REVIEW PAPER

The Effects of DSM5 Autism Diagnostic Criteria on Number of Individuals Diagnosed with Autism Spectrum Disorders: A Systematic Review

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Received: 19 March 2014 / Accepted: 25 March 2014 / Published online: 21 June 2014 © Springer Science+Business Media New York 2014

Abstract A systematic review of empirical papers comparing the application of DSM-IV and DSM5 diagnostic criteria for Autism Spectrum Disorders identified 12 papers. The application of DSM5 diagnostic criteria resulted in an approximately one third reduction in Autism Spectrum Disorders. The reduction was approximately two thirds for mild forms of Autism. The implications for practice and research are discussed.

Keywords Autism · Asperger syndrome · DSM-IV · DSM5 · Diagnosis

The American Psychiatric Association (2013) published revised diagnostic criteria for autism spectrum disorders (ASD) which involved substantial changes. These changes included (1) elimination of individual categories of ASD, such as autistic disorder, pervasive developmental disorder not otherwise specific, and Asperger syndrome; (2) introduction of coding of disorder severity; (3) reduction in the number of symptom domains from three to two by collapsing DSM-IV's domains of communication and social behavior into one domain; (4) changes in the number of symptoms required for diagnosis in each of the domains; and (5) introduction of a new disorder, social (pragmatic) communication disorder. Although these new criteria have been approved, they were not subject to extensive empirical evaluation prior to development.

Between 2012 and 2013, a number of studies have been reported that appear to demonstrate that DSM5 criteria reduce the diagnosis of ASD, which has raised considerable concern (Tsai and Ghazuiddin 2013); however, there has been no

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P. Sturmey (⊠) · S. Dalfern Queens College, City University of New York, New York, NY, USA e-mail: Peter.Sturmey@qc.cuny.edu systematic review and synthesis of this literature. Further, it is unclear if this possible reduction of ASD diagnoses will be uniform across all individuals with ASD or whether it will impact certain subgroups of individuals formerly diagnosed with DSM-IV ASD. Therefore, the aim of this paper was to systematically search the literature to locate all current empirical papers comparing DSM-IV and DSM5 criteria for ASD and to synthesize their findings.

Method

Procedure

On 24 June 2013, the second author searched PschINFO© using the keywords "Autism" and "DSM 5". The inclusion criterion was that the article had to compare DSM-IV and DSM5 diagnostic criteria for autism spectrum disorders empirically. Articles that reported sensitivity and specificity, factor analytic studies, reviews, and editorials were excluded. The search identified 995 articles of which 13 studies met inclusion criteria. On 15 July 2013, the second author gathered studies using the terms "Autism AND DSM 5 AND DSM 4" on PschINFO©. The second search identified 436 articles of which 19 articles met inclusion criteria. These searches yielded 32 non-overlapping articles that met inclusion criteria. On 29 July 2013, the second author searched Google Scholar© for all articles that cited these 32 articles that also met the inclusion criteria. This search identified 24 articles that met criteria. These searches yielded 56 nonoverlapping articles of which 27 reported data and 29 did not. Finally, the second author categorized the 27 empirical articles into those that (a) directly compared DSM-IV and DSM5 criteria; (b) evaluated the psychometric properties of DSM5 criteria, such as specificity and specificity; and (c) other empirical studies. The study by Huerta et al. (2012) was excluded since although, this study's abstract reported a reduction of 9 % in the number of

Article	Population	Method	
Mattila et al. (2011)	5,484 children aged 8 years	Instruments were the Autism Spectrum Screening Questionnaire, ADI-R, ADOS, and WISCR by a clinical and a research psychologist. School-day observations of 24 children were undertaken. Patient records were studied	
Gibbs et al. (2012)	N=132; aged 2 to 16 years ($M=6.06$ years)	Informal observations, ADOS, ADI-R, and an ADI-R were completed. Information gathered from background reports, information from teachers, or other professionals.	
Matson et al. (2012a)	227 adults aged 18–88 years with ID: those who only met criteria for DSM-IV-TR ASD, those who met DSM5 criteria, and controls with ID not meeting either criteria for autism	Participants were all residents of developmental centers previously diagnosed with ID. Professionals and doctoral students administered both the ASD-DA and DSM-IV-TR/ICD-10 checklist	
Matson et al. (2012b)	2,493 caregivers and their toddlers with varied biomedical conditions, developmental delays, mild ID, ASDs, PDD-NOS; mean age=26 months (<i>M</i> =25.88)	BISCUIT-part 1, M-CHAT, and BDI-2. All ASD diagnoses were made by a licensed doctoral level psychologist with over 30 years of experience who had access to the information gathered from the assessment administrations and a record review, and made diagnoses based on clinical judgment using the DSM-IV-TR and the DSM-5 criteria	
Matson et al. (2012c)	2,721 toddlers at risk for DDs	Pediatricians and healthcare professionals previously identified the toddlers as at risk for DD. A licensed doctoral psychologist administered the BISCUIT-part 1, the M-CHAT, and BDI-2°	
Taheri and Perry (2012)	131 children previously diagnosed with either autistic disorder or PDD-NOS aged 2–12 years	Mullen, S-BIS, WPPSI or WPPSI, WISC, VABS, and CARS	
Worley and Matson (2012)	208 children with ASD aged 3–16 years	Participants were recruited though advocacy groups, support groups, schools, and through an outpatient clinic. The ASD-DC and the DSM-IV-TR/ICD-10 checklist were completed by the parents or caregivers	
Mayes et al. (2013)	n=125 psychiatric sample: 25 children with high-functioning autism, 25 children with low-functioning autism, 25 children with a clinical diagnosis of PDDNOS, 25 children with diagnoses other than autism (ADHD, ODD, MA, etc.); 1–16 years of age ($M=6.7$); pediatric sample, $n=25$ —9 children have LFA, 4 have HFA, 5 have PDDNOS, and 7 have developmental delays without autism. The children are 1–3 years of age ($M=2.6$)	Completion of the CASD, a review of early intervention records, and standardized developmental and language test scores	
Mazefsky et al. (2013)	498 high-functioning verbal participants with ASD mean IQ 105; 5–61 years; mean age of 21.8	Certified and trained diagnosticians administered the ADOS and ADI-R. All participants had a clinical diagnosis of an ASD which was confirmed by expert clinical opinion and supported by standard DSM-IV-based cut-offs on the ADOS and ADI-R	
Rieske et al. (2013)	424 children and adolescents aged 2–18 years, a control group (124), a DSM-V diagnosis group (192), a DSM-IV diagnosis group (108)	The DSM-IV-TR/ICD-10 checklist caregivers were told to complete the measure based on their behavioral observations of the child, the ASD-CC	
Wilson et al. (2013)	150 intellectually able adults; 18–65 years, mean age, 32	Instruments included the ADI-R, ADOS-G, OCI-R, the Barkley's Current and Childhood Symptom Scales, and HADS. Information was compiled by a consultant psychiatrist, junior doctor, and ADI-R/ADOS-G administrator (nurse or psychologist)	
Young and Rodi (2013)	233 participants; 23 participants did not meet criteria for a PDD diagnosis based on DSM-IV-TR criteria (age range, 12 months–54 years, <i>M</i> =9 years, SD=8.89 years)	Participants were either self-referred or referred by a medical or allied health practitioner for a diagnostic assessment conducted by a single clinician or a dual team, which was comprised of a speech pathologist and psychologist. The ADI-R and the CARS were used. Information was obtained from parent questionnaires, the social communication questionnaire, and structured observation	

searched PubMed[©] and Google Scholar[©] using the terms "Autism" AND "DSM5". This search located an additional

two articles of which one met inclusion criteria. Twelve articles that met the inclusion criterion were retained.

The second author then tabulated features of the retained studies including sample characteristics and methods and instruments used for diagnosis. We distinguished two types of studies. Studies using method 1 compared diagnostic criteria in heterogeneous populations, such as large samples of individuals who may or may have ASD or clinic samples of individuals who may or may not have a variety of disorders, including ASD who were subsequently evaluated for DSM-IV and DSM5 ASD. Studies using method 2 compared diagnostic criteria in homogenous populations of individuals who had or were very likely to have ASD, such as studies of individuals all of whom met DSM-IV or clinical criteria for ASD who were subsequently evaluated for DSM5 ASD.

The authors then identified or calculated the percentage change in the proportion of individuals who met diagnostic criteria for ASD. The percentage change in the proportion of individuals with ASD was calculated by subtracting the number of individuals with DSM-IV ASD from the number of individuals with DSM5 ASD and expressing it as a percentage of the number of individuals with DSM-IV ASD. Where data were reported for subgroups, such as individuals with autism, pervasive developmental disorders not otherwise specified (PDD-NOS), Asperger syndrome, higher functioning ASD, etc.; the second author also calculated these data separately for each group. paper included a wide range of ages with participants ranging in age from 17 months to 88 years. Seven studies included children aged 5 or younger and four included participants aged over 21 years. Studies also included the full range of degrees of ASD from individuals at risk for various disorders, PDD-NOS, and autism disorder. The samples were also varied in the degree of intellectual disability (ID) from average intellect to profound ID. Finally, the sample came from a wide range of settings including preschoolers living at home, school, special education, community, and institutional populations. Thus, the samples in these papers were highly varied. Sample sizes were adequate to large ranging from 131 to 5,484 participants. Only one paper (Wilson et al. 2013) used method 1 (heterogeneous sample), and the remaining 11 papers used method 2 (homogenous samples).

The median overall change in diagnosis of ASD from all papers was -36.97 % = (range, -7 to -54 %). When changes were compared between less impaired subgroups (Asperger disorder, high functioning autism, and PDDNOS) and more impaired subgroups (AD, low IQ), there were large differences. The median reductions in the more impaired subgroups were -19.35 % (range, 0 to -26.3 %) and in the less impaired subgroups were -71.27 % (range, -16.6 to -100 %; Table 2).

Discussion

Results

Table 1 displays the methodological features and percentage change in the diagnosis of DSN-IV and DSM5 ASD. The

This systematic review of 12 empirical papers comparing the application of DSM-IV and DSM5 diagnostic criteria for ASD found consistent data across studies showing a median reduction of about a third. Reductions were not uniform, but were much smaller in individuals with more severe forms of ASD, but were approximately two thirds for less severe forms of

Table 2 The overall reduction in diagnosis of autism spectrum disorders from 12 studies

Article	Overall results	Less Impaired	More Impaired
Mattila et al. (2011)	54 % Reduction <i>N</i> =12/26	AD: 100 % reduction <i>N</i> =0/11 High IQ: 63.6 % reduction <i>N</i> =8/22 IQ >70	Low IQ: 0 % reduction <i>N</i> =4/4 IQ=50–69
Gibbs et al. (2012)	23 % Reduction N=85/111	AD: 16.6 % reduction <i>N</i> =15/18 PDDNOS: 50 % reduction <i>N</i> =17/34	ASD:10.2 % reduction N=53/59
Matson et al. (2012a)	36.53 % reduction N=99/156		
Matson et al. (2012b)	47.74 % reduction N=404/773		
Matson et al. (2012c)	47.79 % reduction N=415/795	PDDNOS: 78.94 % reduction <i>N</i> =72/342	ASD: 24.28 % reduction N=343/453
Taheri and Perry (2012)	37.40 % reduction N=82/131	PDDNOS: 83.33 % reduction <i>N</i> =6/36	ASD: 19.35 % reduction N=75/93
Worley and Matson (2012)	32.3 % reduction N=121/180		
Mayes et al. (2013)	14 % reduction N=70/83		
Mazefsky et al. (2013)	7 % reduction N=463/498		
Rieske et al. (2013)	36 % reduction <i>N</i> =192/300		
Wilson et al. (2013)	24 % reduction <i>N</i> =61/80		
Young and Rodi (2013)	42.86 % reduction N=120/210	AD: 38.6 % reduction <i>N</i> =64/114 PDDNOS: 100 % reduction <i>N</i> =0/20	ASD: 26.3 % reduction <i>N</i> =56/76

ASD. We can be confident that these findings are robust because they were robust across populations, ages, methods, and researchers. Other data sources confirm that DSM5 criteria exclude individuals with mild forms of ASD. First, Mayes et al. (2014) found that DSM5 criteria were less likely to identify individuals with ASD compared to two psychometric measures and clinical diagnoses of ASD. Second, several studies have found that individuals who meet DSM5 criteria have more severe disabilities than individuals that meet DSM-IV, but not DSM5 criteria for ASD (Matson et al. 2012a; McPartland et al. 2012).

These findings have important implications for services and research. If services apply DSM5 criteria, this may lead to exclusion of many individuals with mild forms of ASD. It is unclear if this will actually happen since practitioners routinely apply diagnoses inaccurately based more on unmet child need than truth (MacMillan et al. 1996). Although practitioners and family members may be eager to diagnostically avoid labels such as ID in preference for more acceptable labels such as specific learning disabilities or emotional disorders, they may be less motivated to avoid diagnostic labels such as ASD, which sometimes give access to better services and are less stigmatizing that other labels. Only future monitoring of patterns of service use will show what may happen.

The American Psychiatric Association partially acknowledged the exclusion of mild forms of ASD when the added rider to the diagnostic criteria was that "Individuals with a well-established DSM-IV diagnosis of autistic disorder, Asperger's disorder, or pervasive developmental disorder not otherwise specified should be given the diagnosis of autism spectrum disorder", but this approach to diagnosis of autism is strange. It enables individuals who previously were diagnosed with ASD to continue to be eligible for services, but results in two odd consequences. First, consider two individuals with identical symptoms who meet DSM-IV criteria for ASD, one of whom was diagnosed before the publication of DSM5 and one diagnosed after: The first will be diagnosed with ASD and eligible for services, but the second will not. The second consequence is that the meaning of the DSM5 term ASD is obfuscated as it refers to different entities using the same term.

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