## ADMINISTRATIVE RULES Fiscal Estimate & Economic Impact Analysis

1. Type of Estimate and Analysis	2. Date	
Original Updated Corrected	3/28/24	
3. Administrative Rule Chapter, Title and Number (and Clearinghouse Number if applicable) $CSB\ 4$		
4. Subject Monitored Precription Drug History Reports		
5. Fund Sources Affected □ GPR □ FED ⊠ PRO □ PRS □ SEG □ SEG-S	6. Chapter 20, Stats. Appropriations Affected s. 20.165 (1) (hg)	
7. Fiscal Effect of Implementing the Rule         Image: No Fiscal Effect       Increase Existing Revenues         Indeterminate       Image: Decrease Existing Revenues	☐ Increase Costs ☐ Decrease Costs ☐ Could Absorb Within Agency's Budget	
Local Government Units     Pub	cific Businesses/Sectors lic Utility Rate Payers	
Small Businesses (if checked, complete Attachment A) 9. Estimate of Implementation and Compliance to Businesses, Local Governmental Units and Individuals, pers. 227.137(3)(b)(1). \$0		
<ul> <li>10. Would Implementation and Compliance Costs Businesses, Local Governmental Units and Individuals Be \$10 Million or more Over Any 2-year Period, pers. 227.137(3)(b)(2)?</li> <li>Yes X No</li> </ul>		
11. Policy Problem Addressed by the Rule The objective of the proposed rule is to allow an authorized patient representative to request monitored prescription drug history reports on behalf of a patient both in person and via mail.		
12. Summary of the Businesses, Business Sectors, Associations Representing Business, Local Governmental Units, and Individuals		
that may be Affected by the Proposed Rule that were Contacted for Comments. The rule was posted on the Department of Safety and Professional Service's (DSPS) website for 14 days to solicit public comment on economic impact, including how the proposed rules may affect businesses, local government units, and individuals. No comments were received.		
13. Identify the Local Governmental Units that Participated in the Development of this EIA. None.		
14. Summaryof Rule's Economic and Fiscal Impact on Specific Businesses, Business Sectors, Public Utility Rate Payers, Local Governmental Units and the State's Economyas a Whole (Include Implementation and Compliance Costs Expected to be Incurred)		
DSPS estimates a total of \$3,200 in one-time costs for implementing the provisions of this rule to support the equivalent of a 0.1 limited term employee for activities including rulemaking and coordination with PDMP program staff. The one-time costs cannot be absorbed in the currently appropriated agency budget.		
15. Benefits of Implementing the Rule and Alternative(s) to Implementing the Rule The benefit of implementing the rule is that patients authorized representatives will have the option to request monitored presciption drug history reports via mail, in addition to in-person requests.		
16. Long Range Implications of Implementing the Rule The long range implications of implementing this rule is greater patient and customer satisfaction with the Prescroption Drug Monitoring Program (PDMP) through increased accessability to monitored prescription drug history reports where allowed by law. 17. Compare With Approaches Being Used by Federal Government		
None.		

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18. Compare With Approaches Being Used by Neighboring States (Illinois, Iowa, Michigan and Minnesota) Illinois: The Illinois Prescription Monitoring Program allows patients access to their personal prescription history based on a validation process established by administrative rules [720 Illinois Compiled Statutes Chapter 570 Section 318 (m)]. The administrative rules governing patient access to their prescription history require that the patient, parent, or guardian complete a notarized request for a personal information report of a patient's prescription history, and submit it by mail to the Illinois Prescription Monitoring Program [Illinois Administrative Coder Title 77 Chapter X Subchapter e Part 2050 Section 2080.190 (a)].

Iowa: The Iowa Prescription Monitoring Program allows patients or a patient's agent to request that individual patient's own prescription history report by submitting a request form. Request forms may be submitted in-person with a government issued photo identification or via mail if the request form is notarized and sent with a certified copy of the patient's government issued identification. A patient's agent may sign the request form in lieu of the patient if a copy the legal document establishing the agency relationship is provided. The patient's agent must also present a government issued identification for in-person requests or a certified copy of a government issued identification for mailed requests. [657 Iowa Administrative Code Chapter 37 Section 37.16 (7)].

Michigan: The administrative rules that govern the Michigan Automated Prescription System, the states electronic system for monitoring schedule II to V controlled substances, does not specify whether a report of a patient's prescription history can be disclosed, nor how a report may be obtained by a patient. [Michigan Administrative Rules R 338.3162b].

Minnesota: The Minnesota Prescription Monitoring Program allows a patient who has been prescribed a controlled substance to access the program's database to obtain information on users who have access to that patient's data records. A patient may submit a request for this information on a notarized form from the Minnesota State Board of Pharmacy's website.[Minnesota Statutes Chapter 152 Section 152.126 Subdivision 11].

19. Contact Name	20. Contact Phone Number
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## ATTACHMENT A

1. Summaryof Rule's Economic and Fiscal Impact on Small Businesses (Separatelyfor each Small Business Sector, Include Implementation and Compliance Costs Expected to be Incurred)

2. Summary of the data sources used to measure the Rule's impact on Small Businesses

3. Did the agency consider the following methods to reduce the impact of the Rule on Small Businesses?

Less Stringent Compliance or Reporting Requirements

Less Stringent Schedules or Deadlines for Compliance or Reporting

Consolidation or Simplification of Reporting Requirements

Establishment of performance standards in lieu of Design or Operational Standards

Exemption of Small Businesses from some or all requirements

Other, describe:

4. Describe the methods incorporated into the Rule that will reduce its impact on Small Businesses

5. Describe the Rule's Enforcement Provisions

6. Did the Agency prepare a Cost Benefit Analysis (if Yes, attach to form)

🗆 Yes 🛛 No