Medication ReconciliatioN

&

Polypharmacy Committee

Annual Report and Recommendations Submitted to: Connecticut Health Information Technology Advisory Council [DATE OF SUBMISSION]

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Acknowledgements:

On behalf of the State of Connecticut, the Executive Director of the Office of Health Strategy (OHS) and the Health Information Technology Officer express their sincerest gratitude to all those who participate in the Medication Reconciliation and Polypharmacy (MRP) Committee. The safe and effective use of medications is essential for the health of the citizens of Connecticut. The insights and perspectives of participants results in meaningful, practical, and realistic recommendations that will no doubt lead to improvements in the health and wellbeing of the citizens of Connecticut.

Executive Summary:

The Medication Reconciliation and Polypharmacy Committee (MRPC) was officially established as a standing committee of the Health Information Technology Advisory Council (HITAC) on September 19th, 2019. The MRPC was established to continue the work of the Medication Reconciliation Polypharmacy Working Group, the major output of which was a report with eleven recommendations related to medication reconciliation and polypharmacy¹.

During the past year the MRPC has organized and met to prioritize and advance the working group recommendations. The committee and its members were impacted by the COVID-19 pandemic – both personally and professionally in some cases. After a brief hiatus during the early part of the pandemic, the committee continued to meet in a fully remote format and redesigned its activities to meet social distancing requirements. This report summarizes the progress made and work planned for next year.

Notable achievements in the first year of committee activity include:

- Establishing a public-private cross-functional membership to ensure representation of all voices in committee matters and to garner support for committee outputs and decisions. Membership includes two patient advocates;
- Advancing the work identified in the recommendations of the working group. The committee made progress with deprescribing, CancelRx, medication reconciliation process improvement, and supported related professional education events.
- Developing a robust set of business and functional requirements for a best possible medication history (BPMH) across all residents of the State of CT. These requirements will provide a strong framework in developing a technical solution towards a BPMH.

A summary of the committee activities is presented in Table 4.

The main body of this report presents a high-level summary of the committee's activity and progress during the past year. Appendices are used to document outputs as stand-alone sections that can be easily referenced and made available to subsequent efforts.

¹ https://portal.ct.gov/-/media/OHS/Health-IT-Advisory-Council/MRP/Final-Recoms-Med-Rec.pdf

Background:

Medication Reconciliation and Polypharmacy

The MRP Working Group report notes that medications are a large and growing component of the prescriber's armamentarium and are the first line treatment for 88% of chronic diseases.² As of the publication of that report, the percentage of patients taking multiple prescription medications is also increasing. According to the most recent data (2011-2014) from the Centers for Disease Control and Prevention (CDC), 40.7% of seniors (65 years or older) and 10.9% of the total population were taking five (5) or more prescription medications within the past 30 days. For seniors, the 40.7% represents almost a three-fold increase from the period of 1988-1994 (13.8%). More recent data (2015-2016³) show 22.4% of the population, aged 40-79, taking five or more prescription medications within the past 30 days.

Because a patient's medication regimen is the basis for many treatment decisions, it is extremely important that medication lists are accurate to maximize therapeutic impact and prevent potentially life-threatening patient safety events.

Continuing the work of the MRP Work Group, Medication Reconciliation and Polypharmacy Committee was chartered as a standing committee of the Health IT Advisory Council. The Health IT Advisory Council was established per Public Acts 16-77 and 17-2. These acts recognized a need within the State of Connecticut for a designated Health Information Technology Officer (HITO), and the Health IT Advisory Council was established to advise the HITO and coordinate health IT activities for health reform initiatives in Connecticut. This standing committee, the Medication Reconciliation and Polypharmacy Committee (MRPC) was officially established on September 19th, 2019. As described in the MRPC Project Charter⁴, the purpose of the MRPC is:

to provide strategic guidance, recommendations and ongoing support to the Health IT Advisory Council and the Office of Health Strategy (OHS) for the development and maintenance of a best possible medication history (BPMH), supported by communication, education, and user-friendly digital tools.

To that end, the MRPC has five main goals through September 2021:

- 1. Develop an approach for the creation of a BPMH
- 2. Create an online directory of relevant tools and solutions for communication to key stakeholders
- 3. Support⁵ development and implementation related to key medication reconciliation activities⁶
- 4. Develop an implementation plan for the eleven recommendations from the working group
- 5. Actively monitor and support funding opportunities related to the purpose and goals of the MRPC

These five goals are closely tied to the eleven recommendations from the working group, and the activities that have occurred since the establishment of the committee are summarized in **Error! Reference source not found.**. The remainder of this report will focus on the progress toward the five goals and eleven recommendations.

² Ekstrand, MJ. *Transforming "Med Wreck" into "Med Rec:" One Health System's Journey*. Webinar presentation: Pharmacy Quality Alliance; July 2017.

³ https://www.cdc.gov/nchs/data/databriefs/db347-h.pdf

⁴ See Appendix A of this report, or visit the OHS website

⁵ The MRPC is to support: The Office of Health Strategy, Health Information Alliance, Inc., the Department of Social Services, and the Department of Consumer Protection

⁶ Activities related to deprescribing and CancelRx described in Appendix C.

Committee Membership and Goals

Membership of the MRPC broadly represents stakeholders involved in matters of medication reconciliation and polypharmacy. Per the charter for the group, experience and expertise is to involve, at a minimum a diverse group of healthcare professionals, consumer advocates, industry representatives, informaticians, and subject matter experts, as detailed in Table 1 below. The committee meeting schedule for the past year is presented in Table 2 below. All meeting materials including agendas, minutes, and recordings can be found on the OHS website⁷.

Table 1: MRPC Members, Affiliations, and Membership Categories

Member Name	Membership Category
Sean Jeffery, Co-Chair	Expert in medication reconciliation
Nitu Kashyap, Co-Chair	Expert in medication reconciliation
Alejandro Gonzalez-Restrepo	Expert in psychopharmacology
Amy Justice	Expert in polypharmacy
Anne VanHaaren	Pharmacist
Diana Mager	Represents LTPAC/Hospice
Ece Tek	Prescribing practitioner
Elizabeth Taylor	Represents a state agency
Jason Gott	Represents a state agency
Jennifer Osowiecki	Represents expertise in law
Jeremy Campbell	Represents pharmaceuticals
Kate Sacro	Expert in medication reconciliation
Lesley Bennett	Represents consumers
Margherita Giuliano	Pharmacist
Marie Renauer	Represents hospitals
Mark Silvestri	Represents a FQHC
MJ McMullen	Represents expertise in CancelRx
Nate Rickles	Expert in polypharmacy
Patricia Carroll	Represents consumers
Rachel Petersen	Represents expertise in CancelRx
Rod Marriott	Represents a state agency
Stacy Ward-Charlerie	Represents expertise in CancelRx
Valencia Bagby-Young	Represents a state agency

Table 2: MRPC Planned 2020 Meetings

Planned 2020 Meeting Schedule of the MR	
January 13, 2020	July 20, 2020*
February 27, 2020	August 17, 2020
March 16, 2020 *	September 24, 2020 *
April 20, 2020	October 19, 2020 *
May 18, 2020 *	November 16, 2020 *
June 25, 2020 *	December 21, 2020 *

Note: In light of Covid 19, in March 2020 the committee moved to virtual meetings. All virtually held meetings are marked with an (*). Additionally, two planned meetings were cancelled, those dates are identified with a strike through (-).

As shown in Figure 1, the MRPC has continued the work of the MRP Working Group, with great success toward the main goal of developing requirements for the best possible medication history⁸. The remainder of this report will give more detail into the work done to date toward the eleven recommendations of the MRP Workgroup.

⁷ https://portal.ct.gov/OHS/HIT-Work-Groups/Medication-Reconciliation-and-Polypharmacy-Committee/MRPC-Meeting-Agenda

⁸ https://ctvideo.ct.gov/ohs/MRPC Brown Presentation.mp4

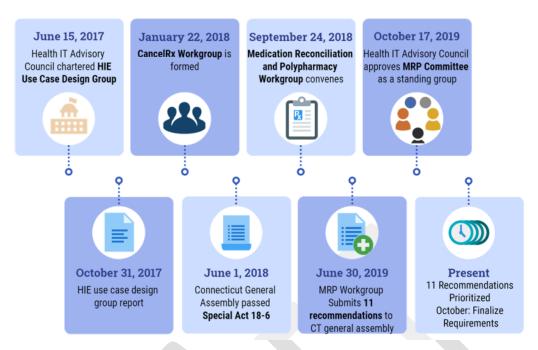


Figure 1: MRPC Evolution and Activities

Committee Progress

The committee was approved in October 2019 and convened its first session in November 2019. Members elected two co-chairs to serve. With general agreement on committee objectives and goals for 2020, the group started with reviewing and prioritizing tasks. We used several tools to elicit member comments and generate a healthy discussion. Subsequently focusing on some time-sensitive tasks before launching work on some foundation tasks. The key phases of work are summarized below.

- 1. Identify goals and objectives: A charter with overall goals, objectives and deliverables was created and approved. See Appendix A for charter
- 2. Prioritize tasks: The committee members unanimously agreed to pursue the recommendations summarized in the Medication Reconciliation Polypharmacy Workgroup report. An electronic survey was created to elicit impact and effort assessments from all members. Results of the survey are presented in this impact-effort grid. From this grid, low effort items without any dependencies were selected as work streams in addition to a single foundational element that was deemed as the cornerstone for any medication reconciliation at any level of care. This critical element was the Best Possible Medication History (BPMH) that is defined in Figure 3.



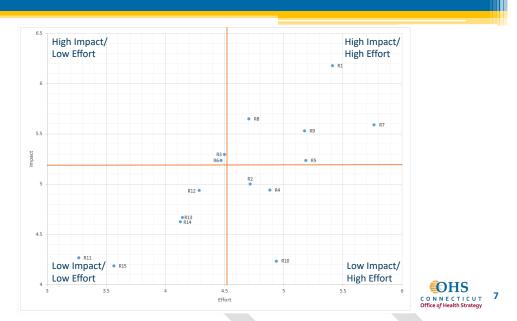


Figure 2: Impact Effort Grid

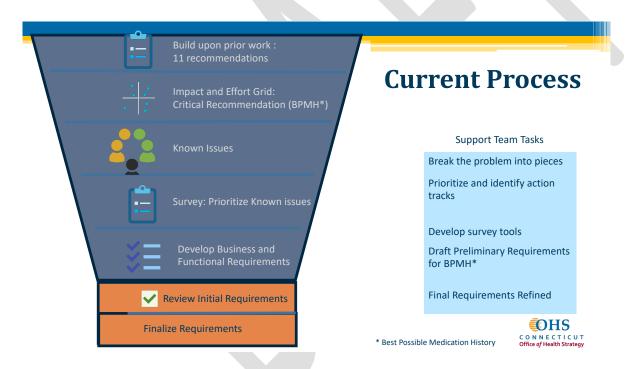


Figure 3: BPMH Recommendation Development

3. Timelines and Deliverables: Priority tasks were next plotted on a timeline to account for any deliverable deadlines or collisions with other important deadlines. For instance, new SCRIPT standards for e-prescribing afforded an opportunity to promote the CancelRx message standard for electronic deprescribing of medications that are discontinued in an electronic prescribing system. This work built upon prior work done by the CT State Cancel Rx subgroup. The overall initial timeline for tasks is presented in Figure 4 below. The timelines needed to be revised due to two unexpected natural events. In April, the State of CT saw its peak of COVID-19 cases during the first wave of the pandemic. Several committee members were pulled to alternate duties and the April meeting was suspended. Similarly, in August, storm damage across the state required us to cancel the meeting. Despite these natural disasters, the committee members with support from the administrative team were able to iteratively work on and collate BPMH requirements. These are summarized below under the BPMH section.

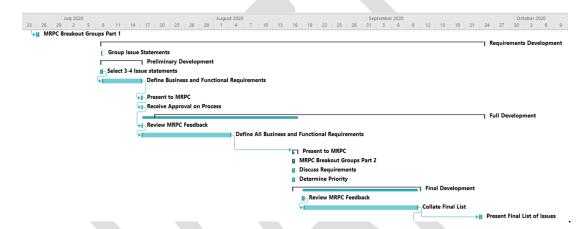


Figure 4: MRPC Abbreviated Activities Timeline

Finalize and Plan Next Steps: At the close of fiscal year 2020, the committee drafted its final report which is presented here. In addition, a roadmap for next steps was reviewed and is presented in the Future work section below. We recognize that medication safety is an important public health mission and needs broad participation across stakeholders. During the committee meetings, we have attempted to stay abreast of developments in this space across the state and the country. We have had periodic presentations from the Health Information Technology Advisory Council (HITAC), which was established to advise the State of Connecticut's Health Information Technology Officer and coordinate health IT activities for health reform initiatives in Connecticut. Additionally, the MRPC has kept informed on the development of the Health Information Exchange, Connecticut Information Exchange (ConnIE), a nonprofit, nongovernmental entity established to enable the development and operation of health information exchange (HIE) services for the state of Connecticut. Finally, the committee has met with other organizations to understand pros and cons of an approach to expand the Prescription Drug Monitoring Program (PDMP), which collects prescription data for Schedule II through Schedule V drugs into a centralized database. In Connecticut the Prescription Monitoring and Reporting System (CPMRS) is used by healthcare providers and pharmacists in the active treatment of their patients. The purpose is to present a complete picture of a patient's-controlled substance use, including prescriptions by other providers. The future work section encompasses this situational awareness into a vision of possible ways to bring BPMH to the citizens and clinicians across the State. The committee membership and support

teams have made consistent efforts to stay tied into state agency activities, and maintain collaborative working relationships, with the realization that the work of this committee is a part of the larger state Health IT landscape.

Committee Tasks

Key tasks and activities related to each task are summarized in this section. These tasks were identified and prioritized as outlined in committee progress section above.

Best Possible Medication History (BPMH)

In keeping with the MRP workgroup recommendation for near-term efforts focused on making tangible progress toward an enhanced and uniform best possible medication list, the committee made a concerted effort to move forward with the development of the BPMH. Efforts began with a curated a statement to capture the spirit of a BPMH:

Safe, quality and timely delivery of healthcare requires access to the 'Best-Possible Medication History". The BPMH should include all prescription and non-prescription medications, supplements and herbal products. The BPMH should be accurate, up-to-date and accessible to stakeholders (including but not limited to patients, caregivers and health care providers) at the point of decision making. Access to the BPMH will support collaborative care, reduce medication costs and errors and improve clinical outcomes.

This statement embodies the objectives listed in the charter while recognizing that in the short term the focus will be on identifying critical features and input sources for the BPMH. To identify and refine the requirements statements⁹, we first identified roles that would be consumers of the BPMH. Roles were then grouped into logical cohorts such as clinicians, prescribers and health systems in one group and patients, care givers and home health agencies in another group. Thereafter using breakout sessions, each group identified gaps in medication history

that exists today in their respective areas. These gaps or "known issues" were statements for answering the question "What problem are we trying to solve?" These issues were then distilled into features that would be most useful in a BPMH as business and functional requirements by the administrative team and endorsed by the committee. A timeline of activities can be seen in Figure 4, and a list of known issue statements, business requirements, functional requirements are detailed in Appendix B.

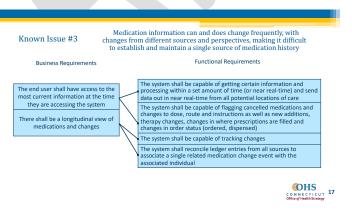


Figure 5: Development of the BPMH Requirements

⁹ Business requirements define the 'why' for the BPMH: they provide the scope of needs to be addressed. Functional requirements define the 'what' for the BPMH: they provide the specific steps the MRPC will need to take to achieve the business requirements. These requirements are more explicitly described in Appendix B.

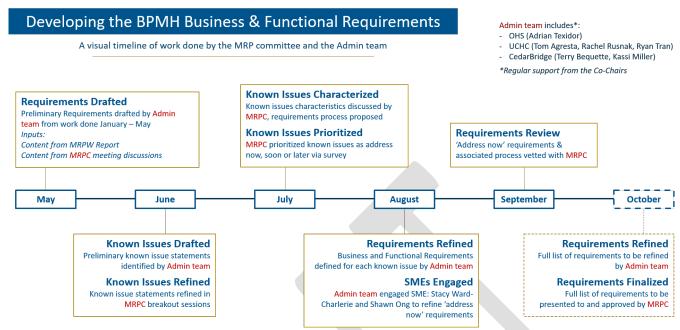
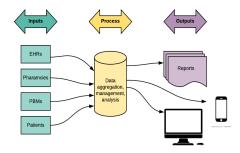


Figure 6: Development of the BPMH Business & Functional requirements

Next Steps

The business and functional requirements as outlined in Appendix B provide a strong foundation for informing a technical solution for medication history regardless of the scope or tool used. For instance, sources of discrete medication history that are available in a standardized and shareable digital format are identified. Critical data elements needed for medication history to be useful such as dose, strength, last dispense date are identified. The need for near-real-time updates is clearly evident from these requirements. Any technical solution, whether designed as a dynamic clearinghouse or repository would need to meet these requirements. Furthermore, some

elements of a user interface by role are also described in these requirements. For example, clinicians would need to see the latest information by default and would like access to dispense history which can be a click or two away. The MRPC will continue to expand on these feature sets and technical specifications and partner with other stakeholders at state or national level.



Implementation and Adoption of CancelRx and promoting Deprescribing

The ability to cancel a prescription medication electronically has existed from a technical perspective for several years through a technical messaging standard (SCRIPT Standard 10.6) developed by the National Council for Prescription Drug Programs (NCPDP) and adopted by the Office of the National Coordinator (ONC). With the implementation of 2017 NCPDP standards, all prescribing systems were required to support sending electronic cancelation messages when a medication is discontinued. This offered an opportunity to promote education, awareness and preparedness amongst end users and pharmacies. Through advocacy efforts, we influenced decision by a major national e-prescribing hub to remove any implementation licensing fees to pharmacies for sending medication cancellation messages using the CancelRx standard. This was identified in previous work as a significant barrier for retail pharmacies to adopt the new standard. We noted in impressive uptake of adoption of this standard by retail pharmacies. See Figure 5, which describes Surescripts trends in Connecticut. A statewide survey of all prescribers was conducted in partnership with the State PDMP program. This was followed by a statewide webinar to address common misconceptions and promote CancelRx adoption.

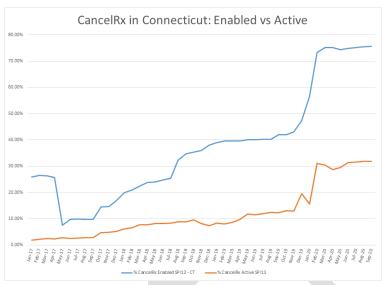


Figure 7: Surescripts Trends in Connecticut

Committee members also developed and presented a webinar for providers across the state on deprescribing. Deprescribing, the planned and supervised process of dose reduction or stopping of medication, is an essential practice of good prescribing, and is closely tied to CancelRx, an electronic tool that facilitates communication around deprescribing. CancelRx allows prescribers to send electronic cancellation messages to pharmacies when medications are discontinued. One committee served as a moderator and two additional members served as panelists at a successful continuing medical education event hosted in June of this year. The webinar had 165 attendees with learning objectives that included the impacts of polypharmacy, the challenges associated with deprescribing, the role of Health IT in medication management, CancelRx workflow and best practices to implement and apply CancelRx in clinical practice.

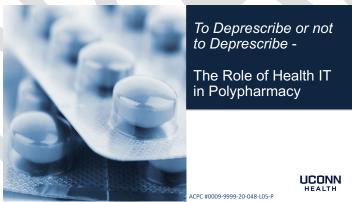


Figure 6: Image from the Deprescribing webinar.

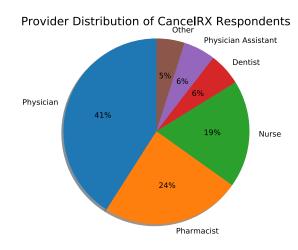


Figure 7: Distribution of responding provider types.

Next Steps

In addition to ongoing education to improve the effectiveness of deprescribing practices the committee will review the relationship of deprescribing with activity associated with other recommendations. For example, deprescribing has overlap with team-based medication reconciliation and with reports that might be developed for the BPMH. With the close ties of deprescribing and CancelRx in mind, the MRPC considers adoption of CancelRx to be a critical component of the committee's purpose and goals. The committee will continue to monitor the implementation of CancelRx in Connecticut and will consider policy initiatives, including legislation, to ensure this standard is fully adopted in the state and that measures of this tool's effectiveness are in place.

Medication Reconciliation Resources

Medication reconciliation resources have been identified and produced. These resources are expected to be available via a web repository in Winter 2020. Preliminary resources, including peer reviewed literature, as well as documentation around standards, recommendations and best practices have also been collated and will be included in the repository. This work is ongoing.

SUPPORT Act and IAPD funding

A number of timing factors, including the startup of Connie as the state's HIE and the availability of resources to pursue activities related to the SUPPORT Act in the short window of time available through the IAPD, precluded making much progress with these considerations. However, the fundamental interest of leveraging the Prescription Drug Monitoring Program (PDMP) database and processes to support the BPMH is still keen. The Director of the Department of Consumer Protection (DCP) Drug Control Unit, who oversees and administers the State's PDMP, is an active member of the MRPC. Recently awarded Support Act funds, have resulted in plans that will provide integration for EHRs with Connecticut's Prescription Drug Monitoring Program. This integration will occur at no cost for providers and health systems and will facilitate easier access to Connecticut's PDMP for providers. Ongoing close coordination with the DCP is planned to ensure harmonization between the agency and the MRPC. A potential area for collaboration for the MRPC could include identifying how to best facilitate the adoption of the PDMP/EHR integration opportunity and driving adoption and use throughout the state. The MRPC could offer as educational opportunities to showcase how other HIEs and healthcare systems across the country have done this in implemented successfully.

Privacy and confidentiality of medication-related information was frequently mentioned in the course of discussing other topics at MRPC meetings during the past year. Operation of the BPMH, when it becomes an implemented resource for providers and prescribers, must align with HIPAA. Depending on the data sources, the BPMH may also be impacted by alignment with 42 CFR Part 2¹⁰.

Due to the timing of IAPD submission and approval the MPRC was not able to access certain planning funding available through the IAPD. However, these funds should now be available in FFY 2021 and the committee's ideas for utilizing these funds to advance the work are being finalized.

Next Steps

The MRPC will determine how best to align the activity and projects of the committee with the Medicaid enterprise, to the fullest extent possible. Initiatives such as the BPMH, CancelRx, and best practices for deprescribing and polypharmacy have benefits for Medicaid populations and providers and the associated Medicaid funding can help to secure implementation and ongoing support.

As the MRPC wraps up the requirements for a BPMH it will continue to work on policy alignment including medication quality measures, incentives for medication management and reconciliation, assessing a policy mandate for the CancelRx standard, privacy and confidentiality of medication-related information, and introducing team-based medication reconciliation into healthcare provider workflows.

The committee will continue to monitor activities related to the integration of the Prescription Drug Monitoring Program, updates, enhancements and to evaluate potential collaboration opportunities with DCP, in efforts to support the BPMH and medication reconciliation in general.

¹⁰ https://www.hhs.gov/about/news/2020/07/13/fact-sheet-samhsa-42-cfr-part-2-revised-rule.html

Table 4: Overview of MRPC Accomplishments on MRPW Recommendations

Recommendation	Task	Activity
Best Possible Medications History (BPMH)	Near-term efforts (1-2 years) should be focused on making tangible progress toward an enhanced and uniform best possible medication list.	The committee has engaged in a thoughtful process to identify known concerns and translated them in to business and functional requirements.
Patient Engagement	Develop a process for patient and family/caregiver engagement.	The development of requirements for the BPMH was undertaken with the patient at the forefront. Several of the identified requirements pertain to patient and family engagement. Activities in this area will continue in 2021.
Medication Reconciliation Process Improvements	Develop a repository of evidence-based, best practice medication tools, technical advisories, subject matter experts, and policy and regulatory guidance documents should be developed.	Surveys have been deployed, a webinar series has been initiated, and the development of the repository is in progress, and expected to launch in Winter 2020.
Team Approach	Mission critical team members, whose participation in medication reconciliation is essential for success, should be identified.	Key roles were identified in the process of the BPMH requirements development process.
Implementation and Adoption of CancelRx	Complete a formal assessment of the return on investment (ROI) for the CancelRx standard and other medication reconciliation recommendations to support the widespread adoption by pharmacies should be conducted.	CancelRx adoption in Connecticut is relatively high. Alternative approaches to increase adoption are under consideration.
Deprescribing	Identify evidence-based best-practices for deprescribing and added to the repository of medication reconciliation tools.	Surveys were developed and deployed to providers and pharmacists in Connecticut. Additionally, a continuing education webinar on deprescribing was hosted in June 2020. The materials developed for the webinar, as well as survey results, and materials identifying best practices will be included in the forthcoming repository.
Technology	Focus on developing a best possible medication list, leveraging existing data resources that include community pharmacies, PBMs, and EHRs.	The committee has discussed technology and potential solutions regularly and has identified some technical suggestions for the BPMH. Technology will remain a consideration for the committee.
SUPPORT Act Funding and Planning/Design Process	Develop a process for communication and coordinated planning between the SUPPORT Act activities and the initiatives and future planning activities recommended by the MRP Work Group.	In progress
Aligned Policy	Identify and align work with medication quality measures that align clinically meaningful outcomes.	The committee has considered policy issues and keeps them in mind the work completed thus far in the development of the BPMH, and work in deprescribing and CancelRx. Further work is expected in this area.
Planning/Design Process and Use of IAPD Funding	Funds from the current IAPD should be utilized to finalize planning, design and requirements for the projects and services recommended in this report.	These plans were delayed due to the Covid 19 public health crisis. The Committee is planning to pursue work in this arena in 2021.
Continuation of the MRP Workgroup	The MRP Work Group should be constituted as a Standing Committee of Health IT Advisory Council.	The workgroup was transitioned into a standing committee that has an approved charted, meets monthly, and is composed of diverse membership.

Next Steps and Future Work

As it pertains to the original five goals outlined in the MRPC charter, the committee considers Goal 1, the development of a detailed strategic approach for the creation of a BPMH, and Goal 4, the development of and implementation plan for deprescribing transaction standards to be completed. The requirements and recommendations of the committee for the BPMH address these goals and lay the foundation for future development at the State level. Progress towards Goal 2, providing support Implementation Advance Planning Document (IAPD) and Substance Use-Disorder Prevention that Promotes Opioid Recovery and Treatment for Patients and Communities (SUPPORT) Act funded initiatives as related to the MRPC are in progress, and the committee's ideas for utilizing these funds to advance the work are being finalized.

The committee proposes to focus future work on Goals 2 and 3: the development and launch of the online directory of medication management and medication reconciliation tools and solutions and continuing to serve as a resource and collaborative partners across Connecticut. Partners include the Office of Health Strategy (OHS), whose mission is to promote equal access to high quality health care, control costs and ensure better health for the people of Connecticut, the Connecticut Information Exchange (ConnIE), Connecticut's fledgling, non-profit health information exchange, with the mission of enabling the exchange of clinical information across all health care settings, the Department of Social Services (DSS), who administers the State's Medicaid program, the Department of Consumer Protection (DCP), who administers the prescription drug monitoring program, and others. The MRPC plans to continue this work through the original deadline of September 2021. Continued engagement of the MRPC and its membership with Connecticut State Agencies, non-profits, and national entities will strengthen the position of Connecticut and its journey towards medication safety. The Committee recognizes the value that these collaborations bring, and that these partners are those who will implement the recommendations and work we produce, moving them through the next stages of development, prototyping, and application.



Figure 8: An Illustration of MRPC Connection to State Entities

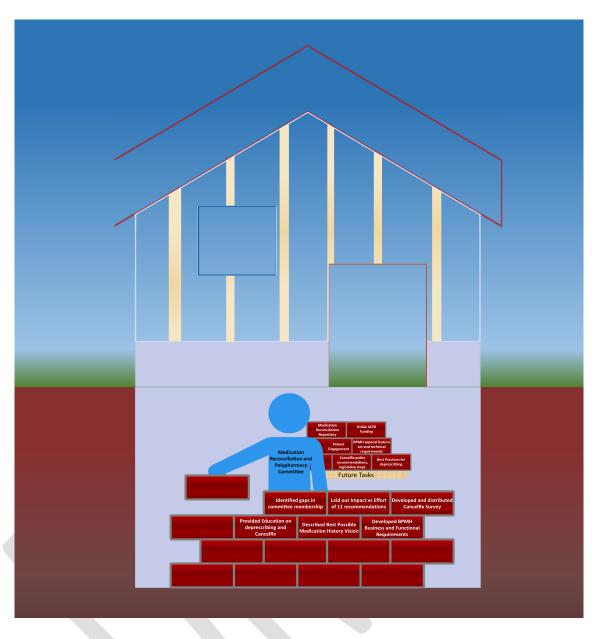


Figure 9: Foundational Activities of the MRPC

Project Charter

Medication Reconciliation & Polypharmacy Committee (MRPC) of the Health IT Advisory Council

September 24, 2019

Article 1: Name

Section 1: The name of this entity shall be the Medication Reconciliation & Polypharmacy Committee (MRPC), established by the Health Information Technology (IT) Advisory Council on September 19, 2019.

Article 2: Purpose

Section 1: The purpose of the MRPC is to provide strategic guidance, recommendations, and ongoing support to the Health IT Advisory Council and the Office of Health Strategy (OHS) for the development and implementation of patient-centered and evidence-based best practices necessary to contribute to the development and maintenance of a best possible medication history (BPMH), supported by communication, education, and user-friendly digital tools. The MRPC will build upon the approved recommendations and areas of focus identified by the Medication Reconciliation & Polypharmacy Work Group (found in Appendix A: MRP Work Group Recommendations). Through September 2021, the MRPC will focus on the following project goals:

- **Goal 1:** Develop a detailed strategic approach for the creation of a BPMH, supported by active patient engagement, that results in near-term value for stakeholders while laying the foundation for a longer-term, more extensive and integrated solution.
- Goal 2: Create an online directory of medication management and medication reconciliation tools and solutions for communication of evidence-based, best practice medication tools; patient engagement strategies; technical advisories; subject matter experts; and policy and regulatory guidance documents.
- Goal 3: Serve as a resource to OHS, Health Information Alliance, Inc., Department of Social Services, and Department of Consumer Protection to support development and implementation related to: technical solutions and use cases; workflow integration; medication reconciliation pilot activities; stakeholder engagement; and measurement and evaluation.
- Goal 4: Develop an implementation plan for the Medication and Polypharmacy Work Group recommendations related to deprescribing transaction standards, including CancelRx.
- Goal 5: Support Implementation Advance Planning Document (IAPD) and Substance Use-Disorder Prevention that Promotes Opioid Recovery and Treatment for Patients and Communities (SUPPORT) Act funded initiatives and actively monitor funding opportunities related to the stated purpose and goals of MRPC.

Other goals may be considered to support the purpose and goals of the MRPC, as needed. **Section 2:** The MRPC will not endorse or recommend any specific software solutions as part of its work; however, they may evaluate and review functionality for the purposes of achieving the above stated purpose and goals.

Article 3: Membership

Section 1: Membership in the MRPC shall be broadly representative of stakeholders involved in the matters of medication reconciliation and polypharmacy. Experience and expertise represented will include, at a minimum, the following:

- Patients, consumers, and caregivers
- Subject matter expertise in medication reconciliation
- Subject matter expertise in polypharmacy
- Community pharmacy
- Payers / pharmacy benefit managers
- Hospitals and health systems
- Providers / prescribers
- Long-term post-acute care, including skilled nursing and home health
- State agencies, including, but not limited to:
 - Department of Social Services
 - Department of Consumer Protection
 - Department of Mental Health and Addition Services
 - Department of Developmental Services
- Representation from the Health IT Advisory Council
- Health IT technology development and implementation

Section 2: Members of the MRPC shall initially be appointed by the Co-Chairs of the Health IT Advisory Council. Thereafter, the Chair, or Co-Chairs, in consultation with OHS, shall appoint members of the MRPC.

Section 3: Membership recruitment should take into consideration such factors as the geographic residence, race, ethnicity, and language of potential candidates to ensure that the group is as representative of as many perspectives and experiences as possible. This group will also include patient advocates to consult on patient engagement.

Section 4: As determined by the Chair, or Co-chairs, of the MRPC, additional subject matter experts (SMEs) may be sought on a permanent or periodic basis for the areas identified in the MRP Work Group's recommendations (see Appendix A), including, but not limited to: software development; electronic health record (EHR) and pharmacy information systems; and policy and regulations.

Section 5: Although this is not a time-limited group, membership will be reviewed annually by the Health IT Advisory Council and OHS to determine if membership is adequate to support the above stated purpose and goals of the MRPC. Recognizing that consistent participation in MRPC meetings is critical for success, failure by any member to attend at least 66% of meetings (within a given calendar year), or members who are absent for three consecutive meetings, shall result in consideration of termination from the MRPC. Members should notify the Chair, or Co-chairs, if they will be absent for any meeting. Members serve on a voluntary basis and without compensation.

Article 4: Officers

Section 1: The Chairperson shall be chosen by the members of the MRPC during the first scheduled meeting. The MRPC may also choose to elect Co-Chairs if Co-Chairs (rather than a single Chair) are deemed a better structure to support the stated purpose and goals.

Section 2: As Chair, or Co-chairs, the selected individual(s) will be responsible for setting meeting agendas, establishing regular meeting schedules, appointing subcommittees as needed, and

acting as liaison between the MRPC, OHS, the Health IT Advisory Council, and the Health Information Alliance, Inc.

Article 5: Subcommittees

Section 1: Subcommittees of the MRPC may be formed as needed by the Chair, or Co-chairs, in collaboration with OHS designated staff. Subcommittee leaders will be appointed by the Chair, or Co-chairs, in collaboration with OHS designated staff. The subcommittee lead member is responsible for organizing subcommittee meetings, with assistance from OHS staff, as necessary. The subcommittee lead member will report subcommittee findings and recommendations to the full MRPC for their information or action.

Article 6: Operating Procedures

Section 1: The MRPC operates as a standing committee of the Health IT Advisory Council. All records of the MRPC will be transmitted as soon as practical to OHS for inclusion in Health IT Advisory Council matters as appropriate.

Section 2: The MRPC is initially chartered for the period November 2019 through September 2021. At the conclusion of this initial period, the MRPC may decide to continue its work if deemed valuable at that time. If so, the MRPC will make a request for continuation to the Health IT Advisory Council. Otherwise, the MRPC shall document the basis for the conclusion of its work.

Section 2: The Office of Health Strategy (OHS) may establish procedures to allow members to participate in meetings by videoconference or teleconference.

Section 3: Meetings will be governed by Robert's Rules of Order, Abbreviated. One half of the membership will constitute a quorum. Action on agenda items may be taken by no less than a majority of members present at the meeting.

Section 4: The Chair, or Co-chairs, may solicit agenda items from members in advance of a meeting and establish agendas in collaboration with the OHS designated staff. Items may be added to the agenda on the day of the meeting if approved by the Chair, or Co-Chairs.

Section 5: All meeting information will be published on the Connecticut Public Notice web site and on the OHS web site. Meeting changes will be sent by email to members no later than 9 AM the day of the scheduled meeting.

Article 7: Duties of OHS

- OHS will provide the MRPC and the Chair, or Co-chairs, with support in the areas of meeting
 facilitation, the development of agenda and meeting materials, logistical planning and
 scheduling, research and analysis and stakeholder engagement. This support will be provided
 by OHS personnel or through engagement of professionals with required expertise.
- OHS shall inform the MRPC about all known changes in federal and state policy as well as rules and regulations that impact its work and the stated purpose and goals.
- OHS will consult with ongoing committees and advisory bodies in the state, maintain familiarity of the subject and purpose of the MRPC, and communicate perceived areas of opportunity for collaboration.
- OHS will ensure ongoing communication between the MRPC and relevant OHS staff and leadership as well as communication with the Health IT Advisory Council.
- OHS staff assigned to the MRPC will attend all meetings and inform its members of timely developments relevant to its work.
- An OHS administrative support member(s) will assist the MRPC's Chair, or Co-chairs, as needed, to maintain membership and interested parties with information, distribute meeting

agendas and notices to the membership and interested parties, and record the meeting minutes of the MRPC's meetings, including attendance.

Article 8: Duties of Health IT Advisory Council

- The Health IT Advisory Council shall approve the MRPC Project Charter and any updates.
- The Health IT Advisory Council shall regularly review work products of the MRPC and provide feedback as requested and appropriate.
- The Health IT Advisory Council will take action on MRPC recommendations as necessary to support the work of the MRPC and the purpose of the Health IT Advisory Council, consistent with enhancing the health and healthcare of CT and its residents.
- The Health IT Advisory Council will consider requests from the MRPC for resources and support as needed to support the Purpose and Goals of the MRPC.
- The Health IT Advisory Council will consider requests from the MRPC for extension of its activities beyond the initial period that concludes September 2021.



Appendix B: Medication Reconciliation and Polypharmacy Work Group Recommendations

1. Best Possible Medications History (BPMH)

The committee developed business and functional requirements for the proposed BPMH solution via work sessions, surveys, and environmental scan research.

2. Patient Engagement

To support this goal, the committee maintains active membership of patient advocates on the committee, has encouraged patient involvement in webinars and always keep the patient perspective as the north star in the development of the requirements for recommendation 1 above and the impetus behind work supporting recommendations 5 and 6 that follow.

3. Medication Reconciliation Process Improvements

The committee provided input on surveys and a webinar series that created information which will eventually be stored in an online repository. Additionally, several of the requirements for recommendation 1 above are written specifically for improvements to workflow.

4. Team Approach

During the development of the requirements to support recommendation 1, the committee identified three main roles involved in medication reconciliation: health systems & organizations, prescribers & clinicians, and patients & home health. The existence and the functioning of the committee itself supports this recommendation.

5. Implementation and Adoption of CancelRx

The committee assisted with surveys to assess the overall CancelRx adoption within the state. CancelRx was featured in a webinar discussing deprescribing. Continuing education credits (CME) were available for medical and pharmacist professionals.

6. Deprescribing

The committee supported the creation of content for a CME webinar on deprescribing.

7. Technology

The business and functional requirements developed for recommendation 1 include language about possible technical characteristics to support the best possible medication history, but technical requirements are not being addressed by the committee.

8. SUPPORT Act Funding and Planning/Design Process

Members of the committee supported the SUPPORT Act funding and planning/design process between February and September. There was no clear immediate impact of SUPPORT Act activity and that of the MRPC although the MRPC continues to track the progress of the Prescription Drug Management Program and considers that to be an information source for the BPMH. The committee receives periodic updates on advancements with the PDMP from the Director of the DCPs Drug Control Unit, an active member of the MRPC.

9. Aligned Policy

Operation of the BPMH, when it becomes an implemented resource for providers and prescribers, must of course align with HIPAA. Depending on the data sources it may also be impacted by alignment with 42 CFR Part 2. As the MRPC wraps up the requirements for a BPMH it will do additional work on policy alignment including aligned medication quality measures, incentives for medication management and reconciliation, assessing a policy mandate for the CancelRx standard, privacy and confidentiality of medication-related information, and introducing team-based medication reconciliation into healthcare provider workflows.

10. Planning/Design Process and Use of IAPD Funding

The MRPC intends to utilize planning funding that has been proposed for FFY 2021 in the current IAPD-U. More significantly the MRPC will determine how best to align the activity and projects of the committee with the Medicaid enterprise, to the fullest extent possible. Initiatives such as the BPMH, CancelRx, and best practices for deprescribing and polypharmacy have benefits for Medicaid populations and providers and the associated Medicaid funding can help to secure implementation and ongoing support.

11. Continuation of the MRP Work Group

The committee has had nine successful regular meetings as a continuation of the MRP work group. This recommendation is complete for the current year. The committee will continue to meet and develop initiatives through September 30, 2021 as currently funded and documented in the charter document.

Appendix C: BPMH Requirements

Following the guidance of the Medication Reconciliation Polypharmacy Work Group Final Report¹¹, the Medication Reconciliation Polypharmacy Committee (MRPC) has developed a robust set of recommendations to support the decision-making process around a potential solution for the development of a Best Possible Medication History (BPMH). The vision for the BPMH was presented at the June 25th, 2020 meeting of the committee:

Safe, quality and timely delivery of healthcare requires access to the 'Best-Possible Medication History". The BPMH should include all prescription and non-prescription medications, supplements and herbal products. The BPMH should be accurate, up-to-date and accessible to stakeholders (including but not limited to patients, caregivers and health care providers) at the point of decision making. Access to the BPMH will support collaborative care, reduce medication costs and errors and improve clinical outcomes.

This vision statement, along with the guiding principle that the patient is the north star, lead to the creation of twenty-two known issue statements and their associated business and functional requirements. The known issues were determined by considering the three perspectives involved in medication reconciliation: health systems & organizations, prescribers & clinicians, and patients & home health. These requirements have been delineated into five high-level categories, simply to easily organize the thought process and recommendations. In the case of the perspectives as well as the categories, there is overlap and any assumptions made in that regard have been explicitly outlined. This report explains the process and rationale behind the requirements and provides high-level recommendations for next steps toward a successful BPMH.

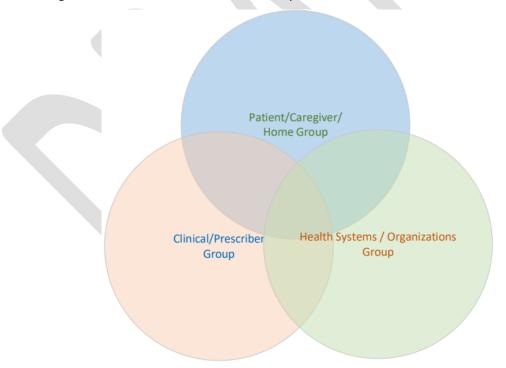


Figure 2: Breakout Session Groups/BPMH Perspectives

22

¹¹ https://portal.ct.gov/-/media/OHS/Health-IT-Advisory-Council/MRP/Final-Recoms-Med-Rec.pdf

Known Issues

During the June 25th, 2020 virtual meeting, the committee was divided into three separate virtual breakout rooms aligned with the perspectives as indicated in Figure 2 (Table 1 shows where each committee member participated). These groups discussed the drafted known issue statements, which were intended to fully represent the list of issues a patient or caregiver may encounter related to a lack of a best possible medication list.

Health Systems & Organizations	Clinicians & Prescribers	Patients & Home Health
Anne VanHaaren	Nitu Kashyap (facilitator)	Sean Jeffery (facilitator)
Elizabeth Taylor	Alejandro Gonzalez- Restrepo	Diana Mager
Jameson Reuter	Margherita Giuliano	Nate Rickles
Jennifer Osowiecki	Marie Renauer	Pat Carroll
Rod Marriott	Stacy Ward-Charlerie	
	Leslev Bennett	

Table 1: Committee Members Present in Each Breakout Session

The committee curated a list of twenty-two known issues, which can be summarized by the overarching known issue statement (known issue #17):

Accurate medication information is imperative for all patients and is particularly challenging for vulnerable populations or others who receive care in multiple settings and/or have complex treatment regimens.

The known issues can be loosely organized into five categories: General; Patient-Centric; Provider/Caregiver-Centric; Technical; Policy and Workflow Related. The final category includes known issues that would require modifications to existing policy and workflows, and is unlikely to be solved by any system, however robust. The following sections describe the known issues at a high level, using summary language to keep the discussion concise and some known issues are combined into a single statement for simplicity¹².

General Known Issues

This subset of known issues has implications across all settings:

- There is no single accurate source of medications available to all those involved in a patient's care issue 1
- Medication information changes frequently issue 3
- There is a lack of clarity on which medications or supplements to include in a medication listissue 4,8,16

Patient-Centric Known Issues

This subset of known issues most significantly impacts the individual receiving care:

- Patients may not trust having their information shared^{issue 12}
- Patients may not know when or how their medication was changed issue 14
- Accurate medication information is imperative to all patients issue 17

Provider-Centric Known Issues

This subset of known issues most significantly impacts the individuals involved in a patient's care:

- Patients and those involved in their care may not understand why a medication has been ordered issue 2,7
- Caregivers outside of an EHR may not have access to view real-time medication history issue 10

¹² Full List of Numbered Known Issue Statements

• Caregivers outside of an EHR may not have ability to contribute to a medication list issue 11

Technical Known Issues

This subset of known issues is most directly related to the limits present in existing systems, necessary roles and responsibilities for a system, or access to various types of relevant data:

- Medication histories may be presented differently in different systems issue 5, 15
- Changes to medication history do not occur in real-time issue 9
- EHRs lack access to real-time pharmacy data necessary for medication reconciliation issue 20
- Medication reconciliation requires defined roles and responsibilities issue 21

Policy and Workflow Related Known Issues

This subset of known issues is most directly related to the limits in existing policies, education, consent, or workflow specific issues that impact medication reconciliation:

- Accurate and timely medication reconciliation requires provider and caregiver educationissue 6
- Users accessing a medication history may not have proper consent to view/edit information^{issue 13}
- Consistently accepted workflows for medication reconciliation do not exist across all settings issue 18
- Automatic refill programs are not in sync with medication history changes issue 19
- Successful adherence to medication reconciliation processes is not measured/enforced^{issue 22}

Business and Functional Requirements

From the full, comprehensive list of known issues, the committee drafted associated business and functional requirements, which are detailed below. Much like the known issues, the business and functional requirements have undergone much discussion and refinement. To best assess how to address the known issue statements, the following guidance was used for the business requirements:

Business requirements define the 'why' for the BPMH:

- How does the BPMH relate to the objectives, vision, and goals of the MRPC?
- These requirements provide the scope of the needs to be addressed, and
- These requirements are high-level and can be broadly defined.

And, similarly, for the functional requirements – which define the 'what' for the BPMH:

- What are the specific behaviors of the BPMH (actions, processes, data, interactions, users)?
- What are the specific steps the MRPC and its support will take to achieve the business requirements?
- These requirements are specific and well defined.

Keeping this in mind, the committee, with support from the Admin team, defined a preliminary set of business and functional requirements, to support each of the known issues. This list is long, and while some redundancies occur in the list, it is meant to be exhaustive: all business and related functional requirements are explicitly defined. The figures below are designed to help the reader understand the delineation between the known issues, business requirements, and associated functional requirements.

Taking Figure 2 as an example, the first business requirement for known issue 1 is:

(1.1) A single list of all current medications for a patient shall be available 13

To support this business requirement, the MRPC recommends the following three functional requirements:

 $^{^{13}}$ Note that the numbering (1.1), (1.1.1), (1.1.2), etc. is consistent with the table containing the business and functional requirements, which can be found in Appendix B.

- (1.1.1) The medication list may be static with periodic updates or it can be created when requested from component information systems,
- (1.1.2) Individuals with records on the medication list shall be uniquely identified, and
- (1.1.3) Medication history shall be available to all physicians, providers, health systems, home health and pharmacists involved in a patient's care in accordance with governance rules, as well as to patients and caregivers.

It's important to note that the second two functional requirements *also* serve to support the second business requirement which is

(1.2) The medication list shall be accessible through a query and meet compliance requirements.

The remainder of this section shows these relationships for the full list of business and functional requirements associated with the known issues, as described above. Following the full list is a description of proposed next steps to best leverage this large body of work.

General Known Issues: Requirements

Business Requirements

These business and functional requirements address the known issues most accurately described as having implications across all settings. The first known issue in this category:

• There is no single accurate source of medications available to all those involved in a patient's care issue 1

Has the associated business and functional requirements shown in Figure 3.

The medication list may be static with periodic updates or it can be created A single list of all current medications when requested from component information systems for a patient shall be available Individuals with records on the medication list shall be uniquely identified The medication list shall be accessible Medication history shall be available to all physicians, providers, health through a query and meet compliance systems, home health and pharmacists involved in patient's care in accordance requirements with governance rules, as well as to patients and caregivers The medication list shall be in a Individuals accessing the medication list shall be validated using identity standardized electronic format management (MPI or Provider Directory) The medication list shall adhere to a The medication list shall accept updates from Health Information Technology standardized vocabulary or (HIT) systems nomenclature The medication list shall interact with HIT systems The medication list shall be compiled Medication history disambiguation shall be facilitated through software and from multiple sources human readable interfaces (deduplication, modifications) Sources shall be vetted for accuracy of the information submitted

Functional Requirements

Figure 3: Known Issue 1 Business and Functional Requirements

The second known issue in this category:

Medication information changes frequently issue 3

Has the associated business and functional requirements shown in Figure 4.

Business Requirements

Functional Requirements

The end user shall have access to the most current information at the time they are accessing the system

There shall be a longitudinal view of medications and changes

The system shall be capable of getting certain information and processing within a set amount of time (or near real-time) and send data out in near real-time from all potential locations of care

The system shall be capable of flagging cancelled medications and changes to dose, route and instructions as well as new additions, therapy changes, changes in where prescriptions are filled and changes in order status (ordered, dispensed)

The system shall be capable of tracking changes

The system shall reconcile ledger entries from all sources to associate a single related medication change event with the associated individual

Figure 4: Known Issue 3 Business and Functional Requirements

The third known issue in this category:

• There is a lack of clarity on which medications or supplements to include in a medication list issue 4,8,16

Has the associated business and functional requirements shown in Figure 5.

Business Requirements

Functional Requirements

There shall be education material promoted and provided to individuals about Medication History, including clarity of what is/is not included

Non-prescription substances being taken or used by a patient (OTC, supplements, herbals, etc.) contribute to a medication history and this information should be sought and added to the medication history

Users shall have access to the full array of prescription & non-prescription drugs through the BPMH

In the first phase of implementation, Medication History shall include prescribed medications (other OTC medications, vitamins, dietary supplements, etc. will be included in future updates)

Information about Medication History shall be available in understandable language and in the more common non-English languages spoken in Connecticut

Telephone interpretation services shall be available for non-English speaking individuals not covered by the non-English language documentation available

The BPMH shall have the capability to add, store and identify/tag OTC and supplements as a part of the medication list

The medication history shall tag information about other substances to qualify the level of confidence in the accuracy of the information (e.g. confirmed by pt, occasional use, anecdotal 2nd party report, other qualifying parameters)

When available, a standardized nomenclature such as NDC codes, RxNorm or other nomenclature shall be used

OTC medications, supplements and herbals entered by patients and caregivers shall be appropriately tagged to indicate the data entry source

The BPMH shall have the capability to add, store and identify and tag OTC and supplements as a part of the medication list

Figure 5: Known Issue 4, 8, 16 Business and Functional Requirements

Patient-Centric Known Issues: Requirements

These business and functional requirements address the known issues which most significantly impact the individual receiving care. The first known issue in this category:

Patients may not trust having their information shared issue 12

Has the associated business and functional requirements shown in Figure 6.

Patients (and/or their proxy, conservator, legal guardian, etc.) shall be informed of health information sharing intentions and possibilities, including medication information, and meaningful, informed consent shall be obtained from patients when appropriate

Consent rules/regulations shall be adhered to for all endusers/systems of the BPMH

The medication history solution shall leverage Connecticut's existing consent management system

Figure 6: Known Issue 12 Business and Functional Requirements

The second known issue in this category:

Patients may not know when or how their medication was changed^{issue 14}

Has the associated business and functional requirements shown in Figure 7.

Business Requirements

Functional Requirements

Medication history shall be presented in various levels and forms of healthcare literacy

The system shall be capable of alerting the patient to a medication change

The information shall be presented in common understandable terms

The information shall be presented in medical language

The information shall be presented in multiple languages

The information shall be presented visually

Alerts shall include details about who made changes and the details of the change

Patients shall be able to confirming they have seen the alert

Patients shall be able to incorporate changes into their own medication list via download or a mobile app

Figure 7: Known Issue 14 Business and Functional Requirements

The third known issue in this category:

Accurate medication information is imperative to all patients issue 17

Has the associated business and functional requirements shown in Figure 8.

Business Requirements

Functional Requirements

Design of the Medication History shall consider complexities associated with a full spectrum of patient circumstances, including a multiplicity of care settings, complex treatment regimens, and the impacts of social determinants of health that can impact access or adherence to medication (e.g., poverty; homelessness)

The most current Medication History must be available to patients. In addition to digital access, the BPMH should be printable in an easy-to-read and logical format so that even if the patient cannot read/understand, other people or caregivers should be able to easily understand all information about current medications

For patients receiving care in multiple settings the Medication History shall reflect those settings and the associated providers, for purposes of medication reconciliation

Complex medication regimens shall be accurately represented and understood (such as different dosages at different times of day, tapering dosages, alternate day dosages, altered dosages for renal or other problems, chemotherapy dosages that may be weight or exposure based)

Figure 8: Known Issue 17 Business and Functional Requirements

Provider-Centric Known Issues: Requirements

These business and functional requirements address the known issues which most significantly impact the individuals involved in a patient's care. The first known issue in this category:

Patients and those involved in their care may not understand why a medication has been ordered issue 2,7

Has the associated business and functional requirements shown in Figure 9.

Business Requirements

Functional Requirements

The medication list shall accommodate the inclusion of medication instructions There shall be a clear description of The reason the medication was ordered shall be available in why a medication has been ordered a language the patient/caregiver can understand for an individual (by providers and clinicians) Medication history shall include a field or fields to describe reason medication was ordered Pharmacists and dispensers shall have Each medication on the list shall have an accompanying information as to why a medication is diagnostic code to explain why the medication was ordered being ordered when permissible by Accurate medical technical language shall be used for law or patient consent pharmacist and provider users

Figure 9: Known Issue 2 and 7 Business and Functional Requirements

The second known issue in this category:

Caregivers outside of an EHR may not have access to view real-time medication history issue 10

Has the associated business and functional requirements shown in Figure 10.

Business Requirements

Functional Requirements

Medication histories shall be available for patient engagements in all care settings, including home health care

Medication histories shall be available through a variety of mechanisms to support workflows in different care settings, including via a website/portal, mobile app or appropriately connected HIT source

Intermediate records prepared to support care settings and workflows shall include the full Medication History (this includes paper records as well as electronic records)

Changes to or discrepancies in the medication history, verified in a care encounter, must be able to be submitted to the system for notation and resolution

Figure 10: Known Issue 10 Business and Functional Requirements

The third known issue in this category:

• Caregivers outside of an EHR may not have ability to contribute to a medication list issue 11

Has the associated business and functional requirements shown in Figure 11.

Business Requirement

Functional Requirement

A variety of mechanisms to update a medication list shall be supported, including a minimum acceptable mechanism that can be used in the most limited circumstances

The medication list shall accept input through electronic and non-electronic mechanisms, including telephone reporting and mailed notifications

Figure 11: Known Issue 11 Business and Functional Requirements

Technical Known Issues: Requirements

These business and functional requirements address the known issues most directly related to the limits present in existing systems, necessary roles and responsibilities for a system, or access to various types of relevant data. The first known issue in this category:

Medication histories may be presented differently in different systems^{issue 5, 15}

Has the associated business and functional requirements shown in Figure 12.

Business Requirements

Functional Requirements

Medication history shall leverage available technology in all steps of the data stream

BPMH data shall be made available through standard APIs to mobile technology vendors

The system shall have the capability to dynamically query source systems to present the most up to date information

Medication history information shall be available in a vendor agnostic format, using standards to support subsequent parsing for vendor system compatibility Prescribing information and reason medications were ordered/modified/canceled shall be captured from prescribers using EHR technology in all settings (incl. modifications for dosage/pause/etc in hospital)

Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy)

The pharmacy shall update medication history if a prescription is returned to stock

The prescriber shall update medication history with cancelled prescription information, and the pharmacy shall provide acknowledgement of cancellation

Data shall be made available utilizing ONC standards for API development for consumer apps (such as FHIR 4.0 and USCDI – updated as required)

User centered design shall be incorporated into the development of the GUI

Upon accessing a medication history from an EHR or mobile app, the system shall present a dynamically updated BPMH in near real-time

Information from vendor systems shall follow an agnostic format used by the Medication history solution (with standards to support parsing as needed)

Medication history information shall be available through a portal

Medication history individual reports shall be available to a variety of output mechanisms (such as fax or e-mail) which can be directed by an authorized user with established consent to share the information with a recipient

Figure 12: Known Issue 5 and 15 Business and Functional Requirements

The second known issue in this category:

Changes to medication history do not occur in real-time^{issue 9}

Has the associated business and functional requirements shown in Figure 13.

Business Requirements

Functional Requirements

Additions to a med list from one HIT system shall be available in an electronic format to all users and the system shall process input from all HIT systems, and push to all systems – in as real-time as possible (The BPMH is the source of truth)

Changes to a medication (dosage, reason for changing dose, therapy, cancellation, cancellation reason, route, instructions) shall be available in the source system and other systems accessed by end users in near real-time and should have date/time stamp

If additions and changes cannot be reflected in near realtime, and indicator in the user's system shall reflect that a change has occurred or will occur

A change log and process shall be in place for the medication list

Clear identification of the source systems for changes updates or delete shall exist in an audit log or similar functionality

Standardized nomenclature for changes shall be implemented

A (centralized?) system that monitors changes shall be implemented and available to end-systems to alert them to the changes

Figure 13: Known Issue 9 Business and Functional Requirements

The third known issue in this category:

EHRs lack access to real-time pharmacy data necessary for medication reconciliation issue 20

Has the associated business and functional requirements shown in Figure 14.

Business Requirement

Functional Requirements

The Medication History shall accommodate medication claims data

Consideration of whether the medication history shall be populated with APCD medication claims data on a regular basis will be made with design and accuracy as decision points

The Medication History shall support a query to pharmacy benefits information for an individual

Figure 14: Known Issue 20 Business and Functional Requirements

The fourth known issue in this category:

Medication reconciliation requires defined roles and responsibilities issue 21

Has the associated business and functional requirements shown in Figure 15.

Business Requirement

Functional Requirements

The system shall permit various roles to carry out various functions of medication reconciliation The system shall permit reconciliation guidance/governance to be specific to institutions, care settings, and caregiver types

The system shall be configurable based on state, organization, and/or legislative guidance

The system shall have roles based on access and control

Figure 15: Known Issue 21 Business and Functional Requirements

Policy and Workflow Related Known Issues: Requirements

These business and functional requirements address the known issues most directly related to the limits in existing policies, education, consent, or workflow specific issues that impact medication reconciliation. The first known issue in this category:

Accurate and timely medication reconciliation requires provider and caregiver education issue 6

Has the associated business and functional requirements shown in Figure 16.

Business Requirements

Functional Requirements

Medication reconciliation shall be a recognized quality aspect of every patient encounter

The system shall have the capability to assign role-based access and to adapt to more than one workflow for medication reconciliation as needed for organizational structure and requirements

The system shall have the capability to design a user-training module for various stakeholders

Education with CME shall be available to providers and caregivers on medication reconciliation processes and procedures

A maturity model for medication reconciliation shall be developed, with an established minimum expectation for reconciliation effort and results

Medication discrepancies that could have harmful outcomes if not addressed should be reconciled during the patient encounter (Patient safety shall be the primary consideration)

The system shall have the capability to allow medical assistants or others involved in a patients care to begin the mark-up process of a medication reconciliation, with the option of a separate user given permissions to finalize the process

Figure 16: Known Issue 6 Business and Functional Requirements

The second known issue in this category:

Users accessing a medication history may not have proper consent to view/edit information issue 13

Has the associated business and functional requirements shown in Figure 17.

Business Requirement

Functional Requirements

Consent rules/regulations shall be adhered to for all end-users/systems of the BPMH

The Medication History solution shall implement criteria for who can view medication history information and for who can make changes to the information based on industry standards/current regulations

They system shall account for patient consent as well as guardian, POA, conservator consent to share medication information

The system shall verify that the end-user has access and provide necessary audit logs

If more than one consent policy is involved in the domains involved in a BPMH use case (viewing a record; changing a record; etc.) the more restrictive policy shall prevail

Figure 17: Known Issue 13 Business and Functional Requirements

The third known issue in this category:

• Consistently accepted workflows for medication reconciliation do not exist across all settings issue 18

Has the associated business and functional requirements shown in Figure 18.

Business Requirement

Functional Requirements

Workflow templates supporting medication reconciliation shall be available in a variety of formats that support healthcare setting variations and encounter modalities

Medication reconciliation templates shall be available in an electronic format that supports interactive editing

The system shall have the ability to receive reconciliation templates with changes being integrated into the medication list

Reconciliation templates shall also be available in a paper format for use by home health or family caregivers

Figure 18: Known Issue 18 Business and Functional Requirements

The fourth known issue in this category:

• Automatic refill programs are not in sync with medication history changes issue 19

Has the associated business and functional requirements shown in Figure 19.

Business Requirement

Functional Requirements

Automatic refill programs at pharmacies shall verify the status of a prescription with the Medication History before refilling When a prescriber decides to discontinue a medication there shall be an update made to the medication history

The system shall be capable of receiving notifications for cancellations and sending those notifications to the pharmacy, in a format the pharmacy can receive and process

Figure 19: Known Issue 19 Business and Functional Requirements

The fifth known issue in this category:

• Successful adherence to medication reconciliation processes is not measured/enforced^{issue 22}

Has the associated business and functional requirements shown in Figure 20.

Business Requirement

Functional Requirements

The Medication History solution shall establish metrics to track the successful usage of the system

Metrics for the Medication History solution shall measure utilization, system performance, activities by type, and outcomes

Reports for the Medication History solution shall provide utilization histories by time period at the individual provider or caregiver level and at the practice or higher ordered entity level

An individual whose Medication History is in the system shall have access to a report of the individual's Medication History detail and a record of changes made to the individual history over time (including what changes were made, by whom, and when)

Figure 20: Known Issue 22 Business and Functional Requirements

Next Steps and Closing Thoughts

The committee recommends using the known issues, and business and functional requirements described in this report to inform the decision-making process around the possible procurement of a best possible medication history solution for the state of Connecticut. While these requirements are meant to be complete, the list does not include any recommended technical requirements, as those are out of scope for the MRPC.

Full List of Numbered Known Issue Statements

- 1. There is no single accurate source and list of medications concurrently available to all the physicians, providers, health systems, pharmacists, and patients (and their designated caregivers), involved in a patient's care
- 2. Patients may not understand why they are taking a particular medication, and/or providers may not understand why a medication has been ordered for their patient
- 3. Medication information can and does change frequently, with changes from different sources and perspectives, making it difficult to establish and maintain a single source of medication history
- 4. Patients are unaware of what should be included in a medication list such as OTC medications, vitamins and dietary supplements, creams and ointments
- 5. The use of technology to improve medication history, including EHRs, smartphone apps, and repositories of information, presents new problems of interoperability and data currency (user adaptation to new technologies is a related additional problem)
- 6. Medication reconciliation is not a simple process and provider education is an ongoing problem systematic approaches to medication reconciliation can vary depending on the size/type of organization, staffing, and technology capabilities
- 7. Pharmacists/dispensers are not always aware of the condition being treated by a prescription
- 8. Information on OTC, supplements and herbals does not originate in provider or prescriber systems and is challenging to capture from patients and caregivers
- 9. Additions and changes to medication history do not occur in real-time from all sources
- 10. Home care nurses are not provided with accurate real-time medication histories prior to engaging with the patient, nor do the paper records they prepare include the full medication history
- 11. Family caregivers may not have access or mechanisms to contribute information to a medication list
- 12. Patients may not be adequately informed or trust having their information shared, including through technology

- 13. People accessing a medication history may not have proper consent to view the information or to make changes to the information
- 14. Patients may not know when their medication was changed or who might have made the changes
- 15. Medication histories may be presented differently in different systems or settings, creating challenges in interpreting or understanding the information
- 16. Despite front line responsibility for prescribing, coordinating, and reconciliation patients' medications, primary care providers often lack complete information on the full array of medications that a patient is using (e.g. Rx, OTC, supplements)
- 17. Accurate medication information is particularly challenging for vulnerable populations who receive care in multiple settings and have complex treatment regimens
- 18. There is a need for consistently accepted workflows that adapt medication reconciliation to various sizes and types of healthcare settings, encounters (in person, remote), and to the most effective time during different types of encounters
- 19. Automatic refill programs at pharmacies may not be in sync with medication history changes or provider decisions to discontinue a medication
- 20. Some EHRs lack access to real-time pharmacy benefits data and/or pharmacy claims data to be used in medication reconciliation
- 21. Institutions do not define the roles and responsibilities necessary to perform medication reconciliation
- 22. No metrics exist for the measurement of adherence to medication reconciliation processes

	Table of Known Issues, Business Requirements and Functional Requirements	
	ere is no single accurate source and list of medications concurrently available to all the pholers, health systems, pharmacists, and patients (and their designated caregivers), involve	-
1.1	A single list of current medications for a patient shall be available	business
1.1.1	The medication list may be static with periodic updates or it can be created when requested from component information systems	functional
1.1.2	Individuals with records on the medication list shall be uniquely identified	functional
1.1.3	Medication history shall be available to all physicians, providers, health systems, home health and pharmacists involved in patient's care in accordance with governance rules, as well as to patients and caregivers	functional
1.2	The medication list shall be accessible through a query and meet compliance requirements	business
1.2.1	See 1.1.2 (Individuals with records on the medication list shall be uniquely identified)	functional
1.2.2	See 1.1.3 (Medication history shall be available to all physicians, providers, health systems, home health and pharmacists involved in patient's care in accordance with governance rules, as well as to patients and caregivers)	functional
1.2.3	Individuals accessing the medication list shall be validated using identity management (MPI or Provider Directory)	functional
1.3	The medication list shall be in a standardized electronic format	business
1.3.1	The medication list shall accept updates from HIT systems	functional
1.3.2	The medication list shall interact with HIT systems	functional
1.4	The medication list shall adhere to a standardized vocabulary or nomenclature	business
1.5	The medication list shall be compiled from multiple sources	business
1.5.1	Medication history disambiguation shall be facilitated through software and human readable interfaces (deduplication, modifications)	functional
1.5.2	Sources shall be vetted for accuracy of the information submitted	functional
	ients may not understand why they are taking a particular medication, and/or providers m stand why a medication has been ordered for their patient.	nay not
	There shall be a clear description of why a medication has been ordered for an	
2.1	individual (by providers and clinicians)	business
2.1.1	Medication history shall include a field or fields to describe reason medication was ordered	functional
2.1.2	Each medication on the list shall have an accompanying diagnostic code to explain why the medication was ordered	functional
2.1.3	The medication list shall accommodate the inclusion of medication instructions	functional
2.1.4	The reason the medication was ordered shall be available in a language the patient/caregiver can understand	functional
2.1.5	Accurate medical technical language shall be used for pharmacist and provider users	functional
	dication information can and does change frequently, with changes from different sources ectives, making it difficult to establish and maintain a single source of medication history.	
3.1	The end user shall have access to the most current information at the time they are accessing the system	business
3.1.1	The system shall be capable of getting certain information and processing within a set amount of time (or near real-time) and send data out in near real-time from all potential locations of care	functional
3.1.2	The system shall be capable of flagging cancelled medications and changes to dose, route and instructions as well as new additions, therapy changes, changes in where prescriptions are filled and changes in order status (ordered dispensed)	functional

2

3

3.2

3.2.1

(ordered, dispensed)

The system shall be capable of tracking changes

There shall be a longitudinal view of medications and changes

business

functional

	3.2.2	The system shall reconcile ledger entries from all sources to associate a single related medication change event with the associated individual	functional
		ients are unaware of what should be included in a medication list such as OTC medicatio ietary supplements, creams and ointments.	ns, vitamins
	4.1	There shall be education material promoted and provided to individuals about Medication History, including clarity of what is included in Medication History and what is not included	business
4	4.1.1	In the first phase of implementation, Medication History shall include prescribed medications (other OTC medications, vitamins, dietary supplements, etc. will be included in future updates)	functional
	4.1.2	Information about Medication History shall be available in understandable language and in the more common non-English languages spoken in Connecticut	functional
	4.1.3	Telephone interpretation services shall be available for non-English speaking individuals not covered by the non-English language documentation available	functional
	inform	e use of technology to improve medication history, including EHRs, smartphone apps, and nation, presents new problems of interoperability and data currency (user adaptation to ne plated additional problem).	•
	5.1	Medication history shall leverage available technology in all steps of the data stream to create and maintain the information in medication history	business
	5.1 5.1.1		business functional
		Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in	
5	5.1.1	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data	functional
5	5.1.1	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy)	functional functional
5	5.1.1 5.1.2 5.1.3	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy) The pharmacy shall update medication history if a prescription is not picked up and is returned to stock The prescriber shall update medication history with cancelled prescription information, and the pharmacy shall	functional functional functional
5	5.1.1 5.1.2 5.1.3 5.1.4	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy) The pharmacy shall update medication history if a prescription is not picked up and is returned to stock The prescriber shall update medication history with cancelled prescription information, and the pharmacy shall provide acknowledgement of cancellation BPMH data shall be made available through standard APIs to mobile technology vendors Data shall be made available utilizing ONC standards for API development for consumer apps (such as FHIR 4.0 and USCDI – updated as required)	functional functional functional
5	5.1.1 5.1.2 5.1.3 5.1.4 5.2	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy) The pharmacy shall update medication history if a prescription is not picked up and is returned to stock The prescriber shall update medication history with cancelled prescription information, and the pharmacy shall provide acknowledgement of cancellation BPMH data shall be made available through standard APIs to mobile technology vendors Data shall be made available utilizing ONC standards for API development for consumer apps (such as FHIR 4.0 and USCDI – updated as required) User centered design shall be incorporated into the development of the GUI where possible	functional functional functional functional business
5	5.1.1 5.1.2 5.1.3 5.1.4 5.2 5.2.1	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy) The pharmacy shall update medication history if a prescription is not picked up and is returned to stock The prescriber shall update medication history with cancelled prescription information, and the pharmacy shall provide acknowledgement of cancellation BPMH data shall be made available through standard APIs to mobile technology vendors Data shall be made available utilizing ONC standards for API development for consumer apps (such as FHIR 4.0 and USCDI – updated as required) User centered design shall be incorporated into the development of the GUI where possible The system shall have the capability to dynamically query source systems to obtain and present the most up to date information	functional functional functional functional business functional
5	5.1.1 5.1.2 5.1.3 5.1.4 5.2 5.2.1 5.2.2	Stream to create and maintain the information in medication history Prescribing information and reason medications were ordered/modified/ canceled shall be captured from prescribers using EHR technology (regardless of setting – including modifications for dosage/pause/etc. in hospital) Prescription information including date filled and date dispensed shall be captured from pharmacy systems data (including delivery by pharmacy) The pharmacy shall update medication history if a prescription is not picked up and is returned to stock The prescriber shall update medication history with cancelled prescription information, and the pharmacy shall provide acknowledgement of cancellation BPMH data shall be made available through standard APIs to mobile technology vendors Data shall be made available utilizing ONC standards for API development for consumer apps (such as FHIR 4.0 and USCDI – updated as required) User centered design shall be incorporated into the development of the GUI where possible The system shall have the capability to dynamically query source systems to	functional functional functional functional business functional functional

Medication reconciliation is not a simple process and provider education is an ongoing problem - systematic approaches to medication reconciliation can vary depending on the size/type of organization, staffing, and technology capabilities

6.1	Medication reconciliation shall be a recognized quality aspect of every patient encounter	business
6.1.1	Education with CME shall be available to providers and caregivers on medication reconciliation processes and procedures	functional
6.1.2	A maturity model for medication reconciliation shall be developed, with an established minimum expectation for reconciliation effort and results	functional
6.1.3	Medication discrepancies that could have harmful outcomes if not addressed should be reconciled during the patient encounter (<i>Patient safety shall be the primary consideration</i>)	functional

		The system shall have the capability to assign role-based access and to adapt	
	6.2	to more than one workflow for medication reconciliation as needed for	business
	6.2.1	organizational structure and requirements The system shall have the capability to allow medical assistants or others involved in a patient care to begin the mark-up process of a medication reconciliation, with the option of a separate user given permissions to finalize the process	functional
	6.3	The system shall have the capability to design a user-training module for various stakeholders	business
-			
	Pha	rmacists/dispensers are not always aware of the condition being treated by a prescription	า
	7.1	Pharmacists and dispensers shall have information as to why a medication is being ordered when permissible by law or patient consent	business
7	7.1.1	See 2.1.5 (Accurate medical technical language shall be used for pharmacist and provider users)	functional
	7.1.2	See 2.1.1 (Medication history shall include a field or fields to describe the reason(s) medication was ordered)	functional
	7.1.3	See 2.1.2 (Each medication on the list shall have a field for an accompanying standardized diagnostic code or meta-data to explain why the medication was ordered)	functional
		ormation on OTC, supplements, and herbals does not originate in provider or prescriber s nging to capture from patients and caregivers	systems and is
	8.1	Other non-prescription substances being taken or used by a patient (OTC, supplements, herbals, etc.) contribute to a medication history and this information should be sought and added to the history	business
8	8.1.1	The BPMH shall have the capability to add, store and identify/tag OTC and supplements as a part of the medication list	functional
	8.1.2	The medication history shall tag information about other substances to qualify the level of confidence in the accuracy of the information (e.g. confirmed by pt., occasional use, anecdotal 2nd party report, other qualifying parameters)	functional
	8.1.3	When available, a standardized nomenclature such as NDC codes, RxNorm or other nomenclature shall be used	functional
	8.1.4	OTC medications, supplements and herbals entered by patients and caregivers shall be appropriately tagged to indicate the data entry source	functional
	Ada	litions and changes to medication history do not occur in real-time from all sources	
9	9.1	Additions to a med list from one HIT system shall be available in an electronic format to all users and the system shall process input from all HIT systems, and push to all systems – in as real-time as possible (<i>The BPMH is the source of truth</i>)	business
	9.1.1	If additions and changes cannot be reflected in near real-time, and indicator in the user's system shall reflect that a change has occurred or will occur	functional
	9.2	Changes to a medication (dosage, reason for changing dose, therapy, cancellation, cancellation reason, route, instructions) shall be available in the source system and other systems accessed by end users in near real-time and should have date/time stamp	business
	9.2.1	A change log and process shall be in place for the medication list	functional
	9.2.2	Clear identification of the source systems for changes updates or delete shall exist in an audit log or similar functionality	functional
	9.2.3	Standardized nomenclature for changes shall be implemented	functional
	9.4.5	A (centralized?) system that monitors changes shall be implemented and available to end-systems to alert them to the changes	functional

	Hon	ne care nurses are not provided with accurate real-time medication histories prior to enga	aging with the		
		t, nor do the paper records they prepare include the full medication history.	ignig with the		
	•	Medication histories shall be available in advance of patient engagements in all			
	10.1	care settings, including home health care	business		
10	10.1.1	Medication histories shall be available through a variety of mechanisms to support workflows in different care settings, including via a website/portal, mobile app or appropriately connected HIT source	functional		
	10.1.2	Intermediate records prepared to support care settings and workflows shall include the full Medication History (this includes paper records as well as electronic records)	functional		
	10.1.3	Changes to or discrepancies in the medication history, verified in a care encounter, must be able to be submitted to the system for notation and resolution	functional		
	Fan	nily caregivers may not have access or mechanisms to contribute information to a medica	ation list.		
11	11.1	A variety of mechanisms to update a medication list shall be supported, including a minimum acceptable mechanism that can be used in the most limited circumstances	business		
	11.1.1	The medication list shall accept input through electronic and non-electronic mechanisms, including telephone reporting and mailed notifications	functional		
	Pati techno	ents may not be adequately informed or trust having their information shared, including to Blogy.	hrough		
12	12.1	Patients (and/or their proxy, conservator, legal guardian, etc.) shall be informed of health information sharing intentions and possibilities, including medication information, and meaningful, informed consent shall be obtained from patients when appropriate	business		
	12.1.1	Consent rules/regulations shall be adhered to for all end-users/systems of the BPMH	functional		
	12.1.2	The medication history solution shall leverage Connecticut's existing consent management system	functional		
	People accessing a Medication History may not have proper consent to view the information or to make changes to the information.				
	13.1	Consent rules/regulations shall be adhered to for all end-users/systems of the BPMH	business		
13	13.1.1	The Medication History solution shall implement criteria for who can view medication history information and for who can make changes to the information based on industry standards/current regulations	functional		
	13.1.2	They system shall account for patient consent as well as guardian, POA, conservator consent to share medication information	functional		
	13.1.3	The system shall verify that the end-user has access and provide necessary audit logs	functional		
	13.1.4	If more than one consent policy is involved in the domains involved in a BPMH use case (viewing a record; changing a record; etc.) the more restrictive policy shall prevail	functional		
<u> </u>					
	Pati	ents may not know when their medication was changed or who might have made the cha	anges.		
	14.1	Medication history shall be presented in various levels and forms of healthcare literacy	business		
	14.1.1	The information shall be presented in common understandable terms	functional		
14	14.1.2	The information shall be presented in medical language	functional		
	14.1.3	The information shall be presented in multiple languages	functional		
	14.1.4	The information shall be presented visually	functional		
	14.2	The system shall be capable of alerting the patient to a medication change	business		
	14.2.1	Alerts shall include details about who made changes and the details of the change	functional		
	14.2.2	Patients shall be able to confirm they have seen the alert	functional		

	1.4		, ,				
		dication histories may be presented differently in different systems or settings, creating ch	nallenges in				
	merpi	reting or understanding the information.					
	454	Medication History information shall be available in a vendor agnostic format,	husinasa				
	15.1	using standards to support subsequent parsing by vendor systems for compatibility with their systems	business				
4-		Medication History information submitted from vendor systems shall follow an agnostic format used by the					
15	15.1.1	Medication History solution, using standards to support parsing by the solution system	functional				
	15.1.2	Medication History information shall be available through a portal	functional				
	15.1.3	Medication History individual reports shall be available to a variety of output mechanisms (such as fax or e-mail) which can be directed by an authorized user with established consent to share the information with a recipient	functional				
	15.1.4	The Medication History solution shall support multiple languages as more commonly used in the state of Connecticut, and shall be aligned with Section 508, as revised	functional				
	Des	spite front line responsibility for prescribing, coordinating, and reconciling patients' medica	ntions, primary				
	care p	providers often lack complete information on the full array of medications that a patient is	using (e.g. Rx,				
	OTC,	supplements).					
16	16.1	Users shall have access to the full array of prescription & non-prescription drugs through the BPMH	business				
	16.1.1	The BPMH shall have the capability to add, store and identify and tag OTC and supplements as a part of the medication list	functional				
	Acc	urate medication information is particularly challenging for vulnerable populations who re	ceive care in				
	multip	le settings and have complex treatment regimens.					
	17.1	Design of the Medication History shall consider complexities associated with a full spectrum of patient circumstances, including a multiplicity of care settings, complex treatment regimens, and the impacts of social determinants of health that can impact access or adherence to medication (e.g., poverty; homelessness)	business				
17		The most current Medication History must be available to patients. In addition to digital access, the BPMH					
	17.1.1	should be printable in an easy-to-read and logical format so that even if the patient cannot read/understand, other people or caregivers should be able to easily understand all information about current medications	functional				
	17.1.2	For patients receiving care in multiple settings the Medication History shall reflect those settings and the associated providers, for purposes of medication reconciliation	functional				
	17.1.3	Complex medication regimens shall be accurately represented and understood (such as different dosages at different times of day, tapering dosages, alternate day dosages, altered dosages for renal or other problems, chemotherapy dosages that may be weight or exposure based)	functional				
		re is a need for consistently accepted workflows that adapt medication reconciliation to v					
	_	rpes of healthcare settings, encounters (in person, remote), and to the most effective time	during				
	aiπere	ent types of encounters.					
18	18.1	Workflow templates supporting medication reconciliation shall be available in a variety of formats that support healthcare setting variations and encounter modalities	business				
	18.1.1	Medication reconciliation templates shall be available in an electronic format that supports interactive editing	functional				
	18.1.2	The system shall have the ability to receive reconciliation templates with changes being integrated into the medication list	functional				
	18.1.3	Reconciliation templates shall also be available in a paper format for use by home health or family caregivers	functional				

Patients shall be able to incorporate changes into their own medication list via download or a mobile app

14.2.3

functional

		omatic refill programs at pharmacies may not be in sync with Medication History changes ons to discontinue a medication.	or provider	
19	19.1	Automatic refill programs at pharmacies shall verify the status of a prescription with the Medication History before refilling	business	
	19.1.1	When a prescriber decides to discontinue a medication there shall be an update made to the medication history	functional	
	19.1.2	The system shall be capable of receiving notifications for cancellations and sending those notifications to the pharmacy, in a format the pharmacy can receive and process	functional	
		ne EHRs lack access to real-time pharmacy benefits data and/or pharmacy claims data to ation reconciliation.	be used in	
20	20.1	The Medication History shall accommodate medication claims data	business	
20	20.1.1	Consideration of whether the medication history shall be populated with APCD medication claims data on a regular basis will be made with design and accuracy as decision points	functional	
	20.1.2	The Medication History shall support a query to pharmacy benefits information for an individual	functional	
	Institutions do not define the roles and responsibilities necessary to perform medication reconciliation			
	21.1	The system shall permit various roles to carry out various functions of medication reconciliation	business	
21	21.1.1	The system shall permit reconciliation guidance/governance to be specific to institutions, care settings, and caregiver types	functional	
	21.1.2	The system shall be configurable based on state, organization, and/or legislative guidance	functional	
	21.1.3	The system shall have roles based on access and control	functional	
	No	metrics exist for the measurement of adherence to medication reconciliation processes.		
22	22.1	The Medication History solution shall establish metrics to track the successful usage of the system	business	
	22.1.1	Metrics for the Medication History solution shall measure utilization, system performance, activities by type, and outcomes	functional	
	22.1.2	Reports for the Medication History solution shall provide utilization histories by time period at the individual provider or caregiver level and at the practice or higher ordered entity level	functional	
		An individual whose Medication History is in the system shall have access to a report of the individual's		

Appendix D: Deprescribing and CancelRx Update

Deprescribing & the Use of Cancel RX

Deprescribing is the planned and supervised process of dose reduction or stopping of medication that is no longer of benefit to the patient or might be causing harm. Deprescribing is an essential practice of good prescribing – reducing doses that are too high or discontinuing medications that are unnecessary. ¹⁴ CancelRx is an electronic tool that facilitates communication around deprescribing by allowing prescribers to send electronic cancellation messages to pharmacies when medications are discontinued. ¹⁵ CancelRx is used to remove active prescriptions from a patients medication regimen and communicate that to the pharmacy. Due to the intertwined nature of deprescribing and the CancelRx tool, and the overlap of work done in these areas by the Medication Reconciliation and Polypharmacy Committee (MRPC), the decision has been made to combine the work performed in these areas into a single update.

CancelRx

A CancelRx transaction communicates an electronic message to the pharmacy from a provider when the provider discontinues a medication in their EHR system. Providers have the ability to also enter a note in the message to indicate why the medication was discontinued. As of 2010 the *National Council for Prescription Drug Programs* (NCPDP)¹⁶ CancelRx tool was available, however adoption was quite limited. Updates to the Medicare Access and CHIP Reauthorization Act of 2015 (MACRA) introduced requirements that EHRs include capability to send electronic cancellation messaging to pharmacies. Additionally, prescribers and pharmacies are responsible for certifying their capacity to send and consume CancelRx messages. This has led to an increase in adoption of CancelRx. As of September 2020, approximately 78% of Connecticut pharmacies are CancelRx enabled.¹⁷ This statistic puts Connecticut at the forefront of CancelRx adoption, with higher adoption levels than the national average.

Anecdotal reports as well as implementation data reveal that while acceptance has increased, work remains to improve adoption of this technology. Additional outreach and education efforts targeted at informing providers about how the technology works, as well as encourage it's use through incentives and/or requirements could help push Connecticut closer to full adoption.

In the context of the Covid 19 public health crisis it becomes imperative that we consider ways in which deprescribing activities, increased adoption of CancelRx, and education efforts can move forward via telehealth and virtual opportunities. In support of the MRPC and enhanced adoption of Health Information Technology (HIT) in Connecticut, the team at UConn Health, led by Dr. Tom Agresta developed and presented a virtual continuing education event for providers and pharmacists. Additional virtual outreach and education activities should be considered to aid in the increased awareness of these tools, as well as further drive adoption of deprescribing standards within Connecticut.

Activities & Opportunities

¹⁴ https://deprescribing.org/what-is-deprescribing/

¹⁵ Pitts SI, Barasch N, Maslen AT, Thomas BA, Dorissaint LP, Decker KG, Kazi S, Yang Y, Chen AR. Understanding CancelRx: Results of End-to-End Functional Testing, Proactive Risk Assessment, and Pilot Implementation. Appl Clin Inform. 2019 Mar;10(2):336-347. doi: 10.1055/s-0039-1688698. Epub 2019 May 22. PMID: 31117135; PMCID: PMC6531208.

https://www.ncpdp.org

¹⁷ CancelRx and the Medication Management Puzzle: A Pharmacist's Perspective, **October 10, 2017,** Stacy Ward-Charlerie, PharmD, Pharmacist Data Manager, Surescripts

Survey

In early 2020 a survey was developed for providers in the State of Connecticut to assess their current methods of deprescribing, awareness of the CancelRx standard, and to identify areas of interest for education and outreach efforts. The survey also assessed prescriber and pharmacist satisfaction with the standard, if they were currently using it. In March 2020 the survey was shared through a list serve of prescribers in the State, using Qualtrics. Over 500 providers responded to the survey. Respondents spanned a variety of practice settings and provider types, including physicians, pharmacists, advance practice nurses, dentists, and physician assistants. The results determined that more pharmacists than prescribers were familiar with the CancelRx standard and participants demonstrated that there was an interest for educational opportunities on the topic and led to the development of the deprescribing continuing education webinar held in June 2020. The survey questions and a selection of results can be found at the end of this appendix.

Education & Outreach

Education and outreach activities around deprescribing and CancelRx are ongoing. Initial education to providers, pharmacists, and others occurred over the summer, via a panel style continuing education webinar. The event, *To Deprescribe or not to Deprescribe: The Role of Health IT in Polypharmacy* was held in June 2020. This event offered continuing education credits to physicians and pharmacists, and covered the impact of polypharmacy, challenges associated with deprescribing, the role of Health IT in medication management, the SCRIPT standard, CancelRx transaction data flow and best practices to implement and apply deprescribing in clinical practice. This event was recorded and can be shared widely, beyond initial attendees. Additionally, a frequently asked questions (FAQ) document was developed based on attendee input and is available to be shared broadly. Further opportunities for educating at the organization and pharmacy level are being considered. Additional provider focused events are in development, and the development of an opportunity to engage patients and providers to promote shared decision-making around deprescribing is currently under consideration.

Education and outreach materials that have been produced around deprescribing to date and the CancelRx standard are available in digital formats and have been distributed via email. These items are expected to be available via a web repository in Winter 2020. Preliminary resources, including peer reviewed literature, as well as documentation around standards, recommendations and best practices have also been collated and will be included in the repository. This work is ongoing.

Legislative & State Leadership

Opportunities for collaboration with State leadership and the Legislature have been identified in the encouragement and/or incentivizing of participation in the CancelRx standard. At this time formal assessment or activities have occurred in this area. Since currently available data indicate that over 78% of Connecticut pharmacies are already enabled, and federal programs already incentivize adoption, it may be most useful at this time to consider seeking support for assessing which pharmacies are not currently enabled and why and performing targeted outreach to those pharmacies not participating. Alternatively, as the majority of pharmacies are already enabled, an option is to seek legislative support and sponsorship of a bill to require those remaining pharmacies to turn on CancelRx capability.

Health Information Exchange

There are also prospects for a partnership with the State's Health Information Exchange (HIE) to further the work of the MRPC. An area that has been identified as ripe for collaboration is the development of a business case for the sustainability of CancelRx. Under the HIE's Use Case Factory model, CancelRx could be combined with effective deprescribing strategies and packaged as a medication reconciliation use case for the State's HIE. Further, upon the implementation of the HIE there may be an opportunity to utilize risk algorithms to identify population health strategies around deprescribing. This latter opportunity will be practical to explore once the HIE is operational.

Office of Health Strategy

An occasion may also exist for MRPC collaboration work with the Office of Health Strategy's (OHS) Primary Care Modernization and Cost Containment Offices. There may be opportunities for deprescribing recommendations and/or incentives to be incorporated into these ongoing projects at OHS. The goals of these offices align with deprescribing efforts to improve healthcare outcomes and reduce costs.

Department of Consumer Protection

Collaboration efforts with the Department of Consumer Protection (DCP) have helped to align the agency with the activities of the MRPC. The Director of the DCP's Drug Control Unit is an active member of the MRPC. Additionally, recent efforts, aided by the Substance Use-Disorder Prevention that Promotes Opioid Recovery and Treatment (Support Act) funds, have resulted in plans that will provide integration for EHRs with Connecticut's Prescription Drug Monitoring Program. This integration will occur at no cost for providers and health systems and will facilitate easier access to Connecticut's PDMP for providers. Ongoing close coordination with the DCP is planned to ensure harmonization between the agency and the MRPC.

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Status of CancelRx In Connecticut

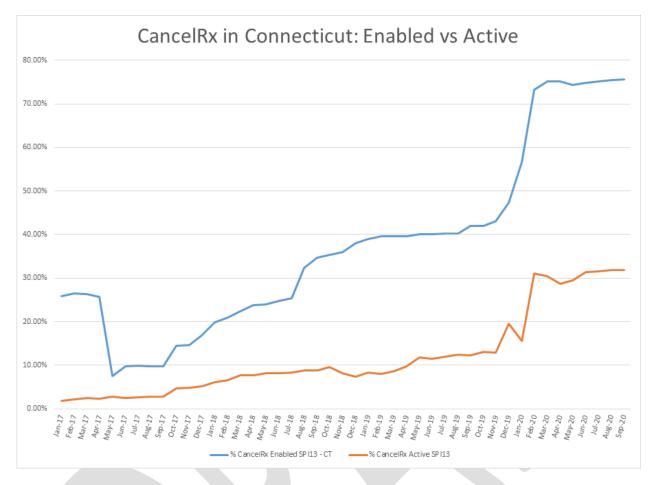


Figure 1 Represents the providers in Connecticut that are enabled to use CancelRx vs. those that are actively using CancelRx.

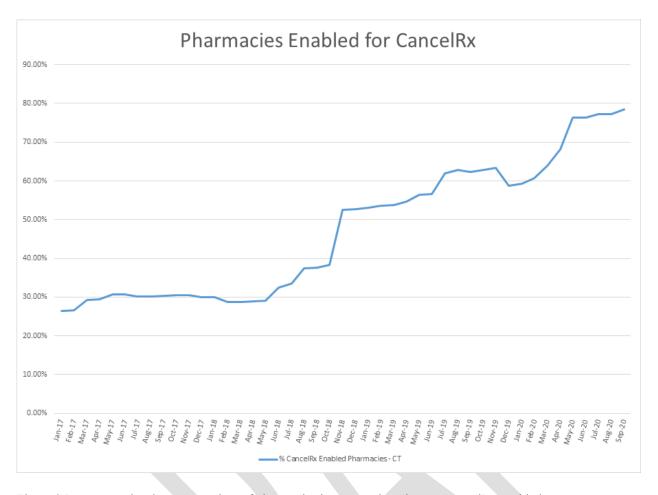


Figure 2 Represents the change over time of pharmacies in Connecticut that are CancelRx enabled.

Cancel RX Survey Questions

The CancelRx survey was designed and administered to gain an understanding of the current levels of familiarity of CancelRx among providers and to identify if training was desired. Additionally, the survey sought to understand which educational methods providers might be most interested in participating in. Below is an outline of the survey questions.

CancelRx Survey

- Q1 The purpose of this survey is twofold: 1) to assess the awareness of and knowledge about CancelRx, which is an electronic messaging standard for deprescribing medications from within an Electronic Health Record and 2) to determine the best method to create and deliver educational materials to facilitate the safe and effective use of this standard by healthcare stakeholders who are involved in prescription cancellations.
- Q2 Please select the option below that best describes your role in healthcare.
- Q3 Which setting(s) do you practice in? Select all that apply.
- Q4 Please state the name of the organization(s) where you are employed.
- Q36 If you are employed in a state other than Connecticut, please indicate below.
- Q5 Which of the following best describes your role (select all that apply):
- Q6 Which of the following best describes your role (select all that apply):
- Q7 How does the pharmacy receive the cancellation? (select all that apply)
- Q8 Prior to this survey, were you familiar with CancelRx or its purpose?
- Q9 Does your organization use CancelRx?
- Q10 Does your pharmacy have CancelRx enabled?
- Q11 How do you notify the appropriate parties a medication has been discontinued? (choose all that apply)
- Q12 Do you know which providers send cancellation transactions using CancelRx?
- Q13 (Click and Drag) Rank the method you use from most to least, where a ranking of 1 represents the most utilized.
- Q14 (Click and Drag) Rank the methods in which the pharmacy receives prescription cancellations from most (1) to least (4)
- Q15 For the following questions, assume your electronic medical record allows you to click on a button labeled "cancel," and once you do, an electronic message is sent to the pharmacy associated with that medication to notify them that the prescriber has discontinued the medication (CancelRx).
- Q16 For the following questions, assume your pharmacy information system allows you to receive an electronic message from the prescriber for a medication cancellation and permits you to respond electronically to confirm the cancellation (CancelRx).
- Q17 CancelRx has eased my everyday workflow
- Q18 In what ways has CancelRx positively impacted your workflow?
- Q19 In what ways has CancelRx negatively impacted your workflow?
- Q20 CancelRx has reduced medication discrepancies.
- Q21 CancelRx has supported patient medication adherence.
- Q22 When I use the CancelRx function, I feel confident that the message was successfully transmitted.
- Q23 I believe CancelRx would ease my everyday workflow
- Q24 I believe CancelRx would reduce medication discrepancies.
- Q25 I believe CancelRx would support patient medication adherence.
- Q26 Do you know who to reach out to if you have any questions about CancelRx?
- Q27 Who do you contact to get help with your electronic prescription system?
- Q28 How interested would you be in receiving a short training session regarding the proper utilization of CancelRx?
- Q29 Rate the level of interest for each of the instructional methods.
- O30 Please provide any comments.
- Q35 Optional.

Provide your email below.

A Selection of CancelRx Survey Results

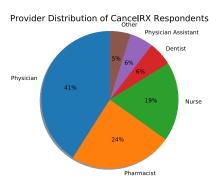


Figure 3 Represents the distribution of provider types responding to the CancelRx survey.

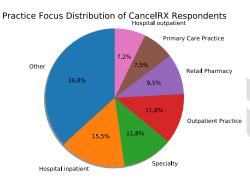


Figure 4 Respresents the distribution of practice settings of respondents to the CancelRx survey.

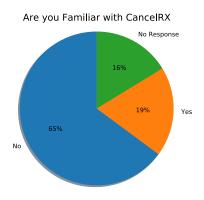


Figure 5 Represents the percentages of respondents with prior knowledge of CancelRx.

Interest in CancelRX Training Among Those Unfamiliar

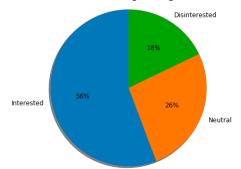


Figure 6 Represents the percentage of respondents interested in CancelRx educational opportunities.

