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Ground Floor Hearing Room, Andrew Jackson Building

Welcome/Introductions/Announcements

- A. Opening remarks by TDMHSAS Commissioner Marie Williams
- B. Tutorial on Emergency Rulemaking Process
- II. SB2009/HB2002-Discussion & Recommendations re: Injectable Buprenorphine Mono Product
- III. SB777/HB717

I.

- A. Discussion & Recommendations re: Requirements of SB777/HB7171 re: changes to the Buprenorphine Treatment Guidelines
  - 1. Initiating periodic prescriber-initiated and prescriber-led discussions with patients re: patient readiness to taper down or taper off opioids employed in treatment.
- B. Discussion & Solicitation of Advisory Comments re: Requirements of SB777/HB717 re: changes to TDMHSAS Rule 0940-05-35 (Office-Based Opiate Treatment Facilities) in order to establish:
  - 1. Standards for determining what constitutes a high dose of a the opioid employed in treatment at a nonresidential office-based opiate treatment facility;
  - 2. Protocols for initiating or switching a patient at a nonresidential officebased treatment facility to a high dose of the opioids employed in treatment;
  - 3. Protocols for initiating periodic prescriber-initiated and prescriber-led discussions with patients re: patient readiness to taper down or taper off the opioids employed in treatment; and,
  - 4. Requirements for approval of dispensing of buprenorphine products at a nonresidential office-based opiate treatment facility, including, but not limited to:
    - a) Reporting dispensing authority to licensing board;

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- b) Checking the CSMD prior to dispensing; and,
- c) Reporting to the CSMD amounts dispensed to the extent permitted by 42 CFR part 2.
- IV. SB2095/HB2510-FYI re: Buprenorphine Prescribing Authority Working Group
- V. Adjourn