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Eisma, Maarten C.; Janshen, Antje; Lenferink, Lonneke I.M.

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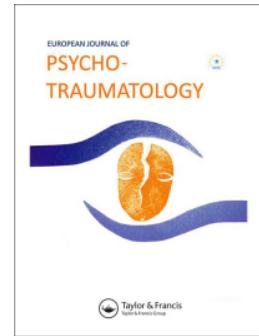
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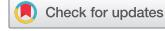
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LETTER TO THE EDITOR

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## Content overlap analyses of ICD-11 and DSM-5 prolonged grief disorder and prior criteria-sets

Maarten C. Eisma <sup>a</sup>, Antje Janshen<sup>a</sup> and Lonneke I. M. Lenferink <sup>a,b,c</sup>

<sup>a</sup>Department of Clinical Psychology and Experimental Psychopathology, Faculty of Behavioral and Social Sciences, University of Groningen, Groningen, The Netherlands; <sup>b</sup>Department of Psychology, Health, & Technology, Faculty of Behavioural, Management, and Social Sciences, University of Twente, Enschede, The Netherlands; <sup>c</sup>Department of Clinical Psychology, Faculty of Social Sciences, Utrecht University, Utrecht, The Netherlands

### ABSTRACT

**Background:** The International Classification of Diseases eleventh edition (ICD-11) has recently included prolonged grief disorder (PGD), a diagnosis characterized by severe, persistent, and disabling grief. The text revision of the Diagnostic and Statistical Manual of Mental Disorders 5 (DSM-5-TR) is scheduled to include a similar but distinct diagnosis, also termed PGD. Concerns have been raised that these new diagnoses are qualitatively different from both prior proposed diagnoses for pathological grief and each other, which may affect the generalizability of findings obtained with different criteria sets.

**Objective:** We conducted a content overlap analysis of PGD<sub>ICD-11</sub>, PGD<sub>DSM-5-TR</sub>, and previous proposals for pathological grief diagnoses (i.e. PGD 2009; complicated grief (CG), PGD ICD-11 beta draft, persistent complex bereavement disorder (PCBD) per DSM-5).

**Methods:** Using the Jaccard's Index, we established the degree of content overlap between core and accessory symptoms of PGD<sub>ICD-11</sub>, PGD<sub>DSM-5-TR</sub>, and prior proposals for pathological grief diagnoses.

**Results:** Main findings are that PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub> showed moderate content overlap with each other and with most prior proposed diagnoses for pathological grief. PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub> showed the strongest content overlap with their direct predecessors, PGD<sub>ICD-11</sub> beta draft and PCBD, respectively.

**Conclusions:** Limited content overlap between PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub> and preceding criteria sets may threaten generalizability of past research on phenomenological characteristics of pathological grief to current criteria sets. Similarly, findings obtained with instruments to assess PGD<sub>ICD-11</sub> may not generalize to PGD<sub>DSM-5-TR</sub> and vice versa. Researchers should aim to determine under which circumstances criteria sets for PGD yield similar or distinct characteristics. Convergence of criteria sets for PGD remains an important goal for the future.

### Análisis de solapamiento de contenido del Trastorno por Duelo

### Prolongado del CIE-11 y el DSM-5 y criterios diagnósticos previos

**Antecedentes:** La Decimoprimera Clasificación Internacional de Enfermedades (CIE-11) ha incluido recientemente el Trastorno Por Duelo Prolongado (PGD por sus siglas en inglés), un diagnóstico caracterizado por un duelo severo, persistente e incapacitante. La versión revisada del Manual Diagnóstico y Estadístico de los Trastornos Mentales (DSM-5-TR) tiene agendado incluir un diagnóstico similar pero diferente, también llamado PGD. Ha existido preocupación de que ambos diagnósticos sean cualitativamente diferentes de aquellos propuestos previamente para duelo patológico y también entre sí, lo que puede afectar la posibilidad de generalización de los hallazgos obtenidos con cada conjunto de criterios diagnósticos.

**Objetivo:** Conducimos un análisis de solapamiento de contenido de los criterios diagnósticos del PGD de acuerdo a la CIE-11, del PGD de acuerdo al DSM-5-TR y de propuestas previas para diagnósticos de duelo patológico [como el PGD de Prigerson y colaboradores, publicado el 2009, Duelo complicado (CG por sus siglas en inglés) del borrador beta de la CIE 11, el Trastorno por Duelo Complejo Persistente (PCBD por sus siglas en inglés) del DSM-5].

**Métodos:** Usando el Índice de Jaccard, establecimos el grado de solapamiento del contenido entre los síntomas principales y accesorios de los criterios diagnósticos del PGD de acuerdo a la CIE-11, del PGD de acuerdo con el DSM-5-TR y de propuestas previas para diagnósticos de duelo patológico.

**Resultados:** Los resultados principales son que los criterios diagnósticos del PGD de acuerdo a la CIE-11 y PGD de acuerdo al DSM-5-TR mostraron un solapamiento de contenido moderado entre ellos y también con la mayoría de los diagnósticos de duelo patológico previamente propuestos. Ambos diagnósticos mostraron el mayor solapamiento de contenidos con sus

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### PALABRAS CLAVE

Duelo; Pérdida; Diagnóstico;  
DSM-5 TR; CIE-11;  
Solapamiento de Contenido;  
Trastorno por Duelo  
Prolongado

### 关键词

哀伤; 丧亲; 诊断; DSM-5-TR;  
ICD-11; 内容重叠; 延长哀  
伤障碍

### HIGHLIGHTS

- Content overlap analyses showed moderate overlap between symptoms of PGD per DSM-5-TR and ICD-11 and between these diagnoses and prior criteria sets.
- We should establish when new criteria sets for PGD behave similarly or differently.
- Convergence of PGD criteria sets is needed.

predecesores directos, el Duelo Complicado del borrador beta de la CIE-11 y el PCBD respectivamente.

**Conclusiones:** el solapamiento limitado de contenidos entre los criterios diagnósticos del PGD de acuerdo a la CIE-11 y PGD de acuerdo al DSM-5-TR y los criterios precedentes pueden amenazar la generalización de investigación pasada de las características fenomenológicas del duelo patológico en los criterios diagnósticos actuales. En forma similar, los hallazgos obtenidos con instrumentos para evaluar el PGD de acuerdo a la CIE-11 pueden no ser generalizables al PGD de acuerdo al DSM-5-TR. Los investigadores debiesen determinar bajo qué circunstancias los criterios diagnósticos de PGD muestran características distintas o similares. La convergencia de los criterios diagnósticos de PGD sigue siendo una importante meta para el futuro.

## ICD-11 和 DSM-5 延长哀伤障碍和先前标准集的内容重叠分析

**背景:** 国际疾病分类第 11 版 (ICD-11) 最近纳入了延长哀伤障碍 (PGD), 其特征是严重, 持续和致残的哀伤。精神疾病诊断和统计手册 5 (DSM-5-TR) 的文本修订计划纳入类似但不同的诊断, 也称为 PGD。有人担心这些新诊断与先前提出的病理性哀伤诊断和彼此之间存在质的不同, 这可能会影响使用不同标准集获得的结果的推广性。

**目的:** 我们对 PGDICD-11, PGDDSM-5-TR 和先前提出的病理性哀伤诊断 (即依据 Prigerson et al., 2009 的 PGD; 复杂性哀伤 (CG), PGD ICD-11 测试版草案, 依据 DSM-5 的持续性复杂丧亲症 (PCBD)) 进行了内容重叠分析。,

**方法:** 使用 Jaccard 指数, 我们确定了 PGDICD-11, PGDDSM-5-TR 以及先前提出的病理性哀伤诊断的核心和附加症状之间的内容重叠程度。

**结果:** 主要发现是 PGDICD-11 和 PGDDSM-5-TR 显示出中度的内容重叠, 并且与大多数先前提出的病理性哀伤诊断重叠。 PGDICD-11 和 PGDDSM-5-TR 分别与其直接前身 PGDICD-11 测试版草案和 PCBD 显示出最强的内容重叠。

**结论:** PGDICD-11 和 PGDDSM-5-TR 和先前的标准集之间有限的内容重叠可能会威胁到过去关于病理性哀伤现象学特征研究对当前标准集的推广性。同样, 使用评估 PGDICD-11 的工具获得的结果可能无法推广到 PGDDSM-5-TR。研究人员应着眼于确定在何种情况下为 PGD 设定的标准会产生相似或不同的特征。 PGD 标准集的收敛性仍然是未来的一个重要目标。

Dear Editor,

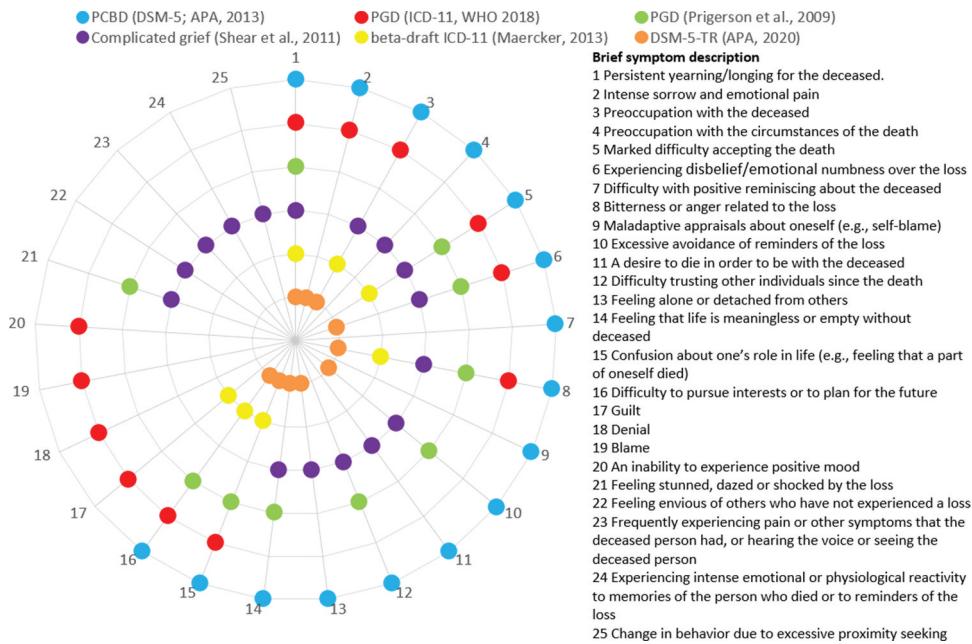
Over the past decades, there have been multiple attempts to define a diagnosis characterized by severe, persistent, and disabling grief, i.e. pathological grief. These proposed diagnoses have received different names, including complicated grief (CG: Shear et al., 2011), persistent complex bereavement disorder (PCBD: American Psychiatric Association [APA], 2013) and, most commonly, prolonged grief disorder (PGD, e.g. Prigerson et al., 2009, PGD<sub>2009</sub>; Maercker et al., 2013; PGDICD-11 beta draft). In 2018, a diagnosis termed PGD was formally added to the International Classification of Diseases eleventh edition (PGDICD-11, ICD-11: World Health Organization [WHO], 2018). A different diagnosis named PGD will be added to the text revision of the Diagnostic and Statistical Manual of Mental Disorders 5 in 2022 (PGD<sub>DSM-5-TR</sub>, DSM-5-TR: Boelen, Eisma, Smid, & Lenferink, 2020; Prigerson, Kakarala, Gang, & Maciejewski, 2021). Figure 1 displays core and accessory symptoms of all mentioned criteria-sets.

A concern regarding the development of new criteria-sets for pathological grief is that they are, as a rule, qualitatively different from preceding criteria-sets (e.g. Boelen & Prigerson, 2012; Djelantik et al., 2021; Eisma & Lenferink, 2017; Stelzer, Zhou, Maercker, O'Connor, & Killikelly, 2020). Criteria-sets differ in number of included symptoms, symptom content, and diagnostic algorithms (Eisma, Rosner, & Comtesse, 2020; Lenferink, Boelen, Smid, & Paap, 2021). Consequently, the phenomenological characteristics of different pathological

grief criteria-sets vary. For example, PGDICD-11 has limited diagnostic agreement with prior proposed criteria-sets, such as PCBD and PGD<sub>2009</sub> (e.g. Boelen & Lenferink, 2020; Boelen, Lenferink, Nickerson, & Smid, 2018; Comtesse et al., 2020; Cozza et al., 2020), although the extent of agreement partially depends on the chosen diagnostic algorithm (Eisma et al., 2020). Therefore, previous findings on important clinical issues, ranging from dimensionality of diagnoses to treatment efficacy, may not generalize to newer criteria-sets. Additionally, since PGDICD-11 and PGD<sub>DSM-5-TR</sub> also differ in symptom count, content, and diagnostic algorithms, findings obtained with one version of PGD may not generalize to the other.

Therefore, clarifying to what extent criteria-sets capture the same content and whether and when criteria-sets of pathological grief yield similar or different results appears important. The aim of the present contribution is to assess the comparability of different criteria-sets using a mathematical approach. Specifically, we will establish the extent to which the content of core and accessory symptoms in PGDICD-11, PGD<sub>DSM-5-TR</sub>, and preceding criteria-sets overlap. We derive these methods from Fried (2017), who used a similar approach to illustrate the limited content overlap between items from seven frequently used self-report measures of depression.

We estimated content overlap between criteria-sets using the Jaccard Index, a similarity coefficient for binary data ranging from 0 (no overlap among criteria-sets) to 1 (complete overlap). It is calculated with the following formula:  $J = s / (u_1 + u_2 + s)$ , where  $J$  is the Jaccard Index,  $s$



**Figure 1.** Similarities and differences between diagnostic criteria sets of pathological grief.

is the number of items that two criteria-sets share, and  $u_1$  and  $u_2$  are the number of symptoms unique to each criteria set. Since there is no established guideline on categorizing the strength of overlap using the Jaccard Index, we will apply the rule by Evans (1996) for the correlation coefficient as an indicator: very weak 0.00–0.19, weak 0.20–0.39, moderate 0.40–0.59, strong 0.60–0.79, and very strong 0.80–1.0 (Fried, 2017).

Supplemental Table S1 shows the results. A first main finding is that there is moderate overlap between the most recent criteria-sets PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub> ( $J = 0.47$ ). PGD<sub>ICD-11</sub> shows the strongest overlap with the PGD<sub>ICD-11</sub> beta draft ( $J = 0.58$ ), whereas PGD<sub>DSM-5-TR</sub> shows the strongest overlap with PCBD ( $J = 0.63$ ), illustrating that they most closely resemble their direct predecessors. Both PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub> show least overlap with CG ( $J = 0.22$  and  $0.37$ , respectively). Overall, the mean overlap between PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub> with all other criteria-sets is moderate ( $J = 0.41$  and  $0.48$ , respectively). CG stands out as the diagnosis showing the least overlap with all other criteria-sets ( $J = 0.35$ ), whereas PCBD shows most content overlap with other criteria-sets ( $J = 0.49$ ).

Overall, our analysis demonstrated modest content overlap between prior proposed criteria-sets and both PGD<sub>ICD-11</sub> and PGD<sub>DSM-5-TR</sub>. Moreover, the two newest criteria-sets showed limited content overlap with each other. These findings complement prior empirical research demonstrating differences between the characteristics of different criteria-sets for pathological grief (e.g. Boelen et al., 2020; Boelen & Lenferink, 2020; Comtesse et al., 2020; Cozza et al., 2020). Using a single validated instrument, such as the recently developed Traumatic Grief Inventory–Self Report Plus, to assess symptoms of different criteria-sets, may be instrumental to further elucidate when criteria-sets behave similarly or differently

(Lenferink, Eisma, Smid, de Keijser, & Boelen, Lenferink, et al., 2022).

Together, results suggest that limited content overlap could partly explain differences in findings across different criteria-sets. Two courses of action may help reduce this problem of generalizability in the future. First, we should strive for greater convergence of future diagnostic criteria-sets with presently used criteria-sets (Lenferink et al., 2021). Second, since PGD<sub>ICD-11</sub> uses a descriptive diagnosis without a formal diagnostic algorithm, we could investigate which PGD<sub>ICD-11</sub> algorithm yields to the greatest convergence with past criteria-sets, and, more importantly, with PGD<sub>DSM-5-TR</sub> criteria (Eisma et al., 2020).

Some limitations warrant mention. First, this work is a mathematical exercise that complements but does not substitute empirical studies of similarities and differences between pathological grief criteria-sets. Second, we only compared core and accessory symptoms of criteria-sets. For example, we did not take into account differences in time criteria or diagnostic algorithms between proposed diagnoses. A third limitation is that for CG we split up some compound symptoms (e.g. ‘Frequent intense feeling of loneliness or like life is empty or meaningless without the person who died’ was separated into ‘loneliness’ and ‘feeling life is empty/meaningless’) because these symptoms were also separated in other criteria-sets (see Lenferink et al., 2021 for details). This may have led us to overestimate content overlap between CG and other criteria-sets. Fourth, one grief researcher assessed overlap between criteria sets (cf. Lenferink et al., 2021). Multiple assessors may have yielded more reliable and replicable classifications of symptoms.

Notwithstanding these limitations, our analyses have demonstrated suboptimal comparability in the

content of past and current pathological grief criteria-sets. We have highlighted how this may result in problems of generalizability of findings obtained with past and current criteria-sets. This work illustrates the need for further convergence of diagnoses and empirical investigations of the similarities and differences between pathological grief diagnoses and related phenomenological characteristics.

## Disclosure statement

No potential conflict of interest was reported by the author(s).

## ORCID

Maarten C. Eisma  <http://orcid.org/0000-0002-6109-2274>  
Lonneke I. M. Lenferink  <http://orcid.org/0000-0003-1329-6413>

## Data availability statement

The authors confirm that the data supporting the findings of this study are available within the article and its supplementary materials.

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