

<h1>Regulatory Analysis Form</h1> <p>(Completed by Promulgating Agency)</p> <p>(All Comments submitted on this regulation will appear on IRRC's website)</p>		<p>INDEPENDENT REGULATORY REVIEW COMMISSION</p> <p>RECEIVED</p> <p>Independent Regulatory Review Commission November 13, 2025</p>	
<p>(1) Agency Department of Human Services</p>		<p>IRRC Number: 3466</p>	
<p>(2) Agency Number: 14 Identification Number: 558</p>			
<p>(3) PA Code Cite: 55 Pa. Code Chapter 5100</p>			
<p>(4) Short Title: Mental Health Procedures</p>			
<p>(5) Agency Contacts (List Telephone Number and Email Address): Primary Contact: Barry Decker (717) 772-7640, bdecker@pa.gov Secondary Contact: Tara Pride (717) 346-8116, tpride@pa.gov</p>			
<p>(6) Type of Rulemaking (check applicable box):</p> <p><input checked="" type="checkbox"/> Proposed Regulation <input type="checkbox"/> Final Regulation <input type="checkbox"/> Final Omitted Regulation</p>		<p><input type="checkbox"/> Emergency Certification Regulation; <input type="checkbox"/> Certification by the Governor <input type="checkbox"/> Certification by the Attorney General</p>	
<p>(7) Briefly explain the regulation in clear and nontechnical language. (100 words or less)</p> <p>The purpose of this proposed rulemaking is to amend Chapter 5100 to align with Act 32 of 2022 and Health Insurance Portability and Accountability Act of 1996 (HIPAA) requirements. Act 32 of 2022 requires the Department of Human Services (Department) to promulgate regulations reflecting changes to the Mental Health Procedures Act, which include the addition of covered entities and business associates to the list of entities to which the Department may disclose confidential information.</p>			
<p>(8) State the statutory authority for the regulation. Include <u>specific</u> statutory citation.</p> <p>The Department has authority to promulgate this regulation under section 112 of the Mental Health Procedures Act (50 P. S. § 7112) (act) and section 201(2) of the Mental Health and Intellectual Disabilities Act of 1966 (50 P. S. § 4201(2)). In addition, section 3 of the act of July 7, 2022 (P.L. 428, No. 32) (Act 32) requires the Department to promulgate regulations for the purpose of implementing the amendment of section 111(a) of the act regarding access to confidential documents of persons in treatment for a covered entity or a covered entity's business associate.</p>			

(9) Is the regulation mandated by any federal or state law or court order, or federal regulation? Are there any relevant state or federal court decisions? If yes, cite the specific law, case or regulation as well as, any deadlines for action.

Yes, Act 32 of 2022 requires the Department to promulgate regulation in order to implement the amendments to the act. Further, to conform with HIPAA, the Department is proposing changes under this chapter.

(10) State why the regulation is needed. Explain the compelling public interest that justifies the regulation. Describe who will benefit from the regulation. Quantify the benefits as completely as possible and approximate the number of people who will benefit.

The proposed revisions to Chapter 5100 are needed to align with the requirement of Act 32 of 2022. Additionally, the proposed regulation aligns with the confidentiality requirements under HIPAA. This will benefit over 720,000 individuals receiving Department funded behavioral health services in Pennsylvania. This proposed rulemaking will also benefit at least 1,732 providers and 5,655 satellite sites in following a standard process for confidentiality.

(11) Are there any provisions that are more stringent than federal standards? If yes, identify the specific provisions and the compelling Pennsylvania interest that demands stronger regulations.

No. The proposed revisions to Chapter 5100 are needed to align with the requirement of Act 32 of 2022 and are being made to be consistent with Federal HIPAA requirements.

(12) How does this regulation compare with those of the other states? How will this affect Pennsylvania's ability to compete with other states?

The proposed rulemaking provides consistency with Federal HIPAA requirements. As such, the proposed rulemaking will not affect Pennsylvania's ability to compete with other states.

(13) Will the regulation affect any other regulations of the promulgating agency or other state agencies? If yes, explain and provide specific citations.

No, the proposed rulemaking will not affect any other state regulations.

(14) Describe the communications with and solicitation of input from the public, any advisory council/group, small businesses and groups representing small businesses in the development and drafting of the regulation. List the specific persons and/or groups who were involved. ("Small business" is defined in Section 3 of the Regulatory Review Act, Act 76 of 2012.)

In response to the Act 32 of 2022, the Department issued bulletin OMHSAS-23-05 titled, "Confidentiality of Records Changes Due to Act 32 of 2022 and Aligning with Health Insurance Portability and Accountability Act of 1996" on May 12, 2023. Under the bulletin, the Department informed stakeholders of Act 32 of 2022 changes to the list of permissible disclosures in 50 P.S. § 7111(a), and new definitions in 50 P.S. § 7103.1 and announced that the Department would be amending 55 Pa. Code Chapter 5100 to align with Act 32 and federal HIPAA requirements. Additionally, OMHSAS informed stakeholders of a pending OMHSAS bulletin and changes to the Chapter 5100

regulations during a stakeholder meeting on March 21, 2023 that was open to the public and advertised via the OMHSAS public listserv. Link to [Bulletin OMHSAS-23-5_May2023.pdf \(pa.gov\)](#).

(15) Identify the types and number of persons, businesses, small businesses (as defined in Section 3 of the Regulatory Review Act, Act 76 of 2012) and organizations which will be affected by the regulation. How are they affected?

This regulation will benefit over 720,000 individuals who receive Department-funded behavioral health services in Pennsylvania. This regulation will also benefit at least 1,732 providers and 5,655 satellite sites in following a standard process for confidentiality. The Department does not have access to information on all revenue generated by all licensed providers and is therefore unable to determine how many providers or licensed locations would qualify as small businesses. The revision to Chapter 5100 is needed to align with the requirements of Act 32 of 2022. Additionally, the proposed regulation aligns with the confidentiality requirements under HIPAA, allowing for consistency.

(16) List the persons, groups or entities, including small businesses that will be required to comply with the regulation. Approximate the number that will be required to comply.

All 1,732 providers and 5,655 licensed locations of behavioral health services will need to comply with this regulation.

(17) Identify the financial, economic and social impact of the regulation on individuals, small businesses, businesses and labor communities and other public and private organizations. Evaluate the benefits expected as a result of the regulation.

The impact of the regulations on individuals and providers will allow for the alignment of regulations with Federal HIPAA requirements and Act 32 of 2022. This will allow for greater access of information for individuals to a covered entity or business associate.

(18) Explain how the benefits of the regulation outweigh any cost and adverse effects.

The revision to Chapter 5100 is needed to align with the requirement of Act 32 of 2022. Additionally, the proposed regulation aligns with the confidentiality requirements under HIPAA. This will benefit individuals who receive behavioral health services in Pennsylvania for greater access to their own information and proposes a standard confidentiality process for providers.

(19) Provide a specific estimate of the costs and/or savings to the **regulated community** associated with compliance, including any legal, accounting or consulting procedures which may be required. Explain how the dollar estimates were derived.

There are no costs or savings anticipated to the regulated community.

(20) Provide a specific estimate of the costs and/or savings to the **local governments** associated with compliance, including any legal, accounting or consulting procedures which may be required. Explain how the dollar estimates were derived.

There are no anticipated costs or savings to the local governments.

(21) Provide a specific estimate of the costs and/or savings to the **state government** associated with the implementation of the regulation, including any legal, accounting, or consulting procedures which may be required. Explain how the dollar estimates were derived.

There are no anticipated costs or savings to the state government.

(22) For each of the groups and entities identified in items (19)-(21) above, submit a statement of legal, accounting or consulting procedures and additional reporting, recordkeeping or other paperwork, including copies of forms or reports, which will be required for implementation of the regulation and an explanation of measures which have been taken to minimize these requirements.

There are no legal, accounting or consulting procedures or additional reporting, recordkeeping or other paperwork which will be required for implementation of the regulation.

(22a) Are forms required for implementation of the regulation?

No new forms are required for implementation.

(22b) If forms are required for implementation of the regulation, **attach copies of the forms here**. If your agency uses electronic forms, provide links to each form or a detailed description of the information required to be reported. **Failure to attach forms, provide links, or provide a detailed description of the information to be reported will constitute a faulty delivery of the regulation.**

No new forms are required for implementation.

Gloria Gilligan

01/02/2025

(23) In the table below, provide an estimate of the fiscal savings and costs associated with implementation and compliance for the regulated community, local government, and state government for the current year and five subsequent years.

	Current FY Year 2024-2025	FY +1 Year 2025-2026	FY +2 Year 2026-2027	FY +3 Year 2027-2028	FY +4 Year 2028-2029	FY +5 Year 2029-2030
SAVINGS:	\$	\$	\$	\$	\$	\$
Regulated Community	0	0	0	0	0	0
Local Government	0	0	0	0	0	0
State Government	0	0	0	0	0	0
Total Savings	0	0	0	0	0	0
COSTS:						
Regulated Community	\$0	\$0	\$0	\$0	\$0	\$0
Local Government	0	0	0	0	0	0
State Government	0	0	0	0	0	0
Total Costs	0	0	0	0	0	0
REVENUE LOSSES:						
Regulated Community	0	0	0	0	0	0
Local Government	0	0	0	0	0	0
State Government	\$0	\$0	\$0	\$0	\$0	\$0
Total Revenue Losses	0	0	0	0	0	0

(23a) Provide the past three-year expenditure history for programs affected by the regulation.

Program	FY -3 2020-2021	FY -2 2021-2022	FY -1 2022-2023	Current FY 2023-2024
Mental Health	\$ 822,470,000	\$ 866,093,000	\$ 855,567,000	\$ 956,535,000
MA Capitation	\$4,557,295,000	\$3,418,498,000	\$3,594,065,000	\$3,945,519,000
MA Fee-for-Service	\$644,059,000	\$589,137,000	\$697,345,000	\$686,639,000

(24) For any regulation that may have an adverse impact on small businesses (as defined in Section 3 of the Regulatory Review Act, Act 76 of 2012), provide an economic impact statement that includes the following:

- (a) An identification and estimate of the number of small businesses subject to the regulation.

RESPONSE: The Department does not have access to information on all revenue generated by all licensed providers and is therefore unable to determine how many providers or licensed locations would qualify as small businesses. All 1,732 providers and 5,655 licensed locations of behavioral health services will need to comply with this regulation, which proposes to codify the statutory changes under Act 32 and Federal HIPAA requirements.

- (b) The projected reporting, recordkeeping and other administrative costs required for compliance with the proposed regulation, including the type of professional skills necessary for preparation of the report or record.

RESPONSE: There are minimal to no additional costs anticipated.

- (c) A statement of probable effect on impacted small businesses.

RESPONSE: Because this proposed rulemaking codifies existing requirements under Act 32 of 2022 and Federal HIPAA requirements, there is no impact on small businesses.

- (d) A description of any less intrusive or less costly alternative methods of achieving the purpose of the proposed regulation.

RESPONSE: Because this proposed rulemaking codifies existing requirements under Act 32 of 2022 and Federal HIPAA requirements, there are no other methods available for achieving the purpose of the proposed regulation.

(25) List any special provisions which have been developed to meet the particular needs of affected groups or persons including, but not limited to, minorities, the elderly, small businesses, and farmers.

This regulation relates to the confidentiality of records in compliance with Federal and State law. No special provisions are included for specific populations, such as minorities, the elderly, small businesses or farmers.

(26) Include a description of any alternative regulatory provisions which have been considered and rejected and a statement that the least burdensome acceptable alternative has been selected.

To be consistent with Act 32 of 2022 and Federal HIPAA requirements, no other regulatory provisions were considered given the limited nature and anticipated impact of this proposed regulation.

(27) In conducting a regulatory flexibility analysis, explain whether regulatory methods were considered that will minimize any adverse impact on small businesses (as defined in Section 3 of the Regulatory Review Act, Act 76 of 2012), including:

- a) The establishment of less stringent compliance or reporting requirements for small businesses;

RESPONSE: The revision to Chapter 5100 is needed to align with the requirement of Act 32 of 2022. Additionally, the proposed regulation aligns with the confidentiality requirements under HIPAA. As such, there are not less stringent compliance or reporting requirements for small businesses.

- b) The establishment of less stringent schedules or deadlines for compliance or reporting requirements for small businesses;

RESPONSE: The revision to Chapter 5100 is not anticipated to require hardship on small businesses as the revisions are needed to align with the requirement of Act 32 of 2022 and Federal HIPAA requirements.

- c) The consolidation or simplification of compliance or reporting requirements for small businesses;

RESPONSE: The revision to Chapter 5100 is needed to align with the requirement of Act 32 of 2022. Additionally, the proposed regulation aligns with the confidentiality requirements under HIPAA.

- d) The establishment of performance standards for small businesses to replace design or operational standards required in the regulation; and

RESPONSE: There are no performance standards established since the revisions to the regulations are needed to align with the requirement of Act 32 of 2022 and to align with the confidentiality requirements under HIPAA.

- e) The exemption of small businesses from all or any part of the requirements contained in the regulation.

RESPONSE: Small businesses are not exempt from the requirements of the proposed regulation since the revisions to Chapter 5100 are needed to align with the statutory requirement of Act 32 of 2022. Additionally, the proposed regulation aligns with the confidentiality requirements under HIPAA.

(28) If data is the basis for this regulation, please provide a description of the data, explain in detail how the data was obtained, and how it meets the acceptability standard for empirical, replicable and testable data that is supported by documentation, statistics, reports, studies or research. Please submit data or supporting materials with the regulatory package. If the material exceeds 50 pages, please provide it in a searchable electronic format or provide a list of citations and internet links that, where possible, can be accessed in a searchable format in lieu of the actual material. If other data was considered but not used, please explain why that data was determined not to be acceptable.

No data is the basis for this regulation.

(29) Include a schedule for review of the regulation including:

- A. The length of the public comment period: 30 days.
- B. The date or dates on which any public meetings or hearings will be held: No public meetings or hearings will be held.
- C. The expected date of delivery of the final-form regulation: 4th Quarter 2026.
- D. The expected effective date of the final-form regulation: Upon publication of the final-form regulation in the Pennsylvania Bulletin.
- E. The expected date by which compliance with the final-form regulation will be required: Upon publication of the final-form regulation in the Pennsylvania Bulletin.
- F. The expected date by which required permits, licenses or other approvals must be obtained: There are no required permits, licenses or other approvals needed for this regulation.

(30) Describe the plan developed for evaluating the continuing effectiveness of the regulations after its implementation.

The Department will review the regulations on an ongoing basis to ensure compliance with Federal and State law and to assess the appropriateness and effectiveness of the regulation.

RECEIVED

Independent Regulatory
Review Commission

November 13, 2025

CDL-1

**FACE SHEET
FOR FILING DOCUMENTS
WITH THE LEGISLATIVE REFERENCE BUREAU**

(Pursuant to Commonwealth Documents Law)

DO NOT WRITE IN THIS SPACE

Copy below is hereby approved
as to form and legality.
Attorney General

By: Ay M Elliott
(Deputy Attorney General)
8/29/25
Date of Approval

☐ Check if applicable
Copy not approved.
Objections attached.

Copy below is hereby certified to be a true and correct
copy of a document issued, prescribed or promulgated
by:

DEPARTMENT OF HUMAN SERVICES
(Agency)

LEGAL COUNSEL: [Signature]

DOCUMENT/FISCAL NOTE NO. 14-558

DATE OF ADOPTION: _____

BY: [Signature]

TITLE: SECRETARY OF HUMAN SERVICES
(Executive Officer, Chairman or Secretary)

Copy below is hereby approved as to
form and legality. Executive or
Independent Agencies.

BY: Cynthia K. Montgomery
Digitally signed by Cynthia K. Montgomery
DN: cn=Cynthia K. Montgomery, o=PA, email=cymontgomcspa.gov, c=US
Date: 2025.08.06 07:54:59 -04'00'

8/6/2025

Date of Approval

(Deputy General Counsel)
(Chief Counsel, Independent Agency)
(Strike inapplicable title)

☐ Check if applicable. No Attorney
General approval or objection
within 30 days after submission.

NOTICE OF PROPOSED RULEMAKING

DEPARTMENT OF HUMAN SERVICES

OFFICE OF MENTAL HEALTH AND SUBSTANCE ABUSE SERVICES

[55 Pa. Code Chapter 5100]

Mental Health Procedures

Statutory Authority

Notice is hereby given that the Department of Human Services (Department) under the authority of section 112 of the Mental Health Procedures Act (50 P. S. § 7112) (act) and section 201(2) of the Mental Health and Intellectual Disability Act of 1966 (50 P. S. § 4201(2)), intends to amend Title 55, Chapter 5100 (relating to mental health procedures), to read as set forth in Annex A.

Section 201(2) of the Mental Health and Intellectual Disability Act of 1966 (50 P.S. § 4201(2)) requires the Department to consult with the Advisory Committee for Mental Health and Intellectual Disability when developing regulations authorized under the Mental Health and Intellectual Disability Act. Because the Advisory Committee is no longer active, the Department instead presented this proposed rulemaking to the Mental Health Planning Council. The Mental Health Planning Council advises the Department on issues related to mental health, substance abuse, behavioral health disorders and cross-system disability.

In addition, section 3 of the act of July 7, 2022 (P.L. 428, No. 32) (Act 32) requires the Department to promulgate regulations for the purpose of implementing the amendment of section 111(a) of the act regarding access to confidential documents of persons in treatment for a covered entity or a covered entity's business associate.

Purpose of Regulation

The purpose of this proposed rulemaking is to amend Chapter 5100 to align with the amendments to the act made by Act 32 and requirements of the Health Insurance Portability and Accountability Act of 1996 (Pub. L. No. 104-191, 110 Stat. 1936) (HIPAA). Act 32 requires the Department to promulgate regulations reflecting changes made by Act 32 to the act, which include: (1) the new definitions of "business associate" and "covered entity"; and (2) the addition

of covered entities and their business associates to the list of entities to which the Department may disclose confidential information.

Background

On July 7, 2022, Act 32 was signed into law, amending the act by updating definitions and confidentiality requirements. As noted previously, Act 32 also required the Department to promulgate regulations in accordance with the act.

On May 12, 2023, the Department issued bulletin OMHSAS-23-05, titled “Confidentiality of Records Changes Due to Act 32 of 2022 and Aligning with Health Insurance Portability and Accountability Act of 1996” to inform stakeholders of changes to the list of permissible disclosures under section 111(a) of the act (50 P.S. § 7111(a)), and new definitions under section 103.1 of the act (50 P.S. § 7103.1). Additionally, the bulletin announced that the Department would amend Chapter 5100 to align with Act 32 and HIPAA requirements. See: https://www.dhs.pa.gov/docs/Publications/Documents/FORMS%20AND%20PUBS%20OMHSAS/Bulletin%20OMHSAS-23-5_May2023.pdf

Requirements

The following is a summary of the specific substantive provisions in the proposed regulation.

Scope and policy (§ 5100.31). The proposed rulemaking aligns requirements in the chapter with Federal law. Specifically, the proposed rulemaking updates provisions under subsection (b) to specify either the required or permitted disclosure of information, which aligns with HIPAA requirements. The proposed deletion of the second sentence in subsection (f) affords individuals greater right to access their own information. Subsections (g) and (h) are

proposed to be removed because they are not in compliance with HIPAA requirements regarding release of protected health information. Also, subsection (j) is added to specify that to the extent there are any conflicts, Federal law will control.

Nonconsensual release of information (§ 5100.32). The proposed rulemaking adds a new subsection (a.1) to provide that records may be released to a covered entity or a covered entity's business associate in accordance with Federal regulations under 45 CFR Part 164, Subpart E (relating to privacy of individually identifiable health information). This proposed provision fulfills the revisions identified in Act 32 of 2022. Additionally, HIPAA and Act 32 requirements apply to disclosures to health care providers for treatment purposes and disclosures to third party payors. Subsection (a)(2) is proposed to be removed because its provisions are not consistent with the requirements of Act 32 and section 111 of the act. The proposed rulemaking clarifies information exchanged with those other than the provider shall be limited to the relevant information that is requested. The request for information and the action that was taken shall be recorded in the patient's records.

Patient's access to records and control over release of records (§ 5100.33). The proposed rulemaking directs that an individual's access to and control over the disclosure of the patient's record and information be in accordance with Federal regulations under 45 CFR Parts 160 and 164 (relating to general administrative requirements; and security and privacy). Deletions of subsections (a) through (e) are being proposed because they are contrary to HIPAA requirements concerning a patient's right to access their own records and information; therefore, a new subsection (e.1) is being added to reflect that a patient's access to and control over disclosure of the patient's record and information shall be in accordance with 45 CFR Parts 160 and 164. Subsection (h) is being revised to identify the new criminal rule under Pa.R.Crim. P.

No. 703 that governs pre-sentence reports. Additionally, the first sentence of subsection (j) is proposed to be deleted because it is contrary to HIPAA requirements due to its reference to the exception in (c), which also is proposed to be deleted.

Consensual release to third parties (§ 5100.34) (proposed amendment to authorized release to third parties). In accordance with HIPAA, the proposed rulemaking permits access to records upon written authorization in accordance with 45 CFR Part 164. These provisions are being revised in subsection (a). The last three sentences in subsection (a) are proposed to be deleted as they have been addressed and are not necessary with the provision being added to subsection (a). The proposed rulemaking also proposes to delete subsections (b) and (c) because they differ from Act 32 and HIPAA regarding third party access and access to one's own records. Subsection (d) and (e) are also proposed to be deleted because the statement in subsection (d) is no longer required. In addition, the Department proposes to revise the language provisions for voluntary disclosures. Under the proposed rulemaking, a facility shall use an authorization form for voluntary disclosures that complies with HIPAA, and other applicable Federal and State requirements. These provisions are addressed in subsection (f). Specifically, subsection (f) requires a facility to use an authorization form that complies with HIPAA, other applicable laws, and also contains a place for the signature of a staff person who witnesses the consent and a place to record a verbal authorization, if applicable. The remaining paragraphs are proposed to be deleted since they are specifically addressed under HIPAA, and they are not completely consistent with HIPAA requirements. For example, HIPAA does not require certain specificity regarding the purpose or type of information. To maintain consistency with federal requirements regarding voluntary disclosures that require written authorization, the proposed rulemaking cites HIPAA and its corresponding federal requirements and state law.

Lastly, the department proposes to amend to the section title to “authorized release to third parties” to accurately reflect the proposed amendments in the section.

Records relating to drug and alcohol abuse or dependence (§ 5100.37). The proposed rulemaking aligns with the act of July 7, 2022 (P.L. 430, No. 33) (Act 33 of 2022), which amended the Pennsylvania Drug and Alcohol Abuse Control Act (71 P.S. §§ 1690.101—1690.115) . As such, the Department proposes to delete this cross-reference to 4 Pa. Code § 255.5 (relating to projects and coordinating bodies; disclosure of client-oriented information) due to the controlling nature of Act 33 of 2022.

Notice of withdrawal (§ 5100.76). The proposed rulemaking proposes to correct an error under subsection (b). Under the proposed rulemaking, the “...withdraw from involuntary treatment” is corrected to say “...withdraw from voluntary treatment”.

In addition to the proposed revisions discussed previously, the proposed rulemaking also proposes to make technical edits, correct citations and revise outdated terminology, such as the references to the Mental Health and Mental Retardation Act of 1966 and the Regional Commissioner of Mental Health. Under the proposed rulemaking, these are referred to as the Mental Health and Intellectual Disabilities Act of 1966 and a departmental designee, respectively. The proposed rulemaking also updates the reference to the Professional Standards Review Organization and the citations to the Child Protective Services Law and the Human Services Code.

Affected Individuals and Organizations

The proposed rulemaking will benefit over 720,000 individuals receiving Department - funded behavioral health services in Pennsylvania. This proposed rulemaking will also benefit at least 1,732 providers and 5,655 satellite sites by following a standard process for confidentiality.

Accomplishments and Benefits

Revisions to Chapter 5100 are needed to align with the requirement of Act 32 of 2022. The proposed rulemaking will also align with HIPAA, providing for consistency. These proposed amendments will allow for greater access for individuals to their own information.

Fiscal Impact

No cost to the Commonwealth, local government, service providers or individuals is anticipated as a result of this proposed rulemaking.

Paperwork Requirements

There are no legal, accounting or consulting procedures or additional reporting, recordkeeping or other paperwork required to comply with this proposed rulemaking. The proposed rulemaking codifies existing requirements under Act 32 of 2022 and Federal HIPAA requirements. Under the proposed rulemaking, the written authorization shall comply with HIPAA and other Federal and State requirements.

Effective Date

The proposed regulation will be effective upon final publication in the *Pennsylvania Bulletin*.

Public Comment

Interested persons are invited to submit written comments, suggestions or objections regarding the proposed rulemaking to the Department at the following address: Barry Decker, Office of Mental Health and Substance Abuse Services (OMHSAS), Commonwealth Tower, 303 Walnut Street, 11th Floor, Harrisburg, PA. 17105 and RA-PWMHPROCREGS@pa.gov, within 30 calendar days after the date of publication of this proposed rulemaking in the *Pennsylvania Bulletin*. Reference **Regulation No. 14-558** when submitting comments.

Persons with a disability who require an auxiliary aid or service may submit comments by using the Pennsylvania Hamilton Relay Service at 1-800-654-5984 (TDD users) or 1-800-654-5988 (voice users).

Regulatory Review Act

Under § 5(a) of the Regulatory Review Act (71 P.S. § 745.5(a)), on November 13, 2025 the Department submitted a copy of this proposed rulemaking and a copy of a Regulatory Analysis Form to the Independent Regulatory Review Commission (IRRC) and to the Chairpersons of the House Committee on Human Services and the Senate Committee on Health and Human Services. A copy of this material is available to the public upon request.

Under § 5(g) of the Regulatory Review Act, if the IRRC has any comments, recommendations or objections to any portion of the proposed regulation, it may notify the Department and the Committees within 30 days after the close of the public comment period. Such notification shall specify the regulatory review criteria that have not been met. The Regulatory Review Act specifies detailed procedures for review by the Department, the General

Assembly and the Governor, of any comments, recommendations or objections raised, prior to final publication of the regulation.

ANNEX A

TITLE 55. HUMAN SERVICES

PART VII. MENTAL HEALTH MANUAL

CHAPTER 5100. MENTAL HEALTH PROCEDURES

GENERAL PROVISIONS

§ 5100.1. Legal base.

The legal base for this chapter is section 112 of the Mental Health Procedures Act (50 P. S. § 7112), section 201 of the Mental Health and [Mental Retardation] Intellectual Disability Act of 1966 (50 P. S. § 4201), and section 1021 of the [Public Welfare] Human Services Code (62 P. S. § 1021).

§ 5100.2. Definitions.

The following words and terms, when used in this chapter, shall have the following meanings, unless the context clearly indicates otherwise:

* * * * *

Administrator—The person appointed to carry out the duties specified in section 305 of the Mental Health and [Mental Retardation] Intellectual Disability Act of 1966 (50 P. S. § 4305).

* * * * *

Director of treatment team—A physician or licensed clinical psychologist designated by the facility director to assure that each patient receives treatment under the act and this chapter and that the facility's treatment responsibility to the patient, as defined in this chapter, the [Mental Health/Mental Retardation] Mental Health and Intellectual Disability Act of 1966 and the act, are discharged. The director of the treatment team is responsible for implementing and reviewing the

individualized treatment plan, for participating in the coordination of service delivery between other service providers, and for insuring that the unique skills and knowledge of each team member are utilized. The director of the treatment team is responsible for encouraging the person in treatment to become increasingly involved in decisions regarding the treatment planning process.

* * * * *

Mental Health and [Mental Retardation] Intellectual Disability Act of 1966—The act of October 20, 1966 (P. L. 96, No. 3) (50 P. S. §§ 4101—4704).

* * * * *

MENTAL HEALTH REVIEW OFFICER AND PROCEEDINGS

* * * * *

§ 5100.22. Consultation and education.

The administrator shall, in discharging his duties under the Mental Health and [Mental Retardation] Intellectual Disability Act of 1966, provide the court or mental health review officer with:

* * * * *

§ 5100.23. Written application, petitions, statements and certifications.

* * * * *

(f) Submission to county administrator:

* * * * *

(5) Mental health facilities shall file such statistical reports of activities and services required by the act and the Mental Health and [Mental Retardation] Intellectual Disability

Act of 1966 as the Department from time to time may require, so long as the data does not identify individual patients.

* * * * *

CONFIDENTIALITY OF MENTAL HEALTH RECORDS

§ 5100.31. Scope and policy.

* * * * *

(b) Persons seeking or receiving services from a mental health facility are entitled to do so with the expectation that information about them will be treated with respect and confidentiality by those providing services. Confidentiality between service providers [of services] and their [clients] patients is necessary to develop the trust and confidence important for therapeutic intervention. While full confidentiality cannot be guaranteed to everyone as a result of Federal and State [statutes] laws which require and permit disclosure of information for specific purposes, it remains incumbent upon service providers to inform each current [client/patient] patient of the specific limits upon confidentiality which affect [his] the patient's treatment when these limits become applicable. When facilities are required by Federal or State [statutes] law or by order of a court to release information regarding a discharged patient, a good faith effort shall be made to notify the person by certified mail to the last known address.

(c) As used in this chapter, “records” includes, but is not limited to, all written clinical information, observations and reports or fiscal documents, relating to a prospective, present or past [, client or] patient, which are required or authorized to be prepared by the act or by the Mental Health and [Mental Retardation] Intellectual Disability Act of 1966. This includes any central file of [client/patient] patient records and reports which are required to be maintained by

the Department's regulations or other [statutes and regulations] Federal and State law regarding service content for mental health programs. Every [therapist] service provider who reports objective findings must carefully consider the impact of placing in the records statements made privately in [therapy] clinical sessions.

* * * * *

(f) Records of a person receiving mental health services are the property of the [hospital or] mental health facility in which the person is receiving or has received services. [The person who is or was receiving services shall exercise control over the release of information contained in his record except as limited by § 5100.32 (relating to nonconsensual release of information), and be provided with access to the records except to the limitations under § 5100.33 (relating to patient's access to records and control over release of records).]

(g) [The presence or absence of a person currently involuntarily committed at a mental health facility is not to be considered a record within the meaning of subsection (c) and such information may be released at the discretion of the director of a facility in response to legitimate inquiries from governmental agencies or when it is clearly in the patient's best interest to do so] {Reserved}.

(h) [No document which was a public record prior to the person's treatment shall become confidential by its inclusion in the facility's records] {Reserved}.

(i) When information and observations regarding [clients or] patients are not made part of a record, there remains a duty and obligation for staff to respect the patient's privacy and confidentiality by acting ethically and responsibly in using or discussing such information.

(j) To the extent that any requirements in this chapter conflict with Federal law requiring or precluding disclosure of information, Federal law will control.

§ 5100.32. Nonconsensual release of information.

(a) Records concerning persons receiving or having received treatment shall be kept confidential and shall not be released nor their content disclosed without the [consent] written authorization of a person given under § 5100.34 (relating to consensual release to third parties), except that relevant portions or summaries may be released or copied as follows:

(1) To those actively engaged in treating the individual, or to persons at other facilities, including professional treatment staff of State Correctional Institutions and county prisons, when the person is being referred to that facility and a summary or portion of the record is necessary to provide for continuity of proper care and treatment.

(2) [To third party payors, both those operated and financed in whole or in part by any governmental agency and their agents or intermediaries, or those who are identified as payor or copayor for services and who require information to verify that services were actually provided. Information to be released without consent or court order under this subsection is limited to the staff names, the dates, types and costs of therapies or services, and a short description of the general purpose of each treatment session or service] {Reserved}.

(3) To reviewers and inspectors, including the Joint Commission on the Accreditation of Hospitals (JCAH) and Commonwealth licensure or certification, when necessary to obtain certification as an eligible provider of services.

(4) To those participating in [PRSO] Professional Standards Review Organization or Utilization Reviews.

(5) To the county administrator, under [his] the county administrator's duties under applicable [statutes and regulations] State law.

(6) To a court or mental health review officer, in the course of legal proceedings authorized by the act or this chapter.

(7) In response to a court order, when production of the documents is ordered by a court under § 5100.35(b) (relating to release to courts).

(8) To appropriate Departmental personnel under § 5100.38 (relating to child or patient abuse).

(9) In response to an emergency medical situation when release of information is necessary to prevent serious risk of bodily harm or death. Only specific information pertinent to the relief of the emergency may be released on a nonconsensual basis.

(10) To parents or guardians and others authorized to consent when necessary to obtain [consent] written authorization to medical treatment.

(11) To attorneys assigned to represent the subject of a commitment hearing.

(a.1) Records may also be released to a covered entity or a covered entity's business associate, as defined under 45 CFR 164.103, that makes the use, disclosure or request for disclosure in accordance with 45 CFR Part 164 Subpart E (relating to privacy of individually identifiable health information).

(b) Current patients [or clients] or the parents of patients under the age of 14 shall be notified of the specific conditions under which information may be released without their consent.

(c) [Information] Except for disclosures between those providing treatment to the patient, information made available under this section shall be limited to that information relevant and necessary to the purpose for which the information is sought. The information may not, without the patient's consent, be released to additional persons or entities, or used for additional

purposes. Requests for information and the action taken [should] shall be recorded in the patient's records.

§ 5100.33. Patient's access to records and control over release of records.

(a) [When a client/patient, 14 years of age or older, understands the nature of documents to be released and the purpose of releasing them, he shall control release of his records. For a client who lacks this understanding, any person chosen by the patient may exercise this right if found by the director to be acting in the patient's best interest. In the event that the client/patient is deceased, control over release of records may be exercised by the client's/patient's chosen executor, administrator or other personal representative of his estate, or, if there is no chosen personal representative, by a person otherwise empowered by court order to exercise control over the records. In the event that the client/patient is less than 14 years of age or has been adjudicated legally incompetent, control over release of the client's/patient's records may be exercised by a parent or guardian of the client/patient respectively] {Reserved}.

(b) [The term "access" when used in this section refers to physical examination of the record, but does not include nor imply physical possession of the records themselves or a copy thereof except as provided in this chapter] {Reserved}.

(c) [A person who has received or is receiving treatment may request access to his record, and shall be denied such access to limited portions of the record only:

(1) Upon documentation by the treatment team leader, it is determined by the director that disclosure of specific information concerning treatment will constitute a substantial detriment to the patient's treatment.

(2) When disclosure of specific information will reveal the identity of persons or breach the trust or confidentiality of persons who have provided information upon an agreement to maintain their confidentiality] {Reserved}.

(d) [A patient may obtain access to his records through the facility, or in the case of those records kept by the county administrator, through the physician or mental health professional designated by the administrator. Any third parties who are granted access to records may discuss this information with the patient only insofar as necessary to represent the patient in legal proceedings or other matters for which records have been released. Discussion of records with patients should be part of the therapeutic process and is not to be undertaken by other than mental health professionals] {Reserved}.

(e) [The limitations in subsection (c) are applicable to parents, guardians, and others who may control access over records as described in subsection (a) except that the possibility of substantial detriment to the parent, guardian, or other person may also be considered] {Reserved}.

(e.1) A patient's access to and control over disclosure of the record and information contained therein shall be in accordance with 45 CFR Parts 160 and 164 (relating to general administrative requirements; and security and privacy).

(f) If a person wishes to enter a written reaction qualifying or rebutting information in their records which they believe to be erroneous or misleading, they shall have the right to prepare such statement for inclusion as part of their record. The patient's written reaction shall accompany all released records.

(g) The director of the treatment team or the facility director may require that a mental health professional, who is a member of the treatment team, and who has reviewed the record in advance, be present when the patient or other person examines the record to aid in the

interpretation of documents in the record. If the records pertain to a former patient, an appropriate mental health professional may be designated by the facility director.

(h) Access to presentence reports, which may be part of the persons' records, is governed by Pa.R.Crim.P. No. [1404] 703 (relating to disclosure of pre-sentence reports), and the patient may have access to these records only upon order of the sentencing judge or as otherwise required by law. Any conditions of confidentiality imposed by the sentencing judge must be complied with. [Similarly] Similarly, parole and probation reports shall be released or access to them given only in accordance with 37 Pa. Code Part II (relating to Board of Probation and Parole) or as otherwise required by law.

(i) If a person is denied access to all or part of [his] the person's record, this fact and the basis for the denial shall be noted in the person's record.

(j) [When records or information have been forwarded from one agency to another agency, the receiving agency may not refuse the client or patient access to the records received except in accordance with subsection (c).] Records received from other agencies become part of the [client/patient's] patient's active record and are subject to the controls exercised over them by the [client,] patient [,] or those with authority over records as [defined in § 5100.31 (relating to scope and policy)] set forth in this chapter.

§ 5100.34. [Consensual] Authorized release to third parties.

(a) Access to records [, as defined in § 5100.33(b) (relating to patient's access to records and control over release of records) will] not authorized for release under § 5100.32 (relating to nonconsensual release of information) may be granted to persons other than the patient upon written [consent] authorization of the [client/patient] patient or the patient's personal

representative in accordance with 45 CFR Part 164 Subpart E (relating to privacy of individually identifiable health information). [With the consent, copies of excerpts or a summary of a record may be provided to specific persons at the discretion of the director. If copies of excerpts or summaries are provided, a charge may be made against the patient or person receiving the record for the cost of making the copies. The facility may require payment for the copies in advance.]

(b) [When a patient designates a third party as either a payor or copayor for mental health services, this designation carries with it his consent to release information to representatives of that payor which is necessary to establish reimbursement eligibility. Unless otherwise consented to by the patient, information released to the third-party payors shall be limited to that necessary to establish the claims for which reimbursement is sought] {Reserved}.

(c) [Clients, patients, or other persons consenting to release of records are to be informed of their right, subject to § 5100.33 to inspect material to be released] {Reserved}.

(d) [When records are released or disclosed under § 5100.32 (relating to nonconsensual release of information) or subsections (a) and (b) the written or oral disclosure shall be accompanied by a written statement which reads as follows:

“This information has been disclosed to you from records whose confidentiality is protected by State statute. State regulations limit your right to make any further disclosure of this information without prior written consent of the person to whom it pertains.”] {Reserved}.

(e) [The limitation in subsection (d) does not prohibit the re-release of information in accordance with § 5100.32] {Reserved}.

(f) [Each facility shall prepare a form for use in the voluntary release of records which shall meet the following requirements] For voluntary disclosures that require written authorization,

each facility shall use an authorization form that complies with the Health Insurance Portability and Accountability Act of 1996 (Public Law 104-191, 110 Stat. 1936) (HIPAA), other applicable Federal and State requirements, and contains, at a minimum, the following:

- (1) [A time limit on its validity which shows starting and ending dates] {Reserved}.
- (2) [Identification of the agency or person to whom the records are to be released] {Reserved}.
- (3) [A statement of the specific purposes for which the released records are to be used] {Reserved}.
- (4) [A statement identifying the specific relevant and timely information to be released] {Reserved}.
- (5) [A place for the signature of the client/patient or parent or guardian and the date, following a statement that the person understands the nature of his release] {Reserved}.
- (6) A place for the signature of a staff person [obtaining] who witnesses the consent of the [client/patient or parent or guardian] patient or the patient's personal representative in accordance with 45 CFR Part 164 Subpart E (relating to privacy of individually identifiable health information) and the date.
- (7) A place to record a verbal [consent] authorization to release of information given by a person physically unable to provide a signature and a place for the signatures of two responsible persons who witnessed that the person understood the nature of the release and freely gave [his] verbal [consent] authorization.
- (8) [Indication that the consent is revocable at the written request of the person giving consent, or oral request as in paragraph (7)] {Reserved}.
- (g) A mental health facility receiving a request for information from a governmental agency may accept that agency's release of information form if the form complies with HIPAA, other

applicable Federal and State requirements and is signed by the patient[/client] or [the person legally responsible for the control of information] the patient's personal representative unless the patient has specifically expressed opposition to that agency receiving information.

* * * * *

§ 5100.36. Departmental access to records and data collection.

(a) Notwithstanding any part of this chapter to the contrary, [employees] employees of the Department shall not be denied access to any patient records where such access is necessary and appropriate for the [employee's] employee's proper performance of [his] the employee's duties. The facility director or designee shall make such decision, and shall be responsible for limiting access to those portions which are relevant to the request.

(b) Any conflict as to access by an [employee] employee to patient records at State hospitals shall be resolved by [the Regional Commissioner of Mental Health] a departmental designee.

(c) Collection and analysis of clinical or statistical data by the Department, the administrator, or the facility for administrative or research purposes may be undertaken [as] so long as the report or paper prepared from the data does not identify any individual patient without [his consent] the individual patient's written authorization.

§ 5100.37. Records relating to drug and alcohol abuse or dependence.

Whenever information in a patient's records relates to drug or alcohol abuse or dependency, as defined in 71 P.S. § 1690.102, those specific portions of the patient's records are subject to the confidentiality provisions of section 8(c) of the Pennsylvania Drug and Alcohol Abuse Control

Act (71 P.S. § 1690.108(c))[, and the regulations promulgated thereunder, 4 Pa. Code § 255.5 (relating to projects and coordinating bodies: disclosure of client-oriented information)].

§ 5100.38. Child or patient abuse.

Nothing in this chapter shall conflict with the mandatory statutory or regulatory requirements of reporting suspected or discovered child abuse or patient abuse. Whenever a conflict exists between the reporting requirements of the Child Protective Services [Act (11 P.S. §§ 2201—2224),] Law (23 Pa. C.S. §§ 6301 – 6388) and the confidentiality of mental health records, the reporting requirements shall govern.

* * * * *

VOLUNTARY TREATMENT

* * * * *

§ 5100.76. Notice of withdrawal.

* * * * *

(b) The person receiving a signed Form MH 781-F from a patient shall immediately examine the patient's record to determine whether the patient has previously agreed to remain in treatment for a specified period not to exceed 72 hours after having given written notice of intent to withdraw from [involuntary] voluntary treatment. If no such consent has been given, the patient may immediately withdraw from treatment unless an application for emergency involuntary treatment is executed under section 302 of the act (50 P. S. § 7302), and the patient is advised accordingly.

* * * * *

PERSONS CHARGED WITH A CRIME OR UNDER SENTENCE

* * * * *

§ 5100.92. Voluntary examination and treatment of a person charged with a crime or serving a sentence.

* * * * *

(r) Liability for treatment of an individual admitted to a State mental health facility shall be assessed pursuant to section 505 of the [Mental Health/Mental Retardation] Mental Health and Intellectual Disability Act of 1966 (50 P. S. § 4505), and section 408 of the act (50 P. S. § 7408).

* * * * *



COMMONWEALTH OF PENNSYLVANIA
DEPARTMENT OF HUMAN SERVICES

November 13, 2025

Mr. David Sumner, Executive Director
Independent Regulatory Review Commission
555 Walnut Street, Suite 804
Harrisburg, Pennsylvania 17101

Dear Executive Director Sumner:

Enclosed is a proposed regulation that will align departmental regulations with Act 32 of 2022 and the Health Insurance Portability and Accountability Act of 1996 (HIPAA) requirements. Act 32 of 2022 required the Department of Human Services (Department) to promulgate regulations reflecting changes to the Mental Health Procedures Act, which include the new definitions of “business associate” and “covered entity” and the addition of covered entities and business associates to the list of entities to which the Department may disclose confidential information.

This proposed regulation, which amends the ***Pennsylvania Code***, Title 55, amends Chapter 5100 (relating to mental health procedures) and is submitted for review pursuant to the Regulatory Review Act.

The Department of Human Services will provide the Commission with any assistance required to facilitate a thorough review of this proposal.

Sincerely,

A handwritten signature in blue ink, appearing to read 'Val Arkoosh'.

Valerie A. Arkoosh, MD, MPH
Secretary

Enclosure

OFFICE OF THE SECRETARY

RECEIVED

From: [Annmarie Robey](#)
To: [Curley, Maeve](#)
Cc: [Stein, Marianne](#); [Fischer, Kendrick](#)
Subject: Re: [EXTERNAL]: DHS Proposed Regulation #14-558 Mental Health Procedures
Date: Thursday, November 13, 2025 9:51:17 AM
Attachments: [image001.jpg](#)

Independent Regulatory
Review Commission

November 13, 2025

Good morning:

The regulation has been received.

Thank you and have a good day. Annmarie

Annmarie K Robey | Executive Director/Legal Counsel (R)

Pennsylvania House of Representatives

Aging and Older Adult Services Committee

Human Services Committee

(p): 717-772-9842

From: Curley, Maeve <macurley@pa.gov>
Sent: Thursday, November 13, 2025 9:48 AM
To: Annmarie Robey <Arobey@pahousegop.com>
Cc: Stein, Marianne <maristein@pa.gov>; Fischer, Kendrick <kendfische@pa.gov>
Subject: [EXTERNAL]: DHS Proposed Regulation #14-558 Mental Health Procedures

Good morning,

DHS is submitting Reg. No. 14-558, Mental Health Procedures (Proposed Rulemaking) to the Senate Health and Human Services Committee and the House Human Services Committee.

Please provide written (email) confirmation that this rulemaking was received by the Committee chair's office.

Best,
Maeve



Maeve Curley
Pronouns: She/Her
Regulatory Coordinator

PA Department of Human Services | Office of Policy Development
macurley@pa.gov
<https://www.dhs.pa.gov>

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Independent Regulatory
Review Commission

November 13, 2025

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Review Commission

November 13, 2025

From: [Freeman, Clarissa](#)
To: [Curley, Maeve](#)
Cc: [Stein, Marianne](#); [Fischer, Kendrick](#)
Subject: RE: DHS Proposed Regulation #14-558 Mental Health Procedures
Date: Thursday, November 13, 2025 9:52:03 AM
Attachments: [image001.jpg](#)

Confirmed!

Thank you,

Clarissa L. Freeman
Deputy Chief Counsel | Senate Democratic Caucus
Executive Director-Health and Human Services Committee
717-783-1220

From: Curley, Maeve <macurley@pa.gov>
Sent: Thursday, November 13, 2025 9:49 AM
To: Freeman, Clarissa <clarissa.freeman@pasenate.com>
Cc: Stein, Marianne <maristein@pa.gov>; Fischer, Kendrick <kendfische@pa.gov>
Subject: DHS Proposed Regulation #14-558 Mental Health Procedures
Importance: High

EXTERNAL EMAIL

Good morning,

DHS is submitting Reg. No. 14-558, Mental Health Procedures (Proposed Rulemaking) to the Senate Health and Human Services Committee and the House Human Services Committee.

Please provide written (email) confirmation that this rulemaking was received by the Committee chair's office.

Best,
Maeve



Maeve Curley
Pronouns: She/Her
Regulatory Coordinator
PA Department of Human Services | Office of Policy Development
macurley@pa.gov
<https://www.dhs.pa.gov>

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From: [Wright, Imogen L.](#)
To: [Curley, Maeve](#)
Cc: [Stein, Marianne](#); [Fischer, Kendrick](#)
Subject: RE: DHS Proposed Regulation #14-558 Mental Health Procedures
Date: Thursday, November 13, 2025 9:55:46 AM
Attachments: [image001.jpg](#)

Independent Regulatory
Review Commission
November 13, 2025

Good morning Maeve,

This is to confirm receipt of the proposed rulemaking - Reg. No. 14-558.

Thank you,
Imogen

Imogen Wright | Executive Director

House Human Services Committee (D)

36 East Wing, Harrisburg PA

Office: (717) 705-1925 | Cell: (717) 317-2197

iwright@pahouse.net

From: Curley, Maeve <macurley@pa.gov>
Sent: Thursday, November 13, 2025 9:48 AM
To: Wright, Imogen L. <IWright@pahouse.net>
Cc: Stein, Marianne <maristein@pa.gov>; Fischer, Kendrick <kendfische@pa.gov>
Subject: DHS Proposed Regulation #14-558 Mental Health Procedures
Importance: High

Good morning,

DHS is submitting Reg. No. 14-558, Mental Health Procedures (Proposed Rulemaking) to the Senate Health and Human Services Committee and the House Human Services Committee.

Please provide written (email) confirmation that this rulemaking was received by the Committee chair's office.

Best,
Maeve



Maeve Curley

Pronouns: She/Her

Regulatory Coordinator

PA Department of Human Services | Office of Policy Development

macurley@pa.gov

<https://www.dhs.pa.gov>

From: [Burnett, David](#)
To: [Curley, Maeve](#)
Cc: [Stein, Marianne](#); [Fischer, Kendrick](#)
Subject: RE: DHS Proposed Regulation #14-558 Mental Health Procedures
Date: Thursday, November 13, 2025 9:56:04 AM
Attachments: [image001.jpg](#)

RECEIVED

Independent Regulatory
Review Commission
November 13, 2025

Good morning Maeve,

This email is to confirm receipt of the proposed regulation.

Regards,
-David

David Burnett

*Counsel and Executive Director
Senate Health & Human Services Committee
Harrisburg, PA 17120*

From: Curley, Maeve <macurley@pa.gov>
Sent: Thursday, November 13, 2025 9:49 AM
To: Burnett, David <dburnett@pasen.gov>
Cc: Stein, Marianne <maristein@pa.gov>; Fischer, Kendrick <kendfische@pa.gov>
Subject: DHS Proposed Regulation #14-558 Mental Health Procedures
Importance: High

⦿ CAUTION : External Email ⦿

Good morning,

DHS is submitting Reg. No. 14-558, Mental Health Procedures (Proposed Rulemaking) to the Senate Health and Human Services Committee and the House Human Services Committee.

Please provide written (email) confirmation that this rulemaking was received by the Committee chair's office.

Best,
Maeve



Maeve Curley

Pronouns: She/Her
Regulatory Coordinator
PA Department of Human Services | Office of Policy Development
macurley@pa.gov
<https://www.dhs.pa.gov>

From: [Bulletin](#)
To: [Curley, Maeve](#)
Cc: [Whare, Jennifer \(GC\)](#); [Duckett, Danielle A.](#); [Dietrich, Dawn](#); [Serafin, Kenneth](#); [Madden, Victoria](#)
Subject: [External] Re: DHS Proposed Regulation #14-558 Mental Health Procedures
Date: Thursday, November 13, 2025 11:57:54 AM
Attachments: [image001.jpg](#)

November 13, 2025

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Hello Maeve!

Thank you for submitting proposed rulemaking 14-558. Someone from our office will be in touch regarding the December 6 publication in the Bulletin.

Have a great day!

Leah

From: Curley, Maeve <macurley@pa.gov>
Sent: Thursday, November 13, 2025 11:39 AM
To: Bulletin <bulletin@palrb.us>
Cc: Whare, Jennifer (GC) <jwhare@pa.gov>; Duckett, Danielle A. <dduckett@pa.gov>; Dietrich, Dawn <dadietrich@pa.gov>; Serafin, Kenneth <kserafin@pa.gov>; Madden, Victoria <vmadden@pa.gov>
Subject: DHS Proposed Regulation #14-558 Mental Health Procedures

Good morning,

DHS is submitting Reg. No. 14-558, Mental Health Procedures (Proposed Rulemaking) to the Senate Health and Human Services Committee, the House Human Services Committee, LRB, and IRRC.

DHS has requested a publication date of December 6th in the Pennsylvania Bulletin. Please let me know if there are any questions regarding this.

Please provide written (email) confirmation that this rulemaking was received by LRB.
Thank you!

Best,
Maeve



Maeve Curley

Pronouns: She/Her

Regulatory Coordinator

PA Department of Human Services | Office of Policy Development

macurley@pa.gov

<https://www.dhs.pa.gov>