatment provider to share information about your 42 CFR Part 2 - Substance Use Disorder treatment with CRISP who may share it with other members of your health care team for purpose of treatment, payment, and health care operations (TPO).

Examples of who may see your information include, but may not be limited to, your primary care provider, hospital and emergency providers, case managers or care coordinators, your insurance company or payer, and other individuals who are involved in coordination or payment of your care. The information will be shared with members of your healthcare team who participate with the CRISP Shared Services affiliate HIEs including Maryland, DC, West Virginia, Connecticut, Alaska and any HIE affiliates in the future.

Anyone receiving your information must follow all state and federal laws to keep your information private; however, there is the potential for the records used or disclosed pursuant to the consent to be redisclosed by the entities receiving the information and the information may no longer be protected by 42 CFR Part 2 (the federal regulation which protects the privacy of substance use disorder (SUD) information). Once your SUD information is shared with members of your health care team for purposes of treatment, payment, or operations, they may incorporate it into their records and further share it with other health care providers, payers, or organizations that provide services for them. Your information may be redisclosed or shared in accordance with HIPAA regulations, except for uses and disclosures for civil, criminal, administrative, and legislative proceedings against you, the patient.

You can request a list of organizations who have received your information by completing an accounting of disclosures requests at https://disclosures.crisphealth.org. A list of Frequently Asked Questions (FAQ) about sharing 42 CFR Part 2 -- Substance Use Disorder treatment data through CRISP can be found here and at https://www.crisphealth.org/consenttool/#:~:text=Sharing%20Your%20Substance%20Use%20Disorder%20(SUD)%20or%20Mental%20Health%20(MH)%20Treatment%20Data%20Through%20CRISP%20FAQ.

CRISP does not require you to sign this consent, and it will not impact the sharing of any of your health information through the HIE, except for your 42 CFR Part 2 —Substance Use Disorder information. If you do not consent to the disclosure of your SUD information, it may not be readily available through CRISP to those who need the information to give you appropriate care, especially in an emergency.

Patient must elect to share ALL SUD information with this form



CRISP Consent

Consent History

Consent Status: Opted to Disclose All SUD Treatment Data, Expiration Date: Does Not Expire

Dismiss

Next

Type and Amount of Data and Purpose of Disclosure

Purpose

The information shared will be used for purposes of treatment, payment, and health care operations as defined by HIPAA. The information to be shared could include but may not be limited to clinical documents, lab results, hospital discharge summaries, medication information, and claims data relating to my Substance Use Disorder treatment.

Consent Options

Disclose All Substance Use Disorder and Mental Health Data for TPO Purposes

This information could include my treatment plan, medications, laboratory results, clinical notes, health care encounters, claims information, and other data about my Substance Use Disorder and/or Mental Health care.

Review Submission Instructions CRI System for one Section



- Please review instructions for each type of visit carefully.
- For telehealth visits, or consents that are otherwise obtained and on file, please make sure to have the CRISP 42 CFR Part 2 SUD Consent Form (or a substantially similar form) signed and completed by the patient before attesting to having the consent on file in the tool.

Review the revocation and expiration sections

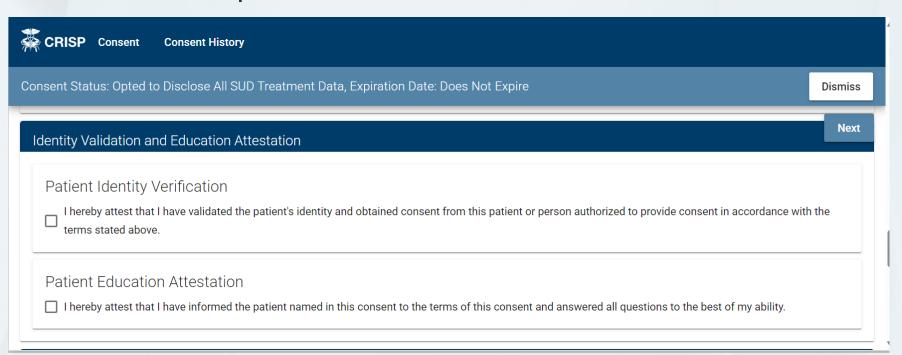


- These sections explain the process and result of revoking this consent, which the patient may do at any time.
- The expiration date will automatically be blank. In the Choose a
 Date field, the patient may select whichever expiration date they
 would like. If they would prefer that this registration does not
 expire, providers may use the toggle in this section to indicate
 this preference.
- For telehealth patients, this must be edited to match the expiration date on the SUD Consent Form, as indicated by the patient
- For in-person visits, this date may be updated to anything in the future based on discussions with your patient

Complete Provider Attestations



- Providers/staff obtaining patient consent must attest that they have:
 - (1) Verified the patient identity and;
 - (2) Informed the patient of all terms of the consent.







- For in-person registration:
 - Patient enters electronic signature using a mouse, stylus pen, or finger via touchscreen/ signature pad.
- For registrations of consents on file:
 - Check the box under "Attestation for Consent on File."
 - Note: The CRISP SUD Consent Form, or a substantially similar form, must be completed by the patient before attesting.

Legal guardian, parent, or legally authorized representative signature (as applicable)

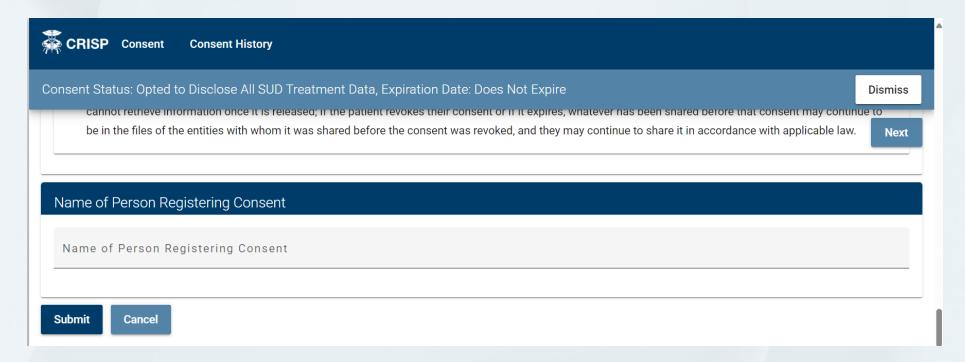


- First checkbox only required if the person signing the consent is the patient's legal guardian, parent, or legally authorized representative and has the legal authority to consent on the patient's behalf.
- Second checkbox allows for capture of both patient's AND legal representative's signatures.
- The person signing on behalf of a patient MUST enter their name into the form and electronically sign.

Submit Consent



- Enter the name of the person registering this consent.
- Click "Submit" once to avoid duplicate entries.
- Click "Print and Exit" or "Exit."



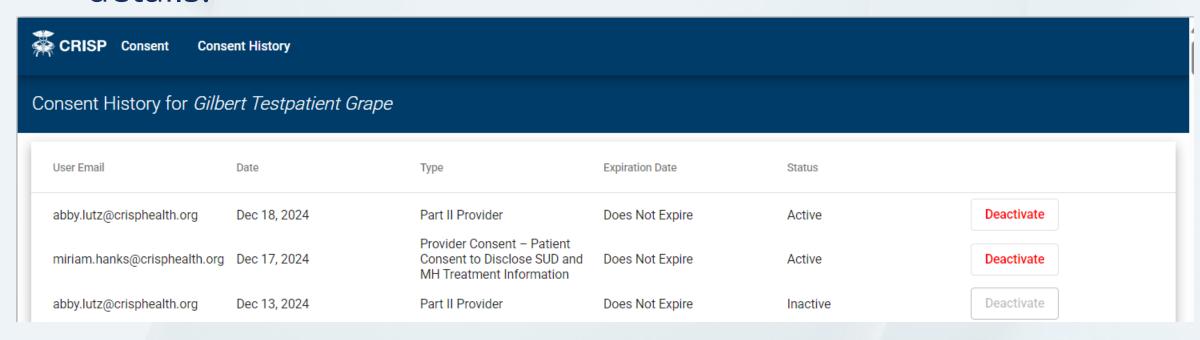


Additional Functions in the Consent Tool

How to view consent history



- After searching for your patient, click "Consent History."
- Click on a row to open that consent.
- A pop-up window will appear with the consent registration details.



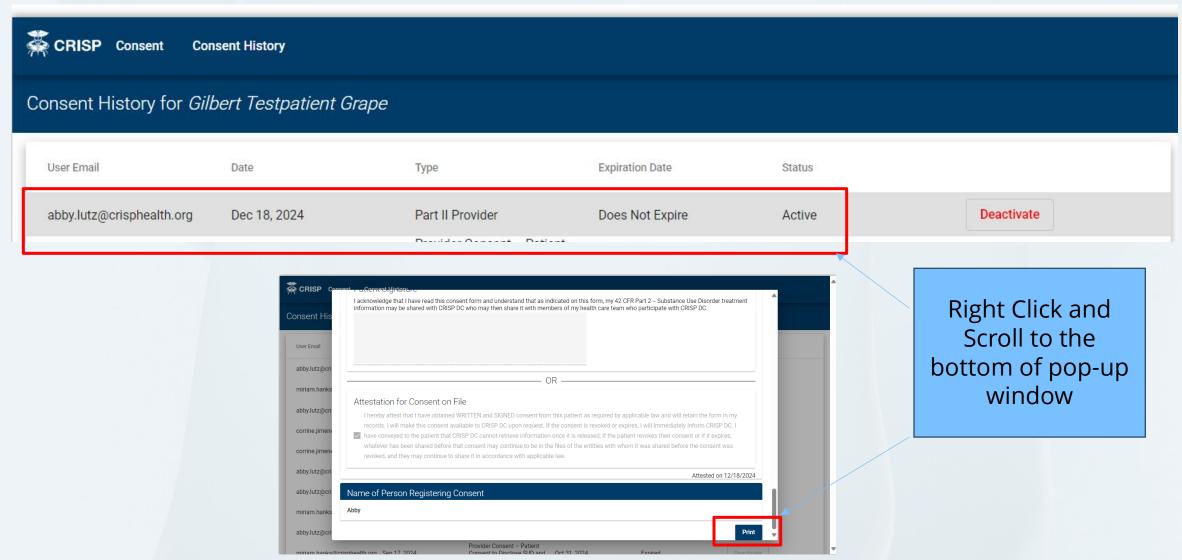


How to print a consent registration

- Providers may review, print, or save a registered consent as a file.
- Search for a patient and open a record in their consent history.
- Scroll to the bottom of the window and click "Print."
- A print preview will appear where providers can make selections for how to print the file.



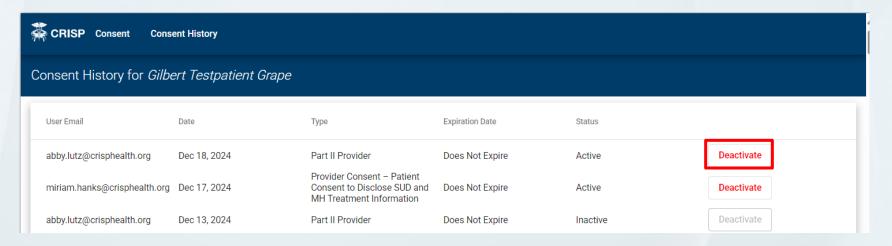




How to deactivate a consent registration



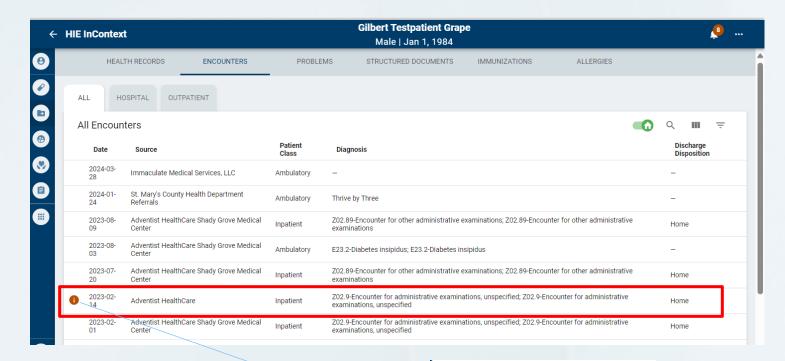
- Search for a patient and locate the "Active" record in their consent history.
- Click "Deactivate" on the record.
- Then click "Deactivate" on the prompt
- The record's status will then update as "Inactive."







- Once consent is submitted for a patient their SUD data covered by 42 CFR Part 2 will be identified within the CRISP HIE with an orange 'i'
- This makes the data easily identifiable amongst other clinical data within the HIE.



42 CFR Part 2 prohibits unauthorized redisclosure of this information. A provider that receives 42 CFR Part 2 protected SUD information from the HIE may record information about the patient's SUD treatment in their medical record for clinical purposes, and in most cases, that would not cause the record to be subject to 42 CFR Part 2 restrictions, unless the provider is already subject to 42 CFR Part 2.

