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## Exempt Action: Final Regulation Agency Background Document

<b>Agency name</b>	Board of Pharmacy, Department of Health Professions
<b>Virginia Administrative Code (VAC) Chapter citation(s)</b>	18VAC110-20
<b>VAC Chapter title(s)</b>	Regulations Governing the Practice of Pharmacy
<b>Action title</b>	December 2022 scheduling of chemicals in Schedule I
<b>Final agency action date</b>	December 6, 2022
<b>Date this document prepared</b>	December 6, 2022

This information is required for executive branch review pursuant to Executive Order 19 (2022) (EO 19), any instructions or procedures issued by the Office of Regulatory Management (ORM) or the Department of Planning and Budget (DPB) pursuant to EO 19. In addition, this information is required by the Virginia Registrar of Regulations pursuant to the Virginia Register Act (§ 2.2-4100 et seq. of the Code of Virginia). Regulations must conform to the Regulations for Filing and Publishing Agency Regulations (1 VAC 7-10), and the *Form and Style Requirements for the Virginia Register of Regulations and Virginia Administrative Code*.

### Brief Summary

*Provide a brief summary (preferably no more than 2 or 3 paragraphs) of this regulatory change (i.e., new regulation, amendments to an existing regulation, or repeal of an existing regulation). Alert the reader to all substantive matters. If applicable, generally describe the existing regulation.*

As specified in Va. Code § 54.1-3443, the Virginia Department of Forensic Science (“DFS”) has identified five (5) compounds which it recommended the Board of Pharmacy place into Schedule I in the Virginia Administrative Code. The placement by regulatory action will remain in effect for 18 months or until the compounds are placed in Schedule I by legislative action of the General Assembly. This action is exempt in accordance with Va. Code § 2.2-4006 of the Virginia Administrative Process Act.

### Mandate and Impetus

*Identify the mandate for this regulatory change and any other impetus that specifically prompted its initiation (e.g., new or modified mandate, internal staff review, petition for rulemaking, periodic review, or board decision). For purposes of executive branch review, “mandate” has the same meaning as defined*

*in the ORM procedures, “a directive from the General Assembly, the federal government, or a court that requires that a regulation be promulgated, amended, or repealed in whole or part.”*

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The impetus of this action is identification by DFS of five compounds which DFS recommends be included in Schedule I under the Virginia Administrative Code pursuant to Va. Code § 54.1-3443(D).

**Statement of Final Agency Action**

*Provide a statement of the final action taken by the agency including: 1) the date the action was taken; 2) the name of the agency taking the action; and 3) the title of the regulation.*

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On December 6, 2022, the Board of Pharmacy amended 18VAC110-20-322 of the Regulations Governing the Practice of Pharmacy to place chemicals specified by DFS into Schedule I in accordance with Va. Code § 54.1-3443(D).