

Policy Number: 500.210

Title: Tardive Dyskinesia Monitoring

Effective Date: 12/3/19

PURPOSE: To provide a systematic way of monitoring offenders/residents for the symptoms of tardive dyskinesia (TD).

APPLICABILITY: All adult and juvenile facilities.

DEFINITIONS:

<u>Abnormal Involuntary Movement Scale (AIMS)</u> – a standardized instrument used to rate offenders/residents for TD.

<u>Neuroleptic medications</u> – prescription drugs classified as anti-psychotics or similar drugs known to be associated with the potential side effects of TD.

<u>Practitioner</u> – physician/psychiatrist, nurse practitioner, physician's assistant, or other person licensed and authorized to prescribe medications.

<u>Tardive dyskinesia</u> – an involuntary movement disorder associated with the long term use (generally greater than six months) of neuroleptic medication.

PROCEDURES:

- A. The psychiatric provider must:
 - 1. Obtain written informed consent. This must occur prior to initiating neuroleptic medications, or continuing them upon intake, and within 14 days of emergency neuroleptic medication administration.
 - 2. Complete the AIMS evaluation prior to initiating neuroleptic medication(s), and/or continuing neuroleptic medication(s) upon intake, and within 14 days of emergency administration of neuroleptic medication(s).
 - 3. Complete the AIMS evaluation at least bi-annually (every six months).
 - 4. Document the course of action in a psychiatric case note including any required follow-up for AIMS evaluation.
 - 5. Return the consent form and AIMS evaluation to the offender's/resident's mental health file.
 - 6. For a tardive dyskinesia (TD) diagnosis:
 - a) Document in the psychiatric note and medical record.
 - b) Inform the offender or legally authorized representative of the diagnosis and risks.
 - c) Obtain an amended written informed consent from the offender, or the legally authorized representative, indicating that they have been informed of the potential risks.

- d) Inform the department of corrections (DOC) medical director.
- 7. For the discontinuation of neuroleptic medications:
 - a) Complete the AIMS evaluation two months after the discontinuation of neuroleptic medication(s).
 - b) If a positive AIMS score is determined, complete the AIMS evaluation again within three months, and as warranted thereafter.

B. Mental health staff must:

- 1. File the consent form in the legal section of the offender's/resident's mental health file.
- 2. File the AIMS evaluation in the clinical section of the offender's/resident's mental health chart.
- 3. Forward a copy of the AIMS evaluation to medical staff for the psychiatric section of the medical file.
- 4. Track offenders requiring AIMS follow-up on the facility Master Client List (MCL), which includes scheduling offenders for follow-up appointments.

C. Nursing staff must:

- 1. File a copy of the consent for the neuroleptic medications in the legal section of the offender's/resident's medical file.
- 2. File a copy of the AIMS evaluation in the psychiatric section of the offender's/resident/s medical file.

INTERNAL CONTROLS:

A. AIMS evaluations and consent forms are filed in the medical and mental health records.

ACA STANDARDS: 4-4397

REFERENCES: Diagnostic and Statistical Manual of Mental Disorders (DSM) IV Criteria 33.82

Neuroleptic-Induced Tardive Dyskinesia

Munetz, Benjamin, "How to Examine Patients Using the Involuntary Movement

Scale," Hospital and Community Psychiatry (1988) 1172-1177.

Policy 500.321, "Administration of Neuroleptic (Antipsychotic) and Non-

Neuroleptic, Psychotropic Medications"

REPLACES: Policy 500.210, "Tardive Dyskinesia Monitoring," 5/1/18.

All facility policies, memos, or other communications whether verbal, written, or

transmitted by electronic means regarding this topic.

ATTACHMENTS: None

APPROVALS:

Deputy Commissioner, Community Services Deputy Commissioner, Facility Services Assistant Commissioner, Operations Support Assistant Commissioner, Facility Services