Diagnostic classification of irritability and oppositionality in youth: a global field study comparing ICD-11 with ICD-10 and DSM-5

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Background: Severe irritability has become an important topic in child and adolescent mental health. Based on the available evidence and on public health considerations, WHO classified chronic irritability within oppositional defiant disorder (ODD) in ICD-11, a solution markedly different from DSM-5's (i.e. the new childhood mood diagnosis, disruptive mood dysregulation disorder [DMDD]) and from ICD-10's (i.e. ODD as one of several conduct disorders without attention to irritability). In this study, we tested the accuracy with which a global, multilingual, multidisciplinary sample of clinicians were able to use the ICD-11 classification of chronic irritability and oppositionality as compared to the ICD-10 and DSM-5 approaches. **Methods:** Clinicians ($N = 196$) from 48 countries participated in an Internet-based field study in English, Spanish, or Japanese and were randomized to review and use one of the three diagnostic systems. Through experimental manipulation of validated clinical vignettes, we evaluated how well clinicians in each condition could identify chronic irritability versus nonirritable oppositionality, episodic bipolar disorder, dysthymic depression, and normative irritability. **Results:** Compared to ICD-10 and DSM-5, ICD-11 led to more accurate identification of severe irritability and better differentiation from boundary presentations. Participants using DSM-5 largely failed to apply the DMDD diagnosis when it was appropriate, and they more often applied psychopathological diagnoses to developmentally normative irritability. **Conclusions:** The formulation of irritability and oppositionality put forth in ICD-11 shows evidence of clinical utility, supporting accurate diagnosis. Global mental health clinicians can readily identify ODD both with and without chronic irritability. **Keywords:** International Classification of Diseases (ICD-11); oppositional defiant disorder; mood dysregulation; irritability; child and adolescent mental health.

Introduction

Irritability in children and adolescents (herein 'youth') has emerged as a significant clinical and public health concern for which assessment and treatment options are limited. Defined as an elevated proneness to anger, irritability has manifestations ranging from normative emotions to chronically irritable mood and aggressive outbursts (Stringaris et al., 2018). Irritability occurs in over a dozen mental disorders including oppositional defiant disorder (ODD), where it is chronic and often a core part of the presentation (Evans et al., 2017). Longitudinal studies show that chronic irritability does not specifically predict bipolar disorder (BD) or conduct disorder (CD), but it does predict ODD, depression, anxiety disorders, suicidality, and poor functional outcomes (Brotman, Kircanski, & Leibenluft, 2017; Evans et al., 2017; Vidal-Ribas et al., 2016). Thus, it is important that diagnostic classification systems accurately identify and characterize severely irritable youth in need of clinical care. Chronic irritability

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was therefore given close attention in the development of the Mental, Behavioural, and Neurodevelopmental Disorders (MBND) chapter in the Eleventh Revision of the World Health Organization's (WHO's) *International Classification of Diseases and Related Health Problems* (ICD-11). The purpose of the present study was to evaluate the utility of the ICD-11 formulation of irritability and oppositionality for global mental health applications.

In May 2019, the World Health Assembly approved ICD-11 as the global standard for collection and reporting of health information by WHO's 194 member nations (Reed et al., 2019). The development of ICD-11 MBND was a massive and globally influential undertaking coordinated by the WHO Department of Mental Health and Substance Abuse (First et al., 2015; International Advisory Group, 2011; Reed, 2010; Reed et al., 2019). As a part of this effort, the Department developed detailed diagnostic guidelines for clinician use and implemented a program of global field studies to assess the reliability and clinical utility of these guidelines (First et al., 2015; Keeley et al., 2016). At the time of this study, the ICD-11 draft guidelines had not yet been finalized, allowing a window for potential refinement based on field study results.

Early in ICD-11 development, clinical utility and global applicability were identified as guiding principles based on the idea that improvements in these domains would foster improvements in clinical service delivery and resulting health data (International Advisory Group, 2011; Reed, 2010). To this end, an international group of experts in disruptive mood and behavior problems (e.g. aggression, oppositionality, mood disorders, irritability) reviewed the literature and proposed changes intended to (a) improve communication among users (e.g. practitioners, patients, families, administrators); (b) foster conceptualization and understanding of disruptive behavior and dissocial disorders (DBDDs); (c) accurately and easily describe actual clinical presentations; (d) assist with clinical management; (e) enhance clinical outcomes (Reed, 2010); and (f) enhance applicability of these changes by multidisciplinary clinicians in diverse settings across the globe.

It was the view of this and other ICD-11 expert working groups that existing evidence argued against introducing chronic irritability as a new standalone disorder. Instead, it was proposed that irritability be identified as a feature of ODD (Evans et al., 2017; Lochman et al., 2015). This decision was justified in part by evidence showing that irritability is a major subdimension of ODD symptoms (e.g. Burke et al., 2014; Ezpeleta et al., 2012; Rowe et al., 2010; Stringaris & Goodman, 2009). Converging research on ODD irritability and alternative models of chronic irritability (e.g. severe mood dysregulation [SMD]; Leibenluft, 2011) showed that irritability can be significantly impairing and predictive of anxiety and depression (Evans et al., 2017; Vidal-Ribas

et al., 2016). The irritable dimension of ODD is also recognized in the *Diagnostic and Statistical Manual of Mental Disorders* (DSM-5; American Psychiatric Association [APA], 2013). The evidence and rationale behind ICD-11's conceptualization of irritability have been detailed by Evans et al. (2017) and Lochman et al. (2015).

ICD-11's interpretation of the evidence base contrasted with DSM-5's. For DSM-5, findings from research on chronic irritability (Brotman et al., 2017; Leibenluft, 2011, 2017) were cited as justification for creating a category named disruptive mood dysregulation disorder (DMDD), a new childhood depressive disorder (APA, 2013; Roy, Lopes, & Klein, 2014). This line of research began as a response to observed upticks in pediatric BD diagnoses in the United States in the 1990s–2000s, apparently driven by some clinicians' interpretation of chronic irritability as a form of pediatric mania. In response, work pioneered by Leibenluft (2011, 2017) and others (for broad reviews, see Brotman et al., 2017; Evans et al., 2017) operationalized chronic irritability as a research syndrome, severe mood dysregulation (SMD), and generated a compelling body of evidence showing that SMD is distinct from BD. Thus, this research helped clarify the boundaries of pediatric BD; it also demonstrated chronically irritable youth were severely impaired and at increased risk for an array of negative affective, behavioral, and functional outcomes. In consideration of the clear clinical need of this population, SMD was adapted, with modifications, to form DMDD in DSM-5.

As a diagnostic category, however, DMDD lacks compelling evidence for its validity, reliability, and clinical utility (Axelson et al., 2012; Evans et al., 2017; Lochman et al., 2015). Research and clinical advances in this area are sorely needed, but efforts to understand, assess, and treat severe irritability are hindered by inconsistency in how it has been measured and classified. Developing an empirically supported, clinically useful, and globally applicable classification is essential for improving care for youth with severe irritability and related problems. The vast majority of youth with DMDD would already (empirically, if not by definition) receive a diagnosis of ODD, and the DMDD diagnosis shows no incremental validity or utility beyond ODD (Freeman et al., 2016). Clinically, the best available treatment options are not different from those for ODD (Stringaris et al., 2018), but DMDD has raised concerns about the possibility of treatment with psychoactive medications with limited evidence for their efficacy or safety in this population (Tourian et al., 2015).

The revised classification of chronic irritability and DBDDs in ICD-11 must also be evaluated in relation to ICD-10, the previous official system for health information and reporting and the most widely used classification system globally (Reed et al., 2011). Given that nearly three decades have passed since the last major revision, it is not surprising that ICD-

11 (WHO, 2020) represents a significant reformulation from ICD-10 (WHO, 1992). For example, ICD-10's F90-F98 section (disorders of childhood and adolescence) has been dissolved, replaced by developmental organization and guidance throughout ICD-11 MBND (Reed et al., 2019). Further, whereas ICD-10 counted ODD as one of several 'Conduct disorders', ICD-11 reflects current evidence by recognizing ODD and CD as the two distinct, major categories of DBDDs. Thus, an important component of the present study was to test whether ICD-11's treatment of DBDDs and irritability, in the context of these other major changes, represents an improvement from ICD-10.

The present study

Despite the sound empirical rationale behind these ICD-11 proposals for ODD and irritability, open questions remain regarding their clinical utility and whether they can be used accurately and consistently by clinicians. We sought to investigate these issues in an ICD-11 global field study on DBDDs in youth. In this report, we focus on research questions examining the performance of ICD-11 guidelines as compared to ICD-10 and DSM-5 in the assessment of severe irritability versus boundary clinical presentations (ODD and bipolar and dysthymic/depressive disorders) and developmentally normative irritability. Specifically, we investigated the extent to which each classification system could assist clinicians in making four key challenging diagnostic distinctions:

1. *Differentiating chronically irritable from nonirritable oppositional behavior.* Most chronically irritable youth, under any diagnostic system, meet the requirements for ODD; yet, many with ODD do not display severe irritability (Evans et al., 2017). This reflects the reality that ODD is heterogeneous, always including disruptive behavior and sometimes including prevailing negative affect dysregulation (Burke & Loeber, 2010). Thus, it is important to test whether clinicians can accurately identify ODD both with and without severe irritability.

2. *Differentiating chronic irritability from episodic BD irritability.* Chronic irritability initially attracted interest because it was purportedly being misdiagnosed as a 'broad phenotype' of bipolar disorder (BD)—that is, some clinicians viewed it as a childhood form of mania (Leibenluft, 2011). If this were the case, certain BD diagnoses (e.g. BD type II, other specified BD) would likely be given. Thus, we investigated this question: Can clinicians differentiate chronic irritability from an episodic BD presentation with irritability?

3. *Differentiating chronic irritability from depressive disorders.* Whereas the preceding question focused on the key distinctions of chronic versus episodic, it is also important to assess whether clinicians can differentiate among two types of nonepisodic, chronic presentations including irritability: ODD versus chronically depressed mood (i.e. dysthymic/persistent depressive disorder). This distinction is clearly supported by research, with different treatment implications for the two conditions (Stringaris et al., 2018).

4. *Differentiating chronic irritability from normative irritability.* Much of the controversy surrounding DMDD has involved concern about pathologizing developmentally normative irritability, such as temper tantrums in young children (Axelson et al., 2011; Wakefield, 2016). Secondary analyses do not show alarmingly high rates of DMDD, but prevalence varies greatly depending on assessment methods (Axelson et al., 2012; Copeland et al., 2013) and effects on the diagnosis of individual children are unclear. It is possible that including a novel 'high severity' disorder may effectively lower the diagnostic threshold for ODD, if the latter is perceived as a 'low severity' disorder by comparison.

These questions 1–4 correspond to comparisons 1–4 in our experimental design in the Methods, Results, and Discussion that follow. While each specific question is relatively narrow in scope, they collectively served two overarching goals: to both *evaluate* and *refine* the proposed ICD-11 guidelines for better utility in global mental health applications.

Methods

This project was part of the larger program of ICD-11 field trials for which the broad methods and justification have been published elsewhere (Evans et al., 2015; Keeley et al., 2016). Below, we focus only on the design, methodology, and sample for the data reported here.

Participants

A global, multilingual, multidisciplinary sample of mental health professionals ($N = 196$) participated in English, Japanese, or Spanish (see Table 1). Clinicians were from 48 countries, with the highest proportions being from India (19.4%), Japan (10.2%), the USA (7.7%), Spain (7.1%), the UK (6.6%), Mexico (5.1%), Nigeria (4.6%), Australia (4.1%), and Canada (4.1%). Participants were identified via the Global Clinical Practice Network (GCPN; https://gcp.network/en/). Beginning in 2011, clinicians from all regions of the globe were invited to register with the GCPN to participate in Internet-based field studies for the ICD-11 (Reed et al., 2015). To be eligible for the present study, clinicians had to (a) be a mental health or primary care professional qualified to work with persons with mental disorders; (b) be currently seeing patients or directly supervising services, including (c) at least some services to children; and (d) have proficiency in one of the study languages. Individuals were ineligible to participate if they had contributed to the ICD-11 MBDN revision process or to the DBDD field study development in any capacity. Informed consent was obtained from all participants. Procedures were exempted from review by the WHO Ethics Review Committee and approved by the Human Subjects Committee of the University of Kansas, where the servers used for data collection were housed.

Development of case vignettes

Five vignettes (labeled A–E; see Table 2, Table S1, and Appendix S1) were developed to help answer the study questions. Each vignette was designed, refined, and validated to match a target clinical presentation. This work was carried out by DBDD working group members and consultants, following best practices for vignette development (Evans et al., 2015). These experts were asked to write vignettes to meet specified requirements, drawing from actual patient presentations. Draft vignettes were edited for internal consistency, clinical clarity, and cross-cultural applicability, and then were sent to a separate group of experts who conducted confirmatory evaluations of each vignette to ensure diagnostic agreement. The task of editing and reviewing vignettes involved coding key clinical features (e.g. symptoms, severity, impairment) and identifying any potentially ambiguous aspects, which were then clarified. By the end of this process, each vignette had been vetted to meet the diagnostic requirements for the target disorder under ICD-11, ICD-10, and DSM-5. Any disagreements were resolved by consensus. On this basis, each vignette was designated as having a single 'correct' diagnosis per each diagnostic system. Vignettes and other materials were developed in English and translated into Spanish and Japanese using a forward- and back-translation process carried out by native speakers with relevant clinical expertise.

Procedures

The study was programmed and hosted in Qualtrics. E-mail invitations were sent to all registered clinicians in each language group who met the participation criteria. When the study was launched, 3,274 GCPN members across all three languages qualified and were contacted via personalized email to invite them to participate. Of these, 686 (21.0% response rate) clicked the link and began the study, of whom 493 (71.9% completion rate) finished the study. The sample of interest here is the subset of 196 participants (39.8% of the total sample) who were allocated to and finished one of the four comparisons (of 13 total comparisons) pertaining to oppositionality and irritability.

Upon following the link and providing consent, clinicians were randomly allocated to the ICD-11, ICD-10, or DSM-5 condition and to a paired comparison condition. Block and stratified randomization were used to ensure a balance of participant numbers and self-reported DBDD expertise levels across cells. Diagnostic system was masked, although ICD-10 or DSM-5 materials could have been recognized by those familiar with them. After reviewing the diagnostic guidelines, clinicians followed prompts to assess the two assigned vignettes, one at a time (order counterbalanced). To help control for potential gender-related biases, vignette character gender was manipulated such that half the participants assessed two male characters and the other half two female characters (see Appendix S1). The characters' age (range 6–11 years) was selected based on clinical and developmental considerations and held constant within vignette. Clinicians could select from broad disorder clusters and specific diagnostic categories, spanning DBDDs, habit/impulse control, bipolar, and depressive disorders. They could review the diagnostic guidelines while deciding and could also enter a different diagnosis or indicate that no diagnosis was warranted. After selecting an initial diagnosis, participants were shown its essential features, one by one, and asked to indicate whether each was present in the vignette case. Next, participants could change or re-confirm their diagnosis. These

Table 1 Participant characteristics

	English f (%)	Spanish f (%)	Japanese f (%)	Total f (%)
Region				
AFRO	17 (12.6)			17 (8.7)
AMRO North	22 (16.3)	1 (2.4)		23 (11.7)
AMRO South	4 (3.0)	26 (63.4)		30 (15.3)
EMRO	9 (6.7)			9 (4.6)
EURO	32 (23.7)	14 (34.1)		46 (23.5)
SEARO	40 (29.6)			40 (20.4)
WPRO Asia	2 (1.5)		20 (100)	22 (11.2)
WPRO Oceania	9 (6.7)			9 (4.6)
Gender				
Male	66 (48.9)	22 (53.7)	15 (75.0)	103 (52.6)
Female	69 (51.1)	19 (46.3)	5 (25.0)	93 (47.4)
Discipline				
Counseling	7 (5.2)			7 (3.6)
Medicine	67 (49.6)	11 (26.8)	16 (80.0)	94 (48.0)
Psychology	51 (37.8)	27 (65.9)	4 (20.0)	82 (41.8)
Social work	4 (3.0)	1 (2.4)		5 (2.6)
Occupational therapy	3 (2.2)			3 (1.5)
Other	3 (2.2)	2 (4.9)		5 (2.6)
Relevant specializations[a]				
Child/adolescent	62 (45.9)	14 (34.1)	7 (35.0)	83 (42.3)
ADHD/DBDD	41 (30.4)	12 (29.3)	4 (20.0)	57 (29.1)
Total N finished/analyzed	135	41	20	196
	M (SD)	*M (SD)*	*M (SD)*	*M (SD)*
Age	49.83 (11.06)	47.88 (10.89)	50.90 (10.17)	49.53 (10.92)
Years of experience	17.19 (10.46)	19.17 (9.95)	18.30 (9.67)	17.72 (10.26)

Data were self-reported from GCPN registration.
[a]Specialization variables represent how many clinicians reported spending at least 50% of their clinical time working with individuals 0–18 years of age (child/adolescent) and how many identified ADHD and conduct disorders as one of their top three areas of specialization (ADHD/DBDD).

Table 2 Comparisons, vignettes, and correct[a] diagnoses

	First[b] vignette	Second[b] vignette
Comparison 1	*Vignette B* An 8-year-old [boy/girl][c] with predominately headstrong/defiant oppositional behavior. • ICD-11: ODD • ICD-10: ODD • DSM-5: ODD	*Vignette C* A 9-year-old [boy/girl][c] with oppositional behavior and chronic irritability, including mood and outburst features. • ICD-11: ODD • ICD-10: ODD • DSM-5: DMDD
Comparison 2	*Vignette C* (See top right)	*Vignette D* An 11-year-old [boy/girl][c] with a history of depressive and hypomanic episodes, including irritable features. • ICD-11: Bipolar type II disorder • ICD-10: Bipolar affective disorder • DSM-5: Bipolar II disorder
Comparison 3	*Vignette C* (See top right)	*Vignette E* A 10-year-old [boy/girl][c] with persistently depressed mood, including irritable features. • ICD-11: Dysthymic disorder • ICD-10: Dysthymia • DSM-5: Persistent depressive disorder
Comparison 4	*Vignette A* A 6-year-old [boy/girl][c] with developmentally appropriate tantrums. • ICD-11: no diagnosis • ICD-10: no diagnosis • DSM-5: no diagnosis	*Vignette C* (See top right)

See Appendix S1 for full vignettes.
[a]'Correct' diagnoses are those that the vignettes were developed and validated to represent.
[b]Order of presentation for a random ~50% of participants (reversed for the other ~50%).
[c]A random ~50% of participants within each comparison viewed two male vignettes (the other ~50% viewed two female).

procedures helped ensure that diagnostic decisions were made with due attention to the diagnostic guidelines and to the clinical features of the vignette.

Analytic plan and power

Analyses used a mixed factorial design involving three diagnostic systems (ICD-11, ICD-10, DSM-5) × four paired vignette comparisons (oppositionality, bipolar, dysthymia, normality; each compared to chronic irritability) as between-participant factors, with a within-participant factor comparing diagnoses of the two paired vignettes. Two-way chi-square statistics were used for bivariate comparisons and the G-square statistic for three-way interactions (Rao & Scott, 1984). Data from all three study languages were combined for analysis. With an average of $n = 32.1$ per pairwise comparison, sensitivity power analyses showed adequate power $(1-\beta = .81)$ to detect large effects $(w = 0.5)$ at standard thresholds $(\alpha = .05)$.

Results

The 196 participating clinicians were randomly allocated to the ICD-11 $(n = 61)$, ICD-10 $(n = 72)$, and DSM-5 $(n = 63)$ conditions and then to four paired-comparison conditions. The results for each comparison are described below. See Figure 1 for a visualization of the results across all four comparisons.

Comparison 1: Can clinicians accurately diagnose irritable versus nonirritable oppositional behavior?

Comparison 1 evaluated clinicians' assessment of two cases: (1) a child with predominately headstrong/defiant behavior (Vignette B), who would receive ODD under any system; and (2) a child with predominately irritable and angry disruptive behavior (Vignette C), who would be best identified as ODD with chronic irritability—anger under ICD-11, ODD under ICD-10, and DMDD under DSM-5.

Participants using ICD-11 selected the correct diagnosis (ODD) for both vignettes 66.7% of the time overall (80.0% and 53.3%, respectively), with no difference between vignettes, $\chi^2(1) = 2.40$, *ns*. In contrast, participants using both ICD-10 $(\chi^2(1) = 7.20, p < .01)$ and DSM-5 $(\chi^2(1) = 13.33, p < .001)$ were less accurate in their diagnosis of Vignette C compared to Vignette B. Specifically, participants in the ICD-10 condition diagnosed
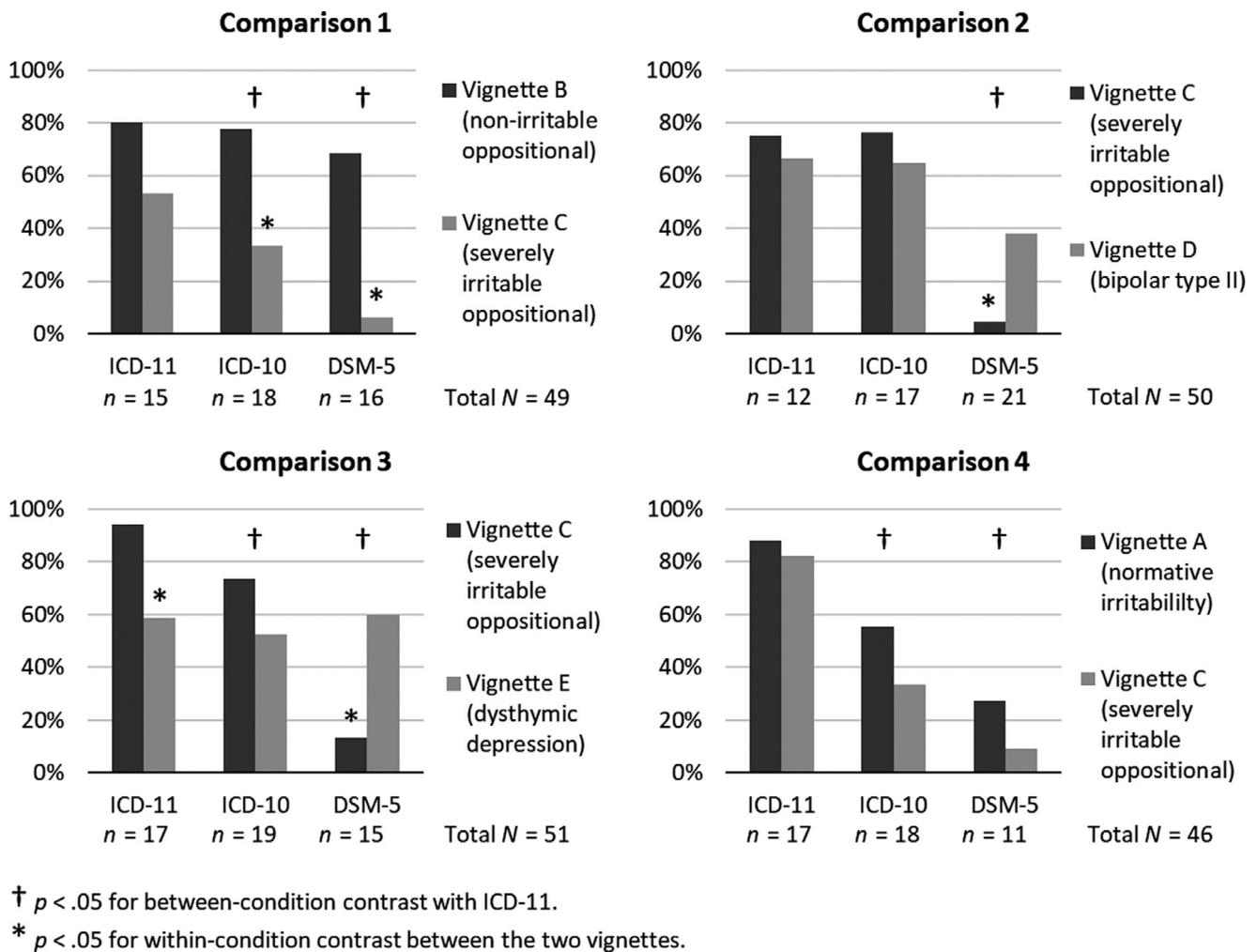
Figure 1 Percentage of participating clinicians assigning the correct diagnosis under ICD-11, ICD-10, and DSM-5 conditions

Vignette B with ODD at 77.8% accuracy compared to Vignette C with ODD at 33.3% accuracy. Those using DSM-5 correctly diagnosed Vignette B with ODD at 68.8% accuracy and Vignette C with DMDD with only 6.3% accuracy. The most frequently selected incorrect answers included unsocialized CD (38.9%) for ICD-10 and ODD (50.0%) for DSM-5. Overall, participants using ICD-11 were more accurate in their diagnostic assignments than those using either ICD-10 ($G^2(4) = 10.78$, $p < .05$) or DSM-5 ($G^2(4) = 22.78$, $p < .001$).

Comparison 2: Can clinicians accurately diagnose chronic irritability versus episodic BD irritability?

Participants assigned to this condition viewed two cases: (1) a child with chronic, severe irritability, anger, and outbursts (Vignette C); and (2) a child with irritable/hypomanic mood and depressive episodes, characteristic of BD type II (Vignette D). Clinicians using ICD-11 ($\chi^2(1) = 0.20$, ns) and ICD-10 ($\chi^2(1) = 0.57$, ns) were equally correct for their diagnoses of Vignette C (75.0% and 76.5%, respectively) and Vignette D (66.7% and 64.7%, respectively). Those using DSM-5 showed lower accuracy in

assessing Vignette C (4.8%) than Vignette D (38.1%), $\chi^2(1) = 6.93$, $p < .01$. Again, those in the DSM-5 condition most often selected ODD (81.0%) for Vignette C, although DMDD was correct. Overall, participants in the ICD-11 condition were more accurate than participants using DSM-5 ($G^2(4) = 23.78$, $p < .001$), but not different from those using ICD-10 ($G^2(4) = 0.80$, ns), in identifying episodic bipolar irritability and chronic oppositional irritability.

Comparison 3: Can clinicians accurately diagnose chronic irritability versus dysthymic depression?

Comparison 3 examined this question with two vignettes: (1) a child presenting with chronic irritability and disruptive behavior, characteristic of severe ODD/DMDD (Vignette C); and (2) a child presenting with chronically depressed mood with irritability, representing persistent depression/dysthymia (Vignette E). Participants using ICD-11 were less accurate in diagnosing Vignette E (58.8%) than they were for Vignette C (94.1%), $\chi^2(1) = 5.88$, $p < .05$; the specific difficulty was in differentiating among depressive disorders (when all unipolar

depressive diagnoses were counted as correct, accuracy was 87.5% and no longer differed from Vignette C, $\chi^2(1) = 0.32$, *ns*). Participants using ICD-10 were equally correct for Vignette C (73.7%) and Vignette E (52.6%), $\chi^2(1) = 1.81$, *ns*. Those using DSM-5 again had difficulty recognizing DMDD as the correct diagnosis for Vignette C (13.3%) and did better diagnosing Vignette E (60.0%), $\chi^2(1) = 7.03$, $p < .01$. The most commonly selected answer for Vignette C for DSM-5 was again ODD (86.7%). Participants in the ICD-11 condition were more accurate overall than those in the ICD-10 ($G^2(4) = 9.82$, $p < .05$) and DSM-5 ($G^2(4) = 24.54$, $p < .001$) conditions.

Comparison 4: Can clinicians accurately diagnose severe versus normative irritability?

Two vignettes were presented: (1) a typically developing child with irritable tantrum behavior within normal limits (Vignette A) and (2) a child with a more severe, chronically irritable presentation (Vignette C). Participants using ICD-11 accurately classified Vignette A as normative (88.2%) and Vignette C as clinical (82.4%), with no difference in accuracy between vignettes, $\chi^2(1) = 0.23$, *ns*. Unlike previous comparisons, participants using ICD-10 had some difficulty diagnosing Vignette C (33.3%) but this was not worse than their performance on Vignette A (55.6%), $\chi^2(1) = 1.80$, *ns*. Vignette C was most often diagnosed as unsocialized CD (38.9%). Participants using DSM-5 had little accuracy in identifying Vignette A (27.3%) and Vignette C (9.1%), with equally poor accuracy on both, $\chi^2(1) = 1.22$, *ns*. In terms of incorrect answers, participants using DSM-5 most often selected ODD for both Vignette A (54.5%) and Vignette C (45.4%). Overall accuracy was greater for participants in the ICD-11 condition compared to those using ICD-10 ($G^2(4) = 15.44$, $p < .01$) and DSM-5 ($G^2(4) = 28.10$, $p < .001$). Specifically, for typical irritability (Vignette A), those using ICD-11 more often correctly selected 'no diagnosis' (88.2%) than those using either ICD-10 (55.6%; $\chi^2(1) = 4.58$, $p < .05$) or DSM-5 (27.3%; $\chi^2(1) = 10.81$, $p < .01$).

Discussion

We examined the clinical utility of the ICD-11 diagnostic classification of youth irritability and oppositionality, as compared to ICD-10 and DSM-5, among a global, multilingual, and multidisciplinary sample of clinicians. Participants assigned to use ICD-11 generally made more accurate diagnostic judgments about various presentations of irritability, oppositionality, and mood disturbance as compared to those using the other diagnostic systems. Notably, clinicians using DSM-5 largely failed to assign DMDD when it was appropriate and

more often applied psychopathological diagnoses to a case with developmentally typical irritability.

Results suggest that clinicians may struggle to accurately diagnose presentations of chronic irritability using ICD-10 (where answers were scattered across ODD/CD types) and DSM-5 (where they tended to select ODD rather than the correct DSM-5 diagnosis, DMDD). In contrast, ICD-11 may lead to better accuracy overall, and for both irritable and nonirritable presentations of oppositionality. This pattern of results was largely consistent across all comparisons. It may be that when assessing a child with prevailing chronic irritable mood and aggressive temper outbursts, clinicians tend to assign a DBDD diagnosis (i.e. ODD) rather than conceptualize it as a mood disorder (depressive or bipolar).

Regarding specific boundary presentations, comparisons 2 and 3 reflect favorably upon clinicians' performance across systems in *identifying* episodic BD with irritability and nonepisodic depression with irritability. However, when it comes to *differentiating* these presentations from chronic irritability and oppositionality, those using DSM-5 had more difficulty than those using ICD-11. Although this pattern was most clear and pronounced with DSM-5, ICD-11 also outperformed ICD-10 in Comparison 3, with results suggesting ICD-11 may be especially effective at steering clinicians to the correct overarching category (i.e. differentiating ODD/DBDDs from depressive disorders) as compared to ICD-10 and DSM-5.

Comparison 4 showed that ICD-11 may lead to greater overall accuracy differentiating typical and atypical irritability, compared to both ICD-10 and DSM-5. Within each of these diagnostic systems, participants showed similar levels of accuracy for typical versus atypical irritability, with ODD being frequently (incorrectly) selected under DSM-5. Notably, clinicians using ICD-11 were more likely to correctly ascertain the *absence* of pathology and indicate that a diagnosis was not warranted. Results suggest ICD-10 and, especially, DSM-5 could possibly lead to more false positive diagnoses in terms of children receiving *any* diagnosis in response to normative irritable and disruptive behavior.

Study strengths include its global, multilingual, multidisciplinary clinician sample and well-validated and controlled experimental methodology. Some limitations should be noted. Although vignette-based designs can be valid and reliable research proxies for clinical decision-making (Evans et al., 2015), the use of vignettes remains a limitation. One implication of the vignette approach is that only a select few types of cases can be represented (5 vignettes in this study) and contrasted (4 comparisons in this study) to investigate central study questions. Our vignettes were not designed to disentangle other important questions, such the diagnostic assessment of youth with severe, explosive temper outbursts but not chronically irritable mood,

based on the identification of the phasic versus tonic components of irritability—an important direction for future work (Brotman et al., 2017; Carlson & Klein, 2018). Researchers should also consider using actual or simulated patients, either through live clinical interviews or through video-recordings. Additionally, sample size was relatively small per cell; however, power was adequate to detect the large effects observed. Despite the global representation of the sample, it was not possible to test for language or geographical differences in results due to small cell sizes. Moving forward, research evidence generated through a variety of methodologies is needed to better understand and evaluate the ICD-11 formulations of oppositionality and irritability in relation to alternatives. These include epidemiological, clinical (assessment and intervention), behavioral, neuroscientific, and genetic approaches.

Overall, findings support the adequacy and improvement (relative to ICD-10 and DSM-5) of the ICD-11 diagnostic classification and guidance for irritability and oppositionality in youth. Clinicians appear to conceptualize presentations of disruptive mood and behavior problems in accordance with longstanding clinical and scientific understandings of psychopathology, irrespective of which diagnostic guidelines they are asked to use. Indeed, ODD is a well-established heterogenous disorder, which global clinicians showed they can readily diagnose both with and without chronic irritability. This has important clinical implications because effective treatment and management of a condition is predicated on its accurate identification. The literature's recognition that irritability is transdiagnostic and that chronic irritability predicts *subsequent* internalizing outcomes (Vidal-Ribas et al., 2016) is very useful, and this was part of the basis for DSM-5 locating DMDD within the depressive disorders section. In providing care to a young patient, however, prediction does not trump current presentation. Chronic irritability nearly always occurs within ODD; in this context, it can be severe and impairing, warranting clinical attention and predicting homotypic and heterotypic outcomes (Burke & Loeber, 2010; Evans et al., 2017).

Whether chronic irritability is conceptualized as ODD or as DMDD, the best available evidence supports virtually identical treatment recommendations: behavioral parent training and/or cognitive behavior therapy (Kircanski et al., 2018; Stringaris et al., 2018). These well-established psychosocial interventions may serve as first-line treatments, accompanied by a thorough clinical assessment and personalization to address any co-occurring concerns (Evans et al., 2020; Stringaris et al., 2018). This study suggests that ICD-11's coverage of irritability and oppositionality within the DBDD section can be implemented accurately by clinicians and will provide appropriate guidance for clinical decision-making.

Supporting information

Additional supporting information may be found online in the Supporting Information section at the end of the article:

Appendix S1. Vignettes used in the study (full text of vignettes A–E).
Table S1. Summary of correct/target diagnoses for each vignette per ICD-11, ICD-10, and DSM-5.

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Key points

- The WHO's ICD-11 classified chronic irritability in oppositional defiant disorder (ODD), a formulation differing from ICD-10 and DSM-5.
- In this vignette-based study, we tested the clinical utility of ICD-11's classification of oppositionality and irritability in youth among 196 global clinicians.
- Participants using ICD-11 more accurately identified youths' clinical presentations, while those using DSM-5 made mistakes concerning normative irritability and disruptive mood dysregulation disorder.
- Results support the clinical utility of ICD-11's ODD formulation and current scientific thinking of ODD as a behavioral disorder that may be accompanied by chronic irritability.
- Careful attention is needed for accurate assessment of youth irritability, with the differential diagnosis likely including ODD, mood disorders, and normative irritability.

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