

2020 NEMSIS v3 Annual Implementation Meeting

Raw Discussion Notes from Breakout Sessions





What characteristics of the epidemic were not adequately addressed by custom additions?

- Extend ICD-10 to utilize U codes
- More specific impression code that directly pertains to current issue
- Some states averse to using Custom Elements, some poor implementation at State level. Many states chose not to use Custom Elements, and then they had to be removed after adding.
- Put codes into Standard
- States already tracking ILI, did not utilize many of Custom Elements
- Reuse of PPE element was needed, international less relevant, contact tracing, administrative addendum added to collect pt data post-call
- Speed of establishing Custom Elements have them ready sooner
- Series of different kinds of elements, trial runs for upcoming seasons
- Watch supplies being used, surveillance trends
- Repetitive questions/elements
- Cost of process implementation cadence of release, grouping, get out ahead, be more agile
- Selection lists became difficult with dev
- Billing overlap with U codes
- Prep time
- List of States implementing Custom Elements would have been helpful





What characteristics of the epidemic were not adequately addressed by custom additions? (con't)

- ILI Testing involvement of EMS, vaccination in field (data collection, implementation), vaccine admin date, S/S post vaccine, vaccine Lot # would be needed, consider 2-cycle dosing, documentation for pt education
- ICD-10 Codes for Telemedicine, add appropriate values
- Standardize COVID custom elements
- NFIRS was capturing data as well, double documentation for some
- Importance of putting Custom Elements into the next version or CP to current standard
- Definition for eCustomConfig for geo-related questions defined as type OTHER but difficult to implement in software
- International questions in relation to regions specific location as opposed to general areas. Implementation is an issue
- Use of general symptoms over COVID-specific impressions more valuable
- Define documentation for PUIs, inconsistency in criteria for PUI
- Family members with ILI S/S, current immunization status
- Charting is already too complex for field clinicians
- Surveillance delay with waiting for ePCRs to finish/submit
- Region/city: the Cities traveled Custom Element is a GNIS code only allows for US cities. What about specific international cities that do not fit GNIS.
- Custom Elements designed for states with multiple vendors to facilitate data exchange. When an area uses a custom element and the state doesn't use or accept how is that useful to statewide analysis.





Could the NEMSIS DEM section be used to address resource-related issues?

- Agencies aren't making DEM changes often, could produce stale data. Agencies just not updating DEM frequently enough.
- Pop-up data in field is an option but would engage field clinician not agency level, agency report could be an option
- Already difficult to keep DEM updated. Already complicated.
- Financial concerns may not be appropriate for field API, staffing may not in field PCR either, does an agency WANT to send financial info up the chain.
- Possible but should is the question. May not be fresh enough for decision making. How would the data be used. Do we already collect other places?
- Only states with mandate for DEM may only find it useful. Chicken-egg —if it's added would more states do DEM.
- Where does the benefit come from it is a state need?
- Potential for API, or does this require its own dataset
- States would have to implement work flow to get timely and updated DEM. Some states cannot enforce
 ePCR within 72 hours.
- This set doesn't really fit in the DEM file. Is this too much info specific to an agency as a business.
- Not all states collect DEM files. Field level collection could be skewed based on who is entering data.
- Where does the external COVID Resource Reporting Tool fall into this? This would replace that external tool.





Could the NEMSIS DEM section be used to address resource-related issues? (con't)

- Administrators don't all support updating DEM regularly. PPE is difficult to track per incident in some software.
- Generic data isn't always actionable. Consider burn rates.
- Where is the benefit to the agency?
- Some issues may apply more to clinicians and some to administrators.
- May be better placed within ePCR data pertaining to exposures. Exposure information may not be available when completing the PCR.
- Duplicate data entry other areas of data collection.
- Question of real-time application. DEM is not as real-time as needed.
- Other issues such as Drug shortages, mixing pt care and operational data, may benefit from its own dataset.
- ePCR should not be used as supply chain database.
- All changes incur implementation expenses. Data integrity also a concern.
- Use own dataset. May be better suited as more abstract and not as specific to COVID.
- Hospitals have a different system can current process be leveraged.
- Is NEMSIS the right collection tool?





DAY 2





Day 2: HIPAA & Data Exchange

What additional legal (instructional) documents would be helpful in facilitating bi-directional data exchange?

- Know how opting out for patients will work
- Business agreements might convolute rather than help
- It's not just legal political, financial, operational barriers
- Need more meeting and educating of hospitals, can't just be solved with documents; clinical staff are scared to give any information to EMS (lacking knowledge of what information can be shared)
- Identify what data would be exchanged and under what circumstances
- Federal government give their stance it is easy to find an attorney with the opposite stance
- Website with .gov at the end pushing the conversation about sharing data
- Standard for bi-directional data sharing follows a specific standard to show all the laws are being followed
- Case study highlighting benefits to the hospital





Day 2: HIPAA & Data Exchange

What additional legal (instructional) documents would be helpful in facilitating bi-directional data exchange? (con't)

- Facilities don't want to be the ones to go first a standard would be a great place to start
- Sense of value for sharing the data bi-directionally
- Recognize the other considerations and finding ways to incorporate them all ensure none of us get it wrong
- Vendor to vendor data exchange document
- Document outlining compliance with a cloud provider vs a brick and mortar provider
- Central data agreement that includes various health information death records
- Checklist or flow chart to see what data can and cannot be shared hospital and EMS
- Multiple laws that overlap research at federal and state level
- When and how to stop sharing what happens when you need to stop sharing or are there circumstances when you have to share
- How do we share this information with the field level providers in the agencies checklist of educational material to be covered with HIPAA or SAMSHA trainings





Day 2: Tools for v3.5

Schedule, State Support, Tracking Transition

- Personal attention was successful in the past agency by agency (virtually and physically)
- Not too aggressive timeline
- Video from NEMSIS to start educating agencies to give a heads up that the changes are coming changes that are in the revision
- State information and timelines published on the NEMSIS site like the map
- More discussion on the timeline not a lot of input from states
- Implementation burden eDisposition changes will be disruptive to vendors and agencies
- Timeline for states with heavy customization
- What Schematron does the state accept or use
- Uptick in custom element at a certain percentage roll into national
- Include in state GIT repo what versions the state is using, when they will stop using them
- Like the fact sheets, code translations
- Custom field catalog shared between states
- US map for 3.4 is helpful, also see in Excel to track
- Weekly emails for state resource updates are helpful
- Make sure any version of Schematron update is reliant on date of the chart vendor and state sides
- Timeline is too unrealistic certain set of steps that need to take place (domino effect, not one timeline for everyone)
- If not fulfilling 3.4 but are looking to transition to 3.5 additional burden to get agencies to submit data (can agencies submit 3.4 and 3.5 to the repo)
- States decide when to go to 3.5





Day 2: Tools for v3.5

Schedule, State Support, Tracking Transition (con't)

- Vendors have to comply with the states and not NEMSIS timeline
- State have statutes about NEMSIS versions have to go to new version if not ready
- Agencies will be on software that are not compliant and have to search for a new one long process (6 months)
- State releases data dictionary with plain English Schematron rules for their state
- What version are the states accepting may say they are on 3.4 but are also accepting 3.3.4
- Break information into 3.4 and 3.5 on the website like v2 to v3
- Beta version of what the states are working on
- Push out timeline a little
- States should tell vendors when to meet the requirement
- States can't commit until the window is no longer moving, no additional elements
- Get ET3 released then move to 3.5
- COVID-19 enhancements and ET3 that pushes out vendor timeline
- Change log that is easier to read expand on data behind the map (transition timeline)
- Training burden all the way down to the agency level eDisposition
- State testing outside of the TAC determines the timeline number of agencies and vendors
- The longer on 2 versions, the harder it may be to transition to 3.5
- Best practices for states on how to get information out to their agencies not be dependent on the vendors (some states to webinars). Something for data managers that they can use
- Change log not so technical so that agencies can understand what's going on
- Document for 3.5 spreadsheet of what is collected nationally, what is shared publically with all the states listed
- Data dictionary with rules and examples of what wording states can use and copy to provide to their agencies
- States budget years in advance, transition might not be in their budget yet could be a few years before it is





Day 2: Improving HIE

Decisions, eOutcomes, FHIR, XSD, CCDA R2 D/C

Alerting the hospital to ensure data flows into the ED system

- Consolidated ePRC short form
- Separate application to get the information into the EMR system
- Not everyone can take FHIR
- Make use of the California model element level data is exchanged
- Need to look at FHIR vs. Core because of the Cures Act that is requiring FHIR use
 - 2022 timeline for change to implementing FHIR
 - What infrastructure is needed, especially in rural areas need solutions to handle the standards
 - There is more funding for rural areas
- Already decided to only use FHIR
- Need a unified approach to send it to hospitals and get the hospital buy-in
- Easier for vendors to connect to an HIE
- What ethical considerations need to be made with the large amount of data being shared who accesses and uses the information in the HIE
- HIE connect a thon
 - Working to get adoption for specific use case for EMS vendors to work with CERNER and EPIC can work with them to help create a standard
 - Andrea Fourquet IHE USA informatics consultant





Day 2: Improving HIE

Decisions, eOutcomes, FHIR, XSD, CCDA R2 D/C (con't)

- How will HL7 map/align with NEMSIS 3.5
- Consider:
 - FHIR v4 that moves data from ePCR to EHR or HIE
 - HIE vendors are part of a work group
 - Reach out to Juan Esparza to learn more about what he is doing in FL
 - Have this conversation at a state level to brainstorm how to enforce use of this
 - Funding for development and sustainment of this project

Hospital EHR push/pull to EMS record (file)

- A separate document with separate lifecycle makes sense for outcomes
- Get data back to end user (EMS clinicians and agencies), may not need to build a new process if the hospital sends back to the ePCR that then goes to NEMSIS (repeat NEMSIS file back to agency after the outcomes information gets populated)
 - Run into challenges of validation data quality standards
 - A separate document would allow for data quality standards
- Need some integration with each hospital and same with the hospital back to EMS
 - Need one endpoint at the TAC that all patient outcomes data connect to and the TAC populates the outcomes data and send it back to agencies
 - eOutcomes is a separate document
- Separate document for eOutcomes
 - Reluctance to receive data from a non-EMS provider back into an ePCR not modify the document originally created by the EMT
 - There is concern from agencies for liability reasons but do need the outcome information but scary to have it in the record if the original is modified by someone else
 - Maybe include a data source element not currently an option
 - Could have multiple outcome set because a patient went to multiple doctors
 - Separate document = separate outcomes
- Whitepaper if outcome data is documented from a different crew (legal precedent for outcomes data populated from the hospital instead of the agency)
 - Is it listed in the ePCR or the hospital record and how does that work there is some legal standing on this already
 - Check with NJ for what they do





DAY 3





Day 3: ET3

Tools, Support, Timeline, etc.

- Webinar for participants, will there be one for the vendors would be helpful for the vendors
- Allow vendors to participate in the data submission webinar as well as state data managers
- User meetings specific to vendors
- Build scenario based use cases for the vendors
- Make criteria available for new records vs. resent records
- 48 hours is too narrow of a window to resubmit records
- Telehealth video recording for HIPAA?
- Federal definition for telemedicine?
- Testing everything out with the vendors TAC will coordinate
- Vendor documentation for the onboarding process
- Jan 2021 start feels aggressive with lack of information available at this time





Day 3: Defined Lists

How can we help States limit additions?/How do we implement analytical roll-ups at the State level?

- Make national level contain everything states filter out codes they don't want
- Control the roll-up and map to existing codes that are more relevant to the state
- Provide templates for different types of users complexity when needed to adjust on the dev side
- Create a working group to determine when and why codes should be added
- Store the actual value and the roll-up value as 2 separate data points
- Allow editing of the basic list that vendors supply
- Meet with agencies and prune down the list of codes
- Roll up needs to be built into reporting at the state level
- 2 different sets one more general and one more granular
- Labeling of codes
- Indicator based roll-ups not mutually exclusive roll-ups





Day 3: Defined Lists

How can we help States limit additions?/How do we implement analytical roll-ups at the State level? (con't)

- Can be solved between the default list and the analytical roll up
- Orientation for new clients about the codes/lists
- Shouldn't have to control as tightly if do the roll-up correctly
- More general set of codes for crews to use
- Be as specific as you can on the front end, roll-up is on the reporting side
- Need to maintain flexibility existing structures continue; QI initiatives that rely on existing structures; rolling up codes in a structure that works at a higher level
- Roll up at the national level, not the state level
- Pathway for EMS agencies to provide insight on the lists
- List come from states, vendors comply
- If any changes needed do a request with the vendor
- State level ability to sync the lists that gets passed down to the local level
- Verbiage needs to be more clinical
- Verbiage needs to be less clinical and more user friendly





Day 3: Data Submission Lag

How can we improve timely record completion?/How do we facilitate immediate submission to NEMSIS TAC?

- What is the state requirement for submission; state mandates and requirements
- QA process after record is submitted how long does that take/what is the work flow
- Schematron rules that are too complex
- Consider a different element than eTimes.03 maybe 05
- Dedication to submit the data needs to start at the field
- Standards by state are reasonable and include buy in from EMS agencies
- Need for state and local regulatory agencies to create standards
- Efficient ways for agencies to submit records easily
- Target by NEMSIS for timeliness of data reception
- QA and amendment process at the agency level before it goes to the state
- Make flow better don't show irrelevant fields for the specific call
- Custom required fields more info for provider to remember and fill out before completing report
- Is the goal of NEMSIS broadened to include surveillance
- Shifts that are 48-72 hours long could delay closing records anything longer is overtime
- State systems don't process immediately or in a timely manner customers modify workflow to not be real time/immediate submission to the state





Day 3: Data Submission Lag

How can we improve timely record completion?/How do we facilitate immediate submission to NEMSIS TAC? (con't)

- Incentive at the agency level to get reports closed faster
- Agency workflow with supervisor/QA review this takes more time; could unreviewed runs go up?
- Timeliness and relaying errors to the medics how quickly is that being relayed back to get fixed?
- Pushing out Schematron without heads up to agencies that could cause more errors and increase time
- Streamline update process at repo level UUID to submit multiple times to the same record and link up correctly
- Relaxing web services to accept incomplete records
- Quality score associated with records that are not well defined fear of not meeting that when submitting quickly
- State awards/recognition for timely submissions
- Collect and Receive vendors offer option to send right away and not give option for batch sending
- Identify and quantify where the issues are
- Encourage faster submissions during certain time points COVID
- Indicate in the dataset when a record is incomplete
- Adding date/time chart was locked on the vendor side between the time the state received it
- States provide digest on number of charts received weekly to help identify any Q sizes or differentials
- Which codes mean you should retry sending later vs a permanent failure code
- Promoting a greater understanding of resubmission for all stakeholders holding and settling periods
- New target for real-time data collection formalize a plan
- Sending reports over and over slows the system down

