INTRODUCTION

The relation between bipolar affective disorder (BAD) and adult attention deficit hyperactivity disorder (ADHD) is not clear (1). Higher rate of ADHD symptoms in patients with BAD and higher rate of BAD in the adulthood in patients with ADHD have increased attention on this subject (2).

The lifelong prevalence of BAD ranges 0.4% to 1.6% (mean 1.2%) (3,4,5). Given that it is considered as bipolar spectrum, this rate exceeds 5% (5,6). The lifelong rate of comorbidity of BAD which is frequently observed with axis I and axis II diagnoses with axis I disorder is 50-70%. When studies about comorbidity of BAD and ADHD are examined, the rates of association for both disorders decrease as the age gets older, but is observed at a rate of 9-35% in the adulthood (7-12).

Although community-based longitudinal follow-up studies investigating the prevalence of adult ADHD are limited and composed of indirect data (13), the symptoms of ADHD diagnosed in the childhood continue at a rate of 50-80% in the adolescence and 30-50% in the adulthood (14). Studies conducted in different countries indicate the prevalence of Adult ADHD as 1-4.4% (15-18). Although discussions about Adult ADHD continue, studies show that this disorder is a frequent diagnosis decreasing the functionality (15,18-20). An important property of ADHD is that it has a comorbidity at a rate of 65-89% in the

ÖZET:
Popüler konu: İki uçlu mizaç bozukluğu ve erişkin dikkat eksikliği hiperaktivite bozukluğu ne kadar birlikte ne kadar ayrı?


Anahtar sözcükler: İki uçlu mizaç bozukluğu, dikkat eksikliği hiperaktivite bozukluğu, ekhastalık

Journal of Mood Disorders 2011;1(3):136-44

ABSTRACT:
Bipolar disorder and adult attention deficit/hyperactivity disorder: the same or different?

The diagnoses of Bipolar Affective Disorder (BAD) and Attention Deficit Hyperactivity Disorder (ADHD) can frequently be present comorbidly. The rates of comorbidity of the both disorders concomitantly have recently arisen the question, whether these two conditions are precursors or precursor symptoms of each other. The possible relationship between them is increasingly drawing attention in all age groups. Comorbidity of BAD/ADHD affects the course of the disease negatively, shortens the periods of well-being, increases exposure to negative life events, and decreases quality of life. It also increases occurrence of anxiety disorder and alcoholism. Currently, studies aimed to understand the relation between these two diseases are inadequate. In this article, similarities and differences of BAD and Adult ADHD are reviewed according to the findings in epidemiologic studies, familial studies and neuroimaging studies.

Key words: Bipolar affective disorder, attention deficit hyperactivity disorder, comorbidity

Journal of Mood Disorders 2011;1(3):136-44
clinical studies conducted (21-23). Studies investigating ADHD comorbidity among these co-diagnoses are limited and their results are controversial. The most common comorbidity is oppositional defiant disorder (24). Also speech and learning disorders and anxiety disorders are frequently observed as comorbidity (7,23,25). The comorbidity rates of ADHD and depression are 10-30% (26,27). This can be unipolar or bipolar depression. Bipolarity is observed at a rate of 5-20% in ADHD (11,28). In addition to comorbidity, BAD and ADHD have similar and differentiating aspects epidemiologically and clinically. The aim of this article is to examine the possible relation between BAD and ADHD in the adults according to the findings in epidemiologic studies, familial studies, clinical studies and neurobiological studies.

**RELATIONSHIP BETWEEN BAD and ADULT ADHD**

**Methodological Difficulties**

Although methodological difficulties are present in all the psychiatric disorders, unanswered questions in the differential diagnosis of these two conditions cause to both diagnostic and therapeutical difficulties. There are many separate studies about BAD and ADHD. Although BAD and ADHD are considered as two different disorders, they show many similar clinical properties including racing thoughts, hyperactivity, increase in the amount of speech, impulsivity, and attention deficit (1,29,30). In adult ADHD, anger and mood changes are frequently observed; increase in the amount of energy and decrease in need of sleep are present (31). In patients with Adult ADHD, decrease in functionality may be observed which appears to be occasionally episodic; this arises from the fact that ADHD is a condition leading to developmental problems, change in performance and reactivity. Overlapping symptoms and the possibility of observing these two disorders concomitantly have recently arisen the question, whether these two conditions are precursors or precursor symptoms of each other. The possible relationship between them is increasingly drawing attention in all age groups (32,33). In addition to overlapping symptoms, these two conditions have also many different properties. As Wingo and Ghaemi (26) recently emphasized, as the disorder is stable BAD shows an episodic course. Furthermore, while increased productivity, increase in self-confidence and the presence of psychotic aspects are observed frequently in euphoric BAD, they are rarely observed in ADHD. Excluding the cases starting in the childhood, adequate function has been proved in cognitive tests including writing, reading and different behavioral measurements in the premorbid period in BAD (35), but decrease in functionality is frequently observed in ADHD starting from the childhood (36). It was found that adults with a diagnosis of ADHD showed lower performance in tests which measured selective attention, continuing attention, and suppression of response compared to normal controls. This shows that attention problems in ADHD continue in the adulthood (37).

It is generally not easy to make retrospective examination and differentiation with our current experience to understand the differences related to the courses of the disorders (32,38). Since clinical course is milder and more disorganized, the risk for misdiagnosis is increased especially for BAD-II (39). Temperament properties of the patient or subsyndromal manifestations also increase the risk of misdiagnosis (40).

According to DSM IV, diagnostic categories of BAD and ADHD seem to be unrelated. There is similarity between some ADHD symptoms and manic symptoms. However, it is not fully clears if this relationship is at a superficial level due to clinical definitions or there is an overlapping at biological level (32).

In developmental point of view, ADHD starts in the childhood and shows different symptoms in the adulthood and the symptoms in the childhood are defined for Adult ADHD. BAD usually starts in the adulthood and the form which starts in the childhood displays different clinical properties. For the diagnosis, symptoms observed in the adulthood are evaluated. The diagnoses of Adult ADHD and childhood BAD do not define developmental changes. Therefore, it is difficult to understand the symptomatic similarities and differences of these two conditions.

Prediction of the prevalence of Adult ADHD comorbidity shows a wide range because of the methods used and sampling differences. Overlapping symptoms of BAD and ADHD constitute the most important diagnostic limitation (32). There may be difficulties in remembering the symptoms which had started in the childhood during the adulthood. In a 16-year follow-up study, only 78% of
the patients who had ADHD in the childhood could remember the symptoms in the adulthood (41). In addition, the frequency of comorbidity might have been overstated, since studies were conducted in patients who seek for help (42). When these are considered together, it can not be accurately evaluated which one affects the prediction regarding the prevalence of ADHD to what extent (11).

Disorder course, core symptoms and phenomenology in patients with a comorbidity of ADHD are different compared to patients without a comorbidity of ADHD (7). The disorder starts at an earlier age in BAD patients with a comorbidity of ADHD, depressive and mixed attacks are observed more frequently, euthymic periods are shorter and anxiety disorders and alcohol-substance addiction are observed more frequently compared to patients without a comorbidity of ADHD (7,11,23,43). Again, in these patients, the quality of life decreases (44), the rates of suicidal attempt and legal problems increase (11) and negative life events are observed more frequently (45).

Despite all these difficulties, three possibilities should be considered, when the literature is examined to understand the nature of the relation between BAD and ADHD; the first possibility is that these two conditions are completely different and have no relation. The second possibility is that these two conditions are two different manifestations of the same condition and the third possibility is that a more complex relation is present as suggested by some authors in familial and phenomenology studies (32).

In this context, review of epidemiological studies, familial studies, neurobiological studies, and clinical studies, which may help us to understand the relation between BAD and ADHD, can be beneficial for us to comprehend the nature of the subject.

**Epidemiological Studies**

Studies show that ADHD in children with BAD is observed at a higher rate (57-100%) as compared to patients with BAD in adolescence (46-49). In a different perspective, the rate of BAD in children with ADHD was found to be 10-fold higher compared to children without ADHD (50). Biederman et al. (51) showed that the diagnostic criteria of both mania and ADHD were maintained in most children with a comorbidity of BAD and ADHD, when overlapping symptoms were excluded. These results noted that there was no diagnostic artifact between ADHD and mania in the childhood despite the same diagnostic criteria. However, this raised controversial subjects. The critics about this subject stated that the diagnoses in this study were not compatible with classical mania and 17% displayed atypical properties (52). This may lead to increase in the rates of artifact related to the symptoms of BAD and ADHD (53). Therefore, repeated studies about the subject are needed. Another reason for the high rate of comorbidity was suggested by Biederman group (51) and West (46,47); DSM III R criteria used for the diagnosis of BAD do not require time for the diagnosis. However, a period of at least one week is needed in DSM IV. Therefore, it is proposed that mood change may not typically last one week and may lead to falsely high rates of comorbidity in children with ADHD. The rate of ADHD in patients with BAD was found to be 9-97% and the rate is higher in cases in whom BAD starts in the childhood as compared to the cases in whom BAD starts in the adolescence (46,48,49). Geller at al (48) assessed 60 BAD and 60 ADHD whose ages are 7-16 years old, using with Washington University at St. Louis Kiddie and Young Adult-Schedule for Affective Disorders and Schizophrenia - Lifetime and Present Episode Version-DSM-IV (WASH-U-KSADS). They divided BAD patients in to the prepubertal and postpubertal groups. They found that ADHD occurred in 97.0% of prepubertal versus 74.1% of postpubertal BAD cases. Nevertheless, bipolar disorder is also diagnosed in approximately 7-23% of the children with a diagnosis of ADHD (54,55).

Faranoe et al. (49) reported that the age of onset of BAD is critically important. It is suggested that a heterogeneous structure showing the properties of BAD and ADHD may be a subgroup in the bipolar spectrum and this may be a different familial subtype which classically starts at the adulthood (43,51,56). When the studies about the association of Adult ADHD and BAD are examined, similar findings are observed, although the rates of association are decreased. In the study performed by Tamam et al., lifelong association with ADHD in patients with BAD was found at a rate of 27% and association of BAD with ADHD continuing in the adulthood was found to be 16% (7). Similar findings were
found in other studies (9-35%) (8-12). In contrast to the other studies, Jaideep et al. (57) found the rate of ADHD to be low (4%) in patients with BAD which started in the childhood in a study they performed in 2006. The difference of our study compared to the other studies may be explained by the fact that it was conducted in a small number of patients with BAD. Again, in a study performed in India, a comorbidity of ADHD was not found in 30 children with a diagnosis of BAD (58). Similarly, Kim Cohen et al. (59) compared the histories of ADHD in children with and without mania and no relation were found between mania and ADHD.

Although few studies state the contrary (58,59), the presence of a strong relation between BAD and a comorbidity of ADHD is noted, when the literature is examined (32). Comorbidity of BAD and ADHD may be a separate subtype of ADHD or BAD. Although a diagnostic artifact is possible (32,46,48,49,54,55), most studies show the strength of the relationship.

**Familial Studies**

Strong familial relationship is known in both BAD and ADHD (28). BAD was found with a 7-8 fold higher rate in the first-degree relatives of patients with BAD compared to the patients without BAD (60).

In a meta-analysis which evaluated the results of 8 studies conducted in parents of children and adolescents with a diagnosis of ADHD, it was shown that the relative risk for a diagnosis of ADHD was increased 2-8 fold in parents of children and adolescents with a diagnosis of ADHD (61). In a familial study conducted by Woźniak et al. (56), a comorbidity of BAD and ADHD was found at a high rate in children younger than 12 years of age and their first-degree relatives. In the meta-analysis of six studies conducted between 1970 and 1990, the prevalence of BAD in the first-degree relatives of children with ADHD (2.6%) was compared with the control group (1.3%) and rates of BAD in the relatives of these children were found to be high. However, statistically significant difference was not found in any of the studies (62). In a study, the rate of ADHD was found to be 28% in 60 children of adult patients with BAD. 9 of these children were reported to have BAD and 88% of these were reported to have ADHD (63).

The prevalence of both BAD and ADHD was found to be high in the first-degree relatives of children with BAD (32,56). In addition, ADHD and BAD were found in association in the relative of children with BAD and ADHD (49).

Persistence of the symptoms of ADHD in the adulthood, high rate of a comorbidity of ADHD in patients with BAD and high rate of mood swings in ADHD may be explained by a genetic relation between BAD and ADHD. Biederman et al. (51) who supported this hypothesis found the rates of ADHD to be also high in families of patients with BAD. However, it is not possible to explain all variations genetically. Many disorders have similar environmental risk factors (64).

In the light of these findings, it can be stated that BAD and ADHD may be disorders related to each other and have similar familial risk factors (32). In addition, the hypothesis that a different type of BAD/ADHD may be present is supported by these studies. However, diagnostic overlaps of these two disorders cause confusion in the evaluation. For instance, it is possible that some of the children diagnosed with ADHD are actually suffering from BAD which would explain the increased rates of BAD within their family members.

**Neurobiological Studies**

Neurocognitive studies of patients with ADHD identified patterns of executive dysfunction in patients with ADHD that are thought to reflect abnormalities in the functioning of the prefrontal cortex, therefore supporting the hypothesis of an alteration of the prefrontal cortex neuroanatomy in ADHD (65-67). The most consistent structural brain imaging findings in children with ADHD have been significantly smaller volumes in the dorsolateral prefrontal cortex, caudate, pallidum, corpus callosum, and cerebellum. Similarly, BAD patients have shown differences in frontal, and temporal regions of the brain, and corpus callosum and basal ganglia, as well (68). Thus, similar areas of the brain were reported to be involved in both conditions with only the temporal lobe seeming to play a role in BAD alone. Although dysfunction of similar brain areas could suggest a relationship between BAD and ADHD, it is important to recognize that these brain areas are not restricted to BAD and ADHD pathologies. In fact, the same areas appear to be activated differentially in a number of psychiatric
conditions including anxiety disorders (69) and depression (69,70). To our knowledge, no neuroimaging study has previously been conducted with ADHD, BAD, and ADHD/BAD patients concurrently.

Most studies have found abnormalities in cerebral activation in ADHD, with a hypoperfusion of frontal and possibly striatal areas (65-67,71-74). Several imaging studies using ligands highly selective for the dopamine transporter sites have studied their density in ADHD subjects compared to controls. Dopamine transporter and the dopamine seem to play important roles in the pathophysiology of ADHD and in response to its treatment. These studies consistently found an increase in binding of dopamine transporter in the striatum of ADHD subjects compared to controls. Several studies also showed a normalization of this brain function following treatment with methylphenidate (75-78). Pharmacological treatments for both ADHD and BAD act on the dopaminergic, noradrenergic and serotonergic systems. ADHD is usually treated with stimulants whereas BAD treatment involves either mood stabilizers, including anticonvulsants, antipsychotics, lithium, and sometimes; antidepressants.

The fact that similar classes of medications can treat both conditions suggests that ADHD and BAD could involve dysfunctions of the same neurotransmitters. However, this finding is not specific to these two conditions as dopamine and serotonin systems are also involved in several other psychiatric disorders (32).

Clinical Conditions

Studies about comorbidity report that the association of BAD/ADHD lead to significant socioeconomic and clinical results (7,34,79,80).

It has been proposed that BAD affects the patient’s marriage and occupational life and leads to severe and rather persistent disorders (79). Coryell et al. (79) followed up 48 patients with BAD for 6 years and compared them with normal controls in their study. They found that these patients could not develop their education and occupational states and were unemployed at a significantly higher rate in the last year of the follow-up. Although coupled with normal controls of the same age, their marriage rates were decreased by half and rates of divorce and separation were found to be 2 fold higher (79). Again, studies have shown that adults with ADHD had a lower socioeconomic status, experienced problems in their occupational life, changed work more frequently, had a shorter educational life and a lower academic success, repeated classes more frequently, matured more lately and experienced more problems in their marriages (49,81,82). Attention deficit, impulse control disorder and disruption in social communication observed in ADHD disrupt academic and professional performance (83,84,85) and driving performance (82,86). Negative life events are observed at a higher rate in individuals with ADHD in all ages compared to controls (87,88). These negative life events were found to be related to the severity of ADHD in adults (45). In addition, these negative life events are also related to other negative results including hopelessness, suicidal thought, incompatibility, loss of environmental control (89).

BAD starts at a significantly earlier age in patients with BAD+ADHD compared to patients without ADHD (7,11,23,43,47,54). In the study performed by Nierenberg et al. (11), 60% of the patients had early onset BAD. Data suggest that early-onset BAD may be related to ADHD (8). Nierenberg et al. (11) report that the age of onset of BAD is critically important and proposed that BAD+ADHD is a subtype starting in the early childhood and this may be a different condition from BAD which classically starts in the adulthood (43). Familial studies conducted on this subject support this (32).

In addition, BAD proceeded more chronically and with disability in individuals with a comorbidity of BAD/ADHD compared to the ones without comorbidity of BAD/ADHD. Shorter time of wellbeing and shorter intervals between attacks were related to higher number of attacks (11).

Tamam et al. (7) found the total number of attacks to be statistically higher in the BAD+ADHD group in the study they performed in 159 bipolar patients. When the patients were examined in terms of the first attack, some studies did not find a difference in terms of the characteristics of the first attack (7,11,23), but some other studies reported that manic and depressive episodes were observed at a higher rate in these patients and the number of days with irritability was higher (10,11).

Suicidal attempts were observed at a higher rate in these patients and legal problems were increased (11). In patients with a comorbidity of BAD and ADHD, increase
in attention deficit was related to anxiety and alcohol and substance abuse and addiction (7,11,44). Alcohol and substance abuse was reported to be high in individuals with both BAD (90,91) and ADHD. Familial studies show that symptoms of ADHD are observed at a high rate in adults with alcohol addiction (92). Symptoms of ADHD were also found at a higher rate in children of individuals with alcohol addiction compared to the control group. In addition, rates of both ADHD and substance abuse were found to be high in adolescent or adult children of individuals with substance abuse (86,92,93).

Association of these two conditions affects the quality of life negatively. Global Assessment of Functioning (GAF) scores are statistically significantly lower in patients with a co-diagnosis of BAD/ADHD compared to patients with only ADHD (44).

CONCLUSION

In conclusion, when taken all together, all these evidences strongly suggest that ADHD and BAD are correlated and even that ADHD/ BAD might constitute a separate condition. Surely, chance alone cannot explain these findings in a satisfactory manner. On the other hand, the results of some studies support the view that these two conditions are interlocked with each other and are the same entity. Methodological difficulties prevent designing a study, which will help us understand, if these two conditions are separate entities or a spectrum. However, additional studies are clearly needed to determine whether the link between ADHD and BAD resides only at the epidemiological level or also at the clinical and biological level.

References:


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