

Change Summary

PSAO Data Submission Guide v3.0 and v4.0

The changes between version 3.0 and 4.0 of the psao' data submission guides (DSG) are indicated in red in the table below.

Type of change	Templat e	Version 3.0 (Cu	rrent)	Version 4.0
Definition Update:		"Current year" means the year 2021.	"Current year"	means the year 2022.
Current year Definition Update:		"Prior Year" means calendar year 2020.	"Prior year" me	ans calendar year 2021.
Prior year				
Update: How to Register and Submit			email address u secondary cont (two) contacts.	pp for a Secure Access Washington Account (SAW). The sed for this SAW account must match the DPT primary or act email address. Please note you are only allowed 2 This is a change from the previous system.
Update Field: Data Validation			be rejected. Each validation to end 43.71C and is considered to cover verification on the cover verification of the community	is a two-step process and at any time submissions may ch submitted file undergoes technical and program sure that the data meets the requirements of RCW compatible with HCA's reporting software. These primarily on of data types (number vs. string) and formats (2023-/2023). The program validation process is performed by fiter technical validation and includes additional checks of plete the data validation process. rejected during Program validation, you will need to ected report within 10 business days of receipt of the submit validation. If your submission passed or failed validation a message indicating your submission was essfully uploaded will appear on the screen. If your mission failed, you would see an error log noting a list of extract must be corrected. All errors must be ected prior to clicking the submit button. If you do not live an email notification of either success or failure in 72 hours of submitting your report, please contact program staff at drugtransparency@hca.wa.gov to irm that your submission was received and processed.

			Step 2 Program validation – An analyst will validate information submitted in ensure it meets program requirements. You will receive an approval email or a rejection email. This email will be sent to the email provided when you registered. If your report is rejected, you will need to resubmit within 10-days. If you need help understanding your error log, the Data Submission FAQ clarifies the meaning of the error and provides guidance on corrections, or you may submit your questions to HCADPTTechSupport@hca.wa.gov for assistance.
Update Field: Correcting Submissions			In the event that you find an error in your approved submission, you will need to fill out the Resubmission form which can be found on our portal prior to resubmitting your report. You will need to let HCA know which
			report you will be resubmitting and the specific reasons why you request to resubmit. HCA will review your request and approve or deny your request within 5 business days. In the event your resubmission is rejected during technical or program validation, you would be subject to the 10-day limit for correcting rejected resubmissions.
Update:		File naming schema:	File naming schema:
Table		psao pharmacy contracted rat	psao_pharmacy_contracted_rates_{YYYY}_{ID}_{YYYYMMDD}.csv
Specifications		es_{YYYY}_{ID}_{YYYYMMDD}.cs	• Example:
		V	psao_pharmacy_contracted_rates2022_S12345_20231001.csv
		Example:	Please use the submission due date not the date the report
		·	was prepared for YYYYMMDD
		psao_pharmacy_contracted_rat es2021_S12345_20211201.cs v (Please use the submission due date not the date the report was prepared)	The submission of this report is due on October 1, 2023, and should include data effective for 2022.
		The submission of this report is	
		due on December 1, 2022, and should include data effective for 2021.	
Update Field:	Pharmacy	Name: Washington DPT Number Type: String	Name: Washington DPT Number Type: String
Washington	Contracte	Max Length: 6 characters	Max Length: 6 characters
DPT Number	d Rates	Format: ABCDE	Format: ABCDEF
Update Field:		Name: Year	Name: Year
Year		Type: Numeric	Type: Numeric
		Format: 9999 Max Length: 4 digits	Format: 9999 Max Length: 4 digits
		Min Length: 4 digits	Min Length: 4 digits
		Rule: 2021	Rule: 2022
		Current year for which the aggregate data is reported.	Current year for which the aggregate data is reported.
Update Field:		Name: Drug Name	Name: Drug Name
Drug Name		Type: String	Type: String
<u> </u>	1		l .

	Max Length: 100 characters Format: ABCDE	Max Length: 1 Format: ABCD				
	Name of the drug for the NDC reported. Only include	Name of the drug for the NDC reported. Only include ingredient name.				
	ingredient name.	For example:				
	For example, if the NDC has a	NDC	Drug Name	Drug Product Name	Label Name	
	Drug Product Name of "fluoxetine HCL 20 mg tablets", then this field should be reported as "fluoxetine". All	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA	
	drug product names with	00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA	
	"fluoxetine" in its name should be reported as a single Drug	00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER	
	Name in this field. Combination	00000000000	AMOXICILLIN	AMOXICILLIN 500 MG TABLET	AMOXICILLIN	
	drug product names should be reported individually as its own	00000000000	AMOXICILLIN	AMOXICILLIN 500 MG CAPSULE	AMOXICILLIN	
	Drug Name instead of by each ingredient.	NOTE: Special	characters, hyp	hens, symbols, or slashes	are allowed.	
	NOTE: Special characters, hyphens, symbols, or slashes are allowed.					
Update Field: Drug Product Name	Name: Drug Product Name Type: String Max Length: 100 characters Format: ABCDE	Name: Drug Product Name Type: String Max Length: 100 characters Format: ABCDE				
	Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage	name as repoi include ingred	rted in standard	the NDC reported, to inclized drug databases. This dosage form, strength, and CC.	name should	
	form, strength, and any other information specific to the NDC.	NDC	Drug Name	Drug Product Name	Label Name	
	For example, "fluoxetine HCL 20 mg tablets" is acceptable.	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA	
		0000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA	
		0000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER	
		0000000000	AMOXICILLIN	AMOXICILLIN 500 MG TABLET	AMOXICILLIN	
		0000000000	AMOXICILLIN	AMOXICILLIN 500 MG CAPSULE	AMOXICILLIN	
Update Field: Label Name	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE	Name: Label N Type: String Max Length: 1 Format: ABCD	.00 characters		•	
	Proprietary or legal name as	Proprietary or	legal name as r	narketed by the manufact	urer.	
	marketed by manufacturer. For example, "fluoxetine HCL",	NDC	Drug Name	Drug Product Name	Label Name	
	"fluoxetine DR" are acceptable.	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA	

			T T	I	1	
				DISOPROXIL FUMARATE		
			0000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA
			0000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER
			0000000000	AMOXICILLIN	AMOXICILLIN 500 MG TABLET	AMOXICILLIN
			0000000000	AMOXICILLIN	AMOXICILLIN 500 MG CAPSULE	AMOXICILLIN
			NOTE: Special	characters, hyp	ohens, symbols, or slas	hes are allowed.
Update Field: General Comments		Name: General Comments Type: String Max Length: 5000 characters Format: ABCDE Nullable Any additional information you would like to submit or provide to explain your responses.	Format: ABCD Nullable Any additional explain your re	000 characters E I information yo	u would like to submit urns.	or provide to
Update: Table Specifications	Pharmacy Year-Over- Year Rate Change	File naming schema: psao_pharmacy_yoy_rate_chan ge_{YYYY}_{ID}_{YYYYMMDD}.cs v Example: psao_pharmacy_yoy_rate_chan ge_2020_S12345_20211201.csv (Please use the submission due date not the date the report was prepared) The submission of this report is due on December 1, 2022, and should include data effective for 2021.	Exa psa csv Ple wa The submissio	cy_yoy_rate_chample: ao_pharmacy_yo asse use the sub s prepared for Y	is due on October 1, 20	_S12345_20231001. he date the report
Update Field: Washington DPT Number Update Field: Year		Name: Washington DPT Number Type: String Max Length: 6 characters Format: ABCDE Name: Year Type: Numeric Format: 9999	Name: Washir Type: String Max Length: 6 Format: ABCD Name: Year Type: Numeric Format: 9999	EF	per	
		Max Length: 4 digits Min Length: 4 digits Rule: 2021	Max Length: 4 Min Length: 4 Rule: 2022	=		

	Current year for which the	Current year f	or which the ag	gregate data is reported.			
Update Field:	aggregate data is reported. Name: Drug Name	Name: Drug N	lame				
opunte ricia.	Type: String	Type: String	idiric				
Drug Name	Max Length: 100 characters	Max Length: 1	.00 characters				
8	Format: ABCDE	Format: ABCD					
	Name of the drug for the NDC	Name of the o	Irug for the NDC	reported. Only include in	gredient name.		
	reported. Only include						
	ingredient name.	For example:					
	For example, if the NDC has a	NDC	Drug Name	Drug Product Name	Label Name		
	Drug Product Name of	0000000000	EFAVIRENZ-	EFAVIRENZ-EMTRICITABINE-	ATRIPLA		
	"fluoxetine HCL 20 mg tablets",	11	EMTRICITABINE	TENOFOVIR DISOPROXIL			
	then this field should be		-TENOFOVIR DISOPROXIL	FUMARATE 10MG TABLET			
	reported as "fluoxetine". All	11	FUMARATE				
	drug product names with	0000000000	ADALIMUMAB	ADALIMUMAB	HUMIRA		
	"fluoxetine" in its name should			PEN INJ 40MG/0.8			
	be reported as a single Drug	0000000000	ADALIMUMAB	ADALIMUMAB	HUMIRA CD/UC/HS		
		00000000000	AMOXICILLIN	PEN INJ CD/UC/HS AMOXICILLIN 500 MG	STARTER AMOXICILLIN		
	Name in this field. Combination	000000000	AIVIOXICILLIN	TABLET	AWOXICILLIN		
	drug product names should be	0000000000	AMOXICILLIN	AMOXICILLIN 500 MG	AMOXICILLIN		
	reported individually as its own			CAPSULE			
	Drug Name instead of by each	NOTE: Special	characters, hyp	hens, symbols, or slashes	are allowed.		
	ingredient.						
	g. ca.c						
	NOTE: Consider the sections						
	NOTE: Special characters,						
	hyphens, symbols, or slashes are	hyphens, symbols, or slashes are					
	allowed.						
Update Field:	Name: Drug Product Name	Name: Drug P	roduct Name				
Update Field:	Name: Drug Product Name Type: String	Name: Drug P Type: String	roduct Name				
	Type: String	Type: String					
Update Field: Drug Product Name	_	_	.00 characters				
Drug Product	Type: String Max Length: 100 characters	Type: String Max Length: 1	.00 characters				
Drug Product	Type: String Max Length: 100 characters Format: ABCDE	Type: String Max Length: 1 Format: ABCD	.00 characters E	the NDC reported, to incl	ude ingredient		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for	Type: String Max Length: 1 Format: ABCD Name of the c	.00 characters E Irug product for	the NDC reported, to incl ized drug databases. This	-		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include	Type: String Max Length: 1 Format: ABCD Name of the coname as repo	.00 characters E Irug product for rted in standard	ized drug databases. This	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in	Type: String Max Length: 1 Format: ABCD Name of the c name as report include ingred	.00 characters E Irug product for rted in standard lient, salt form,	ized drug databases. This dosage form, strength, an	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases.	Type: String Max Length: 1 Format: ABCD Name of the c name as report include ingred	.00 characters E Irug product for rted in standard	ized drug databases. This dosage form, strength, an	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in	Type: String Max Length: 1 Format: ABCD Name of the c name as report include ingred	.00 characters E Irug product for rted in standard lient, salt form,	ized drug databases. This dosage form, strength, an	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases.	Type: String Max Length: 1 Format: ABCD Name of the c name as report include ingred	.00 characters E Irug product for rted in standard lient, salt form,	ized drug databases. This dosage form, strength, an	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include	Type: String Max Length: 1 Format: ABCD Name of the c name as repoi include ingred information sp	.00 characters E Irug product for rted in standard lient, salt form,	ized drug databases. This dosage form, strength, an	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage	Type: String Max Length: 1 Format: ABCD Name of the c name as repoi include ingred information sp	.00 characters E Irug product for rted in standard lient, salt form,	ized drug databases. This dosage form, strength, an	name should		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC.	Type: String Max Length: 1 Format: ABCD Name of the c name as repoi include ingred information sp For example:	OO characters E Irug product for rted in standard lient, salt form, pecific to the NE Drug Name EFAVIRENZ-	ized drug databases. This dosage form, strength, an IC. Drug Product Name EFAVIRENZ-EMTRICITABINE-	name should d any other		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingred information specific for example:	On characters If the product for the instandard lient, salt form, opecific to the NE Drug Name EFAVIRENZ- EMTRICITABINE	ized drug databases. This dosage form, strength, an IC. Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL	name should d any other Label Name		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC.	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingred information specific for example:	Drug Name EFAVIRENZ- EMTRICITABINE -TENOFOVIR	ized drug databases. This dosage form, strength, an IC. Drug Product Name EFAVIRENZ-EMTRICITABINE-	name should d any other Label Name		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingred information specific for example:	On characters If the product for the instandard lient, salt form, opecific to the NE Drug Name EFAVIRENZ- EMTRICITABINE	ized drug databases. This dosage form, strength, an IC. Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL	name should d any other Label Name		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingred information specific for example:	OO characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL	ized drug databases. This dosage form, strength, an IC. Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB	name should d any other Label Name		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingred information specific reasons in the coname as reporting in the conam	OO characters E Irug product for rted in standard lient, salt form, pecific to the NE Drug Name EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8	name should d any other Label Name ATRIPLA HUMIRA		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingredinformation specific for example: NDC 000000000000000000000000000000000	On characters E Irug product for red in standard lient, salt form, pecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIR FUMARATE ADALIMUMAB	ized drug databases. This dosage form, strength, an IC. Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB	name should d any other Label Name ATRIPLA HUMIRA		
_	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the c name as repoi include ingred information sp For example: NDC 000000000000000000000000000000000	On characters E Irug product for red in standard lient, salt form, pecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIR FUMARATE ADALIMUMAB	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CO/UC/HS AMOXICILLIN 500 MG	name should d any other Label Name ATRIPLA HUMIRA HUMIRA		
Drug Product	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting information specific	OO characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting include ingred information specific for example: NDC 000000000000000000000000000000000	On characters E Irug product for reed in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20 mg tablets" is acceptable.	Type: String Max Length: 1 Format: ABCD Name of the coname as report include ingred information specific for example: NDC 000000000000000000000000000000000	On characters E Irug product for reed in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20 mg tablets" is acceptable. Name: Label Name Type: String	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting information specific	On characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB AMOXICILLIN AMOXICILLIN Name	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20 mg tablets" is acceptable. Name: Label Name Type: String Max Length: 100 characters	Type: String Max Length: 1 Format: ABCD Name of the coname as report include ingred information specific for example: NDC 000000000000000000000000000000000	OO characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB AMOXICILLIN AMOXICILLIN Name OO characters	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20 mg tablets" is acceptable. Name: Label Name Type: String	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting information specific	On characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB AMOXICILIN AMOXICILIN Name On characters E	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20 mg tablets" is acceptable. Name: Label Name Type: String Max Length: 100 characters	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting information specific	On characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB AMOXICILIN AMOXICILIN Name On characters E	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN		
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage form, strength, and any other information specific to the NDC. For example, "fluoxetine HCL 20 mg tablets" is acceptable. Name: Label Name Type: String Max Length: 100 characters Format: ABCDE	Type: String Max Length: 1 Format: ABCD Name of the coname as reporting information specific	On characters E Irug product for rted in standard lient, salt form, opecific to the NE Drug Name EFAVIRENZ-EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB AMOXICILIN AMOXICILIN Name On characters E	Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE	name should d any other Label Name ATRIPLA HUMIRA HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN		

		example, "fluoxetine HCL", "fluoxetine DR" are acceptable.	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA
			0000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA
			00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER
			00000000000	AMOXICILLIN	AMOXICILLIN 500 MG TABLET	AMOXICILLIN
			00000000000	AMOXICILLIN	AMOXICILLIN 500 MG CAPSULE	AMOXICILLIN
Update Field: General Comments		Name: General Comments Type: String Max Length: 5000 characters Format: ABCDE Nullable Any additional information you would like to submit or provide to explain your responses.	Format: ABCD Nullable Any additional explain your re	000 characters E I information yo	u would like to submit or urns.	provide to
Update:	PBM Contracted Rates	File naming schema: psao_pbm_contracted_rates_{Y YYY}_{ID}_{YYYYMMDD}.csv Example: psao_pbm_contracted_rates_20 20_512345_20211201.csv (Please use the submission due date not the date the report was prepared) The submission of this report is due on December 1, 2022, and should include data effective for 2021.	Example psao_pb Please u prepared The submission	ntracted_rates_ :: :m_contracted_! se the submissic d for YYYYMMDI	is due on October 1, 2023	31001.csv the report was
Update Field: Washington DPT Number		Name: Washington DPT Number Type: String Max Length: 6 characters Format: ABCDE	Name: Washi Type: String Max Length: 6 Format: ABCD		ber	
Update Field: Year		Name: Year Type: Numeric Format: 9999 Max Length: 4 digits Min Length: 4 digits Rule: 2021 Current year for which the aggregate data is reported.	Name: Year Type: Numeric Format: 9999 Max Length: 4 Min Length: 4 Rule: 2022	digits digits	gregate data is reported.	
Update Field: Drug Name		Name: Drug Name Type: String Max Length: 100 characters	Name: Drug N Type: String Max Length: 1	ame .00 characters		
		Format: ABCDE Name of the drug for the NDC reported. Only include ingredient name.	Format: ABCD Name of the d For example:		reported. Only include in	gredient name.

	For example, if the NDC has a	NDC	Drug Name	Drug Product Name	Label Name
	Drug Product Name of "fluoxetine HCL 20 mg tablets", then this field should be reported as "fluoxetine". All	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA
	drug product names with	00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA
	"fluoxetine" in its name should be reported as a single Drug	00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER
	Name in this field. Combination	00000000000	AMOXICILLIN	AMOXICILLIN 500 MG	AMOXICILLIN
	drug product names should be	00000000000	AMOXICILLIN	TABLET AMOXICILLIN 500 MG	AMOXICILLIN
	reported individually as its own Drug Name instead of by each ingredient.	NOTE: Special	characters, hyp	hens, symbols, or slashes	are allowed.
	NOTE: Special characters, hyphens, symbols, or slashes are allowed.				
Update Field: Drug Product Name	Name: Drug Product Name Type: String Max Length: 100 characters Format: ABCDE	Name: Drug Pr Type: String Max Length: 1 Format: ABCD	00 characters		
	Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include ingredient, salt form, dosage	name as repor include ingred	ted in standard	the NDC reported, to inclized drug databases. This dosage form, strength, and C.	name should
	form, strength, and any other	NDC	Drug Name	Drug Product Name	Label Name
	information specific to the NDC.	0000000000	EFAVIRENZ- EMTRICITABINE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL	ATRIPLA
	For example, "fluoxetine HCL 20 mg tablets" is acceptable.		-TENOFOVIR DISOPROXIL	FUMARATE 10MG TABLET	
	· ·	0000000000	-TENOFOVIR	ADALIMUMAB	HUMIRA
	· ·	0000000000	-TENOFOVIR DISOPROXIL FUMARATE	ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB	HUMIRA CD/UC/HS
	· ·		-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	
	· ·	00000000000	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER
Hadata Field:	mg tablets" is acceptable.	0000000000	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN	ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CO/UC/HS AMOXICILLIN 500 MG TABLET	HUMIRA CD/UC/HS STARTER AMOXICILLIN
Update Field: Label Name	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE	00000000000000000000000000000000000000	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN lame 00 characters	ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INI CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG	HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN
	Name: Label Name Type: String Max Length: 100 characters	00000000000000000000000000000000000000	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN lame 00 characters	ADALIMUMAB PEN INI 40MG/0.8 ADALIMUMAB PEN INI CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE	HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN
	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE Proprietary or legal name as marketed by manufacturer. For	00000000000 00000000000 Name: Label N Type: String Max Length: 1 Format: ABCD Proprietary or	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN lame 00 characters E	ADALIMUMAB PEN INI 40MG/0.8 ADALIMUMAB PEN INI CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE	HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN
	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE Proprietary or legal name as marketed by manufacturer. For example, "fluoxetine HCL",	00000000000 00000000000 Name: Label N Type: String Max Length: 1 Format: ABCD Proprietary or	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN AMOXICILLIN Iame O0 characters E legal name as n Drug Name EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL	ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB	HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN urer. Label Name
	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE Proprietary or legal name as marketed by manufacturer. For example, "fluoxetine HCL",	00000000000 00000000000 Name: Label N Type: String Max Length: 1 Format: ABCD Proprietary or NDC 00000000000	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN Iame O0 characters E legal name as n Drug Name EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	ADALIMUMAB PEN INI 40MG/0.8 ADALIMUMAB PEN INI CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INI 40MG/0.8 ADALIMUMAB	HUMIRA CD/UC/HS STARTER AMOXICILIN AMOXICILIN Urer. Label Name ATRIPLA HUMIRA HUMIRA
	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE Proprietary or legal name as marketed by manufacturer. For example, "fluoxetine HCL",	00000000000 00000000000 Name: Label N Type: String Max Length: 1 Format: ABCD Proprietary or NDC 00000000000	-TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB ADALIMUMAB AMOXICILLIN AMOXICILLIN AMOXICILLIN DISOPROXIL EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8 ADALIMUMAB PEN INJ CD/UC/HS AMOXICILLIN 500 MG TABLET AMOXICILLIN 500 MG CAPSULE Drug Product Name EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA CD/UC/HS STARTER AMOXICILLIN AMOXICILLIN UTER. Label Name ATRIPLA HUMIRA

General Comments	Name: General Comments Type: String Max Length: 5000 characters Format: ABCDE Nullable	Name: General Comments Type: String Max Length: 5000 characters Format: ABCDE Nullable
	Any additional information you would like to submit or provide to explain your responses.	Any additional information you would like to submit or provide to explain your responses.
		Note: Do not include hard returns.
Update: PBM Ye	8	File naming schema:
Rate Change	psao_pbm_yoy_rate_change_{Y YYY}_{ID}_{YYYYMMDD}.csv	psao_pbm_yoy_rate_change_{YYYY}_{ID}_{YYYYMMDD}.csv • Example: Example:
	Example: Example:	psao_pbm_yoy_rate_change_2022_S12345_20231001.csv • Please use the submission due date not the date the report was
	psao_pbm_yoy_rate_change_2	prepared for YYYYMMDD
	020_S12345_20211201.csv (Please use the submission due date not the date the report was prepared)	The submission of this report is due on October 1, 2023, and should include data effective for 20221.
	The submission of this report is due on December 1, 2022, and should include data effective for 2021.	
Delete Field: Washington DPT Number	Name: Washington DPT Number Type: String Max Length: 6 characters Format: ABCDE	Name: Washington DPT Number Type: String Max Length: 6 characters Format: ABCDEF
Update Field: Year	Name: Year Type: Numeric Format: 9999 Max Length: 4 digits Min Length: 4 digits Rule: 2021	Name: Year Type: Numeric Format: 9999 Max Length: 4 digits Min Length: 4 digits Rule: 2022
	Current year for which the aggregate data is reported.	Current year for which the aggregate data is reported.
Update Field: Drug Name	Name: Drug Name Type: String Max Length: 100 characters Format: ABCDE	Name: Drug Name Type: String Max Length: 100 characters Format: ABCDE
	Name of the drug for the NDC reported. Only include ingredient name.	Name of the drug for the NDC reported. Only include ingredient name. For example:
	For example, if the NDC has a	NDC Drug Name Drug Product Name Label Name
	Drug Product Name of "fluoxetine HCL 20 mg tablets", then this field should be reported as "fluoxetine". All	0000000000 EFAVIRENZ- EMTRICITABINE TENOFOVIR DISOPROXIL FLIMARATE 10MG TABLET FLIMARATE
	drug product names with	FUMARATE
	"fluoxetine" in its name should be reported as a single Drug	0000000000 ADALIMUMAB ADALIMUMAB HUMIRA CD/UC/HS PEN INJ CD/UC/HS STARTER

	drug product names should be		1	TABLET	1	
	drug product names should be reported individually as its own	00000000000	AMOXICILLIN	AMOXICILLIN 500 MG	AMOXICILLIN	
	Drug Name instead of by each ingredient.	NOTE: Special characters, hyphens, symbols, or slashes are allowed.				
	NOTE: Special characters, hyphens, symbols, or slashes are allowed.					
Update Field:	Name: Drug Product Name	Name: Drug P	roduct Name			
Drug Product Name	Type: String Max Length: 100 characters Format: ABCDE	Type: String Max Length: 100 characters Format: ABCDE				
	Name of the drug product for the NDC reported, to include ingredient name as reported in standardized drug databases. This name should include	name as repo	rted in standard	the NDC reported, to inc ized drug databases. This dosage form, strength, ar IC.	name should	
	ingredient, salt form, dosage form, strength, and any other	Tor example.				
	information specific to the NDC.	NDC	Drug Name	Drug Product Name	Label Name	
	For example, "fluoxetine HCL 20 mg tablets" is acceptable.	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA	
		00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA	
		00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/H: STARTER	
		0000000000	AMOXICILLIN	AMOXICILLIN 500 MG TABLET	AMOXICILLIN	
		00000000000	AMOXICILLIN	AMOXICILLIN 500 MG CAPSULE	AMOXICILLIN	
Update Field: Label Name	Name: Label Name Type: String Max Length: 100 characters Format: ABCDE Proprietary or legal name as	Name: Label N Type: String Max Length: 1 Format: ABCD	turer.			
	marketed by manufacturer. For	NDC	Drug Name	Drug Product Name	Label Name	
	example, "fluoxetine HCL", "fluoxetine DR" are acceptable.	0000000000	EFAVIRENZ- EMTRICITABINE -TENOFOVIR DISOPROXIL FUMARATE	EFAVIRENZ-EMTRICITABINE- TENOFOVIR DISOPROXIL FUMARATE 10MG TABLET	ATRIPLA	
		0000000000	ADALIMUMAB	ADALIMUMAB PEN INJ 40MG/0.8	HUMIRA	
		00000000000	ADALIMUMAB	ADALIMUMAB PEN INJ CD/UC/HS	HUMIRA CD/UC/HS STARTER	
		0000000000	AMOXICILLIN	AMOXICILLIN 500 MG TABLET	AMOXICILLIN	
		0000000000	AMOXICILLIN	AMOXICILLIN 500 MG CAPSULE	AMOXICILLIN	
Update Field: General Comments	Name: General Comments Type: String Max Length: 5000 characters Format: ABCDE Nullable	Name: Genera Type: String Max Length: 5 Format: ABCD Nullable	6000 characters			
	Any additional information you would like to submit or provide to explain your responses.	Any additiona explain your r	· ·	u would like to submit or	provide to	
		Note: Do not	include hard ret	turns.		