

# Narcissistic Personality Disorder: An Integrative Review of Recent Empirical Data and Current Definitions

Stefan Roepke · Aline Vater

© Springer Science+Business Media New York 2014

**Abstract** Although concepts of pathological narcissism are as old as psychology and psychiatry itself, only a small number of clinical studies are based on the criteria for narcissistic personality disorder (NPD), as defined in the Diagnostic and Statistical Manuals of Mental Disorders (DSM). As a result, NPD appears to be one of the most controversially discussed nosological entities in psychiatry. Whereas the majority of empirical studies used self or other ratings of NPD criteria to address issues of reliability and validity of the diagnostic category (i.e., internal consistency, factor structure, discriminant validity), only recent research has applied experimental designs to investigate specific features of NPD (e.g., self-esteem, empathy, shame). The aim of this review is to summarize available empirical data on NPD and relate these findings to current definitions of NPD (according to the DSM-5, [1]). In order to do so, this review follows the five steps to establishing diagnostic validity proposed by Robins and Guze [2], i.e., (1) clinical description, (2) laboratory studies, (3) delimitation from other disorders, (4) family studies, and (5) follow up studies. Finally, this review suggests pathways for future research that may assist further nosological evaluation of NPD and contribute to the overall goal, the improvement of treatment for patients.

**Keywords** Narcissistic personality disorder · Self-esteem · Empathy · Shame · Stability

## Introduction

Most empirical studies on narcissistic personality disorder (NPD) psychometrically evaluated the diagnostic criteria as defined in editions of the Diagnostic and Statistical Manuals of Mental Disorders (DSM). As a result, controversial discussions on the validity of the NPD construct in preparation for the DSM-5 [1] lacked a key component: laboratory studies. Only very recent research has begun to bridge that gap by applying experimental designs to investigate specific features of NPD (e.g., self-esteem, empathy, shame). The aim of this review is to incorporate those recent findings into the discussion on the validity of NPD. In order to do so, this review follows the five steps for establishing diagnostic validity proposed by Robins and Guze [2], i.e., (1) clinical description, (2) laboratory studies, (3) delimitation from other disorders, (4) family studies, and (5) follow up studies. Furthermore, empirical data are evaluated in relation to the two current definitions of NPD in the DSM-5 (in Sects. II and III). Finally, directions for future research are proposed and implications for treatment are discussed.

---

This article is part of the Topical Collection on *Personality Disorders*

S. Roepke (✉) · A. Vater  
Department of Psychiatry, Charité – Universitätsmedizin Berlin,  
Campus Benjamin Franklin, Eschenallee 3, 14050 Berlin, Germany  
e-mail: stefan.roepke@charite.de

S. Roepke  
Cluster of Excellence “Languages of Emotion”, Freie Universität  
Berlin, Berlin, Germany

A. Vater  
University of Darmstadt, Darmstadt, Germany

## Clinical Descriptions

### Development of NPD as DSM Category

Pathological narcissism was first described by Ellis [3] and further elaborated by psychoanalysts (e.g., [4-6]). Despite its longstanding tradition in psychiatric literature, NPD was not introduced as a psychiatric disorder until the 3rd edition of the DSM in 1980 [7]. Diagnostic criteria in DSM-III were based

on psychoanalytic literature and expert consensus without prior empirical evaluation [8]. In DSM-III-R [9], the polythetic criteria set replaced the mixed polythetic-monothetic model applied in DSM-III. Throughout the revision of NPD criteria for DSM-IV [10], overt grandiose themes were emphasized (see [11]). Additionally, clinically significant distress or impairment in functioning caused by personality disorder symptoms was added as one of the general personality disorder criteria in DSM-IV. The current DSM-5 [1] retained the DSM-IV general personality disorder criteria and the criteria set for NPD in Sect. II. Moreover, DSM-5 proposed an alternative research model for personality disorders in Sect. III. According to Sect. III in DSM-5, NPD is characterized by specific impairments in personality functioning (with characteristic difficulties in areas of identity, self-direction, empathy and intimacy) and pathological personality traits (i.e., grandiosity and attention seeking). Although the new definition of NPD in Sect. III of the DSM-5 is well grounded in clinical descriptive literature of pathological narcissism, it is not backed up by empirical research.

#### Internal Consistency

A number of DSM-based studies assessed internal consistency of the NPD criteria set. In summary, DSM-III and DSM-III-R NPD criteria showed rather low to moderate internal consistency (Cronbach's alpha from .38 to .69; see [11]), whereas the DSM-IV NPD criteria set reached higher internal consistency with acceptable values (Cronbach's alpha from .63 to .88; see [11, 12•]). In sum, NPD (DSM-IV and DSM-5, Sect. II) is characterized by acceptable internal consistency that is comparable to other personality disorders.

#### Typological and Dimensional Structure

Further studies provide conflicting evidence for the factorial structure of NPD. Different studies found evidence for a *one-factor* structure [12•, 13, 14], *two-factor* structure [15] or *three-factor* structure [16] for DSM-IV NPD criteria.

Up to now, only one study assessed the factor structure of a broader spectrum of symptoms based on concepts of pathological narcissism [17]. This study found a more differentiated three-factor structure of NPD (labeled grandiose/malignant, fragile, and high-functioning/exhibitionistic). The authors argue that DSM-IV criteria for NPD are too narrow and under-emphasize central aspects of the construct.

Furthermore, one study used taxometric analyses to analyze whether the latent distribution of the DSM-IV NPD features is discrete or dimensional [15]. Results revealed a latent discontinuity in the distribution of the DSM-IV NPD criteria, indicating a typological model of NPD, rather than a dimensional construct [15].

In summary, current analysis of the factor structure of DSM-IV NPD are not coherent, but suggest that the criteria set that has also been adopted in DSM-5 Sect. II only covers part of the clinical concept of pathological narcissism.

#### Prevalence Rates

Recent studies provide varying prevalence rates of NPD, mainly depending on sample selection. Prevalence of NPD according to DSM-III-R or DSM-IV ranges between 0.0 % and 1.0 % in population-based samples [18–24]. Higher prevalence rates have been reported in psychiatric populations, ranging from 0.8 to 5.8 %, with higher rates in outpatient settings relative to inpatient or day clinic facilities [12•, 25–27].

In addition to sample selection, race, ethnicity, and gender seem to account for variability in prevalence rates. First, NPD rates are higher among African-American men and women and Hispanic women compared to Asian or Pacific Islanders, Native American and Caucasians in a U.S. population-based sample [22]. Second, most (but not all) clinical studies found higher prevalence rates among males e.g., [12•, 15, 28]. This finding is consistent with population-based data reporting higher rates of NPD in men (i.e., 1.2 %) than in women (i.e., 0.7 %) [24].

#### Laboratory Studies

##### Self-esteem

Early psychoanalytic theories developed a self-regulatory model of pathological narcissism that has been further elaborated by theoreticians from social psychology e.g., [29], clinical psychology and psychiatry e.g., [30, 31]. The NPD description in Sect. III of the DSM-5 acknowledges this model and proposes, for example, that NPD patients rely excessively upon others for self-esteem regulation and emotion regulation mirrors fluctuations in self-esteem [1].

Based on the assumptions of the self-regulatory model, the frequently cited “mask model” [32] proposes that trait narcissism is characterized by fragile self-esteem. Fragile self-esteem is characterized by low implicit (i.e., automatic, not necessarily conscious, overlearned) self-esteem and grandiose high explicit (i.e., reflected, conscious) self-esteem compared to nonclinical controls. In order to prevent low implicit self-esteem from becoming more explicit, narcissistic patients may engage in defensive behavior.

Until now, only a few studies analyzed self-esteem in NPD patients. Pincus et al. [33] found that pathological narcissism (assessed with the Pathological Narcissism Inventory, PNI) negatively correlated with explicit self-esteem in patients with mixed psychiatric disorders. Vater et al. [34••] found that NPD

patients scored lower on explicit self-esteem than nonclinical controls, but higher than patients with borderline personality disorder. No significant differences emerged on implicit self-esteem compared to nonclinical controls. Thus, this study contradicts assumptions of unconscious feelings of insecurity in patients with NPD. However, those studies did not analyze short-term or long-term fluctuations in self-esteem that are described in Sect. III of the DSM-5. Thus, future studies should analyze fluctuations in self-esteem with regard to the proposed self-regulatory deficit in NPD.

## Empathy

Lack of empathy is characteristic of NPD in the DSM-5 (Sect. II as a diagnostic criterion, Sect. III as a specific impairment in personality functioning). Ritter et al. [35••] compared NPD patients (N=57), non-clinical controls (N=53), and patients with borderline personality disorder (N=27). This study was based on the multidimensional model of empathy [36, 37] and distinguished between cognitive and emotional empathy. Cognitive empathy [38] refers to the ability to take another person's perspective, and overlaps with the constructs of "Theory of Mind" [39] and "mentalizing" [40]. Emotional empathy [41, 42] refers to the emotional response to another person's emotional state. Both facets of empathy were assessed with the Interpersonal Reactivity Index (IRI, self-report questionnaire) [36], the Multifaceted Empathy Test (MET) [43], and the Movie for the Assessment of Social Cognition (MASC) [44]. Although self-report data ('empathic concern' - subscale of the IRI) suggested no group differences, the more ecologically valid MET task revealed that NPD patients had low emotional empathy scores relative to both control groups. With regard to cognitive empathy, self-report data ('perspective taking' - subscale of the IRI) revealed significant impairment in patients with NPD. On the more ecologically valid MET task, no deficit in cognitive empathy in NPD patients could be detected. Although the assessment of cognitive empathy by means of the sensitive MASC task revealed impairments in NPD patients, those impairments could be explained by cases with comorbid borderline personality disorder. Furthermore, in the NPD sample, the self-report measure of cognitive empathy (IRI subscale 'perspective taking') was negatively correlated with the criterion 'lack of empathy' as measured by the SCID-II interview, indicating that the DSM-IV mainly assesses the subjectively perceived deficit in cognitive empathy. Additionally, the wording for the 'perspective taking' subscale items of the IRI and the SCID-II 'lack of empathy' item seem to indicate that the deficit in cognitive empathy is best characterized as motivational. However, these data suggest that "lack of empathy" in NPD goes beyond the motivational deficit described in Sect. II in DSM-5. With regard to Sect. III of the DSM-5, these data contradict the assumption that NPD patients are not able to

recognize the feelings and needs of others. Instead, the ability to identify feelings, thoughts and intentions of others (i.e., cognitive empathy) is preserved, whereas the emotional response to another person's emotional state (i.e., emotional empathy) is restricted.

Notably, a study by Marissen et al. [45••] was unable to replicate the findings on self-reported empathy (IRI) in outpatients with NPD (N=20) compared to outpatients with cluster C personality disorders (N=20) and nonclinical controls (N=20). Differences between the two studies might account for this contradiction, such as a lower sample size, higher mean age, primarily outpatient status, low comorbidity, or inclusion of only males in the Marissen et al. study. In addition, the authors performed a facial emotion recognition task with pictures from the facial affect series including fear, anger, disgust, happiness, sadness, and neutral expressions. Results revealed that NPD patients were less accurate in facial emotion recognition than both control groups. Analyses of facial emotion expression for individual emotions revealed that differences were due to impaired recognition of fear and disgust in NPD. With regard to Sect. II of the DSM-5, these data suggest that NPD patients have impaired ability to recognize specific emotions (fear and disgust) in others. Thus, these results partially contradict the results of Ritter et al. [35••] and might be explained by the emotional specificity (i.e., only fear and disgust) of this finding, which was not analyzed in the Ritter et al. study. In sum, the NPD criterion lack of empathy which is defined differently in Sects. II and III of the DSM-5 requires further empirical exploration.

## Empathy-related Structural Brain Differences

Following the behavioral finding of impaired emotional empathy in NPD [35••], Schulze et al. [46••] conducted a structural brain imaging study with 17 patients with NPD (DSM-IV) and 17 nonclinical controls. Based on a meta-analysis of studies of empathy in nonclinical samples [47], functional brain imaging data in nonclinical individuals with narcissistic traits [48], and structural brain data from adolescents with conduct disorder [49], the anterior insular cortex was identified as crucial for emotional empathy and defined as a region of interest. Results revealed smaller gray matter (GM) volume in the left anterior insula in patients with NPD than non-clinical controls. Moreover, complementary whole-brain analyses yielded smaller GM volume in additional fronto-paralimbic brain regions comprising the rostral and median cingulate cortex, as well as the dorsolateral and medial parts of the prefrontal cortex [46••]. In sum, the results of Schulze et al. [46••] argue for specific structural alterations in empathy-related brain regions in NPD patients corresponding to deficits in emotional empathy ability as defined in DSM-5 (Sect. III).

## Shame

Following psychoanalytic theories, marked feelings of shame was included as a feature of NPD in the DSM-III. Due to revisions in the DSM-IV, feelings of shame were removed from the main criteria set and listed as one of associated features of NPD. However, clinical conceptualizations of pathological narcissism continued to consider shame as a prominent feature of narcissistic vulnerability e.g., [30, 31, 50]. According to these theories, individuals with pathological narcissism try to avoid or reduce intense feelings of shame and engage in a variety of typical intrapersonal and interpersonal strategies (e.g., aggression, fantasies, perfectionism, diverting attention away from oneself, e.g., [51•]). Furthermore, theoretical conceptualizations assume that NPD patients specifically exhibit high levels of implicit shame compared to non-clinical controls, whereas increased explicit shame might be a more general feature of psychopathology e.g., [52] for discussion see [53••].

Previous research using a small mixed clinical sample (N=26, 24 % NPD) showed a moderately positive correlation between explicit shame, measured with the Experience of Shame Scale [54], and pathological narcissism, measured with the Pathological Narcissism Inventory (PNI, [33]). Another more recent study by Ritter et al. [53••] examined shame in NPD patients without comorbid borderline personality disorder (N=28), patients with borderline personality disorder without comorbid NPD (N=31), and non-clinical controls (N=34). Explicit shame was assessed with self-report inventories (Experiential Shame Scale, ESS, [55]; Test of Self-Conscious Affects version 3, TOSCA-3, [56], while implicit shame was assessed with a modified version of the Implicit Association Task with anxiety as reference category for shame (IAT, [57]). Results revealed that explicit state shame (ESS) and explicit shame-proneness (TOSCA-3) were significantly higher in NPD patients than nonclinical controls, but significantly lower than borderline patients. Most importantly, the IAT revealed that NPD patients carried the highest levels of implicit shame-self associations (relative to anxiety-self associations) compared to both control groups. This study indicates that explicit shame and shame proneness are present in, but not specific to, NPD. Moreover, the interpretation of between-group differences in implicit shame is more challenging: On the one hand, high implicit shame might be specific to NPD. Thus, patients might be characterized by unconscious feelings of shame that they try to avoid by employing defensive behavioral strategies. On the other hand, elevated implicit shame might be less specific to NPD, as the finding can also be explained by strong anxiety-self association counterbalancing strong shame-self associations in borderline personality disorder. Although conclusions from this study must be confirmed by future research, these findings suggest that explicit and implicit shame are relevant features of NPD. Whereas sustained feelings

of shame were adopted as an associated feature of NPD in DSM-5 Sect. II, shame is not mentioned in the Sect. III definition of NPD. Further studies need to re-evaluate the specificity of high implicit shame in NPD and test short-term as well as long-term stability of implicit and explicit shame.

## Delimitation from Other Disorders

### Discriminant Validity

Similar to most personality disorders, NPD has high comorbidity rates with other psychiatric disorders in clinical and nonclinical samples e.g., [22]. Perhaps most extensively studied is the degree to which DSM criteria can be used to distinguish NPD from other personality disorders. Based on DSM-III-R criteria, Morey [58] found that NPD is one of eight personality disorders with more than 50 % overlap with at least one other personality disorder. Furthermore, most NPD criteria loaded on a common factor with antisocial personality disorder [59]. As a consequence of such data, DSM-IV criteria sets were also selected to increase specificity. Early empirical studies indicated that this goal had been achieved. Blais et al. [60] found increased specificity of the DSM-IV NPD diagnosis (compared to DSM-III), with no significant intercorrelation between NPD and other cluster B personality disorders. However, a different picture emerged in item-level analysis. Blais and Norman [61] found moderate correlations of NPD diagnosis with six items of histrionic personality disorder, four items of antisocial personality disorder and three items of paranoid personality disorder. Another study by Gunderson and Ronningstam [62] indicates that only items related to grandiosity discriminated between NPD and patients with antisocial personality disorder. Furthermore, the criterion lack of empathy was even more prevalent in antisocial personality disorder than in NPD. A further study by Holdwick et al. [63] found six out of nine NPD criteria differentiated NPD from borderline patients and six out of nine differentiated NPD from patients with antisocial personality disorder. Another study revealed that NPD criteria (similar to criteria for schizotypal and dependent personality disorder) are as highly correlated with criteria for other personality disorders as they are with each other, casting further doubt on the specificity of the NPD criteria set [64]. However, a follow up study with a larger sample size by Grilo et al. [65] found that NPD criteria correlated better with each other than with those of other personality disorders. Karterud et al. [12•] assessed SCID-II criteria in a large sample of personality disorder patients from day clinics and found that NPD criteria has a low to moderate correlation with NPD diagnosis and borderline personality disorder has the highest number of significant correlations with NPD criteria. In sum, discriminant validity of DSM-IV NPD seems to be limited, yet is quite similar to the

discriminant validity of other personality disorders. Furthermore, the definition of NPD seems to be more valid on a construct level than on a criterion level.

### Family Studies

A clinical study with twins found a heritability rate of 79 % for NPD according to DSM-III-R [66]. Compared to other personality disorders, this rate was among the highest. A non-clinical twin study applying DSM-IV criteria found much lower heritability rates for personality disorders (20–41 %), including NPD (25 %) [67]. A recent nonclinical twin study used data from self-report questionnaires *and* the structured interview for DSM-IV personality disorders [68•]. Results indicate a heritability of 71 % for NPD, which was the highest among the cluster B personality disorders. Another study used a DSM-IV based parent report for personality disorders in a sample of child twins (mean age 9 years) [69]. This study revealed a heritability rate of the NPD criteria set of 66 % (50–81 % for all personality disorders). In sum, empirical data provide strong evidence for the heritability of NPD, although the specific degree of heritability appears to be inconsistent.

### Follow-Up Study

Cross population studies showed that NPD (DSM-IV) is inversely associated with age [22]. Some follow-up studies examined the temporal stability of NPD criteria over time in the general population. However, most individuals included in those studies did not meet the full criteria for NPD (for an overview see [70•]). Ball and colleagues investigated the stability of DSM-III-R NPD items assessed by self- and informant questionnaire in patients with substance dependence [71]. This study indicates a moderately high stability ( $r=.48$ ) between baseline and follow-up after one year. Using individual growth trajectories, Lenzenweger et al. [72] found that the temporal stability of personality disorder features (according to DSM-III-R) varies considerably, as individuals showed trajectories that are stable, increasing, and decreasing. However, remission rates for narcissistic features were comparable to those of other personality disorders ( $r=.39$ ). Samuel and colleagues [73] found evidence that narcissistic criteria exhibit a moderate degree of temporal stability according to the DSM-IV (Kappa  $\kappa=.36$ ) across two years. Finally, Hopwood et al. [74] found rather low temporal stability for DSM-IV NPD criteria ( $r=.24$ ) in patients with mixed diagnoses across ten years.

Presently, only two studies have assessed the temporal stability of NPD diagnosis. Ronningstam et al. [75] assessed three-year stability of NPD using the LEAD diagnostic standard (i.e., longitudinal, expert, all data; [76]). Diagnostic criteria according to DSM-III-R and DSM-IV were applied

using the Diagnostic Interview for Narcissism [77]. According to this study, 50 % of NPD patients still met full criteria for this DSM diagnosis at a three-year follow-up. However, the sample size in this study was rather small. Thus, a recent study by Vater et al., [70•] recruited 96 patients with a diagnosis of NPD according to DSM-IV at baseline. Forty patients participated in the follow-up assessment after two years. The results indicate a moderate remission rate of 53 % for NPD as a categorical diagnosis. However, individual NPD criteria differed in their prevalence and temporal stability, similar to findings for other personality disorders e.g., [78]. Furthermore, the results of Vater et al. [70•] indicate that narcissism as a pathological personality trait (assessed with the Dimensional Assessment of Personality Pathology; DAPP-BQ [79]) did not show significant modification over two years. In sum, these results suggest that NPD, like other personality disorders, has moderate stability over time on the criterion-level, a fact that has not yet been recognized in DSM-5 descriptions of the disorder.

### Methodological Limitations

First, many of the cited studies were not designed specifically for NPD research. Thus, clinical cohorts had other primary diagnoses (e.g., other personality disorders, [80•, 90]) and epidemiological studies often did not account for general personality disorder criteria e.g., [22]. Even in research with a focus on NPD, most studies utilized mixed samples with low number of NPD cases e.g., [12•, 13]. Conclusions from these samples, which primarily consisted of sub-threshold cases with NPD-traits, might be restricted as taxometric analyses do not confirm dimensionality of the NPD construct [15]. Furthermore, assessment of NPD criteria is often critical as retrospective chart reviews or clinicians' retrospective criteria recall is used e.g., [59, 81]). Spitzer [76] proposed longitudinal patient observation, expert interviewers and the inclusion of multiple data sources to assess diagnostic criteria (LEAD standard), a procedure that was rarely adhered to in the aforementioned NPD studies, and if so, it was mostly only utilized on a diagnosis level, not on a criterion level. Furthermore, studies using group comparison designs are limited by comorbid psychiatric disorders and sub-threshold psychopathology. In sum, the existing studies carry several methodological limitations restricting the ability to evaluate the validity of the NPD (DSM-IV) criteria set.

### Conclusions, Future Research and Treatment Implications

To date, most empirical studies investigating NPD assessed the validity and reliability of the DSM criteria set. As a consequence, almost all nosological discourse preceding

DSM-5 is based on these findings e.g., [82]. However, NPD criteria and associated features have not been the subject of systematic empirical evaluation using experimental designs. As a result, DSM criteria might only partially cover the specific features of the disorder. This shortcoming is perhaps best highlighted by the criterion ‘lack of empathy’. Research investigating empathy in NPD patients has shown that deficits in this area are more complex and fine-grained than the current DSM-5 descriptions and are not captured by assessment instruments (e.g., SCID-II interview) [35••].

The inclusion of NPD in DSM-5 provides the opportunity to advance NPD research, especially as experimental psychology and neuroscience are now providing useful methodological tools. However, future NPD research must overcome several challenges. The debate over retaining NPD in the DSM-5 concluded with including the disorder twice. This decision introduced a further layer of complexity, especially with respect to sample selection. DSM-5 Sect. II adopts the DSM-IV criteria set, which captures a restricted picture of pathological narcissism by overemphasizing overt grandiose themes [11]. The NPD definition in Sect. III of the DSM-5 purports to acknowledge grandiose and vulnerable features, overtly and covertly expressed [83]. The Sect. III definition is well grounded in the clinical literature of pathological narcissism, but lacks empirical evaluation. Moreover, DSM-5 Sect. III acknowledges an underlying self-regulatory model that is primarily derived from under-evaluated clinical theories e.g., [29], as well as empirical studies relying on nonclinical samples and the Narcissistic Personality Inventory [84]. However, the Narcissistic Personality Inventory has been found to be invalid in NPD patients [85•] and initial attempts to empirically evaluate aspects of the self-regulatory model have been discouraging [34••]. Furthermore, the self-regulatory model assumes that fluctuations in affect and self-esteem are present in NPD. In the future, longitudinal studies should assess short- and long-term fluctuation of grandiose and vulnerable features of NPD (e.g., self-esteem, grandiosity, envy, shame, rage).

In order to investigate these features of NPD, future studies should adhere to a specific standard. First, the assessment of NPD is important: A diagnosis of NPD should be based on diagnostic interview and informant report, accounting for the strong discrepancy between self and informant ratings of NPD criteria [86, 87••]. Further, quality criterion-level evaluation should be conducted using the LEAD standard [76], which calls for longitudinal assessment, expert interview and inclusion of all available data. Additionally, sample selection issues should be taken into consideration. Patient recruitment methodology must acknowledge that NPD patients are more prevalent in private practice than psychiatric hospitals and day clinics [88], which presents a sample recruitment challenge. Furthermore, NPD cases in psychiatric hospitals will present with higher comorbidity rates relative to other settings, such as community-based samples [22, 89]. Finally, future studies

should use evaluated experimental designs, larger samples with NPD patients meeting full criteria and include both clinical and non-clinical control groups.

The major aim of defining diagnostic entities is to inform treatment prognosis. Predictive validity of the DSM-IV NPD criteria has been shown to be non-informative for long-term outcome under treatment-as-usual conditions [80•, 90]. Furthermore, to date, an evidence-based intervention designed specifically for NPD patients is not available. Randomized intervention studies including only NPD cases do not exist and studies assessing outcome of specific psychotherapeutic e.g., [91] or pharmacological e.g., [92, 93] interventions for personality disorders are non-informative with regard to included NPD cases (mainly due to low number of cases). Nevertheless, the NPD construct has already shown some predictive validity with respect to psychotherapeutic outcome, as trait narcissism has been associated with higher treatment drop out [94]. Thus, the future challenge on one hand is to evaluate extant specific interventions for NPD, and on the other hand, utilize accumulating knowledge of NPD to inform the further development of these interventions, which would ultimately provide empirical support for the utility of the NPD construct.

In summary, most of the extant empirical NPD research psychometrically evaluated DSM criteria assessed by self-report or clinical interview. Only very recent empirical studies apply methods from experimental psychology and neuroscience to evaluate NPD. Given the challenges presented by the two different NPD constructs in DSM-5, future research should continue to evaluate specific features of the disorder despite general psychopathology, especially with regard to the establishment of treatment strategies.

#### Compliance with Ethics Guidelines

**Conflict of Interest** Stefan Roepke and Aline Vater declare that they have no conflict of interest.

**Human and Animal Rights and Informed Consent** This article does not contain any studies with human or animal subjects performed by any of the authors.

#### References

Papers of particular interest, published recently, have been highlighted as:

- Of importance
- Of major importance

1. American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders (DSM-5). 5th ed. Washington: American Psychiatric Press; 2013.

2. Robins E, Guze SB. Establishment of diagnostic validity in psychiatric illness: its application to schizophrenia. *Am J Psychiatry*. 1970;126:983–7.
3. Ellis H. Autoeroticism: A psychological study. *Alienist Neurol*. 1989;19:260–99.
4. Freud S. On narcissism: An introduction. Standardth ed. London: Hogarth Press; 1914.
5. Kernberg O. Borderline conditions and pathological narcissism. New York: Jason Aronson; 1975.
6. Kohut H. Forms and transformations of narcissism. *J Am Psychoanal Assoc*. 1966;14:243–72.
7. American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders (DSM-III). 3rd ed. Washington: American Psychiatric Press; 1980.
8. Frances A. The DSM-III personality disorders section: a commentary. *Am J Psychiatry*. 1980;137:1050–4.
9. American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders (DSM-III-R), 3rd rev ed. Washington: American Psychiatric Press; 1987.
10. American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders (DSM-IV). 4th ed. Washington: American Psychiatric Press; 1990.
11. Cain NM, Pincus AL, Ansell EB. Narcissism at the crossroads: Phenotypic description of pathological narcissism across clinical theory, social/personality psychology, and psychiatric diagnosis. *Clin Psychol Rev*. 2008;28:638–56.
12. Karterud S, Oien M, Pederson G. Validity Aspects of the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition, narcissistic personality disorder construct. *Compr Psychiatry*. 2011;52: 517–26. *A validity study of the NPD construct challenging the notion that the DSM-IV criteria for NPD capture a distinct diagnostic category.*
13. Miller JD, Hoffman BJ, Campbell WK, Pilkonis PA. An examination of the factor structure of Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition, narcissistic personality disorder criteria: one or two factors? *Compr Psychiatry*. 2008;49:141–5.
14. Kubarych TS, Aggen SH, Kendler KS, Torgersen S, Reichborn-Kjennerud T, Neale MC. Measurement non-invariance of DSM-IV narcissistic personality disorder criteria across age and sex in a population-based sample of Norwegian twins. *Int J Methods Psychiatr Res*. 2010;19:156–66.
15. Fossati A, Beauchaine TP, Grazioli F, Carretta I, Cortinovis F, Maffei C. A Latent structure analysis of diagnostic and statistical manual of mental disorders, fourth edition, narcissistic personality disorder criteria. *Compr Psychiatry*. 2005;46:361–7.
16. Blais MA, Hilsenroth MJ, Castlebury FD. Content validity of the DSM-IV borderline and narcissistic personality disorder criteria sets. *Compr Psychiatry*. 1997;38:31–7.
17. Russ E, Shedler J, Bradley R, Westen D. Refining the construct of narcissistic personality disorder: Diagnostic criteria and subtypes. *Am J Psychiatry*. 2008;165:1473–81.
18. Torgersen S, Kringlen E, Cramer V. The prevalence of personality disorders in a community sample. *Arch Gen Psychiatry*. 2001;58: 590–6.
19. Lenzenweger MF, Lane MC, Loranger AW, Kessler RC. DSM-IV personality disorders in the National Comorbidity Survey Replication. *Biol Psychiatry*. 2007;62:553–64.
20. Samuels J, Eaton WW, Bienvenu 3rd OJ, Brown CH, Costa Jr PT, Nestadt G. Prevalence and correlates of personality disorders in a community sample. *Br J Psychiatry*. 2002;180:536–42.
21. Coid J, Yang M, Tyrer P, Roberts A, Ullrich S. Prevalence and correlates of personality disorder in Great Britain. *Br J Psychiatry*. 2006;188:423–31.
22. Stinson FS, Dawson DA, Goldstein RB, Chou SP, Huang B, Smith SM, et al. Prevalence, correlates, disability, and comorbidity of DSM-IV narcissistic personality disorder: results from the wave 2 national epidemiologic survey on alcohol and related conditions. *J Clin Psychiatry*. 2008;69:1033–45.
23. Dhawan N, Kunik ME, Oldham J, Coverdale J. Prevalence and treatment of narcissistic personality disorder in the community: a systematic review. *Compr Psychiatry*. 2010;51:333–9.
24. Trull TJ, Jahng S, Tomko RL, Wood PK, Sher KJ. Revised NESARC Personality Disorder Diagnoses: Gender, Prevalence, and Comorbidity with Substance Dependence Disorders. *J Personal Disord*. 2010;24:412–26.
25. Grilo CM. Are there gender differences in DSM-IV personality disorders? *Compr Psychiatry*. 2002;43:427–30.
26. Zimmerman M, Rothschild L, Chelminski I. The prevalence of DSM-IV personality disorders in psychiatric outpatients. *Am J Psychiatry*. 2005;162:1911–8.
27. Soeteman DI, Hakkaart-van Roijen L, Verheul R, Busschbach JJ. The economic burden of personality disorders in mental health care. *J Clin Psychiatry*. 2008;69:259–65.
28. Lynam DR, Widiger TA. Using a general model of personality to understand sex differences in the personality disorders. *J Personal Disord*. 2007;21:583–602.
29. Morf CC, Rhodewalt F. Unraveling the Paradoxes of Narcissism: A Dynamic Self-Regulatory Processing Model. *Psychol Inq*. 2001;12: 177–96.
30. Horowitz M. Clinical phenomenology of narcissistic pathology. *Psychiatr Ann*. 2009;39:124–8.
31. Ronningstam E. Narcissistic personality disorder: a current review. *Curr Psychiatr Rep*. 2010;12:68–75.
32. Jordan CH, Spencer SJ, Zanna MP, Hoshino Browne E, Correll J. Secure and defensive high self-esteem. *J Pers Soc Psychol*. 2003;85:969–78.
33. Pincus AL, Ansell EB, Pimentel CA, Cain NM, Wright AGC, Levy KN. Initial construction and validation of the Pathological Narcissism Inventory. *Psychol Assess*. 2009;21:365–79.
34. Vater A, Ritter K, Schröder-Abé M, Schütz A, Lammers CH, Bosson J, et al. When Grandiosity and Vulnerability Collide: Implicit and Explicit Self-Esteem in Patients with Narcissistic Personality Disorder. *J Behav Ther Exp Psychiatry*. 2013;44:37–47. *Study providing evidence for low explicit self-esteem in NPD compared to non-clinical controls.*
35. Ritter K, Dziobek I, Preißler S, Rüter A, Vater A, Fydrich T, et al. Lack of empathy in patients with narcissistic personality disorder. *Psychiatry Res*. 2011;187:241–7. *First empirical assessment of the 'Lack of empathy' criterion in NPD patients indicating deficits in emotional empathy with cognitive empathy within normal limits.*
36. Davis MH. Measuring Individual-Differences in Empathy - Evidence for a Multidimensional Approach. *J Pers Soc Psychol*. 1983;44:113–26.
37. Blair RJ. Responding to the emotions of others: dissociating forms of empathy through the study of typical and psychiatric populations. *Conscious Cogn*. 2005;14:698–718.
38. Baron-Cohen S, Wheelwright S. The empathy quotient: an investigation of adults with Asperger syndrome or high functioning autism, and normal sex differences. *J Autism Dev Disord*. 2004;34:163–75.
39. Premack D, Woodruff G. Chimpanzee problem-solving: a test for comprehension. *Science*. 1978;202:532–5.
40. Frith U, Frith CD. Development and neurophysiology of mentalizing. *Philosophical Transactions of the Royal Society of London. Series B. Biol Sci*. 2003;358:459–73.
41. Mehrabian A, Epstein N. A measure of emotional empathy. *J Pers*. 1972;40:525–43.
42. Eisenberg N, Miller PA. The relation of empathy to prosocial and related behaviors. *Psychol Bull*. 1987;101:91–119.
43. Dziobek I, Rogers K, Fleck S, Bahnmann M, Heekeren HR, Wolf OT, et al. Dissociation of cognitive and emotional empathy in adults with Asperger syndrome using the multifaceted empathy test (MET). *J Autism Dev Disord*. 2008;38:464–73.

44. Dziobek I, Fleck S, Kalbe E, Rogers K, Hassenstab J, Brand M, et al. Introducing MASC: a movie for the assessment of social cognition. *J Autism Dev Disord.* 2006;36:623–36.
45. Marissen MAE, Deen ML, Franken IHA. Disturbed emotion recognition inpatients with narcissistic personality disorder. *Psychiatry Res.* 2012;198:269–73. *Experimental facial emotion recognition study in NPD patients indicating deficits in fear and disgust labelling.*
46. Schulze L, Dziobek I, Vater A, Heekeren HR, Bajbouj M, Renneberg B, et al. Gray Matter Abnormalities in Patients with Narcissistic Personality Disorder. *J Psychiatr Res.* 2013;47:1363–9. *First structural brain data from patients with NPD indicating lower gray matter density in empathy-related brain regions such as the anterior insular and further fronto-limbic structures.*
47. Fan Y, Duncan NW, de Greck M, Northoff G. Is there a core neural network in empathy? An fMRI based quantitative meta-analysis. *Neurosci Biobehav Rev.* 2011;35:903–11.
48. Fan Y, Wonberger C, Enzi B, de Greck M, Ulrich C, Tempelmann C, et al. The narcissistic self and its psychological and neural correlates: an exploratory fMRI study. *Psychol Med.* 2011;41:1641–50.
49. Sterzer P, Stadler C, Poustka F, Kleinschmidt A. A structural neural deficit in adolescents with conduct disorder and its association with lack of empathy. *Neuroimage.* 2007;37:335–42.
50. Kernberg O. Narcissistic personality disorders: Part I. *Psychiatr Ann.* 2009;39:105–66.
51. Schoenleber M, Berenbaum H. Shame regulation in personality pathology. *J Abnorm Psychol.* 2012;121:433–46. *Conceptual paper on maladaptive shame regulation strategies in NPD and other personality disorders.*
52. Tracy JL, Robins RW. Putting the self into self-conscious emotions: a theoretical model. *Psychol Inq.* 2004;15:103–25.
53. Ritter K, Vater A, Rüsçh N, Schröder-Abé M, Schütz A, Fydrich T, Lammers CH, Roepke S. Shame in patients with narcissistic personality disorder. *Psychiatry Res.* 2014;215(2):429–37. doi:10.1016/j.psychres.2013.11.019. *First empirical assessment of implicit and explicit shame in patients with NPD indicating the relevance of this emotion.*
54. Andrews B, Qian M, Valentine JD. Predicting depressive symptoms with a new measure of shame: the experience of shame scale. *Br J Clin Psychol.* 2002;41:29–42.
55. Turner JE. An Investigation of Shame Reactions, Motivation, and Achievement in a Difficult College Course (Unpublished Doctoral Dissertation). Austin: University of Texas; 1998.
56. Tangney JP, Dearing RL, Wagner PE, Gramzow R. The Test of Self-Conscious Affect-3 (TOSCA-3). George Mason University; 2000.
57. Rüsçh N, Lieb K, Göttler I, Hermann C, Schramm E, Richter H, et al. Shame and implicit self-concept in women with borderline personality disorder. *Am J Psychiatry.* 2007;164:500–508.
58. Morey LC. Personality disorders in DSM-III and DSM-III-R: convergence, coverage, and internal consistency. *Am J Psychiatry.* 1988;145:573–7.
59. Morey LC. The categorical representation of personality disorder: a cluster analysis of DSM-III-R personality features. *J Abnorm Psychol.* 1988;97:314–21.
60. Blais MA, Hilsenroth MJ, Castlebury FD. Psychometric characteristics of the cluster B personality disorder under DSM-III-R and DSM-IV. *J Personal Disord.* 1997;11:270–8.
61. Blais MA, Norman DK. A psychometric evaluation of the DSM-IV personality disorder criteria. *J Personal Disord.* 1997;11:168–76.
62. Gunderson JG, Ronningstam E. Differentiating narcissistic and antisocial personality disorders. *J Personal Disord.* 2001;15:103–9.
63. Holdwick Jr DJ, Hilsenroth MJ, Castlebury FD, Blais MA. Identifying the unique and common characteristics among the DSM-IV antisocial, borderline, and narcissistic personality disorders. *Compr Psychiatry.* 1998;39:277–86.
64. Grilo CM, McGlashan TH. Convergent and discriminant validity of DSM-IV axis II personality disorder criteria in adult outpatients with binge eating disorder. *Compr Psychiatry.* 2000;41:163–6.
65. Grilo CM, McGlashan TH, Morey LC, Gunderson JG, Skodol AE, Shea MT, et al. Internal consistency, intercriteria overlap and diagnostic efficiency of criteria sets for DSM-IV schizotypal, borderline, avoidant and obsessive-compulsive personality disorders. *Acta Psychiatr Scand.* 2001;104:264–72.
66. Torgersen S, Lygren S, Oien PA, Skre I, Onstad S, Edvardsen J, et al. A twin study of personality disorders. *Compr Psychiatry.* 2000;41:416–25.
67. Kendler KS, Aggen SH, Czajkowski N, Røysamb E, Tambs K, Torgersen S, et al. The structure of genetic and environmental risk factors for DSM-IV personality disorders: A multivariate twin study. *Arch Gen Psychiatry.* 2008;65:1438–46.
68. Torgersen S, Myers J, Reichborn-Kjennerud T, Røysamb E, Kubarych TS, Kendler KS. The heritability of Cluster B personality disorders assessed both by personal interview and questionnaire. *J Personal Disord.* 2012;26:848–66. *Twin study revealing .71 heritability of NPD when including self-report and clinical interview.*
69. Coolidge FL, Thede LL, Jang KL. Heritability of personality disorders in childhood: a preliminary investigation. *J Personal Disord.* 2001;15:33–40.
70. Vater A, Ritter K, Strunz S, Ronningstam E, Renneberg B, Roepke S. Stability of Narcissistic Personality Disorder: Tracking Categorical and Dimensional Rating Systems Over a Two-Year Period. *Personal Disord.* 2014 Feb 10. [Epub ahead of print]. Follow-up study that provides evidence for moderate stability of NPD across two years.
71. Ball SA, Rounsaville BJ, Tennen H, Kranzler HR. Reliability of personality disorder symptoms and personality traits in substance-dependent inpatients. *J Abnorm Psychol.* 2001;110:341–52.
72. Lenzenweger MF, Johnson MD, Willett JB. Individual growth curve analysis illustrates stability and change in personality disorder features: The Longitudinal Study of Personality Disorders. *Arch Gen Psychiatry.* 2004;61:1015–24.
73. Samuel DB, Hopwood CJ, Ansell EB, Morey LC, Sanislow CA, Yen S, et al. Comparing the temporal stability of self-report and interview assessed personality disorder. *J Abnorm Psychol.* 2011;120:670–80.
74. Hopwood CJ, Morey LC, Donnellan MB, Samuel DB, Grilo CM, McGlashan TH, et al. Ten-year rank-order stability of personality traits and disorders in a clinical sample. *J Pers.* 2013;81:335–44.
75. Ronningstam EF, Gunderson J, Lyons M. Changes in pathological narcissism. *Am J Psychiatr.* 1995;152:253–7.
76. Spitzer RL. Psychiatric diagnosis: are clinicians still necessary? *Compr Psychiatry.* 1983;24:399–411.
77. Gunderson JG, Ronningstam E, Bodkin A. The diagnostic interview for narcissistic patients. *Arch Gen Psychiatry.* 1990;47:676–80.
78. McGlashan TH, Grilo CM, Ralevski E, Morey LC, Gunderson JG, Skodol AE, et al. Two-year prevalence and stability of individual DSM-IV criteria for schizotypal, borderline, avoidant, and obsessive-compulsive personality disorders: toward a hybrid model of axis II disorders. *Am J Psychiatry.* 2005;162:883–9.
79. Livesley WJ, Jackson DN. Manual for the dimensional assessment of personality pathology. Port Huron: Sigma Press; 2002.
80. Morey LC, Hopwood CJ, Markowitz JC, Gunderson JG, Grilo CM, McGlashan TH, et al. Comparison of alternative models for personality disorders, II: 6-, 8- and 10-year follow-up. *Psychol Med.* 2012;42:1705–13. *Analysing data from the CLPS study, results reveal that the DSM-IV NPD diagnosis did not predict 10-year outcome under treatment as usual conditions.*
81. Morey LC. A psychometric analysis of five DSM-III categories. *Personal Individ Differ.* 1985;6:323–9.



82. Alarcón RD, Sarabia S. Debates on the narcissism conundrum: trait, domain, dimension, type, or disorder. *J Nerv Ment Dis.* 2012;200:16–25.
83. Skodol AE, Bender DS, Morey LC. Narcissistic Personality Disorder in DSM-5. *Personal Disord.* 2013 Jul 8. [Epub ahead of print].
84. Raskin RN, Hall CS. A narcissistic personality inventory. *Psychol Rep.* 1979;45:590.
85. Vater A, Schröder-Abé M, Ritter K, Schulze L, Renneberg B, Bosson J, et al. The Narcissistic Personality Inventory - a useful tool for assessing pathological narcissism? Evidence from patients with Narcissistic Personality Disorder. *J Pers Assess.* 2013;95:301–8. *Study showing that the Narcissistic Personality Inventory is not useful for clinical research.*
86. Dowson JH. DSM-III-R narcissistic personality disorder evaluated by patients' and informants' self-report questionnaires: relationships with other personality disorders and a sense of entitlement as an indicator of narcissism. *Compr Psychiatry.* 1992;33:397–406.
87. Cooper LD, Balsis S, Oltmanns TF. Self- and informant-reported perspectives on symptoms of narcissistic personality disorder. *Personal Disord.* 2012;3:140–54. *Study showing low self- and informant-report agreement among DSM-IV diagnostic criteria and the categorical diagnosis of NPD.*
88. Westen D. Divergences between clinical and research methods for assessing personality disorders; Implications for research and the evolution of Axis II. *Am J Psychiatry.* 1997;154:895–903.
89. Ritter K, Roepke S, Merkl A, Heuser I, Fydrich T, Lammers CH. Comorbidity in patients with narcissistic personality disorder in comparison to patients with borderline personality disorder. *Psychother Psychosom Med Psychol.* 2010;60:14–24.
90. Hopwood CJ, Malone JC, Ansell EB, Sanislow CA, Grilo CM, McGlashan TH, et al. Personality assessment in DSM-5: empirical support for rating severity, style, and traits. *J Personal Disord.* 2011;25:305–20.
91. Bamelis LLM, Evers SMAA, Spinhoven P, Arntz A. Results of a Multicenter Randomized Controlled Trial of the Clinical Effectiveness of Schema Therapy for Personality Disorders. *Am J Psychiatry.* 2014. doi:10.1176/appi.ajp.2013.12040518.
92. Roepke S, Merkl A, Dams A, Ziegenhorn A, Anghelescu IG, Heuser I, et al. Preliminary evidence of improvement of depressive symptoms but not impulsivity in cluster B personality disorder patients treated with quetiapine: an open label trial. *Pharmacopsychiatry.* 2008;41:176–81.
93. Hollander E, Tracy KA, Swann AC, Coccaro EF, McElroy SL, Wozniak P, et al. Divalproex in the treatment of impulsive aggression: efficacy in cluster B personality disorders. *Neuropsychopharmacology.* 2003;28:1186–97.
94. Ogrodniczuk JS, Piper WE, Joyce AS, Steinberg PI, Duggal S. Interpersonal problems associated with narcissism among psychiatric outpatients. *J Psychiatr Res.* 2009;43:837–42.