Office of Regulatory Management

Economic Review Form

Agency name	DEPARTMENT OF MEDICAL ASSISTANCE SERVICES
Virginia Administrative Code (VAC) Chapter citation(s)	12VAC30-130-1000
VAC Chapter title(s)	Amount, Duration and Scope of Selected Services
Action title	Pharmacy and Therapeutics Committee Increase
Date this document prepared	January 28, 2026
Regulatory Stage (including Issuance of Guidance Documents)	Fast-Track

Cost Benefit Analysis

Complete Tables 1a and 1b for all regulatory actions. You do not need to complete Table 1c if the regulatory action is required by state statute or federal statute or regulation and leaves no discretion in its implementation.

Table 1a should provide analysis for the regulatory approach you are taking. Table 1b should provide analysis for the approach of leaving the current regulations intact (i.e., no further change is implemented). Table 1c should provide analysis for at least one alternative approach. You should not limit yourself to one alternative, however, and can add additional charts as needed.

Report both direct and indirect costs and benefits that can be monetized in Boxes 1 and 2. Report direct and indirect costs and benefits that cannot be monetized in Box 4. See the ORM Regulatory Economic Analysis Manual for additional guidance.

Table 1a: Costs and Benefits of the Proposed Changes (Primary Option)

(1) Direct & Indirect Costs & Benefits (Monetized)	The 2025 Appropriations Act, Item 288, CC.2.a, requires DMAS to increase the maximum number of Pharmacy and Therapeutics Committee members from 12 members to 16 members and also requires DMAS to include one physician from each contracted managed care organization (MCO). As a result, this action increases the maximum number of committee members to 16 and adds the requirement of having one physician from each contracted MCO. There are no direct or indirect costs associated with the regulation changes. The benefit of expanding the P&T Committee is that it broadens the committee membership and aims to improve the quality of the guidance provided by this committee.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits
	(a)	(b)
(3) Net Monetized Benefit		
(4) Other Costs & Benefits (Non-Monetized)		
(5) Information Sources		

Table 1b: Costs and Benefits under the Status Quo (No change to the regulation)

(1) Direct & Indirect Costs & Benefits (Monetized)	No change to the regulation would result in less comprehensive input and the potential that several MCOs may not be represented as part of the P&T Committee.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits
	(a)	(b)
(3) Net Monetized Benefit		

(4) Other Costs & Benefits (Non-Monetized)	
(5) Information Sources	

Table 1c: Costs and Benefits under Alternative Approach(es)

(1) Direct & Indirect Costs & Benefits (Monetized)	There are no alternative approaches that would accomplish the goal of this regulatory action.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits
	(a)	(b)
(3) Net Monetized Benefit		
(4) Other Costs & Benefits (Non-Monetized)		
(5) Information Sources		

Impact on Local Partners

Use this chart to describe impacts on local partners. See Part 8 of the ORM Cost Impact Analysis Guidance for additional guidance.

Table 2: Impact on Local Partners

(1) Direct & Indirect Costs & Benefits (Monetized)	No local partners will incur any direct or indirect costs of the regulatory changes contained in the regulatory action. A direct benefit from the regulatory change is the increased input from the now included MCOs.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits
	(a)	(b)

(3) Other Costs & Benefits (Non-Monetized)	
(4) Assistance	
(5) Information Sources	

Impacts on Families

Use this chart to describe impacts on families. See Part 8 of the ORM Cost Impact Analysis Guidance for additional guidance.

Table 3: Impact on Families

(1) Direct & Indirect Costs & Benefits (Monetized)	There are no direct or indirect costs to families. The direct benefit of this proposed change to families is the potential for improved health outcomes as more expertise and input will be included in the decision-making process regarding medications.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits
	(a)	(b)
(3) Other Costs & Benefits (Non-Monetized)		
(4) Information Sources		

Impacts on Small Businesses

Use this chart to describe impacts on small businesses. See Part 8 of the ORM Cost Impact Analysis Guidance for additional guidance.

Table 4: Impact on Small Businesses

(1) Direct & Indirect Costs & Benefits (Monetized)	There are no direct or indirect costs to small businesses.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits

	(a)	(b)
(3) Other Costs & Benefits (Non-Monetized)		
(4) Alternatives		
(5) Information Sources		

Changes to Number of Regulatory Requirements

Table 5: Regulatory Reduction

For each individual action, please fill out the appropriate chart to reflect any change in regulatory requirements, costs, regulatory stringency, or the overall length of any guidance documents.

Change in Regulatory Requirements

VAC Section(s) Involved*	Authority of Change	Initial Count	Additions	Subtractions	Total Net Change in Requirements
12VAC30-130-1000	**(M/A):**	**30**	2	0	2
	(D/A):	**0**	0	0	0
	(M/R):	**0**	0	0	0
	(D/R):	**0**	0	0	0
				Grand Total of Changes in Requirements:	**(M/A):2**
					(D/A): 0
					(M/R):0
					(D/R): 0

Key:
Please use the following coding if change is mandatory or discretionary and whether it affects externally regulated parties or only the agency itself:
(M/A): Mandatory requirements mandated by federal and/or state statute affecting the agency itself
(D/A): Discretionary requirements affecting agency itself
(M/R): Mandatory requirements mandated by federal and/or state statute affecting external parties, including other agencies
(D/R): Discretionary requirements affecting external parties, including other agencies

Cost Reductions or Increases (if applicable)

VAC Section(s) Involved*	Description of Regulatory Requirement	Initial Cost	New Cost	Overall Cost Savings/Increases

Other Decreases or Increases in Regulatory Stringency (if applicable)

VAC Section(s) Involved*	Description of Regulatory Change	Overview of How It Reduces or Increases Regulatory Burden

Length of Guidance Documents (only applicable if guidance document is being revised)

Title of Guidance Document	**Original Word Count**	**New Word Count**	**Net Change in Word Count**

*If the agency is modifying a guidance document that has regulatory requirements, it should report any change in requirements in the appropriate chart(s).