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S.F. No. 3566 – Criteria for prescribing controlled substances for the treatment of intractable pain

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S.F. 3566 modifies the prescribing and administering criteria for the use of controlled substances when treating a diagnosed condition that cause intractable pain. S.F. 3566 also requires the prescriber and patient to enter into a patient-provider agreement that includes expectation, responsibilities, and rights.

Section 1 (152.125) makes modifications to the prescribing criteria for controlled substances when treating intractable pain.

Subdivision 1 adds definitions for drug diversion, palliative care, and rare disease.

Subd. 1a establishes criteria for the evaluation and treatment of intractable pain when treating a nonterminally ill patient.

Subd. 2, paragraph (a) authorizes advanced practice registered nurses and physician assistants to prescribe or administer a controlled substance to a patient as part of the patient's treatment of a diagnosed condition causing intractable pain. Requires the provider to enter into a patient-provider agreement.

Paragraph (b) states that a prescriber shall not be subject to any civil or criminal actions or any investigation, termination, or disenrollment by either the commissioner of health or human services solely for prescribing a dosage that equates to an upward deviation from morphine milligram equivalent dosage recommendations or thresholds specified in state or federal opioid prescribing guidelines or policies.

Paragraph (c) prohibits a prescriber who is treating intractable pain with a controlled substance from tapering a patient's medication dosage solely to meet a predetermined dosage recommendation or threshold if the patient is stable and complaint with the treatment plan; is experiencing no serious harm from the level of medication prescribed, and is in compliance with the patient-provider agreement.

Paragraph (d) specifies that a prescriber's decision to taper a patient's medication dosage must be based on factors other than a morphine milligram equivalent recommendation or threshold.

Paragraph (e) specifies that no pharmacist, health plan company, or pharmacy benefit manager shall refuse to fill a prescription for an opiate issued by a licensed practitioner authorized to prescribe opiates solely on the prescription exceeding a predetermined morphine milligram equivalent dosage recommendation or threshold.

Subd. 3 and 4 add advanced practice registered nurse and physician assistant to these subdivisions. Make other technical changes.

Subd. 5, paragraph (a) requires the prescriber and patient to enter into an agreement that includes the patient's and prescriber's expectations, responsibilities, and rights according to the best practices and current standard of care.

Paragraph (b) requires that the agreement be signed by the patient and the prescriber, and a copy of the agreement included with the patient's medical record and a copy be provided to the patient.

Paragraph (c) requires the agreement to be reviewed at least annually and if there is a change to the patient's treatment plan, the agreement must be revised and updated and signed by the patient with a copy provided to the patient and included in the patient's medical record.

Paragraph (d) specifies that a patient provider agreement is not required in an emergency or inpatient hospital setting.