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November 5, 2025

Pharmacy Benefit Manager Data Collection Listening Session



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Housekeeping

Recording Notice

 This session will be recorded for internal use and documentation purposes.

Voluntary Participation

 Your participation is completely voluntary. You may choose to engage at your comfort level.

Use of Shared Information

 Insights from this session may be used to inform future decisions, but individual responses will remain confidential.

How to participate

- You can participate by:
 - · Using the chat feature
 - · Unmuting to speak
 - · Responding to polls or prompts
 - Providing written feedback after the session.

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ENGAGE EVERYONE



BE HEARD AND SEEN



ACKNOWLEDGE SPEAKER



MAXIMIZE MICROPHONES



MINIMIZE NOISE



MAXIMIZE VISUAL DISPLAYS

These guidelines are intended to improve the meeting experience for virtual participants, as well as people with hearing loss, visual impairment, and those for whom English is an additional language. Developed by the Access AIR and AIR CREW Employee Resource Groups With Support From the AIR Diversity and inclusion Office.

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Agenda

Background

- · Review of associated legislation and requirements
- · Overview of upcoming engagement opportunities

Material Review

- · Overview of draft materials
- · Content review within submission materials and resources
- · Space for discussion and feedback

Wrap-up

- Next steps
- · Opportunities for feedback

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Objectives

- This session is designed to gather feedback, including insights, concerns, and suggestions to shape final implementation
- Your input helps ensure the reporting templates and procedures are practical, clear, and aligned with stakeholder needs
- Responses will be reviewed by CHIA to:
 - Refine submission templates and manuals
 - Adjust timelines and outreach strategies
 - Develop guidance materials for future reporting cycles

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Legislation Overview

- An Act Relative to Pharmaceutical Access, Costs, and Transparency mandates pharmacy benefit manager (PBM) data reporting to improve transparency
- Signed into law on January 9, 2025 by Governor Maura Healey
- Establishes M.G.L. c.12C § 10A, directing CHIA to collect detailed data from PBMs
- Directs CHIA to promulgate regulations necessary to ensure uniform data reporting from PBMs to analyze rebates, fees, wholesale acquisition costs, formulary, and maximum allowable cost lists, etc.
- Incorporates a listening session as a part of CHIA's implementation

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Future Opportunities for Engagement

- This session will not cover:
 - Feedback related to draft regulation 957 CMR 12.00
 - A public hearing will be held on November 20, 2025 at 10:00 AM
 - Register in advance by emailing Regulations@chiamass.gov
 - Submit written testimony to <u>Regulations@chiamass.gov</u>
 - Must include Sender's full name, mailing address, and organization or affiliation (if applicable).
 - PBM licensure requirements
 - The Division of Insurance has a dedicated email to PBM licensure and registration questions: DOI.PBM@mass.gov

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Draft Reporting Template

Purpose:

- Standardizes PBM data for CHIA
- Provides insight into pharmacy benefits for PBMs
- Collects information for regulatory review

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Draft Formulary Template

Purpose:

- Collects formulary data for each plan option
- Details drug lists, formulary tiers, and utilization management
- Provides information for key points of contact
- Supports regulatory review and transparency

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Overview of the Draft Data Specification Manual



Purpose:

- Provides comprehensive guidelines and requirements for PBM data submission
- Includes instructions, schedules, and clear technical specifications to support consistency and accuracy
- Offers contacts for support and clarification

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Draft Reporting Template

This document includes:

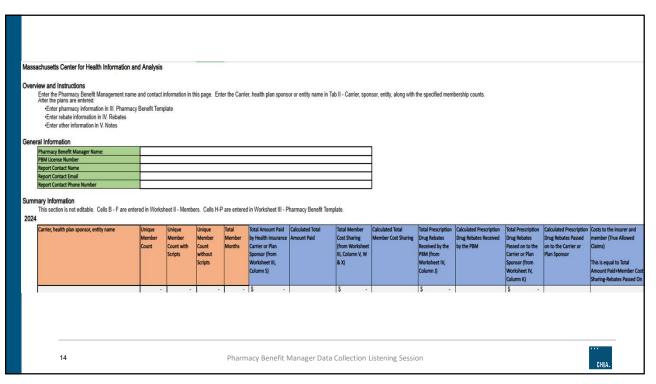
- PBM and contact information
- Carrier, health plan sponsor or entity name, along with the specified membership counts.
- Pharmacy information
- Rebate information
- Required questions

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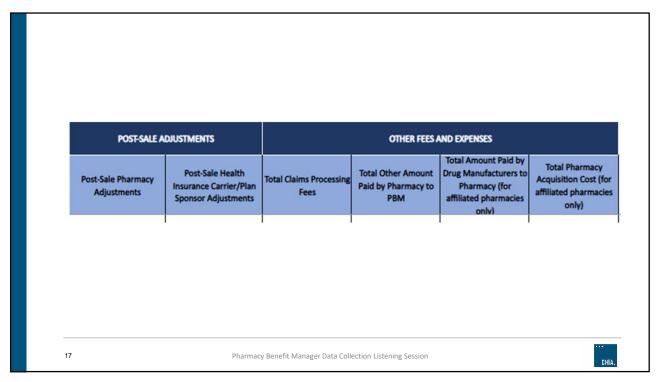


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lease provide an overview of your company ne template. Note that data is requested or	s information a		alam All area	lauara aan k	ambiand and a JJ-J	l to ACO which been	haan ingludad /-						
ere not rejected or denied.													
20-100 (27-0-9 € 10 del 10 20 del 10		Mem	bers			·			Total Fees				
Carrier, health plan sponsor, entity name	Unique Member Count	Unique Member Count with Scripts	Unique Member Count without Scripts	Total Member Months	Total Fees Paid by Drug Manufacturers	Total Fees Paid by Drug Manufacturers to PBM Rebate Aggregator	Total Fees Paid by Drug Manufacturers to PBM	Total Manufacturer Administrative Fees	Total Other Fees Paid by Drug Manufacturers	Total Fees Paid by PBM Rebate Aggregator to PBM	Total Fees Paid by PBM to PBM Rebate Aggregator	Total Fees Paid by Insurance Carrier/Plan Sponsor for Rebate Negotiation Services	Total F Passed Th to Insur Carrier/ Spons

Total WAC	Total AWP	Total Pharmacy Dispensing Revenue	Ingredient Cost Paid	Dispensing Fe	Other Pharmacy res Paid Receivable Amounts Paid	Total Amount Paid by Hea Insurance Carrier or Plat Sponsor			Amount Paid by Member	Total Member Deductibles	Total M Coinsu		al Member Copay ^{Ti}	otal Other Amount P to Pharmacy
							+	-					-	
										,				
YEAR		PLAN INFORMAT	TON			DRUG INFORMATION		PHARMACY	3408		U	ITILIZATION INFO	DRMATION	
Calendar Year	Pharmacy Benefit Manager Name	Carrier, health plan spo	onsor, entity	Risk Type	Drug or Product Name	lational Drug Code Bran	Category (Traditional d, Traditional Generic, ialty Brand, Specialty Generic)	Affiliated Pharmacy Flag	3408 Flag	Total Number of Units	Unit Type	Total Numbe Prescription		Total Members Scripts
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YEAR		PLAN IN	NFORMATION			2 21		INFORMATION				
Calendar Year	Pharmacy Benefit N Name	Manager	Carrier, health plan name	Risk Type	Drug or Product Name	Brand, Trac Specialty E	ory (Traditional litional Generic, grand, Specialty generic)	National Drug Cod	e (NDC)	Total Number of Prescriptions	Total Days Supply	Total WAC
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					REBATES							
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Total Prescriptio	on Drug Rebates	Rebate			Total Prescription D	Orug Base		tion Drug Other	Passed			assed Through at
Total Prescriptic	on Drug Rebates	Rebate	es Paid to PBM		Total Prescription D	Prug Base		tion Drug Other	Passed	Through to Insura		assed Through at
Total Prescriptic	on Drug Rebates	Rebate	es Paid to PBM		Total Prescription D	Orug Base		tion Drug Other	Passed	Through to Insura		assed Through at

	Provide a definition of administrative fees, including distinctions based on payer/payee relationships (e.g., fees paid by health plans to PBMs vs. manufacturers to PBMs or PBM Rebate Aggregators).	
	List each of your affiliated entities that operate in the Commonwealth or provide services that affect insurance carriers, plan sponsors, or members in the Commonwealth, including but not limited to mail order pharmacies, specialty pharmacies, retail pharmacies, insurers, third-party administrators, 340B split billing vendors, copay maximizers/accumulators, discount card programs, drug private labelers, and wholesale distributors.	
	Please describe the services you provide to your clients (health plans, employers, etc.) by selecting all that apply: Formulary management	
	 □ Pharmacy network design □ Rebate negotiation □ Claims processing □ Utilization management and/or other clinical programs 	
	☐ Other: (please specify)	
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	ow you determine the pharmacy reimbursement rate for traditional generic, traditional brand, specialty generic, and ugs including any benchmarks you use (ex. NADAC, WAC, AWP, MAC lists). What proportion of prescriptions for traditional
	brand, specialty generic, and specialty brand drugs are reimbursed using each methodology you employ?
What criteria or per	erformance metrics determine when and how post-sale pharmacy adjustments are applied?
When rebates are	applied, what factors influence whether patients see a direct reduction in their out of pocket costs?
Additional question	ns or miscellaneous items
•	

Thoughts and Feedback for Draft Submission Template

- Are there any fields that would be challenging for your organization to complete?
- Are there any missing data elements that are necessary to contextualize your organization's submission?
- What changes could be made to mirror your organization's internal systems to improve automation?

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Draft Formulary Template

Overview of document:

- Drug list and formulary tiers sheet for each plan option
- Step therapy protocols at the drug-level
- Options to reduce drug list sheets if similarities exist across plan options

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Instructions: Please submit a Drug List and Formulary Tiers sh Tiers sheet for each plan option. However, becau			*			
multiple Formulary Tiers sheets together with the			Contract of the second	A STATE OF THE SECOND STAT		
Important: •The Carrier, health plan sponsor, entity name en II. Carrier, sponsor, entity in the PBM Reporting Te •The Formulary Tier ID in IV. Formulary Drug List r	mplate.			ist match a Carrier, healti	n plan sponsor, en	itity name entry in tab
The Formulary Tier ID must be unique for each fo the template will create this ID. The Formulary Drug List ID must be unique for ea person filling out the template will create this ID.	rmulary tier, startii	ng with the letter "T"	and followed by thr			
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The Formulary Tier ID must be unique for each for the template will create this ID. The Formulary Drug List ID must be unique for each person filling out the template will create this ID. Create a new sheet for each new formulary by: Right-click on the "Formulary Tiers" tab Select "Move or Copy Sheet" Click the "Create a Copy" box Repeat for Formulary Drug List	rmulary tier, startii	ng with the letter "T"	and followed by thr	wed by five numbers betw		

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	art Contact Phone Number
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sponsor, entity name Formulary Tier ID					or entity name in PBM		unique for the set of tiers	described on this
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Drug Tier	Tier Description Required: Enter a description for each							
usually signify less preferred	Required: Enter a description for each tier including cost sharing information. Enter IVIA for any tier that is not associated with the formulary.							
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Carrier, health plan sponsor, entity name				A STATE OF THE PARTY OF THE PAR		ntity name in PBM Reporting template	
ormulary Tier ID ormulary Drug List ID					nulary Tier ID in tab III. Formu		ch drug list will have a unique identifier (E.g. D50123).
Plan Name(s)					ng with 'D' lottowed by 3 hai nal plan names with a comm		unug ust mit nave a unique identine (E.g. D00123).
Formulary URL	-						
	Drug Information			Utilization Manage	ement	Distribution	Notes
Drug Name	Therapeutic Category	Drug Tier	Prior Authorization	Quantity Limits		Limited Distribution Drug	
Enter the brand and generic names on separate rows.		Select NA if this drug is not a part		Select Yes if Quantity Limits	Required	If this is a limited distribution drug, select one of the listed requirements or "Other". If Other is selected, specify	Please enter any notes/comments/clarifications for each row here.
4	categories with a comma.	*	524			in Notes. If the drug is not limited,	
	18		•				

Thoughts and Feedback for Draft Formulary Template

- Do you anticipate any challenges applying the tiering structure outlined in the formulary template?
- What changes would be most useful for providing this information at an individual plan level?
- What system do you use to standardize therapeutic classification?

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Data Specification Manual



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What's inside?

- Contents of the document:
 - Introduction
 - File Submission Instructions and Schedule
 - · Data Submission Guidelines
 - · Data Specifications
 - · a. PBM Reporting Template
 - b. Formulary Template

Data Specification Manual

957 CMR 12.00:

Pharmacy Benefit Manager Reporting

October 2025

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Section: File Submission Instructions

- Details the process for PBMs to submit required data, including instructions for notification if data elements cannot be reported
- Provides guidance on requirements for submission

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Section: Data Submission Guidelines

- Specifies the requirements for submitting pharmacy benefits and rebate information
- Provides instructions for completing both the reporting template and the formulary template

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Section: Data Specifications

- Expands on data elements to guide user submission
- Defines the specific data elements that are necessary for different columns across the templates

IV. Data specifications

PBM Reporting Template

Tab I: Overview

Item	Column	Description	Data Requirements
Pharmacy Benefit Manager Name	C8	Name of the Pharmacy Benefit Manager that is represented in this document. All entries in the other worksheets will be for this PBM.	Data Type: Text
PBM License Number	C9		Data Type: Text
Report Contact Name	C10	Name of the person who is knowledgeable and available to answer questions about entries in this workbook.	Data Type: Text

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Draft Data Specification Manual Review – Data Specifications

Tab III. Pharmacy Benefit Template:

Data in this table should be produced at the PBM, calendar year, carrier [or plan], risk type, affiliated pharmacy status, 340B status (for affiliated pharmacies), and NDC level.

Item	Column	Description	Data Requirements
		Year	
Calendar Year	A	Calendar year in which the drug was dispensed.	Data Type: Integer
		Plan Information	
Pharmacy Benefit Manager Name	В	Name of the pharmacy benefit manager completing the template from I. Overview.	Data Type: Text
Carrier, health plan sponsor, entity name	С	Select from the drop-down list. If the plan is not shown, add it to the list in II. Carrier, health plan sponsor, entity name.	
Risk Type	D	Self-insured or Fully-insured.	

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Thoughts and Feedback on Draft Data Specification Manual

- Are there any missing or unclear field definitions and data requirements outlined in section IV of the Data Specification Manual?
- Is there any missing guidance in the manual to ensure accurate and consistent data entry across all required fields?
- Are there any formatting and data type specifications (e.g., text, max length) that do not seem feasible to implement within your internal systems? And why?
- Have you encountered any challenges interpreting or applying the specifications in the Overview tab? If so, what additional guidance would be helpful?

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Next Steps

- Future users should continue to review the materials and familiarize themselves with the manual and templates
- Written feedback can be provided and sent to the following email: [RxData@chiamass.gov]
 - All participants are highly encouraged to provide any comments and/or feedback regarding the materials discussed in this listening session
 - Please submit written feedback by November 14, 2025
- Please note: Feedback shared will NOT be made publicly available and only CHIA and AIR will have access to this information. The feedback will not be published externally.

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Thank you!