

481—556.4(124) Reporting requirements.

556.4(1) Reportable data. The following will be reported to the PMP in accordance with Iowa Code sections 124.551, 124.552, and 124.554(1) “g”:

- a. Controlled substances dispensed to a patient for self-administration.
- b. Opioid antagonists dispensed or administered by a practitioner, including:
 - (1) To an emergency department patient, and
 - (2) To a patient upon discharge from a hospital, correctional facility or care facility.
- c. If the pharmacy did not dispense or administer any reportable prescriptions during a reporting period, a zero report.

556.4(2) Required data elements.

a. In addition to the information required in Iowa Code section 124.552(1), the following elements will be reported:

- (1) Form of transmission of prescription origin.
 - (2) Refill number.
 - (3) Number of refills authorized.
- b. In an exceptional circumstance, a practitioner may request an extension of time for transmitting program information.

556.4(3) Exemptions—practitioners. The following are exempt from reporting controlled substances:

- a. A licensed veterinarian in the normal course of professional practice.
- b. A DEA-registered narcotic treatment program that is subject to the recordkeeping provisions of 21 CFR 1304.24 as amended on June 28, 2021.

556.4(4) Board notification. A pharmacy that does not engage in any reportable dispensing or administration will notify the board at the time of licensure.

556.4(5) Submission format. Data will be transmitted via the PMP’s current version of data upload or electronic submission.

556.4(6) Submission errors. Upon notification of a potential error in program information, the reporting practitioner will promptly correct the error.

[ARC 9342C, IAB 6/11/25, effective 7/16/25; see Delay note at end of chapter]