Validity of the Korean Version of DIVA-5: A Semi-Structured Diagnostic Interview for Adult ADHD

Minha Hong 1
JJ Sandra Kooij 2,3
Bongseog Kim 4
Yoo-Sook Joung 5
Hanik K Yoo 6
Eui-Jung Kim 7
Soyoung Irene Lee 8
Soo-Young Bhang 9
Seung Yup Lee 10
Doug Hyun Han 11
Young Sik Lee 11
Geon Ho Bahn 10

1Department of Psychiatry, Myongji Hospital, Hanyang University College of Medicine, Goyang, Korea; 2PsyQ, Psycho-Medical Programs, Expertise Center Adult ADHD, The Hague, The Netherlands; 3Department of Psychiatry, Amsterdam University Medical Center, Amsterdam, The Netherlands; 4Department of Psychiatry, Inje University College of Medicine, Seoul, Korea; 5Department of Clinical Psychiatry, Seoul Brain Research Institute, Seoul, Korea; 6Department of Psychiatry, College of Medicine, Ewha Womans University, Seoul, Korea; 7Department of Psychiatry, Sookchunhyang University College of Medicine, Bucheon, Korea; 8Department of Psychiatry, Nowon Eulji University Hospital, Eulji University School of Medicine, Seoul, Korea; 9Department of Psychiatry, Sungkyunkwan University School of Medicine, Seoul, Korea; 10Department of Psychiatry, Kyung Hee University School of Medicine, Seoul, Korea; 11Department of Psychiatry, Chung-Ang University College of Medicine, Seoul, Korea

Correspondence: Geon Ho Bahn Department of Psychiatry, Kyung Hee University School of Medicine, 23 Kyungheedae-Ro, Dongdaemun-gu, Seoul 02447, Korea Tel +82-2-958-8556 Fax +82-2-957-1997 Email mmopeian@khu.ac.kr

Background: This study was to assess the validity of the Korean version of the semi-structured Diagnostic Interview for Adult ADHD, third edition (DIVA-5). The secondary aim was to compare sociodemographic and psychiatric comorbidities in adult patients with and without a diagnosis of ADHD.

Methods: A total of 279 participants were recruited from nine psychiatric outpatient clinics in Korea. All participants were administered the Mini-International Neuropsychiatric Interview (MINI) Plus v.5.0.0, the Adult ADHD Self-Report Scale v1.1 (ASRS-v1.1) Symptom Checklist, and DIVA-5. Diagnosis concordance between two board-certified psychiatrists and DIVA-5 were analysed.

Results: The DIVA-5 showed a diagnostic accuracy of 92%, a sensitivity of 91.30%, and a specificity of 93.62%. Significant clinical and demographic differences between ADHD and control groups were found.

Conclusion: The Korean version of DIVA-5 is a reliable tool for assessing and diagnosing ADHD in adult Korean populations.

Keywords: adult ADHD, validity, DIVA-5, Korean

Introduction
Attention deficit/hyperactivity disorder (ADHD), which was formerly classified under the “disorders usually first diagnosed in infancy, childhood, or adolescence” section, was introduced as one of the neurodevelopmental disorders in the Diagnostic and Statistical Manual of Mental Disorders, fifth edition (DSM-5).1 ADHD prevalence in adulthood is estimated to be between five and eight percent,2 and the persistence of ADHD in children into adulthood varies between 60% and 80%.3,4 Numerous negative outcomes, such as criminality, academic failure, unemployment, behavioural disturbance, addiction, accidents, and premature death, have been reported to be related to ADHD.5,6

Although early diagnosis and early intervention is needed in adults, as well as in children and adolescents, there is no single gold standard tool for the diagnosis of ADHD. A detailed interview by experienced clinicians, and information gathered from informant(s) in addition to the patient are highlighted when diagnosing ADHD.7

To our knowledge, there are only two semi-structured interviews available for the accurate diagnostic assessment of ADHD, based on the DSM criteria in the
adult population: Conners’ Adult ADHD Diagnostic Interview for DSM-IV (CAADID), and the Diagnostic Interview for ADHD in Adults (DIVA-2). However, no official and commercial version of the CAADID is available in the Korean language. The DIVA-2 is a semi-structured diagnostic interview that is available on the DIVA Foundation website. The DIVA has recently been updated according to the DSM-5 criteria, and DIVA-5 is currently available in 12 translated versions, with many more to come. In 2016, the DIVA-2 was revised into DIVA-5 based on DSM-5 criteria by Dr P Asherson of the London Institute of Psychiatry (London, UK), Dr A Ramos-Quirorga of the University of Barcelona (Barcelona, Spain), and S Kooij and A Bron of the DIVA Foundation. The DSM-5 criteria for ADHD, issued with the permission of the American Psychiatric Association, was adopted. The previous version, DIVA-2, has been validated in Swedish and Spanish, but none of the DIVA-5 versions have been validated yet.

The primary aim of the present study was to examine the criterion validity of the Korean version of the DIVA-5 in an adult clinical sample. The secondary aim was to compare sociodemographic and psychiatric comorbidities in adult patients with and without a diagnosis of ADHD.

Methods

Procedures and Participants

Translation of the Diagnostic Interview for ADHD in Adults (DIVA-5) into Korean

The translation of the Dutch version of the DIVA-5 into Korean, as well as its back translation, were carried out according to the standards designed and approved by the DIVA Foundation (The Hague, The Netherlands). The Dutch version was translated into Korean by a professional translator and, where necessary, adjusted to the language and wording used in clinical psychiatry in Korea by clinicians experienced in adult ADHD (DY). The back translation from Korean to Dutch was done by the World Translation Institute (www.worlsys.biz). Expenses were paid by the Adult ADHD Study Group of the Korean Academy of Child and Adolescent Psychiatry. All of these processes were supervised by Professor GHB (Kyung Hee University, Seoul), BSK (Inje University, Seoul), SYB (Eulji University, Seoul), ISL (Soonchunhyang University, Buchun), SWC (Kyung Hee University), MH (Myongji Hospital, Seoul), RH (Kyung Hee University), and JWH (Kangwon National University, Chuncheon, Korea). The adaptation of the Korean version was based on the advice provided by S Kooij and A Bron of the DIVA Foundation; this collaboration ended in January, 2017.

All study participants were recruited from the Outpatient Services of eight university hospitals (Kyung Hee University Hospital, Samsung Seoul Hospital, Inje University Sanggye Paik Hospital, Chung Ang University Hospital, Nowon Eulji Hospital, Ewha Womans University Hospital, Myongji Hospital, and Soonchunhyang Bucheon Hospital) between January 2017 and December 2018. Those who agreed to participate provided written informed consent. The inclusion criteria was a referral for an ADHD assessment, and being between the ages of 17 and 65 years. Exclusion criteria included: treatment with ADHD medication in the previous 3 months; the presence of congenital genetic diseases; organic brain disease; severe physical disease requiring management such as renal failure, liver disease, cancer; a history of schizophrenia, bipolar I disorder, and other psychotic disorders within six months; and autism spectrum disorder; intellectual disability.

The clinical psychiatric assessment was performed by two board-certified psychiatrists. If the diagnosis of ADHD by the two experienced psychiatrist was consistent, patients were classified in the ADHD group; if not, they were classified as controls. All participants in this study were assessed using the Mini-International Neuropsychiatric Interview (MINI) Plus version 5.0.0 (MINI-Plus V.5.0.0) to evaluate comorbidity and were administered the Adult ADHD Self-Report Scale Symptoms Checklist version 1.1 (ASRS-V.1.1). Participants were also asked to fill out a questionnaire designed to collect sociodemographic characteristics. Trained psychiatrists or psychologists then tested all participants using DIVA-5 to confirm an adult ADHD diagnosis according to DSM-5 criteria.

Assessment Tools

Diagnostic Interview for ADHD in Adults (DIVA-5), Korean Version

DIVA is a semi-structured interview tool administered by a clinician. DIVA-2, which was based on DSM-IV criteria for ADHD in adults, was revised into DIVA 5.01 based on the DSM-5 criteria. DIVA-5 was translated into Korean according to the process described above. DIVA-5 consists of three parts that cover the following areas: 1) ADHD symptoms in childhood and adulthood, 2) age of ADHD onset, and 3) areas of impairment due to ADHD. If three or more criteria were met for either inattention and/or hyperactivity/impulsivity in childhood before the age of 12 years old, and five or more criteria were met in adulthood as reported by the patient and collateral informant(s),
the requirements for a clinical diagnosis of lifetime ADHD were considered to have been met.

**Adult ADHD Self-Report Scale (ASRS)**
The ASRS is an 18-item self-report scale based on the DSM-IV symptom criteria developed by the Korean Workgroup on Adult ADHD in conjunction with the World Health Organization.\(^2\) The scale is composed of two parts, parts A and B. The scale focuses on symptom frequency rather than severity to make scale instructions easier for participants to understand. Each ASRS question asks respondents how often a particular ADHD symptom had occurred over the past six months using a 5-point Likert scale ranging from 0 (never) to 4 (very often). Total scores ranged from 0 to 72. The Korean version of the ASRS has been validated.\(^3\) Part A is called the ASRS Screener, which is the short form of the ASRS, and comprises six questions selected based on stepwise-logistic regression.\(^4\) The ASRS Screener score is the sum of the dichotomous responses for the six questions—the scoring approach recommended by the ASRS V1.1 Screener manual—whereby respondents who endorsed at least four out of six items were considered at “elevated” risk for ADHD.\(^5\)

**MINIPLUS V5.0 Structured Interview**
The MINI\(^6\) is a short, structured diagnostic interview developed jointly by psychiatrists and clinicians in the United States and Europe for DSM-IV-TR\(^7\) and ICD-10.\(^8\) It is applied to assess current DSM-IV Axis I psychiatric morbidities.

**Statistical Analysis**
All computations were performed using SPSS software V.21.0 (IBM Corp, Armonk, NY, USA). Group comparisons were performed using the Pearson’s \(\chi^2\) test for categorical variables, and mean differences in continuous variables were evaluated using the independent samples \(t\)-test. A \(p < 0.05\) was used to define statistical significance. The clinical ADHD diagnosis established by two clinical experts was used as the external criterion for the calculation of sensitivity, specificity, positive predictive value (PPV), negative predictive value (NPV), and area under the receiver operating characteristic curve (AUC) of DIVA-5.

**Research Ethics**
This study’s protocol was approved by the Institutional Review Board of the institutes from which participants were recruited, including Kyung Hee University Hospital (KMCIRB 2017–02-054), Sanggye Paik Hospital (SGPAIK-2017-05-012-005), Ewha Womans University Hospital (EUMC 2017–01-013), Chung Ang University Hospital (IRB-No. 1741–005-279), Soonchunhyang University Hospital (SCHBC 2016–11-012-006), Eulji University Hospital (EMCS 2016–07-001), Sungkyunkwan University Samsung Medical Center (SMC 2016–07-061) and Myongji Hospital (MJH 2017–07-013-001); the principal investigator was GHB. Written informed consent was provided by all participants. Some of the data relating to participants in this study were published in a previous paper.\(^9\) The study has been performed in accordance with the ethical standards laid down in the Helsinki II Declaration about informed consent, voluntariness and anonymity.

**Results**
The demographic characteristics of the ADHD and control groups are given in Table 1. There were significant differences between the two groups in terms of educational level and employment status (\(\chi^2 = 24.962, p = 0.000; \chi^2 = 48.796, p < 0.001\)).

The presence of comorbid psychiatric disorders was assessed using MINI-PLUS 5.0 (Table 1). The proportion of participants with a comorbid disorder was higher in the ADHD group (72.4%) than in the control group (14.0%).

The validity of the DIVA-5 was assessed by comparing the scores of the participants, with and without a diagnosis of ADHD, by two independent clinicians. Table 2 presents the sensitivity and specificity of DIVA-5 and ASRS (both Screener and Total scores). In the clinical utility study of the Korean ASRS, a cut-off score of 32 was suggested.\(^10\) DIVA-5 had a sensitivity and specificity of 91.30% and 93.62%, respectively.

**Discussion**
This is the first study to validate the DIVA-5. The results of this study demonstrate DIVA-5 to be a reliable diagnostic tool with good predictive value for the diagnosis of ADHD in adults in a Korean psychiatric population. Given that only two other studies have investigated the validity of the previous version of the DIVA (DIVA-2),\(^11\) to our knowledge this is the first study with the appropriate statistical analysis to explore the psychometric properties of the revised version of DIVA-5 in Korean. The validation of this diagnostic interview provides clinicians with a useful diagnostic tool for assessing ADHD in adult Korean populations.

One of the advantages of DIVA-5 is that it comprehensively assesses symptoms of ADHD, from childhood to
adulthood, by using behavioural examples for each DSM-5 criterion. Moreover, the extensive assessment of lifetime impairment in five areas of psychological functioning is another advantage of the DIVA 5.0 in comparison with other tools. As it is a semi-structured interview for the clinician, it does not have the weaknesses of self-report measures.

Although Kessler et al.\textsuperscript{22} found that the 6-item ASRS Screener, which is based on dichotomized scoring, showed
the best sensitivity and specificity for diagnosing ADHD adults, the Korean version of the full ASRS outperformed the ASRS Screener.\textsuperscript{19} Our study results were in line with Heo et al.\textsuperscript{19} Considering the relatively lower sensitivity the ASRS both Screener and Total compared to the DIVA-5, the ASRS should be used with caution.

We found a higher proportion of unemployment in participants with ADHD compared to controls. This is consistent with the literature.\textsuperscript{23,24} In those who had work, the proportion of simple laborers and skilled workers in participants with ADHD was higher than that of managers and experts compared to controls (data not shown). This is compatible with the fact that ADHD is associated with underachievement, first at school and later at work.\textsuperscript{25,26,27}

Regarding comorbid psychiatric disorders, we found a large difference between the ADHD and control groups. The high comorbidity rate in the ADHD group (72\%) is comparable to previously reported results.\textsuperscript{4} In our study, only 2.1\% (3/141) of controls had psychiatric disorders. We assumed that this was because the major participating institutions were university hospitals and, as university students were more likely to participate in the study as controls, the proportion of “very healthy” participants was higher. This needs to be further verified in a well-designed study, such as a cohort study.

This study had some limitations. First, as most of the subjects—patients and healthy controls—were recruited from metropolitan areas, the study cannot be said to be representative of the general Korean population. Second, male overrepresentation can be partially explained as reflection in the sample recruitment that the prevalence of clinical diagnosis in Korea is almost twice higher in males than females.\textsuperscript{28}

Third, the criterion validity of DIVA-5 in this study only refers to adult symptoms, since ASRS include only symptoms of ADHD in adulthood and no instruments have been included for childhood symptoms. In addition, because there were no valid interview tools specific to adults with ADHD in Korean, DIVA-5 was compared to only a rating scale such as ASRS. Despite these limitations, we found sound sensitivity and specificity values for DIVA-5 and expect the interview to be extensively used in clinical practice and research settings. This may increase the awareness surrounding, and diagnostic assessment of, adult ADHD in the Korean population.

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