

# Cyclothymic Disorder

DSM-5-TR 301.13 | ICD-10-CM F34.0

*Two or more years of fluctuating subthreshold hypomanic and depressive symptoms that never reach full episode criteria.*

**LIFETIME PREVALENCE**  
~0.4-1% (US adults)

**TYPICAL ONSET**  
Adolescence/early adult

**SEX RATIO**  
~1:1 in community samples

**COURSE**  
15-50% convert to BD I/II

## CLINICAL PICTURE

- Mood shifts every few days to weeks with no sustained euthymic baseline, which patients describe as simply being moody by nature.
- Up phases bring energy, sociability, reduced sleep need, and productivity; down phases bring fatigue, withdrawal, and self-doubt.
- Interpersonal instability, impulsive decisions, and erratic work or academic performance follow the shifting affective states.
- Patients typically present during a down phase and describe the up phases as their normal, good, or most effective self.
- Substance use frequently maps onto the cycle, with stimulants or alcohol used to extend or truncate particular mood states.
- Onset in adolescence means the pattern is embedded in identity and is often mislabeled as a personality problem by others.

## DIAGNOSTIC CRITERIA AT A GLANCE

- At least two years in adults, or one year in children and adolescents, of numerous periods of hypomanic and depressive symptoms.
- Symptoms must be present at least half the time with no symptom-free interval longer than two consecutive months during that period.
- No full manic, hypomanic, or major depressive episode may occur during the initial two-year qualifying period of the illness.
- Symptoms cause clinically significant distress or impairment and are not attributable to substances or another medical condition.
- If a full episode later develops, the diagnosis converts to Bipolar I or II, with cyclothymic disorder recorded as the antecedent.

## NEUROBIOLOGY & PHYSIOLOGY

- Familial aggregation with Bipolar I and II is high, supporting cyclothymia as a bipolar-spectrum phenotype rather than mere temperament.
- Circadian rhythm instability and highly variable sleep duration track the state changes and are measurable with actigraphy.
- Reward system reactivity resembles that seen in Bipolar II but with shorter, lower-amplitude oscillations across the affective range.
- Affective temperament research links cyclothymic temperament to elevated emotional reactivity and behavioral activation sensitivity.
- Antidepressant exposure can accelerate cycling and unmask hypomania, revealing the bipolar diathesis beneath the subthreshold picture.
- Comorbid substance use compounds mood instability and independently worsens the long-term neurobiological and functional trajectory.

## PSYCHOLOGICAL MECHANISMS

- Chronic instability prevents formation of a stable self-concept, which is frequently misread by clinicians as personality pathology.
- Reinforcement obtained during productive up phases discourages engagement with treatment that appears to flatten the mood range.
- Interpersonal consequences accumulate as partners and employers respond to unpredictability rather than to a recognized illness.
- Rhythm disruption from irregular sleep schedules, travel, and shift work directly amplifies the amplitude of mood oscillation.
- Mood charting builds the pattern recognition patients need in order to intervene before a swing consolidates into an episode.

## DIFFERENTIAL & COMORBIDITY

- Borderline personality disorder shows interpersonally triggered shifts within hours plus identity disturbance and abandonment fear.
- Bipolar II requires a full hypomanic episode plus a major depressive episode, whereas cyclothymia remains subthreshold on both.
- Rule out substance-induced mood instability, thyroid disease, and the emotional dysregulation that accompanies untreated ADHD.
- Comorbid substance use disorders, anxiety disorders, and ADHD are frequent and each worsens functional and treatment outcomes.
- Suicide risk is meaningful despite subthreshold severity, particularly when impulsivity and active substance use are present.

## Treatment EVIDENCE-BASED MANAGEMENT

### PHARMACOLOGIC

- No treatment is FDA-approved; **lithium 0.5-0.8 mEq/L** and **valproate** are used off label with modest supportive evidence.
- **Lamotrigine 100-200 mg/day** targets depressive predominance and offers a favorable tolerability and metabolic profile.
- Low-dose **quetiapine 50-300 mg/day** may reduce mood lability when affective instability is the most prominent complaint.
- Avoid antidepressant monotherapy, which risks cycle acceleration, emergent mixed states, and unmasking of frank hypomania.
- Address comorbid substance use directly, since ongoing stimulant or alcohol use makes genuine mood stabilization unattainable.

### PSYCHOTHERAPY

- **Psychoeducation** covering bipolar-spectrum illness, prodromes, and cycle triggers is the core and best-supported intervention.
- **IPSRT** stabilizes daily routines and sleep timing, which is the most direct available lever on cycling frequency.
- **CBT** addresses depressive cognition and the risky decisions patients make during elevated or activated phases.
- **DBT** skills in distress tolerance and emotion regulation help when impulsivity and interpersonal chaos dominate the picture.
- Family or couples work reduces conflict driven by unpredictable state changes and meaningfully improves treatment adherence.

### ADJUNCT & SUPPORTIVE

- Daily mood charting across 8 to 12 weeks serves as both the diagnostic instrument and the primary self-monitoring tool.
- A fixed wake time, consistent meal timing, and limited alcohol intake measurably reduce the amplitude of mood oscillation.
- Monitor annually for conversion to Bipolar I or II, since a substantial minority of patients progress to full episodes.
- Screen with the **MDQ** and obtain collateral report, because self-report alone consistently underestimates up phases.
- Vocational counseling helps where erratic performance threatens employment, vocational training, or academic standing.

- ☆ **CLINICAL PEARLS**
- Chronic instability without full episodes; check for symptom-free gaps over two months.
  - A single full episode ends the cyclothymia diagnosis and starts Bipolar I or II.
  - Antidepressants alone can accelerate cycling; cover with a mood stabilizer.

# References

American Psychiatric Association. (2022). *Diagnostic and statistical manual of mental disorders* (5th ed., text rev.).

<https://doi.org/10.1176/appi.books.9780890425787>

National Institute for Health and Care Excellence. (2014). *Bipolar disorder: Assessment and management* (NICE Guideline CG185).

<https://www.nice.org.uk/guidance/cg185>

National Institute of Mental Health. (n.d.). *Bipolar disorder*. U.S. Department of Health and Human Services.

<https://www.nimh.nih.gov/health/topics/bipolar-disorder>

Sadock, B. J., Sadock, V. A., & Ruiz, P. (2021). *Kaplan & Sadock's synopsis of psychiatry* (12th ed.). Wolters Kluwer.

Stahl, S. M. (2021). *Stahl's essential psychopharmacology* (5th ed.). Cambridge University Press.

Van Meter, A. R., Youngstrom, E. A., & Findling, R. L. (2012). Cyclothymic disorder: A critical review. *Clinical Psychology Review*, 32(4), 229-243.

<https://doi.org/10.1016/j.cpr.2012.02.001>

## NOTE

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