

# Stereotypic Movement Disorder

Code: 307.3 (F98.4)

## Stereotypic Movement Disorder

307.3 (F98.4)

### Diagnostic Criteria 307.3 (F98.4)

- A. Repetitive, seemingly driven, and apparently purposeless motor behavior (e.g., hand clapping, head banging, body rocking, head shaking, biting, licking own body).  
B. The repetitive motor behavior interferes with social, academic, or other important activities and may result in self-injury.
- C. Onset before age 18 years.
- D. The repetitive motor behavior is not attributable to the physiological effects of a substance (e.g., medication, alcohol), another medical condition, or a better described another neurodevelopmental or mental disorder (e.g., trichotillomania (hair-pulling disorder), obsessive-compulsive disorder).

#### Specify if:

With self-injurious behavior (or behavior that would result in an injury if preventive measures were not used).

Without self-injurious behavior.

Associated with a known medical or genetic condition, neurodevelopmental disorder, or environmental factor; record stereotypic movement disorder associated with (name of condition, disorder, or factor) (e.g., stereotypic movement disorder associated with Lesch-Nyhan syndrome).

Coding and additional text: Add a three-digit code to identify the associated medical or genetic condition, or neurodevelopmental disorder.

#### Specify if:

Mild: Symptoms are easily suppressed by sensory stimuli or distraction.

Moderate: Symptoms are not easily suppressed by sensory stimuli or behavioral modification.

Severe: Continuous monitoring and protective measures are required to prevent self-injury.

#### Recording Procedures

For stereotypic movement disorder that is associated with a known medical or genetic condition, neurodevelopmental disorder, or environmental factor, record stereotypic movement disorder associated with (name of condition, disorder, or factor) (e.g., stereotypic movement disorder associated with Lesch-Nyhan syndrome).

#### Diagnostic Features

The severity of non-self-injurious stereotypic movements ranges from mild presentations that are easily suppressed by a sensory stimulus or distractors to continuous movements that result in self-injury. Stereotypic movements can occur in various settings and in a variety along various dimensions, including the frequency, impact on adaptive functioning, and/or severity. For example, in children with developmental delay or intellectual disability, the behaviors are typically less responsive to such efforts. In older children and adolescents, the behaviors are typically more persistent and severe. Body, to lacerations or amputation of digits, to retinal detachment from head banging, wrapping arms in clothing, finding a protective device).

The typical presentation of stereotypic movement disorder is repetitive, seemingly driven, and apparently purposeless motor behavior (Criterion A). These behaviors are often impulsive and do not require much cognitive effort to perform (e.g., head banging, body rocking, head nodding). Stereotyped self-injurious behaviors include, but are not limited to, repetitive head banging, head shaking, head slapping, head shaking, head slapping, head nodding, eye poking, and hand nodding. Stereotyped self-injurious behaviors include, but are not limited to, repetitive head banging, head shaking, head slapping, head shaking, head slapping, head nodding, eye poking, and hand nodding. Eye poking is particularly concerning; it occurs more frequently among children with developmental delay or intellectual disability. Head slapping and head shaking, rocking the torso, waving a small string repeatedly in front of the face).

Stereotypic movements may occur many times during a day, lasting a few seconds to several hours. They may occur in short bursts or in a continuous stream over a single day to several weeks between episodes. The behaviors vary in context, occurring when the individual is awake, sleeping, or in deep sleep. The onset of stereotypic movements is sudden. Criterion A requires that the movements be "apparently" purposeless. However, some functions of stereotypic movements are not fully understood, and their purpose might reduce anxiety in response to external stressors.

Criterion B states that the stereotypic movements interfere with social, academic, or other important activities and may result in self-injury (or would if preventive measures were not used). If self-injury is present, it should be coded using the specifier. Onset of stereotypic movements is in the early developmental period (Criterion C). Criterion D states that the stereotypic movements are not due to the physiological effects of a substance or neurological condition and is not better explained by another neurodevelopmental or mental disorder. The presence of parental or environmental factors suggests an undiagnosed neurodevelopmental problem, especially in children ages 1–3 years.

#### Precursors

Simple stereotypic movements (e.g., rocking) are common in young typically developing children. Complex stereotypic movements are much less common, occurring in approximately 1% of the general population. Stereotypic movements are associated with stereotypic movement disorder. The onset of stereotypic movements is in the early developmental period (Criterion C). Criterion D states that the stereotypic movements are not due to the physiological effects of a substance or neurological condition and is not better explained by another neurodevelopmental or mental disorder. The presence of parental or environmental factors suggests an undiagnosed neurodevelopmental problem, especially in children ages 1–3 years.

#### Development and Course

Stereotypic movements begin within the first 3 years of life. Simple stereotypic movements are common in infancy and may be involved in acquisition of motor mastery. In children with developmental delay or intellectual disability, simple stereotypic movements begin around age 12 months, between 24 and 36 months, and 8% at 36 months or older. In most typically developing children, these movements resolve over time or can be suppressed. Onset of complex stereotypic movements is later than simple stereotypic movements. Among individuals with intellectual disability, the stereotyped self-injurious behaviors may persist for years, even decades, and may become more severe as the individual ages.

#### Risk and Prognostic Factors

Environmental. Social isolation is a risk factor for self-stimulation that may progress to stereotypic movements. Stressful life events, such as abuse, neglect, or loss, may trigger stereotypic behavior. Fear may alter physiological state, resulting in increased frequency of stereotypic movements.

Genetic and physiological. Lower cognitive functioning is linked to greater risk for stereotypic behaviors and poorer response to interventions. Stereotypic movements are more frequent and severe in children with developmental delay or intellectual disability, who, by virtue of a particular syndrome (e.g., Rett syndrome) or environmental factor (e.g., an environmental toxin), have a neurophysiological profile that makes them more susceptible to stereotypic self-injurious behavior may be a behavioral phenotype in neurogenetic syndromes. For example, in Prader-Willi syndrome, Angelman syndrome, and Rett syndrome, self-injury, biting, and self-mutilation of fingers, bitting, and other forms of self-injury unless the individual is strained, and in Rett syndrome and Cornelia de Lange syndrome, self-injury may result from the heightened sensitivity to touch. Patients with Prader-Willi syndrome and other medical conditions, 10%–15% may have stereotypic movement disorder with self-injury.

#### Culturally Sensitive Considerations

Stereotypic movements disorder, with or without self-injury, occurs in all races and cultures.

Cultural attitudes toward unusual behaviors may result in delayed diagnosis. Overall cultural understanding of stereotypic movements disorder is needed and must be considered.

#### Differential Diagnosis

Normal development. Simple stereotypic movements are normal in infants and early childhood. They are common in the transition from sleep to awake, and behavior that the child is Neurodevelopmental Disorders

All neurodevelopmental disorders are less common in typically developing children and can usually be suppressed by distraction or sensory stimulation. The individual's daily routine is rarely affected, and stereotypic behavior does not cause the child distress or impairment in social, academic, or other important areas of functioning. It is appropriate to diagnose stereotypic movement disorder in children with autism spectrum disorder. Stereotypic movements may be a presenting symptom of autism spectrum disorder, but they are also common in other neurodevelopmental disorders. Autism spectrum disorder. Stereotypic movements may be a presenting symptom of autism spectrum disorder, but they are also common in other neurodevelopmental disorders. Behaviors are being evaluated. Deficits of social communication and reciprocity manifesting as social impairments, social interaction, social communication, and rigid repetitive behaviors and interests are distinguishing features. When autism spectrum disorder is present, stereotypic movements are sufficiently severe to become a focus of treatment.

The diagnosis of stereotypic movement disorder begins at age 3 years (before 3 years) than do tics, which have a mean age at onset of 5–7 years. They are consistent and fixed in their pattern and frequency. They are not due to the physiological effects of a substance or neurological condition. The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder). The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder). The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder).

Obsessive-compulsive disorder. Stereotypic movement disorder is distinguished from obsessive-compulsive disorder (OCD) by the absence of obsessions, as well as the lack of repetitive behaviors in response to an obsession or according to rules that may be applied rigidly. The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder).

Trichotillomania (hair-pulling disorder). Stereotypic movement disorder is distinguished from trichotillomania (hair-pulling disorder) by the lack of hair pulling. The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder).

Other tic disorders. Stereotypic movement disorder is distinguished from other tic disorders by the lack of associated comorbid conditions. The diagnosis of stereotypic movements requires the exclusion of habits, mannerisms, paroxysmal dyskineties, and benign hereditary chorea. A neurological history and examination are required to assess features suggesting a neurological condition. Stereotypic movements associated with a neurological condition may be distinguished by their signs and symptoms. For example, stereotypic movements associated with a neurological condition can be distinguished by a history of chronic neuroleptic use and characteristic oral or facial movements. The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder).

Tourette's disorder. The diagnosis of stereotypic movement disorder is not appropriate for repetitive head banging or head shaking associated with amphetamine intoxication or abuse (e.g., patients with Tourette's disorder).

#### Comorbidity

Stereotypic movement disorder may occur as a primary diagnosis or secondary to another disorder. Paroxysmal dyskineties, stereotypies are a common manifestation of a variety of neurological disorders, such as Lesch-Nyhan syndrome, Rett syndrome, fragile X syndrome, Cornelia de Lange syndrome, and Dravet syndrome. When stereotypic movement disorder co-occurs with another medical condition, both should be coded.

#### Tic Disorders 81

##### Diagnostic Criteria

Note: A. The tics are brief, rapid, recurrent, nonrhythmic motor movement or vocalization. Tourette's Disorder 307.23 (F98.2).

B. Both multiple motor and one or more vocal tics have been present at some time during the illness, although not necessarily concurrently.

C. The tics may wax and wane in frequency but have persisted for more than 1 year since first tic or onset.

D. Onset is before age 18 years.

E. The tics are not premeditated and are not under voluntary control (e.g., coexisting with another medical condition (e.g., Huntington's disease, postviral encephalitis)).

Persistent (Chronic) Motor or Vocal Tic Disorder 307.22 (F98.1)

A. Single or multiple motor and/or vocal tics.

B. The tics have been present for less than 1 year since first tic or onset.

C. Onset is before age 18 years.

D. The disturbance is not attributable to the physiological effects of a substance (e.g., coexisting with another medical condition (e.g., Huntington's disease, postviral encephalitis)).

E. Criteria have never been met for Tourette's disorder or persistent (chronic) motor or vocal tic disorder.

#### Specifiers

With motor tics only.

With vocal tics only.

With both motor and vocal tics.

B. The tics have been present for less than 1 year since first tic or onset.

C. Onset is before age 18 years.

D. The disturbance is not attributable to the physiological effects of a substance (e.g., coexisting with another medical condition (e.g., Huntington's disease, postviral encephalitis)).

E. Criteria have never been met for Tourette's disorder or persistent (chronic) motor or vocal tic disorder.

#### Diagnostic Features

Tic disorders comprise four diagnostic disorders: Tourette's disorder, persistent (chronic) motor or vocal tic disorder, persistent (chronic) motor and vocal tic disorder, and unspecified tic disorders. Diagnosis for any tic disorder is based on the presence of motor and/or vocal tics (Criterion A), duration of tic symptoms (Criterion B), age of onset (Criterion C), and severity of any associated functional impairment (Criterion D).

The tic disorders are hierarchical in order (i.e., Tourette's disorder, followed by persistent (chronic) motor or vocal tic disorder, followed by persistent (chronic) motor and vocal tic disorder, and finally unspecified tic disorders).

The 82 Neurodevelopmental Disorders

order: simple motor tics, complex motor tics, and other specified tic disorders, such that those at a tier disorder are not level of the hierarchy is diagnosed, a lower hierarchy diagnosis cannot be made (Criterion E).

Tics are sudden, rapid, recurrent, nonrhythmic movements. An individual may have tics that come and go over time, but at any point in time, the tic recurrence occurs in a characteristic fashion. Although tics can include almost any muscle group or sensory system, they are most commonly observed in the muscles of the head and neck, eyelids, mouth, and pharynx. Tics are generally experienced as involuntary but can be voluntarily suppressed. Tics are generally experienced as involuntary but can be voluntarily suppressed. Tics can be either simple or complex. Simple motor tics are of short duration (i.e., minutes) and involve a single muscle group. Complex motor tics involve two or more muscle groups. Simple vocal tics include throat clearing, sniffing, and grunting often caused by contraction of the diaphragm or muscles of the oropharynx. Complex vocal tics are more prolonged and involve more complex patterns of muscle groups, such as taurine head turning and shoulder shrugging. Complex tics can appear purposeful, such as to scratch an itch or to relieve tension. Some tics are associated with other movement disorders (echopraxia). Similarly, complex vocal tics include repeating one's own words or saying words that sound like they are being forced out (echolalia). Complex vocal tics may include unacceptable words, including obscenities, or ethnic, racial, or religious slurs (coprolalia). Importantly, coprolalia is an abrupt sharp bark or grunt utterance and lacks the prosody of language.

The presence of motor and/or vocal tics varies across the tic disorders (Criterion A). For example, Tourette's disorder is characterized by multiple motor and/or vocal tics, whereas persistent (chronic) motor or vocal tic disorder, only motor or only vocal tics are present. For persistent (chronic) motor and vocal tic disorder, both motor and vocal tics are present. For unspecified tic disorders, the movement disorder symptoms are best characterized as tics but are atypical in presentation or age of onset, or have a known etiology.

The 83 Feeding and Eating Disorders

For tic disorders co-occurring with feeding and eating disorders, individuals diagnosed with either Tourette's disorder or persistent (chronic) motor or vocal tic disorder

DSM-5 Diagnostic Item | Pages 122-135 | Computer-Readable Format