Office of Regulatory Management

Economic Review Form

Agency name	Board of Pharmacy
Virginia Administrative Code (VAC) Chapter citation(s)	18VAC110-20
VAC Chapter title(s)	Regulations Governing the Practice of Pharmacy
Action title	June 2024 scheduling of chemicals in Schedule I
Date this document prepared	5/16/2024
Regulatory Stage (including Issuance of Guidance Documents)	Exempt

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)	There are no direct or indirect costs of the change to the agency. There are no direct or indirect benefits of the change. This change solely adds chemicals to Schedule I based on DFS guidance. This change likely adds costs for law enforcement entities and prosecution entities in the Commonwealth, because the number of substances that it is illegal to possess or use increases. The cost for this change is unquantifiable by the Board, however, because the changes are speculative and involve agencies and entities in the Commonwealth with their own financial considerations of which the Board is not aware. Due to this limitation, no costs or benefits are provided below.	
(2) Present Monetized Values	Direct & Indirect Costs	Direct & Indirect Benefits
	(a) 0	(b) 0
(3) Net Monetized Benefit	0	
(4) Other Costs & Benefits (Non-Monetized)	0	
(5) Information Sources		

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

(1) Direct & Indirect Costs & Benefits (Monetized)	There are no direct or indirect costs to status quo. No direct or indirect benefits to status quo.	
(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 to scheduling actions under Virginia Code § 54.1-3443(D).	
(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)	Please see Table 1a for impacts on local partners. This impact is unquantifiable by the Board, however, so no costs or benefits are included below.	
(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 is no impact on families.	
(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 is no impact on 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
	(M/A):				
	(D/A):				
	(M/R):				
	(D/R):	**1**	0	0	0
				Grand Total of Changes in Requirements:	**(M/A): 0**
					(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
18VAC110-20-322	Adds chemicals and drugs to Schedule I.	There is no increase or decrease in the regulatory burden because practitioners are not required to take any action. Additionally, this

		change is only effective for 18 months. If the General Assembly acts to place these scheduled drugs in Code, the Board will delete these additions following the effective date of changes made in the next General Assembly Session. Rarely, the General Assembly will not act to permanently place drugs in Schedule I that the Board has placed there. If that occurs, the Board will delete the changes after the 18 month effective window has closed.

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

Title of Guidance Document	**Original Length**	**New Length**	**Net Change in Length**