

## CONSUMER PERSPECTIVE

Consumer Empowerment	Chronic Care	Biosurveillance
<p><i>Near Term</i>            Lab results as needed by patient            List of conditions &amp; allergies</p> <ul style="list-style-type: none"> <li>• Health problems</li> <li>• Medication allergies</li> <li>• Allergies</li> </ul> <p>Administrative features</p> <ul style="list-style-type: none"> <li>• Appointment scheduling</li> <li>• Demographic profile</li> <li>• Editing account profile</li> <li>• Insurance eligibility &amp; claims</li> <li>• Financial recordkeeping &amp; management</li> <li>• Privacy &amp; access control</li> </ul> <p>Reminders (examples)</p> <ul style="list-style-type: none"> <li>• Annual check-ups</li> <li>• Cancer screening—mammograms</li> <li>• Cancer screening—colonoscopies</li> <li>• Immunizations</li> </ul> <p><i>Mid-Longer Term</i>            Online consultation</p> <ul style="list-style-type: none"> <li>• Structured email</li> </ul> <p>Summaries of healthcare encounters</p> <ul style="list-style-type: none"> <li>• Dates of services</li> <li>• Diagnosis codes</li> <li>• Procedure codes</li> </ul> <p>Educational information</p> <ul style="list-style-type: none"> <li>• Evidence based health information</li> </ul> <p>Decision support</p> <ul style="list-style-type: none"> <li>• Shared decision making</li> <li>• Communications preferences</li> </ul> <p>Patient health outcomes</p> <ul style="list-style-type: none"> <li>• Adverse events</li> <li>• Medical errors</li> <li>• Patient reported health outcomes</li> </ul> <p><i>Workgroup Issues</i>            PHR not integrated with workflow            Limited pre-populated clinical data &amp; limited patient access            Minimal interoperability or portability            Connectivity between physician offices, PHRs, &amp; pharmacies            State laws regarding labs            Policies for consumer entered data            Quality of pre-populated data</p>	<p><i>Near Term</i>            Secure messaging</p> <ul style="list-style-type: none"> <li>• Online consultation</li> </ul> <p>Vital signs</p> <ul style="list-style-type: none"> <li>• Weight</li> </ul> <p>Glucose monitoring            Spirometry</p> <p><i>Mid-Longer Term</i>            Anticoagulation            Vital signs</p> <ul style="list-style-type: none"> <li>• Blood Pressure</li> <li>• Heart rate and rhythm</li> <li>• Pulse oximetry</li> </ul> <p>Fall/motion monitoring            Monitoring of medications</p> <p><i>Other</i>            Vital signs (general)            Labs and pharmacy            Lesion assessment            Remote monitoring for chronic conditions</p> <p><i>Workgroup Issues</i>            HIT use in specific populations</p> <p><b>Limited interoperability</b></p> <p>Product and services certification            Medico-legal liability risks associated with remote care</p> <ul style="list-style-type: none"> <li>• State licensure constraints</li> </ul> <p><b>Need for confidentiality, privacy and security</b></p> <p>Patient identification for authorization and authentication</p>	<p>Case reporting</p> <ul style="list-style-type: none"> <li>• Lab results</li> <li>• Clinical symptomology</li> <li>• Integration with EHRs</li> <li>• Health alerting (HA)/email alerts</li> </ul> <p>Bi-directional communication</p> <ul style="list-style-type: none"> <li>• Collaborative discussions</li> <li>• Web pages</li> </ul> <ul style="list-style-type: none"> <li>• Interoperable minimum data set</li> </ul> <p>Response management</p> <ul style="list-style-type: none"> <li>• Vaccine</li> <li>• Chemoprophylaxis</li> <li>• Treatment</li> <li>• Isolation/quarantine</li> <li>• Resource identification               <ul style="list-style-type: none"> <li>○ Hospital beds</li> <li>○ Medications</li> <li>○ Medical personnel</li> </ul> </li> <li>• EHR data               <ul style="list-style-type: none"> <li>○ Immunization registry</li> <li>○ Disease registry</li> </ul> </li> </ul> <p>Adverse event reporting</p> <ul style="list-style-type: none"> <li>• Devices, drugs, biologic</li> </ul> <p>Nosocomial infections</p> <ul style="list-style-type: none"> <li>• Medication errors               <ul style="list-style-type: none"> <li>○ Ordering/prescribing/dispensing</li> <li>○ Drug-drug, drug-allergy interaction decision support</li> <li>○ Linkage to FDA structured product labeling database results</li> </ul> </li> </ul> <p><i>Workgroup Issues</i>            Lack of EHR case reporting, adverse events, and electronic lab reporting (ELR) integration  <b>Storage, retrieval, and management concerns of large amounts of data</b>            Lack of central dissemination process for public health HIT standards            Lack of interoperable bi-directional communication            Public health information network (PHIN) can be leveraged            Lack of EHR decision support to prompt</p> <ul style="list-style-type: none"> <li>• Immunization reminder</li> <li>• Prevention guidelines</li> </ul> <p>Need to integrate commercial sector supply chain and national stockpile  <b>Privacy/ Security Concerns</b>            National notifiable disease conditions have been identified</p>

**CONSUMER PERSPECTIVE (continued)**

EHR	Quality	AHIC
<p>Patient identification                      Pharmacy/allergy                      • Patient recorder use of pharmacological                      Lab results                      Problem lists                      Clinical/encounter notes                      Anatomic pathology results                      Vital signs                      Radiology reports                      Immunizations</p> <p><i>Workgroup Issues</i></p> <p>Limited interoperability</p> <p>Pharmacy/ medication interoperability</p> <p>Machine readable and interoperable</p> <ul style="list-style-type: none"> <li>• Encounter notes</li> <li>• Radiology reports</li> <li>• Lab results</li> </ul> <p>CLIA and state policies and guidance for secondary data uses</p> <p>Need for confidentiality, privacy and security</p> <p>Accurate and reliable patient identification needed</p>	<p><i>Near Term</i></p> <p>Inpatient quality measures (core set)</p> <ul style="list-style-type: none"> <li>• Labs</li> <li>• Diagnostic procedures</li> <li>• Medications</li> <li>• Vaccines</li> <li>• Treatment</li> </ul> <p>Ambulatory measures (core set)</p> <ul style="list-style-type: none"> <li>• Labs</li> <li>• Diagnostic procedures</li> <li>• Medications</li> <li>• Vaccines</li> <li>• Patient education</li> </ul> <p>Clinicians have access to feedback (self-assessment)</p> <ul style="list-style-type: none"> <li>• Clinical decision support</li> </ul> <p>Public reporting</p> <ul style="list-style-type: none"> <li>• Data aggregation</li> </ul> <p><i>Mid-Longer Term</i></p> <p>Clinical decision support                      Expanded inpatient quality measures                      Expanded ambulatory quality measures</p> <p><i>Workgroup Issues</i></p> <p>Lack of data and technical standards and clinical documentation</p> <p>Data sharing rights and responsibilities</p> <p>Data security and privacy—policies for secondary uses</p>	<p><i>Cross-Cutting Themes</i></p> <p>Labs, medications, allergies, immunizations                      Securing messaging/on-line consultation                      Bi-directional communications                      Adverse event reporting                      Case reporting                      Clinical decision support systems                      Identification/authentication                      Problem lists                      Clinical encounter notes                      Family history/social factors                      Vitals signs                      Population health/conditions                      Minimum data set                      Confidentiality, privacy &amp; security of patient data                      Data access / Data control                      Data aggregation                      Infrastructure areas missing</p> <ul style="list-style-type: none"> <li>• Security, Network, Repositories</li> </ul> <p><i>Missing Areas</i></p> <p><i>Consumer Empowerment</i></p> <p>Vital measurements                      Text documents                      Provider list                      Health literacy (multilingual support)                      Reminders                      Advance directive/living wills                      Social/family history                      PHR portability methods                      Medication history                      E-prescribing</p> <p><i>Chronic care</i></p> <p>Standardization of device interfaces                      Care plans/clinical flowsheets                      Provider list                      Biosurveillance                      Adverse events / Nosocomial infections                      Clinical data storage for surveillance                      Case reporting                      Bi-directional communications</p> <p><i>EHR</i></p> <p>Lab results                      Anatomic pathology results                      Radiology reports                      Social history                      Procedure reports                      Medications                      Referrals</p> <p><i>Population Health</i></p> <p>Dental</p> <p><i>Workgroup Issues</i></p> <p>Workflow integration                      Int'l public health collaboration                      Legal liability &amp; regulatory barriers                      Consumer consent</p>

## CONSUMER PERSPECTIVE (continued)

CCHIT
<p><b>Patient safety</b></p> <p>Transfer of care</p> <ul style="list-style-type: none"><li>• Transmission of patient data (problems, allergies, med list, etc) across providers and settings</li><li>• Ambulatory and emergency referrals, discharge summaries, etc.</li></ul> <p><b>Medication management</b></p> <ul style="list-style-type: none"><li>• Outpatient prescription writing and transmission to pharmacies</li><li>• Ordering</li><li>• Clinical decision support</li><li>• Transmission</li><li>• Dispensing</li><li>• Administering</li><li>• Reconciliation</li></ul>
HITSP
<p><b>EHR/clinical record</b></p> <ul style="list-style-type: none"><li>• Primary data source standards<ul style="list-style-type: none"><li>○ Problem lists</li><li>○ Meds</li><li>○ Allergies</li><li>○ Text reports</li><li>○ Numeric results</li><li>○ Labs/micro</li><li>○ Images</li></ul></li><li>• HIPAA covered entities<ul style="list-style-type: none"><li>○ X12 Claims attachment</li></ul></li></ul> <p>Secondary uses of data</p> <ul style="list-style-type: none"><li>• Clinical research</li><li>• Clinical trials</li><li>• Population health</li></ul> <p>Quality/control measurements</p> <ul style="list-style-type: none"><li>• Consistency across uses</li></ul> <p>Clinical device data</p> <ul style="list-style-type: none"><li>• Glucometers</li><li>• Monitors</li><li>• Smart pump</li></ul> <p><b>Cross use case work on security (standards)</b></p> <ul style="list-style-type: none"><li>• Vocabulary for role-based access control</li><li>• Audit logging and exchange</li><li>• Authentication models to support chain of trust data exchanges</li></ul>

## PROVIDER PERSPECTIVE

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**PROVIDER PERSPECTIVE (continued)**

EHR	Quality	AHIC
<p>Patient identification</p> <p>Pharmacy/allergy</p> <ul style="list-style-type: none"> <li>• Patient recorder use of pharmacological</li> </ul> <p>Lab results</p> <p>Problem list</p> <p>Clinical/encounter notes</p> <p>Anatomic pathology results</p> <p>Vital signs</p> <p>Radiology reports</p> <p>Immunizations</p> <p><i>Workgroup Issues</i></p> <p>Limited interoperability</p> <p>Pharmacy/ medication interoperability</p> <p>Machine readable and interoperable</p> <ul style="list-style-type: none"> <li>• Encounter notes</li> <li>• Radiology reports</li> <li>• Lab results</li> </ul> <p>CLIA and state policies and guidance for secondary data uses</p> <p>Need for confidentiality, privacy and security</p> <p>Accurate and reliable patient identification needed</p>	<p><i>Near Term</i></p> <p>Inpatient quality measures (core set)</p> <ul style="list-style-type: none"> <li>• Labs</li> <li>• Diagnostic procedures</li> <li>• Medications</li> <li>• Vaccines</li> <li>• Treatment</li> </ul> <p>Ambulatory measures (core set)</p> <ul style="list-style-type: none"> <li>• Labs</li> <li>• Diagnostic procedures</li> <li>• Medications</li> <li>• Vaccines</li> <li>• Patient education</li> </ul> <p>Clinicians have access to feedback (self-assessment)</p> <ul style="list-style-type: none"> <li>• Clinical decision support</li> </ul> <p>Public reporting</p> <ul style="list-style-type: none"> <li>• Data aggregation</li> </ul> <p><i>Mid-Longer Term</i></p> <p>Clinical decision support</p> <p>Expanded inpatient quality measures</p> <p>Expanded ambulatory quality measures</p> <p><i>Workgroup Issues</i></p> <p>Lack of data and technical standards and clinical documentation</p> <p>Data sharing rights and responsibilities</p> <p>Data security and privacy—policies for secondary uses</p>	<p><i>Cross-Cutting Themes</i></p> <p>Labs, medications, allergies, immunizations</p> <p>Securing messaging/on-line consultation</p> <p>Bi-directional communications</p> <p>Adverse event reporting</p> <p>Case reporting</p> <p>Clinical decision support systems</p> <p>Identification/authentication</p> <p>Problem lists</p> <p>Clinical encounter notes</p> <p>Family history/social factors</p> <p>Vitals signs</p> <p>Population health/conditions</p> <p>Minimum data set</p> <p>Confidentiality, privacy &amp; security of patient data</p> <p>Data access / Data control</p> <p>Data aggregation</p> <p>Infrastructure areas missing</p> <ul style="list-style-type: none"> <li>• Security, Network, Repositories</li> </ul> <p><i>Missing Areas</i></p> <p><i>Consumer Empowerment</i></p> <p>Vital measurements</p> <p>Text documents</p> <p>Provider list</p> <p>Health literacy (multilingual support)</p> <p>Reminders</p> <p>Advance directive/living wills</p> <p>Social/family history</p> <p>PHR portability methods</p> <p>Medication history</p> <p>E-prescribing</p> <p><i>Chronic care</i></p> <p>Standardization of device interfaces</p> <p>Care plans/clinical flowsheets</p> <p>Provider list</p> <p><i>Biosurveillance</i></p> <p>Adverse events / Nosocomial infections</p> <p>Clinical data storage for continuous surveillance</p> <p>Case reporting</p> <p>Bi-directional communications</p> <p><i>EHR</i></p> <p>Lab results</p> <p>Anatomic pathology results</p> <p>Radiology reports</p> <p>Social history / Procedure reports</p> <p>Medications</p> <p>Referrals</p> <p><i>Population Health</i></p> <p>Dental</p> <p><i>Workgroup Issues</i></p> <p>Workflow integration</p> <p>Int'l public health collaboration</p> <p>Legal liability &amp; regulatory barriers</p> <p>Consumer consent</p>

## PROVIDER PERSPECTIVE (continued)

CCHIT	
Patient safety	
Transfer of care	
<ul style="list-style-type: none"><li>• Transmission of patient data (problems, allergies, med list, etc) across providers and settings</li><li>• Ambulatory and emergency referrals, discharge summaries, etc.</li></ul>	
Medication management	
<ul style="list-style-type: none"><li>• Outpatient prescription writing and transmission to pharmacies</li><li>• Ordering</li><li>• Clinical decision support</li><li>• Transmission</li><li>• Dispensing</li><li>• Administering</li><li>• Reconciliation</li></ul>	
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HITSP	
EHR/clinical record	
<ul style="list-style-type: none"><li>• Primary data source standards<ul style="list-style-type: none"><li>○ Problem lists</li><li>○ Meds</li><li>○ Allergies</li><li>○ Text reports</li><li>○ Numeric results</li><li>○ Labs/micro</li><li>○ Images</li></ul></li><li>• HIPAA covered entities<ul style="list-style-type: none"><li>○ X12 Claims attachment</li></ul></li></ul>	
Secondary uses of data	
<ul style="list-style-type: none"><li>• Clinical research</li><li>• Clinical trials</li><li>• Population health</li></ul>	
Quality/control measurements	
<ul style="list-style-type: none"><li>• Consistency across uses</li></ul>	
Clinical device data	
<ul style="list-style-type: none"><li>• Glucometers</li><li>• Monitors</li><li>• Smart pump</li></ul>	
Cross use case work on security (standards)	
<ul style="list-style-type: none"><li>• Vocabulary for role-based access control</li><li>• Audit logging and exchange</li><li>• Authentication models to support chain of trust data exchanges</li></ul>	

## SECONDARY USE PERSPECTIVE

Consumer Empowerment	Chronic Care	Biosurveillance
<p><u>Near Term</u>            Lab results as needed by patient            List of conditions &amp; allergies</p> <ul style="list-style-type: none"> <li>• Health problems</li> <li>• Medication allergies</li> <li>• Allergies</li> </ul> <p>Administrative features</p> <ul style="list-style-type: none"> <li>• Appointment scheduling</li> <li>• Demographic profile</li> <li>• Editing account profile</li> <li>• Insurance eligibility &amp; claims</li> <li>• Financial recordkeeping &amp; management</li> <li>• Privacy &amp; access control</li> </ul> <p>Reminders (examples)</p> <ul style="list-style-type: none"> <li>• Annual check-ups</li> <li>• Cancer screening—mammograms</li> <li>• Cancer screening—colonoscopies</li> <li>• Immunizations</li> </ul> <p><u>Mid-Longer Term</u>            Online consultation</p> <ul style="list-style-type: none"> <li>• Structured email</li> </ul> <p>Summaries of healthcare encounters</p> <ul style="list-style-type: none"> <li>• Dates of services</li> <li>• Diagnosis codes</li> <li>• Procedure codes</li> </ul> <p>Educational information</p> <ul style="list-style-type: none"> <li>• Evidence based health information</li> </ul> <p>Decision support</p> <ul style="list-style-type: none"> <li>• Shared decision making</li> <li>• Communications preferences</li> </ul> <p>Patient health outcomes</p> <ul style="list-style-type: none"> <li>• Adverse events</li> <li>• Medical errors</li> <li>• Patient reported health outcomes</li> </ul> <p><u>Workgroup Issues</u></p> <p>PHR not integrated with workflow            Limited pre-populated clinical data &amp; limited patient access            Minimal interoperability or portability            Connectivity between physician offices, PHRs, &amp; pharmacies            State laws regarding labs            Policies for consumer entered data            Quality of pre-populated data</p>	<p><u>Near Term</u>            Secure messaging</p> <ul style="list-style-type: none"> <li>• Online consultation</li> </ul> <p>Vital signs</p> <ul style="list-style-type: none"> <li>• Weight</li> </ul> <p>Glucose monitoring            Spirometry</p> <p><u>Mid-Longer Term</u>            Anticoagulation            Vital signs</p> <ul style="list-style-type: none"> <li>• Blood Pressure</li> <li>• Heart rate and rhythm</li> <li>• Pulse oximetry</li> </ul> <p>Fall/motion monitoring            Monitoring of medications</p> <p><u>Other</u>            Vital signs (general)            Labs and pharmacy            Lesion assessment            Remote monitoring for chronic conditions</p> <p><u>Workgroup Issues</u>            HIT use in specific populations</p> <p>Limited interoperability</p> <p>Product and services certification</p> <p>Medico-legal liability risks associated with remote care</p> <ul style="list-style-type: none"> <li>• State licensure constraints</li> </ul> <p><b>Need for confidentiality, privacy and security</b></p> <p><b>Patient identification for authorization and authentication</b></p>	<p><b>Case reporting</b></p> <ul style="list-style-type: none"> <li>• Lab results</li> <li>• Clinical symptomology</li> <li>• Integration with EHRs</li> <li>• Health alerting (HA)/email alerts</li> </ul> <p><b>Bi-directional communication</b></p> <ul style="list-style-type: none"> <li>• Collaborative discussions</li> <li>• Web pages</li> <li>• Interoperable minimum data set</li> </ul> <p><b>Response management</b></p> <ul style="list-style-type: none"> <li>• Vaccine</li> <li>• Chemoprophylaxis</li> <li>• Treatment</li> <li>• Isolation/quarantine</li> <li>• Resource identification               <ul style="list-style-type: none"> <li>○ Hospital beds</li> <li>○ Medications</li> <li>○ Medical personnel</li> </ul> </li> <li>• EHR data               <ul style="list-style-type: none"> <li>○ Immunization registry</li> <li>○ Disease registry</li> </ul> </li> </ul> <p>Adverse event reporting</p> <ul style="list-style-type: none"> <li>• Devices, drugs, biologic</li> </ul> <p>Nosocomial infections</p> <ul style="list-style-type: none"> <li>• Medication errors               <ul style="list-style-type: none"> <li>○ Ordering/prescribing/dispensing</li> <li>○ Drug-drug, drug-allergy interaction decision support</li> <li>○ Linkage to FDA structured product labeling database results</li> </ul> </li> </ul> <p><u>Workgroup Issues</u></p> <p><b>Lack of EHR case reporting, adverse events, and electronic lab reporting (ELR) integration</b>  <b>Storage, retrieval, and management concerns of large amounts of data</b>  <b>Lack of central dissemination process for public health HIT standards</b>            Lack of interoperable bi-directional communication            Public health information network (PHIN) can be leveraged            Lack of EHR decision support to prompt</p> <ul style="list-style-type: none"> <li>• Immunization reminder</li> <li>• Prevention guidelines</li> </ul> <p>Need to integrate commercial sector supply chain and national stockpile</p> <p><b>Privacy/ Security Concerns</b>            National notifiable disease conditions have been identified</p>

**SECONDARY USE PERSPECTIVE (continued)**

EHR	Quality	AHIC
<p>Patient identification</p> <p>Pharmacy/allergy</p> <ul style="list-style-type: none"> <li>• Patient recorder use of pharmacological</li> </ul> <p>Lab results</p> <p>Problem lists</p> <p>Clinical/encounter notes</p> <p>Anatomic pathology results</p> <p>Vital signs</p> <p>Radiology reports</p> <p>Immunizations</p> <p><i>Workgroup Issues</i></p> <p>Limited interoperability</p> <p>Pharmacy/ medication interoperability</p> <p>Machine readable and interoperable</p> <ul style="list-style-type: none"> <li>• Encounter notes</li> <li>• Radiology reports</li> <li>• Lab results</li> </ul> <p>CLIA and state policies and guidance for secondary data uses</p> <p>Need for confidentiality, privacy and security</p> <p>Accurate and reliable patient identification needed</p>	<p><i>Near Term</i></p> <p>Inpatient quality measures (core set)</p> <ul style="list-style-type: none"> <li>• Labs</li> <li>• Diagnostic procedures</li> <li>• Medications</li> <li>• Vaccines</li> <li>• Treatment</li> </ul> <p>Ambulatory measures (core set)</p> <ul style="list-style-type: none"> <li>• Labs</li> <li>• Diagnostic procedures</li> <li>• Medications</li> <li>• Vaccines</li> <li>• Patient education</li> </ul> <p>Clinicians have access to feedback (self-assessment)</p> <ul style="list-style-type: none"> <li>• Clinical decision support</li> </ul> <p>Public reporting</p> <ul style="list-style-type: none"> <li>• Data aggregation</li> </ul> <p><i>Mid-Longer Term</i></p> <p>Clinical decision support</p> <p>Expanded inpatient quality measures</p> <p>Expanded ambulatory quality measures</p> <p><i>Workgroup Issues</i></p> <p>Lack of data and technical standards and clinical documentation</p> <p>Data sharing rights and responsibilities</p> <p>Data security and privacy—policies for secondary uses</p>	<p><i>Cross-Cutting Themes</i></p> <p>Labs, medications, allergies, immunizations</p> <p>Securing messaging/on-line consultation</p> <p>Bi-directional communications</p> <p>Adverse event reporting</p> <p>Case reporting</p> <p>Clinical decision support systems</p> <p>Identification/authentication</p> <p>Problem lists</p> <p>Clinical encounter notes</p> <p>Family history/social factors</p> <p>Vitals signs</p> <p>Population health/conditions</p> <p>Minimum data set</p> <p>Confidentiality, privacy &amp; security of patient data</p> <p>Data access / Data control</p> <p>Data aggregation</p> <p>Infrastructure areas missing</p> <ul style="list-style-type: none"> <li>• Security, Network, Repositories</li> </ul> <p><i>Missing Areas</i></p> <p><i>Consumer Empowerment</i></p> <p>Vital measurements</p> <p>Text documents</p> <p>Provider list</p> <p>Health literacy (multilingual support)</p> <p>Reminders / Advance directive/living wills</p> <p>Social/family history</p> <p>PHR portability methods</p> <p>Medication history</p> <p>E-prescribing</p> <p><i>Chronic care</i></p> <p>Standardization of device interfaces</p> <p>Care plans/clinical flowsheets</p> <p>Provider list</p> <p><i>Biosurveillance</i></p> <p>Adverse events</p> <p>Nosocomial infections</p> <p>Clinical data storage for continuous surveillance</p> <p>Case reporting</p> <p>Bi-directional communications</p> <p><i>EHR</i></p> <p>Lab results</p> <p>Anatomic pathology results</p> <p>Radiology reports</p> <p>Social history/ Procedure reports</p> <p>Medications</p> <p>Referrals</p> <p><i>Population Health</i></p> <p>Dental</p> <p><i>Workgroup Issues</i></p> <p>Workflow integration</p> <p>Int'l public health collaboration</p> <p>Legal liability &amp; regulatory barriers</p> <p>Consumer consent</p>

## SECONDARY USE PERSPECTIVE (continued)

CCHIT
Patient safety
Patient safety
Transfer of care
<ul style="list-style-type: none"><li>• Transmission of patient data (problems, allergies, med list, etc) across providers and settings</li><li>• Ambulatory and emergency referrals, discharge summaries, etc.</li></ul>
<b>Medication management</b>
<ul style="list-style-type: none"><li>• Outpatient prescription writing and transmission to pharmacies</li><li>• Ordering</li><li>• Clinical decision support</li><li>• Transmission</li><li>• Dispensing</li><li>• Administering</li><li>• Reconciliation</li></ul>
HITSP
<b>EHR/clinical record</b>
<ul style="list-style-type: none"><li>• Primary data source standards<ul style="list-style-type: none"><li>○ Problem lists</li><li>○ Meds</li><li>○ Allergies</li><li>○ Text reports</li><li>○ Numeric results</li><li>○ Labs/micro</li><li>○ Images</li></ul></li><li>• HIPAA covered entities<ul style="list-style-type: none"><li>○ X12 Claims attachment</li></ul></li></ul>
Secondary uses of data
<ul style="list-style-type: none"><li>• Clinical research</li><li>• Clinical trials</li><li>• Population health</li></ul>
Quality/control measurements
<ul style="list-style-type: none"><li>• Consistency across uses</li></ul>
Clinical device data
<ul style="list-style-type: none"><li>• Glucometers</li><li>• Monitors</li><li>• Smart pump</li></ul>
<b>Cross use case work on security (standards)</b>
<ul style="list-style-type: none"><li>• Vocabulary for role-based access control</li><li>• Audit logging and exchange</li><li>• Authentication models to support chain of trust data exchanges</li></ul>