

The Worldwide Prevalence of ADHD: A Systematic Review and MetaRegression Analysis

Guilherme Polanczyk, M.D.

Maurício Silva de Lima, M.D.,
Ph.D.

Bernardo Lessa Horta, M.D.,
Ph.D.

Joseph Biederman, M.D.

Luis Augusto Rohde, M.D., Ph.D.

Objective: The worldwide prevalence estimates of attention deficit hyperactivity disorder (ADHD)/hyperkinetic disorder (HD) are highly heterogeneous. Presently, the reasons for this discrepancy remain poorly understood. The purpose of this study was to determine the possible causes of the varied worldwide estimates of the disorder and to compute its worldwide-pooled prevalence.

Method: The authors searched MEDLINE and PsycINFO databases from January 1978 to December 2005 and reviewed textbooks and reference lists of the studies selected. Authors of relevant articles from North America, South America, Europe, Africa, Asia, Oceania, and the Middle East and ADHD/HD experts were contacted. Surveys were included if they reported point prevalence of ADHD/HD for subjects 18 years of age or younger from the general population or schools according to DSM or ICD criteria.

Results: The literature search generated 9,105 records, and 303 full-text articles

were reviewed. One hundred and two studies comprising 171,756 subjects from all world regions were included. The ADHD/HD worldwide-pooled prevalence was 5.29%. This estimate was associated with significant variability. In the multivariate metaRegression model, diagnostic criteria, source of information, requirement of impairment for diagnosis, and geographic origin of the studies were significantly associated with ADHD/HD prevalence rates. Geographic location was associated with significant variability only between estimates from North America and both Africa and the Middle East. No significant differences were found between Europe and North America.

Conclusions: Our findings suggest that geographic location plays a limited role in the reasons for the large variability of ADHD/HD prevalence estimates worldwide. Instead, this variability seems to be explained primarily by the methodological characteristics of studies.

(*Am J Psychiatry* 2007; 164:942–948)

Attention deficit hyperactivity disorder (ADHD) is characterized by pervasive and impairing symptoms of inattention, hyperactivity, and impulsivity according to DSM-IV (1). The World Health Organization (WHO) (2) uses a different name—hyperkinetic disorder (HD)—but lists similar operational criteria for the disorder. Regardless of the name used, ADHD/HD is one of the most thoroughly researched disorders in medicine (3). It has been associated with a broad range of negative outcomes for affected subjects (4, 5) and with a serious financial burden to families and society (6), which characterizes it as a major public health problem (7).

An understanding of the epidemiological aspects of ADHD/HD may provide insight into its distribution and etiology as well as information for planning the allocation of funds for mental health services (8). In past decades, investigators from all regions of the world have made substantial efforts to define the prevalence of the disorder. Several literature reviews have reported highly variable

rates worldwide, ranging from as low as 1% to as high as nearly 20% among school-age children (8, 9).

Although the reasons for variability across studies remain poorly understood, it has been hypothesized that geographical and demographic factors are associated with it (10). Several investigators have suggested that prevalence rates in Europe were significantly lower than rates found in North America (8, 11, 12). This hypothesis fueled the concern that ADHD/HD may be a product of cultural factors and promoted an enduring debate in the medical literature (8, 11, 12). Other experts have argued that the variability of ADHD/HD prevalence estimates may be best explained by the use of different case definitions and that no variability of the actual prevalence across geographical sites should be found when case definitions are the same (5, 8, 13). A similar position has been adopted by the American Academy of Child and Adolescent Psychiatry, which states that conflicting results of ADHD/HD prevalence estimates can be explained by methodological dif-

This article is discussed in an editorial by Drs. Moffitt and Melchior on p. 856.

ferences across studies, such as method of ascertainment, diagnostic systems and associated criteria (e.g., situational versus pervasive, degree of impairment), assessment methods, informants, and the population studied (4). Unfortunately, no empirical evidence to date supports these assumptions, which led the American Medical Association (3), the National Institute of Mental Health (6), and the Centers for Disease Control and Prevention (7) to call for further research on this issue.

Therefore, the main purpose of this study was to conduct an extensive review of the literature on the prevalence of ADHD/HD to 1) calculate a worldwide-pooled prevalence estimate and 2) determine factors implicated in the variability of estimates by examining the assumptions of different estimates based on geographic location of studies. Since some studies suggest that ADHD/HD prevalence is higher in North America, cross-national comparisons were conducted to investigate differences between North American studies and those conducted elsewhere. The main hypothesis was that geographic location would not significantly explain the variability of worldwide ADHD/HD prevalence estimates after adjustment for methodological differences across studies.

Method

Data Sources

The following four search strategies were used for this systematic review of the literature: 1) computer search of databases, 2) review of specialized textbooks, 3) review of reference lists of all articles retrieved, and 4) contact with authors and experts on ADHD/HD epidemiology. This search strategy is in accordance with expert recommendations on the subject (14).

Computer search. We searched MEDLINE and PsycINFO from Jan. 1978 to Dec. 2005 for articles in English, German, French, Spanish, and Portuguese. Search words/terms were as follows: child*, adolesc*, epidemiology, prevalence, rate, mental disorder*, psychiatric disorder*, ADHD, ADD, attention-deficit, attention-deficit/hyperactivity disorder, hyperactiv*, overactiv*, inattent*, hyperkinetic disorder, and minimal brain dysfunction. An asterisk after a term means that all terms that begin with that root were included in the search.

Step 1. Abstracts were reviewed by the first author (Dr. Polanczyk) and selected for further review if they met one of the following three criteria: 1) significant reviews of literature that covered ADHD/HD prevalence, 2) major guidelines on ADHD/HD, or 3) original investigation of prevalence based on nonreferred samples (schools or community) of subjects 18 years of age or younger diagnosed according to criteria established by any DSM (III, III-R, or IV) (1) or ICD (9 or 10) versions (2). If a criterion was not met because not enough information was provided, the abstract was set aside for further evaluation.

Step 2. Abstracts were reviewed independently by two authors (Drs. Polanczyk and Rohde) and were selected based on their consensus according to the same criteria used in Step 1. If consensus was not reached, the abstract was set aside for further evaluation.

Step 3. Full-text articles of abstracts selected in Step 2 were retrieved and reviewed by one author (Dr. Polanczyk). Inclusion was based on consensus between two investigators (Drs. Polanczyk and Rohde). Disagreements were discussed with a third author (Dr. de Lima). Studies were included if they met the following cri-

TABLE 1. Methodological Characteristics and Geographic Location of the Studies (N=102)

Variable	No. of Studies
Geographic location	
North America	32
Europe	32
Asia	15
South America	9
Oceania	6
Middle East	4
Africa	4
Origin of sample	
Community	42
School	60
Diagnostic criteria	
ICD-10	13
DSM-IV	44
DSM-III-R	36
DSM-III	9
Impairment criterion ^a	
Yes	44
No	48
Number of stages of evaluation	
One	61
Two	33
Two, only screens positive at first stage	8
Source of information ^a	
Best-estimate procedure	23
"And rule"	6
Parents	33
"Or rule"	9
Teachers	15
Subjects	7

^a For some studies, data were not available for extraction.

teria: 1) original surveys on ADHD/HD prevalence (point prevalence), 2) diagnoses based on any DSM (III, III-R, or IV) or ICD (9 or 10) versions, 3) probabilistic sample from the general population (e.g., households, birth registers) or from schools, and 4) subjects 18 years of age or younger. Twin studies and data from the first evaluation of longitudinal studies were included. The exclusion criteria were 1) studies that did not report confidence intervals (CI) or standard errors (SE) and that did not report data that allowed the calculation of these parameters and whose authors did not provide such data upon request and 2) studies that adopted a retrospective diagnostic approach based on subjects' records. Authors were contacted via e-mail if data were missing.

Review of specialized textbooks. The ADHD/HD sections of five major textbooks on child and adolescent psychiatry and on general psychiatry were reviewed to identify references to original surveys that might meet Step 1 criteria.

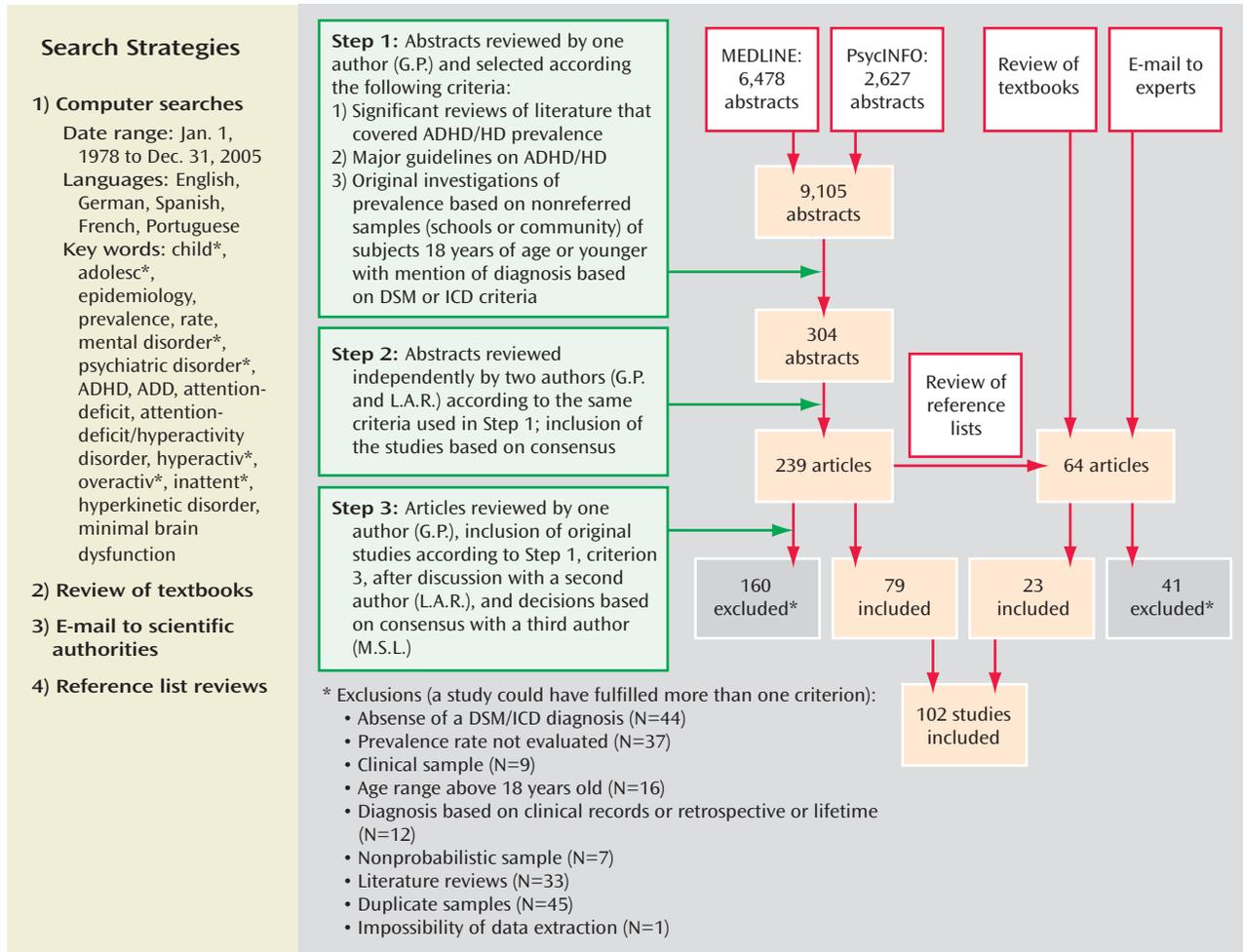
Contact with experts. Forty-five scientific authorities in the field of ADHD/HD from North America, South America, Europe, Africa, Asia, Oceania, and the Middle East were contacted via e-mail and asked whether they were aware of unpublished or ongoing studies, and 28 experts returned responses.

Review of reference lists. The reference lists of all articles selected in Step 3 were reviewed, and the full texts of potentially interesting studies were examined.

Data Extraction

A protocol for data extraction was defined and evaluated by two authors (Drs. Polanczyk and Rohde) (available from authors upon request). Data were extracted from full-text articles by one author (Dr. Polanczyk) and reviewed by a second author (Dr. Rohde). Disagreements were discussed with a third author (Dr. de Lima).

FIGURE 1. Flow Diagram of Study Selection



To determine pooled estimates, each study contributed with only one estimate. Whenever a study generated multiple estimates according to different diagnostic or methodological definitions, one estimate was selected according to a previously defined protocol. This protocol ensured that data on each subject were extracted only once.

The computer program Epi Info version 6.04d (15) was used to calculate CIs if they were not reported in the study.

Data Analyses

The first step of our previously defined strategy of data analysis was to determine an overall pooled-prevalence estimate of ADHD/HD based on all studies. A test of heterogeneity (Q test) was used to determine whether the differences in prevalence estimates across studies were greater than expected by chance. Significant heterogeneity was detected for the pooled estimate. At this point, sensitivity analysis was also performed, and findings showed that no study had skewed the overall result.

Since significant heterogeneity was found across studies, the second step was to conduct univariate analyses to test the individual association of each variable (methodological variables + geographic location of the studies) with the overall pooled-prevalence estimate of ADHD/HD using metaregression analyses (16). This analytical strategy evaluated which variables affected the results. Methodological variables were method of ascertainment of the sample (community or school), sample size, diagnostic crite-

ria (DSM-III, -III-R, or -IV; ICD-9 or -10), source of information (best-estimate procedure, parents, “and rule,” “or rule,” teachers, or subjects), requirement of impairment for the diagnosis (yes or no), number of stages, and response rate (1-attrition rate).

We also tested the association between geographic location and variability of the overall pooled-prevalence estimate of ADHD/HD. Studies were divided into seven groups according to their geographic location: Africa, Asia, Europe, the Middle East, North America, Oceania, and South America. Three studies were fully conducted in Central America (Costa Rica), and one was a multisite study conducted in the United States and Costa Rica. These studies were grouped with those from North America.

In the third step, a random-effects regression model was used to evaluate sources of variability in the overall pooled-prevalence estimate of ADHD/HD. All covariates associated with ADHD/HD prevalence rates for a flexible $p \leq 0.2$ (17) in univariate analyses were included in the final multivariate metaregression model. For these analyses, a significance level of 5% was established.

Age and gender were not included in the final multivariate metaregression model because less than 50% of the studies reported findings stratified by these variables, but individual estimates were computed according to these strata (age was stratified into the following ranges: 6–11 and 12–18). At this stage, we also determined pooled estimates for each geographic region. All analyses were performed using Stata 9.0 (Stata Corp, College Station, Tex.).

TABLE 2. Association Between Methodological Covariates and Geographic Area With ADHD/HD Prevalence Estimates

Variable	Univariate Model	Metaregression (Multivariate Model) ^a
	p	p
Origin of sample	<0.001	
Community		index
School		0.38
Source of information	<0.001	
Best-estimate procedure		index
“And rule”		0.04
Parents		0.03
“Or rule”		0.003
Teachers		<0.001
Subjects		0.46
Impairment criterion	<0.001	
Yes		index
No		0.001
Diagnostic criteria	<0.001	
DSM-IV		index
DSM-III-R		0.02
DSM-III		0.69
ICD-10		0.005
Number of stages of evaluation	<0.001	
One		index
Two		0.25
Two, only screens positive at first stage ^b		0.31
Response rate	0.25	—
Sample size	<0.001	0.81
Geographic area	0.009	
North America		index
Europe		0.40
Oceania		0.45
South America		0.83
Asia		0.85
Africa		0.03
Middle East		0.01

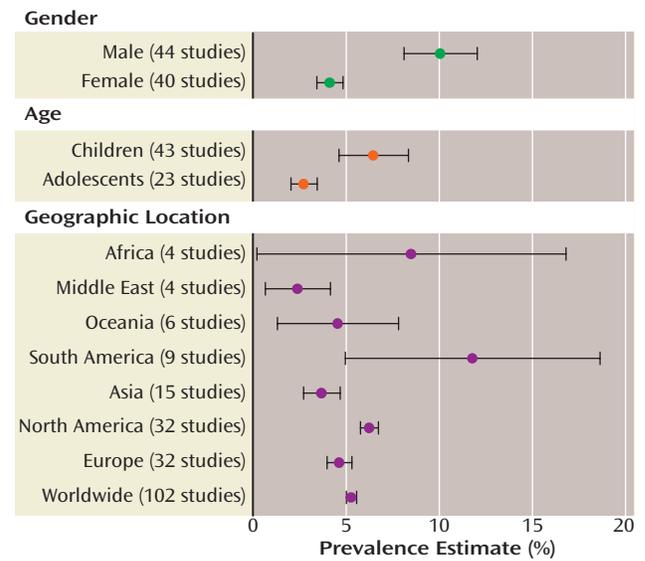
^a Between-study variance assessed by moment-based estimate ($\tau^2=7.815$).

^b Studies using two-stage sampling where only screening positives were assessed in the second stage (for details see the data supplement of the online version of this article).

Results

We screened 9,105 abstracts published in the last 27 years. A total of 303 full-text articles were reviewed, and 102 studies comprising 171,756 subjects were included in this systematic review (Figure 1). The list of reviewed references is available at <http://www.ufrgs.br/psiq/prodah-e.html>. Studies from all continents were retrieved. Of the 102 studies included, 32 were conducted in North America and 32 in Europe. The most frequent diagnostic system was DSM-IV. Characteristics of studies included in the analyses are described in Table 1.

Overall, the pooled prevalence of ADHD/HD was 5.29% (95% CI=5.01–5.56). This analysis revealed significant heterogeneity across studies ($p<0.001$). In univariate metaregression analyses (Table 2), all methodological variables evaluated were significantly associated with ADHD/HD prevalence rates, except response rate ($p=0.25$). The geographic location of studies was also significantly associated with the variability of ADHD/HD prevalence rates and was included in the final metaregression model.

FIGURE 2. ADHD/HD Pooled Prevalence According to Demographic Characteristics and Geographic Location

In the final multivariate metaregression model (Table 2), the methodological variables that remained significantly associated with the prevalence rates were the requirement of impairment for the diagnosis, diagnostic criteria, and source of information. As expected, studies without a definition of impairment had significantly higher ADHD/HD prevalence rates than those with a definition of impairment ($p<0.001$). Studies based on DSM-III-R or ICD-10 criteria, respectively, had significantly lower ADHD/HD prevalence rates than those using DSM-IV criteria ($p=0.02$ and $p=0.005$, respectively). Studies that relied on information provided by parents, teachers, and “or rule,” respectively, were associated with significantly higher ADHD/HD prevalence rates than those relying on a best-estimate procedure ($p=0.02$, $p<0.001$, and $p=0.003$, respectively), whereas those relying on information provided using the “and rule” criterion were associated with significantly lower ADHD/HD prevalence estimates ($p=0.04$) (Table 2).

Geographic location was associated with significant variability between estimates from North America and both Africa ($p=0.03$) and the Middle East ($p=0.01$). Estimates from these areas were significantly lower than estimates from North America. No significant differences were found in prevalence rates between North America and Europe ($p=0.40$), South America ($p=0.83$), Asia ($p=0.85$), or Oceania ($p=0.45$) (Table 2).

The multivariate metaregression model using Europe for the comparison yielded similar findings. Impairment, diagnostic criteria, and source of information remained significantly associated with prevalence rates. Significant differences were found only between Europe and both Africa ($p=0.05$) and the Middle East ($p=0.03$).

To increase power by decreasing degrees of freedom, an additional model was run using only methodological vari-

ables initially. Methodological variables associated with the prevalence rate for a $p \leq 0.20$ in univariate analyses were initially included and progressively deleted from the model using a backward procedure. Then, the geographic location of the studies was entered. In this multivariate model, geographic location was not significantly associated with prevalence rates after adjustment for methodological variables (data available from authors upon request).

Finally, since most studies were conducted in Europe and North America ($N=64$), the same data analysis strategy was applied using only these two regions. In the final multivariate metaregression model, the same methodological variables remained significantly associated with prevalence rates: impairment, diagnostic criteria, and source of information. Again, geographical location (Europe versus North America) was not significantly associated with ADHD/HD prevalence rates ($p=0.61$).

ADHD/HD prevalence rates were stratified by age in 43 studies and by gender in 44 studies. In these studies, both age and gender were significantly associated with prevalence rates ($p < 0.001$). The pooled prevalence of ADHD/HD stratified by gender and age can be found in Figure 2. The pooled prevalence in the seven geographic areas assessed is also shown in Figure 2. These analyses revealed significant heterogeneity across studies ($p < 0.001$).

Discussion

We have conducted a comprehensive systematic review of studies addressing prevalence rates of ADHD/HD worldwide and a metaregression analysis to understand the reasons of estimate variability. Our findings show that 1) the ADHD/HD worldwide-pooled prevalence is 5.29% (95% CI=5.01–5.56); 2) the large variability of ADHD/HD prevalence rates worldwide resulted mainly from methodological differences across studies; and 3) adjusting for methodological differences, prevalence rate variability was only detected between studies conducted in North America and those conducted in Africa and the Middle East. Moreover, no significant differences in ADHD/HD prevalence rates between North America and Europe were detected in the overall metaregression analysis and in the analysis restricted to studies conducted on these two continents. To our knowledge, this is the broadest systematic review of this subject to date.

Our search strategy identified original studies of ADHD/HD prevalence conducted on all continents. Despite the higher number of studies conducted in North America and Europe, our literature search showed that ADHD/HD is a well-studied disorder and that its prevalence rate has been estimated in several different cultures.

Although the ADHD/HD worldwide-pooled prevalence is consistent with results reported in most previous reviews (9, 18, 19), our results should be interpreted with caution because of the large variability found in all analyses. Our metaregression analysis reports, for the first time,

on the critical role that methodological variables (i.e., impairment criterion, diagnostic criteria, and source of information) play in the large variability of ADHD/HD prevalence estimates in different geographic locations. This finding empirically supports assumptions previously discussed in the literature (8, 9). For instance, applying the same methodological procedures and diagnostic criterion, very similar rates of ADHD/HD were found in Russia (20) and Britain (21) (1.3% and 1.4%, respectively). However, when the diagnosis of ADHD/HD was made in the same geographic location but according to a different methodological criterion (i.e., with or without the requirement of functional impairment) estimates ranged from 3.7% to 8.9% (22).

ADHD/HD terminology has undergone significant changes over the past decades. The current ICD-10 and DSM-IV criteria provide very similar lists of symptoms but recommend different ways of establishing a diagnosis. The ICD-10 requires a minimum number of symptoms in all three dimensions (inattention, overactivity, and impulsivity). The DSM-IV defines only two dimensions (with hyperactivity and impulsivity symptoms included in the same dimension), and a diagnosis can be made if there is a minimum number of symptoms in only one dimension. The ICD-10 requires that all criteria be met in at least two different situational contexts, whereas DSM-IV requires the presence of some impairment in more than one setting. The ICD-10 includes mood, anxiety, and developmental disorders as exclusion diagnoses. In DSM-IV, these diagnoses may be classified as comorbid conditions. Therefore, ADHD prevalence rates based on DSM-IV are expected to be higher than those based on ICD-10, which was demonstrated by individual studies (23) and corroborated by our analyses.

Our finding of lower ADHD/HD prevalence rates in Africa and the Middle East than in North America should be carefully interpreted. These geographic areas contributed with only a few studies to the overall analysis, and therefore we cannot rule out the possibility of lower accuracy of the adjustment for methodological aspects of these studies. Nevertheless, data concerning the lack of significant differences in ADHD/HD prevalence rates between North America and Europe in all sets of metaregression analyses are extremely relevant. This finding argues against the view that ADHD/HD is a culturally-based construct peculiar to the North American culture (12). It is important to note, however, that we do not suggest that culture has no influence on the prevalence of this disorder in a given population. Environmental and cultural aspects should play a role in the etiology of ADHD/HD, since the estimates of heritability in the disorder are approximately 80% (24, 25).

Our findings should be understood in the context of some limitations. First, characteristics of the subjects (age and gender) were not included in the final metaregression model because less than 50% of the studies reported re-

sults stratified by age or gender or provided data that allowed us to calculate these distributions. Second, other characteristics of the countries, besides geographic location, could not be assessed. However, the geographic location of a country is conceptually relevant for the purpose of this study and is often cited as a factor that may explain the variability of ADHD/HD prevalence rates. Finally, we did not provide estimates of ADHD/HD prevalence among adults, an emerging new area of research (26) that should be addressed in future studies.

Despite these limitations, we found a large variability in ADHD/HD prevalence rates worldwide. Results suggest that geographic location may play a limited role in the explanation of this variability. The many methodological differences in the 102 studies assessed may be the most important source of variability in the pooled estimate of ADHD/HD prevalence. These results indicate that international efforts from institutions such as WHO should be directed to the standardization of epidemiological studies so that comparable epidemiological data from all regions of the world are made available for the study of relevant mental disorders such as ADHD/HD.

Received Aug. 9, 2006; revisions received Nov. 9, 2006 and Jan. 8, 2007; accepted Jan. 8, 2007. From the ADHD Outpatient Program, Child and Adolescent Psychiatric Division, Hospital de Clinicas de Porto Alegre, Federal University of Rio Grande do Sul, Porto Alegre, Brazil; Pediatric Psychopharmacology Unit, Massachusetts General Hospital, Harvard Medical School, Boston; Federal University of Pelotas and Catholic University of Pelotas, Pelotas, Brazil; and the Post-Graduate Program in Epidemiology, Federal University of Pelotas, Pelotas, Brazil. Address correspondence and reprint requests to Dr. Rohde, Child and Adolescent Psychiatric Division, Hospital de Clinicas de Porto Alegre, Rua Ramiro Barcelos, 2350, Porto Alegre, RS, Brazil 90035-003; Irohde@terra.com.br or Dr. Polanczyk at gvp.ez@terra.com.br (e-mail).

The ADHD outpatient program receives research support from the following pharmaceutical companies: Bristol-Myers Squibb, Eli Lilly, Janssen-Cilag, and Novartis. Dr. Rohde is on the speakers' bureau and is a consultant for Bristol-Myers Squibb, Eli Lilly, Janssen-Cilag, and Novartis; he also serves on the advisory board for Eli Lilly. Dr. de Lima is Medical Director at Eli Lilly, Brazil. Dr. Biederman receives research support from Shire Laboratories, Inc., Eli Lilly, Wyeth-Ayerst, Pfizer, Cephalon, Novartis, and Janssen; he also serves on the speakers' bureaus of Eli Lilly, Pfizer, Novartis, Wyeth-Ayerst, Shire Laboratories, Inc., McNeil, and Cephalon and is on the advisory board of Eli Lilly, CellTech, Shire Laboratories, Inc., Novartis, Noven, McNeil, Janssen, Johnson & Johnson, Pfizer, and Cephalon. Drs. Polanczyk and Horta report no competing interests.

Supported in part by research grants from Conselho Nacional de Desenvolvimento Científico e Tecnológico (CNPq, Brazil) (grants 471761/03-6 and 300226/2002-0), Hospital de Clinicas de Porto Alegre, Porto Alegre, Brazil, and Eli Lilly. There was no involvement of any funding source in the study design, data collection, analysis, interpretation of data, and writing of this article or in the decision to submit the article for publication.

The authors thank the following individuals: Glorisa Canino, Val-samma Eapen, Steve Faraone, Martine Flament, Susan Shur-Fen Gau, Robert Goodman, Peter Jensen, Florence Levy, Anneke Meyer, Cecilia Montiel-Nava, Helmut Remschmidt, Russell Schachar, Joseph Sergeant, James Swanson, and Erik Taylor for their opinions on methodological issues of studies; Steven P. Cuffe, Cristiane Duarte, Bacy Fleitlich-Bilik, Sam Goldstein, Richard Hackett, David Hay, Rolf Loeber, Magda Stouthamer-Loeber, Pall Magnusson, Savita Malhotra, Christa Winkler Metzke, Emma Van der Meulen, Irma Moilanen, Rosalind Neuman, Lawrence Scahill, Martin Schmidt, John Seeley, Jakob Smári, Kenneth P. Tercyak, Márcio Vasconcelos, Kenneth Whiting, and Alessandro Zuddas for providing data about their studies; and

Sandra Costa Fuchs, Jair Mari, Carisi Anne Polanczyk, and Luis Eduardo Rohde for their helpful comments.

References

1. American Psychiatric Association: Diagnostic and Statistical Manual of Mental Diseases (DSM-IV), 4th ed. Washington, DC, American Psychiatric Publishing, 1994
2. World Health Organization: The ICD-10 Classification of Mental and Behavioral Disorders: Diagnostic Criteria for Research. Geneva, Switzerland, World Health Organization, 1993
3. Goldman LS, Genel M, Bezman RJ, Slanetz PJ: Diagnosis and treatment of attention-deficit/hyperactivity disorder in children and adolescents: Council on Scientific Affairs, American Medical Association. *JAMA* 1998; 279:1100-1107
4. Dulcan M: Practice parameters for the assessment and treatment of children, adolescents, and adults with attention-deficit/hyperactivity disorder: American Academy of Child and Adolescent Psychiatry. *J Am Acad Child Adolesc Psychiatry* 1997; 36(suppl 10):85S-121S
5. Swanson JM, Sergeant JA, Taylor E, Sonuga-Barke EJ, Jensen PS, Cantwell DP: Attention-deficit hyperactivity disorder and hyperkinetic disorder. *Lancet* 1998; 351:429-433
6. National Institutes of Health: National Institutes of Health Consensus Development Conference Statement: diagnosis and treatment of attention-deficit/hyperactivity disorder (ADHD). *J Am Acad Child Adolesc Psychiatry* 2000; 39:182-193
7. Lesesne C, Abramowitz A, Perou R, Brann E: Attention Deficit/Hyperactivity Disorder: A Public Health Research Agenda. <http://www.cdc.gov/ncbddd/adhd/dadphra.htm>, 2000 (Accessed Aug. 2006)
8. Bird HR: The diagnostic classification, epidemiology and cross-cultural validity of ADHD, in Attention Deficit Hyperactivity Disorder: State of the Science: Best Practices. Edited by Jensen PCJ. Kingston, NJ, Civic Research Institute, 2002
9. Faraone SV, Sergeant J, Gillberg C, Biederman J: The worldwide prevalence of ADHD: is it an American condition? *World Psychiatry* 2003; 2:104-113
10. Rappley MD: Attention deficit-hyperactivity disorder. *N Engl J Med* 2005; 352:165-173
11. Anderson JC: Is childhood hyperactivity the product of Western culture? *Lancet* 1996; 348:73-74
12. Timimi S, Taylor E: ADHD is best understood as a cultural construct. *Br J Psychiatry* 2004; 184:8-9
13. Rohde LA, Szobot C, Polanczyk G, Schmitz M, Martins S, Tramontina S: Attention-deficit/hyperactivity disorder in a diverse culture: do research and clinical findings support the notion of a cultural construct for the disorder? *Biol Psychiatry* 2005; 57: 1436-1441
14. Stroup DF, Berlin JA, Morton SC, Olkin I, Williamson GD, Rennie D, Moher D, Becker BJ, Sipe TA, Thacker SB: Meta-analysis of observational studies in epidemiology: a proposal for reporting: Meta-Analysis of Observational Studies in Epidemiology (MOOSE) group. *JAMA* 2000; 283:2008-2012
15. Center for Disease Control and Prevention, World Health Organization: Epi Info, Version 6.04d, 2001
16. Thompson SG, Higgins JP: How should meta-regression analyses be undertaken and interpreted? *Stat Med* 2002; 21:1559-1573
17. Maldonado G, Greenland S: Simulation study of confounder-selection strategies. *Am J Epidemiol* 1993; 138:923-936
18. Scahill L, Schwab-Stone M: Epidemiology of ADHD in school-age children. *Child Adolesc Psychiatr Clin N Am* 2000; 9:541-555, vii
19. Szatmari P: The epidemiology of attention-deficit hyperactivity disorders. *Child Adolesc Psychiatr Clin N Am* 1992; 1:361-371

20. Goodman R, Slobodskaya H, Knyazev G: Russian child mental health: a cross-sectional study of prevalence and risk factors. *Eur Child Adolesc Psychiatry* 2005; 14:28–33
21. Ford T, Goodman R, Meltzer H: The British Child and Adolescent Mental Health Survey 1999: the prevalence of DSM-IV disorders. *J Am Acad Child Adolesc Psychiatry* 2003; 42:1203–1211
22. Canino G, Shrout PE, Rubio-Stipec M, Bird HR, Bravo M, Ramirez R, Chavez L, Alegria M, Bauermeister JJ, Hohmann A, Ribera J, Garcia P, Martinez-Taboas A: The DSM-IV rates of child and adolescent disorders in Puerto Rico: prevalence, correlates, service use, and the effects of impairment. *Arch Gen Psychiatry* 2004; 61:85–93
23. Goodman R, Ford T, Richards H, Gatward R, Meltzer H: The Development and Well-Being Assessment: description and initial validation of an integrated assessment of child and adolescent psychopathology. *J Child Psychol Psychiatry* 2000; 41:645–655
24. Biederman J, Faraone SV: Attention-deficit hyperactivity disorder. *Lancet* 2005; 366:237–248
25. Sergeant J: Are we ready for endophenotypes in attention deficit hyperactivity disorder? *Rev Bras Psiquiatr* 2005; 27:262–263
26. Wilens TE, Faraone SV, Biederman J: Attention-deficit/hyperactivity disorder in adults. *JAMA* 2004; 292:619–623