National Survey of Health Information Exchange Organizations

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2022 Health Information Organization (HIO) Survey and CIVITAS Member Survey

The nationwide survey of HIOs is being led by CIVITAS in collaboration with Dr. Julia Adler-Milstein at the University of California, San Francisco and is sponsored by the Office of the National Coordinator for Health IT (ONC). As you know, the field continues to change rapidly, and this survey will enable us to focus on new achievements and identify challenges to create a current and accurate picture of SHIEC member efforts. We request a brief amount of your time to complete our survey. Participation is completely voluntary and will contribute to a research study. Thank you in advance for your time.

The survey includes questions in five broad areas:

- (1) Organizational Demographics
- (2) Public Health Reporting
- (3) Implementation/Use of Standards
- (4) Network-to-Network Connectivity and TEFCA
- (5) Information Blocking

There is a sixth section of questions, only asked of CIVITAS members, that cover a range of supplemental topics.

We will not make ANY responses to questions publicly available or attribute responses to any specific organization. These data will only be presented in aggregate and will be published in a peer-reviewed journal (which we will be happy to send to you) and other publicly available publications and presentations. Please see below for more details on data access and data reporting.

Data Access: Who Will Have Access to Individual, Identified Survey Responses

The CIVITAS leadership team and the UCSF research team that are collecting the data will have access to fully identified survey responses. In addition, the Office of the National Coordinator for Health IT (ONC) that is funding the survey will be given a dataset containing identifiable survey responses in the first five sections only. ONC may choose to share all or part of the dataset with ONC contractors only for the purpose of conducting contracted work and abiding by the same reporting/disclosure terms as described below. The sixth section will only be made available to CIVITAS and the UCSF research team.

Data Reporting: What Data & Derivative Results Will be Reported in Journals, Data Briefs, or Public Documents No individual respondents or responses will ever be identified or reported. All data will be reported at an aggregate level (e.g., across all survey responses). For example, we may report that 10% of HIOs in the US have payers as participants. A subset of data may be reported at the regional level (i.e., aggregated by state or healthcare market/HRR). CIVITAS, UCSF, ONC, and any ONC contractors receiving the data will abide by these terms.

If you are involved with multiple efforts, please let us know so that we can send you another link to the survey. This will ensure that you fill out only one response per effort. We also ask that you respond to survey questions only <u>from the perspective of your organization</u>. Please do not attempt to summarize multiple efforts that may be affiliated with your organization (For example, if you are a state-level HIO, please <u>do not</u> respond on behalf of local HIOs with whom you work.)

To thank you for your time, upon completion of the survey you will be offered a \$50 amazon.com gift certificate. If you are not eligible for our survey, you will be offered a \$10 amazon.com gift certificate.

If you have any questions, please contact the project investigator, Dr. Julia Adler-Milstein (Julia.Adler-Milstein@ucsf.edu or 415-476-9562). Questions for CIVITAS may be directed to Lisa Bari (lbari@civitasforhealth.org or 415-680-6921)

Screening Questions

We would first like to ask you about the type of organization for which you are responding:

1. As of March 1, 2022, was your organization: (select one)

Supporting* "live" electronic health information exchange across your network

Building (or planning for) the infrastructure or services to support*, or pilot testing, electronic health information exchange across your network (End of survey)

No longer pursuing or supporting* electronic health information exchange (End of survey)

Never pursued or supported* electronic health information exchange (End of survey)

2. Does electronic health information exchange take place between independent entities**?

Yes No (End of survey)

* Supporting is defined as offering a technical infrastructure that enables electronic health information exchange to take place.

**Independent entities are defined as institutions with different tax identification numbers; HIE between independent entities requires that *at least one* entity is independent of the other(s).

Organizational Demographics

2.

1. Which of the following general categories apply to your organization: (Select all that apply)

	_ Multi-state HIE
	Single, statewide HIE
	Community or local HIE
	Governmental, state-designated HIE
	Non-governmental, state-designated HIE
	Enterprise HIE (i.e. primarily facilitate exchange between strategically aligned organizations)
	Health Information Service Provider (HISP)
	Other (please list):
What	is your legal organizational structure?
	State Government/Agency
	Private Non-Profit 501c3
	Private For-Profit

- 3. Since January 1, 2020, have you merged or are you planning to merge with another HIE?
 - No, not planning to do so

Other (please specify):

- Currently considering
- Yes, plan to merge. If public, with whom:
- Yes, recently merged. If public, with whom:
- 4. *Which state(s) do you consider the <u>primary</u> ones in which you currently have, or are recruiting new, participants in your HIE? This should ***not*** include state(s) that you connect to via regional/national networks, such as Patient Centered Data Home or eHealth Exchange, or state(s) in which you provide technology for other HIEs that are branded under a different name.



5. 5a. *For the state(s) selected in question 4, please select the specific hospital service area(s)⁺ in which you currently have, or are recruiting new, participants in your HIE.

[†] Hospital Service Areas are geographic areas defined by the Dartmouth Atlas.

[Populate list of HSAs for each State reported in prior question and have check all option for HSAs in a given state]

A hospital service area look-up by zip code can be found at: www.dartmouthatlas.org/data/search_zip.php

If you describe your service area differently or have additional comments on geographic area covered, please comment:

5b. If you have participants in other states or connections to HIEs in other states, please list those states here:

6. For the state(s) selected in question 4, what is the state's general approach to consent? [Populate with states from question 4, limiting to those only reporting 1-5 states.]

Opt-in
Opt-out
Other (please specify):

7. Please indicate which of the following options applies to your HIE model:

 Federated
Centralized
Both (Hybrid)

Other (please specify)

8. Which of the following services do you currently offer that are used by participants in your HIE? (Select all that apply)

GENERAL SERVICES	
Provider Directory	
Patient Consent Management	
Community Medical Record: Aggregation of information from across the community served by the HIE, only including health information (e.g., diagnoses, procedures, medications)]
Community Health Record: Aggregation of information from across the community served by the HIE, including health and non-health information (e.g., transportation, education, and/or housing data)	
Record Locator Service	
Messaging using the Direct Protocol	
Transform other document types or repositories into CCDAs (e.g., MDS, OASIS, Community Health Record)]
Data normalization]
Alerting/event notification (e.g., Admit-Discharge-Transfer)	
Results delivery (i.e., uni-directional push)	
Connection to prescription drug monitoring program (PDMP)	
Prescription fill status and/or medication fill history	
Provide data to third party disease registries (e.g., Wellcentive, Crimson)]
Advanced care planning (i.e., POLST/MOLST)	
Sell de-identified data to third parties	
Patient access to immunization history	
Integrating claims data	
Other (please list):]

Services related to VALUE-BASED PAYMENT MODELS	
Activities related to quality measurement (e.g., generating,	

validating, reporting, etc.)	
Closed-loop referrals tracking	
Identification of gaps in care	
Care coordination platform	
Registry services, including operating as a clinical data registry or qualified clinical data registry (QCDR) ¹	
Providing data to allow analysis by networks/providers	
Analytics (e.g., risk stratification)	
Other (please list):	

- 9. Do entities participating in your HIE cover 100% of your operating expenses?
 - Yes
- 10. Have you received HITECH 90/10 funds for implementation either directly as state designated entity, or indirectly through another entity?

	Yes	
	No	
	Don't	know

- 11. Has your state Medicaid organization ever provided funding to support your HIE?
 - Yes initial, one-time funding only
 Yes ongoing funding only
 Yes both initial and ongoing funding
 In the process of obtaining approval for funding
 No
 Other: Please explain:
- 12. Does your HIE formally partner with your state Medicaid organization to provide data for quality reporting?
 - Yes, our HIE provides data for state quality reporting only
 Yes, our HIE provides data for federal quality reporting only
 Yes, our HIE provides data for state and federal quality reporting
 We are in the process of working with state Medicaid to provide data for quality reporting
 No
 Other: Please explain:
- 13. If you have a Master Patient Index (MPI), please ESTIMATE:

Total number of unique (resolved) individuals in your MPI:

Total number of unique individuals in your MPI with more than only demographic data:

14. Within the past year, please estimate **the number of acute care hospitals** (individual facilities both within health systems and independent, including VA, public, and private) that are directly connected (not via another network) to your HIE:

Do not know

HOSPITALS

Provide data

Do not know

¹ A Qualified Clinical Data Registry (QCDR) is a Centers for Medicare & Medicaid Services (CMS) approved vendor that is in the business of improving health care quality. These organizations may include specialty societies, regional health collaboratives, large health systems or software vendors working in collaboration with one of these medical entities. (CMS)

Receive or view data

Do not know

HIE Support for Public Health Reporting

1. Please list up to 5 state or local public health entities that are connected to your HIE (Connected means that the public health entity provides data to your HIE, receives data from your HIE, and/or pays to participate in your HIE):



1a. For the entity(ies) listed, which type is each public health entity?

Answer Options *populate from those listed above*	State Public Health Agency	Local or County Public Health Agency	Other

1b. For the entity(ies) listed, please report whether each public health entity: (Select all that apply)

Answer Options *populate from those listed above*	provides data to your HIE	Your HIE reports data to	pays to participate in your HIE	None of these options

- 1c. Please report whether the Centers for Disease Control and Prevention (CDC): (Select all that apply)
 - Provides data to your HIE Receives reported data from your HIE
 - None of the above

If any option in column 2 of question 1b is selected:

For questions 2-6 <u>please answer for the PRIMARY public health agency (PHA) to which you are currently</u> reporting data or are establishing ability to report data:

- 2. Please indicate which you consider to be the primary public health agency to which you are currently or planning to establish reporting:
 - 2a. Which of the following reporting services to your **primary public health agency (PHA)** do you offer to **your participating healthcare providers**?

	In production	In testing	In planning	Available, but PHA not able/willing	Not available	Don't know
Syndromic surveillance reporting						
Immunization registry reporting						
Electronic case reporting						
Electronic reportable laboratory result reporting						
Public health registry reporting (administered by or for public health agencies for public health purposes)						
Clinical data and/or specialized registry reporting (administered by or for non- public health agency entities for clinical care and monitoring health care quality and resource use)						
Other COVID-19 related reporting (e.g., registry)						
Vital Record System reporting						

3. For each type of reporting to the primary PHA that is in production, are any of the following provider types currently using these services (i.e., at least one organization providing data for reporting)? (Select all that apply)

	Hospitals	Office-based physicians	LTPAC settings	Urgent Care	Other
Syndromic surveillance reporting					
Immunization registry reporting					
Electronic case reporting					
Electronic reportable laboratory result reporting					
Public health registry reporting					
Clinical data registry reporting and/or specialized registry reporting					
Other COVID-19 related reporting (e.g., registry)					
Vital Record System reporting					

3a. For each type of reporting for 'Other' provider types, please indicate which provider types below.

	Other Provider Types Reporting through your HIE
Syndromic surveillance reporting	
Immunization registry reporting	
Electronic case reporting	

Electronic reportable laboratory result reporting	
Public health registry reporting	
Clinical data registry reporting and/or specialized registry reporting	
Other COVID-19 related reporting (e.g., registry)	
Vital Record System reporting	

- 4. Do you receive any of the following funding source(s) to specifically support public health reporting? (Select all that apply)
 - Fees paid by participants
 Fees paid by State health department
 State Medicaid funding
 STAR HIE program
 CARES Act funding
 Other Federal funding
 Other State funding, including from State health department
 Other. Please list:
 - Do not receive any funding to specifically support public health reporting
- 5. To what extent have you experienced the following barriers to public health reporting? This includes both reporting to and receiving from primary PHA.

	To a Great Extent	Somewhat	Very Little	Not at All	N/A
Patient consent model hinders data exchange with PHA					
State statutes/regulations limit PHA participation with HIE					
Need for data use agreements for public health data					
Limited funding from PHA					
Limited funding from your HIE participants					
PHA lacks staffing					
PHA lacks technical capability to receive messages from your HIE					
PHA lacks technical capability to process messages from your HIE					
Other technical limitations on part of PHA					
PHA has other priorities					
Low return on investment to your HIE					
Other (please list):					

6. Since February 2020, have you expanded the number of provider organizations that engage in public health reporting through your HIE?

Yes
No

Don't know

6a. If yes, Which provider types expanded public health reporting through your HIE? (Select all that apply)

Hospitals and Health Systems		
Ambulatory Clinics/Physician Practices		
Long-term Care Facilities		
Correctional Facilities		

Labs (commercial, public health)		
Behavioral Health Providers		
Other (please list):		

If any option in column 1 of question 1b is selected:

7a. Which of the following types of data do you **receive** from public health entities with which you have established connectivity? (Select all that apply)

	001	
		Syndromic surveillance
		Immunization
		Electronic case reports
		Electronic reportable laboratory results
		Data from public health registry (administered by or for public health agencies for public health purposes)
		Data from clinical data and/or specialized registry (administered by or for non-public health agency entities for
	clin	ical care and monitoring health care quality and resource use)
		Data related to COVID-19
		Vital records
		Other. Please list:
		Don't know
76		at is the surpcess of reactiving sublic health date? (Calent all that each)
70.		at is the purpose of receiving public health data? (Select all that apply)
		To identify opportunities to enrich public health data with HIE data
		To make public health data available to your participants
		Other. Please list:

HIE Support for Public Health Exchange Related to the Pandemic

8. What are your current capabilities to electronically receive hospital data on **bed capacity and resource utilization**? Electronic receipt includes standards-based approaches (e.g., SANER, HL7 feed) and does <u>not</u> include spreadsheet submission and/or manual data entry.

Actively electronically receiving production data

In the process of testing and validating electronic receipt of data (Skip to 9) \square

In planning phase to support this reporting (Skip to 9)

Not planning to support this reporting (Skip to 9)

Don't know (Skip to 9)

8a. If actively electronically receiving production data, to what entities are you submitting this data? (Select all that apply)

City or local public health department(s)

State public health department(s)

Federal entities (such as, the CDC or HHS)

- Other. Please list:
- Don't know

8b. How do hospitals transmit hospital capacity and resource utilization data to your HIE? (Select all that apply)

____ADT messages

_ HL7 v2 messages

SANER FHIR Server https://build.fhir.org/ig/HL7/fhir-saner/introduction.html

Other. Please list:

Don't know

- 8c. What <u>terminology standards</u> are used by hospitals to report hospital capacity and resource utilization data? (Select all that apply)
 - NIEM
 LOINC
 Other. Please list:
 Non-standardized codes
 - Don't know
- 9. Does your HIE currently provide data to PHA(s) to fill gaps in their COVID-19-related data (e.g., missing demographic information)?

Yes
No but could do so
No and could not do so
Don't know

9a. If yes or could do so: Please indicate what types of data are or could be provided to fill gaps. (Select all that apply)

Data Type		/ p	provided	-	/ provided ld be
Race/ethnicity					
Other demographics]
Up-to-date contact information (for contact tracing)]	
Hospitalization information					
Health information such as chronic health conditions					
Immunization data					
Commercial lab results					
Hospital lab results					
Other:					

9b. If yes: How often do PHA(s) electronically receive or query these types of data from your HIE?

Often
Sometimes
Rarely
Never
Don't know

9c. If yes: How are PHA(s) accessing these data? (Select all that apply)

- Single patient lookup through a Portal
- Batch query and response
- ___ API
- Aggregate data and/or statistics (e.g., dashboard)
- SFTP/Amazon S3 file transfer
- ___ Other. Please list:
- ___ Not applicable

10. What other services does your HIE provide to PHA(s) to support COVID-19 response: (Select all that apply)

Analytic and Data Quality Support (beyond those reported above)

Dashboarding and Data Visualization Assistance

Process Automation

Bidirectional Data Sharing/Receiving Data from PHAs

Use of HIE MPIs to Support Public Health Deduplication or Other Services

Outbreak Monitoring and Alerting

___ Public Health Policy Impact Monitoring

- Other. Please list:
- None
- 11. Do PHA(s) contribute COVID-19 immunization registry data or make COVID-19 immunization registry data available for query through your HIE?

Yes
No
Don't know

12. Other than PHAs, who are the users of your HIE's COVID-19 data? (Select all that apply)

Healthcare Providers: Administrators	Emergency Preparedness/Response
Healthcare Providers: Frontline Clinicians	School Nurses
Payers	Contact Tracers
Medicaid	
Other. Please list:	
None	

13. If 'Healthcare Providers: Frontline Clinicians' is checked: What COVID-19 data can frontline clinicians access through your HIE: (Select all that apply)

COVID-19 test results/case status
 COVID-19 antibodies
 Other respiratory illness history
 Vaccination Status
 Hospital Status/Capacity Information
 Healthcare utilization (inpatient, outpatient, EHR visits, etc.)
 Demographics (age, race, ethnicity, etc.)

- Other. Please list:
- Don't know
- 14. If 'Healthcare Providers: Frontline Clinicians' is checked: How can frontline clinicians access COVID-19 data through you HIE: (Select all that apply)

	Individual	patient	look-up	via	portal	or	query
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De-identified reports

Bulk query for identified data on populations

Dashboards and interactive reporting

Public or private briefings on community/statewide COVID-19 status

- Secure email notifications
- Other. Please list:
- Don't know

Lab Participation in COVID-19 Relevant HIE

15. Please report whether each type of stakeholder is involved in your HIE in the following ways:

Answer Options	Provide COVID- 19 Test Results	Provide Data Other Than COVID-19 Test	View or Receive Data
	15 ICST ICSUITS		Dulu
		Results	

Hospital-based labs		
Independent labs (including commercial)		
Physician office-based labs		
Mobile labs (e.g., Point of Care Labs for COVID-19)		
Public health labs		
Other:		

16. If 'Provide COVID-19 Test Results' is checked, for the relevant rows: How timely are COVID-19 test results that you typically receive?

	Real Time; Near Real Time	Within 24 hours	Greater than 24 hours but less than 48 hours	Greater than 48 hours	Don't Know	Not applicable
Hospital-based labs						
Independent labs (including commercial)						
Physician office-based labs						
Mobile labs (e.g., Point of Care Labs for COVID-19)						
Public health labs						
Other:						

- 17. In general, have laboratories sought to limit or refused to provide access, exchange, or use of electronic health information (e.g., laboratory results)?
 - Yes
 No (skip to 21)
 - Have not made request (skip to 21)
- 18. What types of laboratories have sought to limit or refused to provide access, exchange, or use of electronic health information? (Select all that apply)
 - Hospital-based labs
 - Independent labs (including commercial)
 - Physician office-based labs
 - Mobile labs (e.g., Point of Care Labs for COVID-19)
 - Public health labs
 - Other. Please list:
- 19. Which of the following reasons have laboratories used as the basis for limiting or refusing to provide electronic health information to your HIE? (Select all that apply)
 - Role of CLIA or other federal regulations in restricting them from sending additional data
 - Fees associated with HIE participation
 - Labs don't derive value as a data contributor only
 - Concerns with HIE's ability to do patient matching
 - Concerns with producing duplicate data
 - Exchanging data with HIEs is not considered related to treatment, payment, or operations and thus would require patient consent
 - Labs reporting obligation ends with returning result to ordering provider
 - Public health agencies (including emergency rules) do not mandate reporting to HIE

Labs need consent from each individual provider, resulting in your HIE having to execute multiple disclosure forms (e.g., for each participating health care provider)

Technological reasons/use of specific standards (convenient reason or wide spectrum of what labs are able to do)

Other. Please list:

20. To what degree have you been able to overcome these difficulties to access data from laboratories?

___ Not at all ___ To a small extent ___ Somewhat

To a great extent

Fully

21. Does your HIE map from non-standard laboratory test/result codes to LOINC® codes?

Yes
No (Skip to next section)
Don't know (Skip to next section)

21a. Within the past year, based upon the volume of test results received (qualitative and quantitative), to what extent did your HIE have to map those results from non-standard codes to LOINC codes?

All or most
Some
Few
None
Don't know

- 21b. Have you experienced any of the following issues related to mapping to LOINC? (Select all that apply)
 - We do not have sufficient expertise to map to LOINC within our organization
 - We find LOINC and LOINC tools too difficult to use
 - We do not have the resources (personnel/time) to map to and/or maintain mappings to LOINC Other issue. Please specify:
 - No, we have not experienced any issues mapping to LOINC
 - Don't know

1. To what extent does your HIE electronically **receive** data from your participants using the following methods listed below? (Select one option across a row)

Please consider the methods used by participant to provide the data to your HIE. Do not include conversions you may do after receipt. With regards to conformance to standards, if the receipt of the data is in partial conformance, please consider that as conformant.

	Routinely/ from most participants	Sometimes/ From some participants	Rarely/ From few participants	Never	Don't know
Care summaries in a structured format (e.g., CDA, CCR, C32)					
HL7 v2 messages (any type)					
ADT messages (for applicable participants)					
HL7 Fast Healthcare Interoperability Resources (FHIR) messages (DSTU2)					
HL7 FHIR Release 3 (STU) messages					
FHIR v.4.0 messages					

- 1a. If care summaries in a structured format "routinely" or "sometimes" is checked above, then ask: Do you parse C-CDAs (i.e., extract and make available discrete data elements):
 - Yes No Don't know
- 2. To what extent is the information that you **receive from your participants** consistent with different versions of the United States Core Data for Interoperability (USCDI)? USCDI is a standardized set of health data classes and constituent data elements for nationwide, interoperable health information exchange.

	Routinely/ from most participants	Sometimes/ From some participants	Rarely/ From few participants	Never	Don't know
USCDI v1 https://www.healthit.gov/isa/united- states-core-data-interoperability- uscdi					
USCDI v2 https://www.healthit.gov/isa/united- states-core-data-interoperability- uscdi#uscdi-v2					

3. To what extent does your HIE electronically **send or make available** data to your participants using the following methods?

	Routinely/ To most participants	Sometimes/ To some participants	Rarely/ To few participants	Never	Don't know
Care summaries in a structured format (e.g., CDA, CCR, C32)					
HL7 v2 messages (any type)					
HL7 Fast Healthcare Interoperability Resources (FHIR) messages DSTU2					
HL7 FHIR Release 3 (STU)					
FHIR v.4.0					

4. Which types of **clinical and other health-related information** are made available by your HIE (as part of a clinical document or as a structured data element)? (Select all that apply)

	Included in your HIE
Data Provenance	
Clinical Information	
Problems	
Prescribed Medications	
Filled Medications	
Medication Allergies	
Non-Medication Allergies & Intolerances	
Functional Status	
Cognitive Status	
Vital Signs	
Pregnancy Status	
Immunizations	
Family Health History	
Health Concerns	
Clinical Notes	
Imaging/Pathology	
Diagnostic Imaging Order	
Radiology Report (narrative)	
Pathology Report (narrative)	
Laboratory-Related Information	
Laboratory Test(s)	
Laboratory Value(s)/Result(s)	
Laboratory report (narrative)	
Team-Based Care	
Care Plan Field(s), including Goals and Instructions	
Care Team Member(s) (Provider ID, Provider Name)	
Assessment and Plan of Treatment	
Encounter-Related Information	
Procedures	
L	

Admission and Discharge Dates and Locations	
Encounters (Encounter type, diagnosis, time)	
Discharge Disposition	
Referrals	
Discharge Instructions	
Reason for Hospitalization	
Health Equity	
Home Address	
Race/Ethnicity	
Preferred Language	
Health-related Social Needs (e.g., housing, food insecurity)	
Substance Use Disorder (as defined in 42 CFR Part 2)	
Gender Identity	
Sexual Orientation	
Other	
Other (please list):	

5. To what extent does your HIE electronically send or make available to participants:

	Routinely/ To most participants	Sometimes/ To some participants	Rarely/ To few participants	Never	Don't know
Care summaries in a structured format (e.g., CDA, CCR, C32)					
Data in a format consistent with USCDI v1					
Data in a format consistent with USCDI v2					

6. Does your HIE **make data available for participants to query?** Note: query refers to a query-and-response exchange, e.g. a request from one participant through an interface that results in a response delivered into an EHR.

Yes
No
Don't Know

1. Does your HIE: (Select all that apply)

Sell/provide your infrastructure to other HIEs	
Buy/use infrastructure from another HIE	
Connect to other HIEs in SAME state	
Connect to other HIEs in DIFFERENT state(s)	
None of the above	

2. Is your HIE currently using the following national networks to exchange data?

	Live Data Exchange (send or receive)	Implementing	Not Using	Other (please specify):
General Purpose Networks:				
CommonWell				
DirectTrust				
Strategic Health Information Exchange Collaborative (CIVITAS)/Patient Centered Data Home				
e-Health Exchange				
Carequality				
Specific Purpose Networks:				
Surescripts				
Patient Ping				
Audacious Inquiry: Pulse/ENS				
Collective Medical Technologies: EDIE				
Social Service Referral Platform(s) (e.g., Aunt Bertha, Unite Us)				
Other (please list):				

2a. If not using any general-purpose networks in prior question: Please select reason(s) for not using any of the general purpose networks: (Select all that apply)

Do not see the value in what they provide (i.e., services not useful or data limited)

Perceive them as competitors

Participation costs too high

Not a priority

Other. Please list:

3. Is your HIE planning to participate in the Trusted Exchange Framework and Common Agreement? Please find definitions of the roles here: <u>https://www.healthit.gov/topic/interoperability/trusted-exchange-framework-and-common-agreement-tefca</u>

Yes, as a QHIN Yes, as a Participant or as a Sub-participant No Don't know

3a. If no: Why are you not planning on participating in TEFCA? (Select all that apply)

Don't have enough information

____ Don't have time/resources to prepare

Concerns about the terms of the Common Agreement (please briefly describe):

Concerns over privacy and/or security of the network

Concerns about the burden associated with participation (e.g., financial, reporting) (please briefly describe):

- Do not perceive sufficient value in participating (please briefly describe why):
- Other (please list):

3b. If don't know: Why are you unsure about participating in TEFCA? (Select all that apply)

Don't have enough information

Don't have time/resources to prepare

Concerns about the terms of the Common Agreement (please briefly describe):

Concerns over privacy and/or security of the network

Concerns about the burden associated with participation (e.g., financial, reporting) (please briefly describe):

Do not perceive sufficient value in participating (please briefly describe why):

- Have not yet developed a strategic plan to participate
- Other (please list):

3c. If yes: As exchange based on the Trusted Exchange Framework and Common Agreement becomes operational, is your HIE planning to change its operations in any of the following ways:

	Yes	No	Don't know	Not Applicable
Changing Types of services offered				
Selling/providing your services to other HIEs				
Buying/using services from another HIE				
Changing technical infrastructure				
Changing legal agreements and/or policies				
Changing other infrastructure (e.g., creating new training, supporting or making process redesigns (e.g., new workflows))				
Partnering with HIEs in SAME region/state				
Partnering with HIEs in DIFFERENT regions/states				
Partnering with an entity that is not an HIE (e.g., Health IT Developer)				
Other (please list):				

4a. For which of the following exchange purposes (which are included in TEFCA), are your participants currently able to make a REQUEST for information?

	Yes	No	Don't Know
Treatment (as defined by HIPAA)			
Payment (as defined by HIPAA)			

Health Care Operations (as defined by HIPAA)		
Individual Access Services		
Public Health		
Government Benefits Determination (as defined by TEFCA)		

4b. For which of the following exchange purposes (which are included in TEFCA), are your participants currently able to RESPOND WITH ADEQUATE DATA to a Request for information?

	Yes	No	Don't Know
Treatment (as defined by HIPAA)			
Payment (as defined by HIPAA)			
Health Care Operations (as defined by HIPAA)			
Individual Access Services			
Public Health			
Government Benefits Determination (as defined by TEFCA)			

Information Blocking

Information blocking practices have been defined in rules that went into effect on April 5, 2021. The following set of questions ask about practices that may constitute information blocking based on your understanding of the rules. Please respond based on your experience since the rules went into effect (April 5, 2021).

- 1. To what extent are you familiar with the information blocking rules, applicable actors, exceptions, and enforcement timeline?
 - Very Familiar Moderately Familiar Somewhat Familiar Not Familiar
- 2. How often have you encountered **each of the following form(s)** of information blocking by **EHR vendors** (and other Developer(s) of Certified Health IT)?

	Rarely /Never	Sometimes	Often/ Routinely	Don't Know
PRICE				
Examples:				
using high fees to avoid granting third-parties access to data stored in the developer's EHR system				
charging unreasonable fees to export data at a provider's request (such as when switching developers)				
CONTRACT LANGUAGE				
Examples:				
using contract terms, warranty terms, or intellectual property rights to discourage exchange or connectivity with third-party				
changing material contract terms related to health information exchange after customer has licensed and installed the vendor's technology				
ARTIFICIAL TECHNICAL, PROCESS, OR RESOURCE BARRIERS				
Examples:				
using artificial technical barriers to avoid granting third-parties access to data stored in the vendor's EHR system				
using artificial reasons to limit the types of information that can be sent/shared or received				
REFUSAL				
Examples:				
refusing to exchange information or establish connectivity with certain vendors or HIOs				
refusing to export data at a provider's request (such as when switching vendors)				
OTHER (please list):				

6)

3. What proportion of EHR Vendors have you encountered engaging in information blocking?

All/Most		
Some		
Few		
None (skip to 6)		
Don't know or N/A (Don't interact with develope	ers) <mark>(skip t</mark> a	C

3a. Among EHR Vendors that engage in information blocking, how often do they do it?

Routinely
Sometimes
Rarely
Don't know

- 4. When you have experienced practices that you believed constituted information blocking by **EHR vendors** in the past year, how often did you report the information blocking to ONC/HHS?
 - Always
 Most of the time
 Sometimes
 Rarely
 Never
- 5. To what extent does information blocking by **EHR vendors** make it more difficult for you to provide HIE services to your participants?
 - Greatly Moderately Minimally/Not at all Don't know
- 6. In what form(s) have you experienced information blocking by hospitals and health systems?

	Rarely /Never	Sometimes	Often/ Routinely	Don't Know
ARTIFICIAL TECHNICAL, PROCESS, OR RESOURCE BARRIERS				
Examples:				
requiring a written authorization when neither state nor federal law requires it				
requiring a patient to repeatedly opt in to exchange for TPO				
REFUSAL Examples: refusing to exchange information with competing providers, hospitals, or health systems refusing to share data with other stakeholders, such as payers or independent labs				
CLOSED NETWORK EXCHANGE Examples: promoting alternative, proprietary approaches to HIE exchanging only within referral network or with preferred referral partners				
OTHER (please list):				

7. What proportion of **hospitals and health systems** have you encountered engaging in information blocking?

Some

- Few
- None (skip to 9)

] Don't know or N/A (Don't interact with developers) (skip to 9)

- 7a. Among hospitals and health systems that engage in information blocking, how often do they do it?
 - Routinely
 - ___ Sometimes
 - Rarely
 - Don't know

- 8. To what extent does information blocking by **hospitals and health systems** lead to missing patient health information?
 - Greatly
 - Moderately
 - Minimally/Not at all
 - Don't know
- 9. Among other types of stakeholders, to what extent have you observed information blocking behaviors?

	are leve	-	Som	neti	imes	O Roi	fte utin		on't now
Commercial Payers]]				
Commercial Laboratories]]				
Commercial Pharmacies									
National Networks (e.g. CommonWell, eHealth Exchange)									
State, regional, and/or local health information exchange									
Other (please list):									

10. Across all types of stakeholders, to what extent has information blocking decreased since the final regulations went into effect in April 2021?

Greatly Moderately

Minimally/Not at all

Don't know or N/A

Additional Information

1. Initiative or Organization Name:

2. We appreciate your participation. Would you like to receive a copy of our results that will enable you to compare your effort to others in the nation?

Yes
No

3. If you would like to receive a \$50 amazon.com gift certificate, please complete the following fields:

Name:

Email: