Data Provenance

Environmental Scan

HITPC Consumer Empowerment Workgroup Meeting
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• **What is “Provenance”?**
  – Origin of clinical information when first created
    • Information about the source of the data
    • Information about processing/transitions the data has undergone

• **Why is it important?**
  – Enables segmentation of information based on source
  – Enhances provider trust in information being exchanged between providers
  – Enhances provider trust in information received from a patient (e.g., data from PHR)
About the Environmental Scan

• Explore how EHRs, PHRs and health information exchanges (HIEs) track data provenance
• Focus on marking and retaining provenance as systems aggregate data from multiple sources and records are exchanged
• Focus on provenance within CDA documents
• Includes landscape analysis, gap analysis
Key Provenance Considerations

• **Who/What to list as source?**
  – Organization
  – Provider
  – Data entry staff
  – Device details

• **At what level?**
  – Entire document (full record has one source listed)
  – Section (ex. medication section lists its own source)
  – Individual data element (ex. each individual medication lists its own source)

• **How to update or modify source when importing/exporting?**
  – List receiving organization as source
  – List original creator as source
• How are systems documenting provenance today?
  – Considerable variability in how provenance marked and retained in EHRs, PHRs, and HIEs
    • Different levels of granularity (document level vs section level vs data element level)
    • Different practices regarding source (organization vs provider vs individual entering the data)
    • Different practices when importing/exporting data (list original source vs modify source to list importing org)
Flow Down of Provenance Data

• Common response to variability is to utilize “flow down” of provenance data
  – Mark provenance at **document level only**
  – Provenance data inherited at data element levels
    • Example: the source for an individual medication will be the same source listed at document level
  – Results in insufficient granularity of detail; creates integrity issues; undermines trust
Implications for Patient-generated Data

- Generally, provenance at data element level is lacking as records are shared with providers
  - EHRs often export with sending organization listed as source on **document level only** (not section or element)
  - PHRs may import with provenance at document, section and/or data element level, but often export with PHR as source on **document level only**
  - HIEs can export with documentation at data element level, but sufficient information often **not available**

- Resulting flow down of document level provenance data may not meet provider need for granular detail
Challenges and Opportunities

• Currently no dominant provenance model within the HIT community
  – No uniform way of handling data provenance when sharing data

• No harmonized standard currently in place
  – Upcoming work includes HL7 data provenance and privacy support in C-CDA initiative

• Current flow down practice does not meet provider need for granularity regarding origin of data