Psychosocial problems and cognitive functions in children with spina bifida

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Objectives: This study aims to assess psychosocial functioning in relation to lesion level and ambulatory status in children with spina bifida (SB) and compare them to their peers.

Patients and methods: Between March 2013 and May 2013, a total of 31 patients with SB (11 males, 20 females; mean age: 9.4 years; range, 6 to 14.7 years) and 36 typically developing peers (16 males, 20 females; mean age: 9.8 years; range, 6.5 to 14.8 years) were included in the study. All participants were assessed using a semi-structured psychiatric diagnostic interview via the Kiddie Schedule for Affective Disorders and Schizophrenia Present and Lifetime Version (K-SADS-PL), Wechsler Intelligence Scale for Children-Revised (WISC-R), Behavioral Rating Inventory of Executive Functions (BRIEF) parent form, Social Responsiveness Scale (SRS), and Aberrant Behavior Checklist (ABC).

Results: In the SB group, the rate of psychiatric disorders was significantly higher (p=0.001) and the SRS scores and the planning and organizational components of the executive function were higher than their peers (p=0.02 and p=0.007, respectively). The psychiatric diagnosis rate, BRIEF, and SRS total scores did not significantly differ according to lesion level and ambulatory status. The BRIEF initiate and organization of materials subtest scores and ABC scores were significantly lower at high lesion levels (p=0.02, p=0.02, and p=0.02, respectively) and non-community walkers (p=0.002, p=0.03, and p=0.003, respectively).

Conclusion: Psychiatric disorders, impairment in social responsiveness, and planning and organization components of the executive function are prevalent in children with SB with no intellectual disabilities, compared to their peers. Therefore, psychosocial counseling and multidisciplinary follow-up for SB patients seem to be beneficial.

Keywords: Children, cognitive impairment, psychiatric symptoms, psychosocial status, spina bifida.
dysgenesis of the corpus callosum), hydrocephalus and its associated complications have been blamed to cause neurocognitive deficits such as attentional problems, memory difficulties, and poor executive function.\cite{3} Executive functioning is an umbrella term which is usually used to describe cognitive abilities including goal-directed problem solving and adequate cognitive functions to adapt to the needs and changes in the environment. Executive functions represent the capability of processing data obtained from the observations of outer world, resulting in showing appropriate responses, when needed.\cite{4,5} Despite their physical problems, their IQs fall within the average range.\cite{6} Associated health complications include weakened or paralyzed lower extremities, urinary and bowel incontinence, hydrocephalus and orthopedic problems. Problems resulting from complications such as assistance need in mobilization, neurogenic bladder and bowel problems, precocious puberty and short stature related to endocrine dysfunctions may result in internalizing disorders, limited participation in activities with their peers and social skill deficits. Therefore, internalizing problems such as anxiety and depression are common among children with SB.\cite{3,7}

Recently, there has been a surge in the literature investigating psychosocial, behavioral, and social adjustment in patients with SB. However, a limited number of studies has examined impairments in this area in relation to the lesion level and ambulatory status. To date, psychiatric diagnoses have been assessed via caregiver-reported outcomes, and no previous research has evaluated psychiatric symptomatology using a gold-standard psychiatric diagnostic interview. In the present study, therefore, we aimed to evaluate psychiatric symptomatology in children with SB using a gold-standard psychiatric diagnostic interview, to compare children with SB and their age peers in terms of psychiatric symptomatology, cognitive functioning, behavioral problems, executive functions and social responsiveness, and to assess whether psychiatric symptomatology, behavioral problems, executive functions, and social responsiveness varied in relation to the lesion level and ambulatory status.

**PATIENTS AND METHODS**

This cross-sectional, case-control study was conducted at Physical Medicine and Rehabilitation and Pediatric Nephrology outpatient clinics of Marmara University School of Medicine between March 2013 and May 2013. A total of 31 patients with SB (11 males, 20 females; mean age: 9.4 years; range, 6 to 14.7 years) and 36 typically developing peers as the control group (16 males, 20 females; mean age: 9.8 years; range, 6.5 to 14.8 years) were included in the study. Exclusion criteria for both groups were as follows: mental retardation according to clinical evaluation; chronic medical illness for the control group; having a history of long-lasting shunt dysfunction or existing shunt dysfunction; primary sensory loss and hand function deficits which could render assessments impossible; and having a history of psychosis or child neglect and abuse. Demographic and clinical data of the participants, including clinical history, musculoskeletal and neurological examinations were recorded. Clinical examinations were conducted by two experienced rehabilitation physicians. The International Myelodysplasia Study Group Criteria assigning motor levels was used for the evaluation of motor status of children with SB.\cite{8} The Mid-lumbar and thoracic level lesions were classified as the upper lesion level, whilst the sacral and lumbosacral lesions were classified as the lower lesion level.\cite{9} All participants and parents were informed about the nature of the study and a written informed consent was obtained. The study protocol was approved by the Marmara University School of Medicine Ethics Committee (No. 09.2013.0057). The study was conducted in accordance with the principles of the Declaration of Helsinki.

Based on the Hoffer criteria, children with SB were classified as community walkers and non-community walkers.\cite{10} Typically developing children were recruited after a routine examination. The Wechsler Intelligence Scale for Children-Revised (WISC-R) and Kiddie Schedule for Affective Disorders and Schizophrenia Present and Lifetime Version (K-SADS-PL), a semi-structured evaluation procedure for clinical psychiatric diagnoses, were utilized by the child psychiatrist. The parents were asked to complete the Behavioral Rating Inventory of Executive Functions (BRIEF),\cite{11} Social Responsiveness Scale (SRS),\cite{12} and Aberrant Behavior Checklist (ABC).\cite{13}

**Kiddie Schedule for Affective Disorders and Schizophrenia Present and Lifetime Version (K-SADS-PL)**

The K-SADS-PL is a semi-structured interview form for collecting signs and symptoms of psychiatric disorders in children and adolescents.\cite{14} Validity and reliability studies of the Turkish version were carried out for diagnosing childhood psychiatric disorders.\cite{15} The K-SADS-PL is considered the gold standard among diagnostic interview instruments.
for clinical psychiatric diagnoses in children and adolescents.

**The Wechsler Intelligence Scale for Children-Revised (WISC-R)**

The WISC-R is used to measure the intellectual capacity in children with adequate speech and language skills. It has been adapted to Turkish culture and was standardized in Turkish. In our study, we used the subtests of information, vocabulary, picture completion, and picture arrangement for comparison of the groups.

**The Behavioral Rating Inventory of Executive Functions (BRIEF)**

The BRIEF is a caregiver report rating scale developed to assess the behaviors related to executive functions of children ages 5 to 18 years. The Parent Form contains 86 items of behavior as occurring “Never (1),” “Sometimes (2),” or “Often (3).” The BRIEF consists of eight clinical scales (Inhibit, Shift, Emotional Control, Initiate, Working Memory, Plan/Organize, Organization of Materials, Monitor), and two indexes (Behavioral Regulation Index [BRI] and the Metacognition Index [MI]). The BRI is comprised of three subscales (Inhibit, Shift, Emotional Control). The MI is comprised of five subscales (Initiate, Working Memory, Plan/Organize, Organization of Materials, and Monitor), and The Global Executive Composite is a summary score which incorporates all eight clinical scales. Higher scores indicate a greater impairment for each scale and index. Clinical validity has been supported with myelomeningocele study groups. The scale was shown to be valid and reliable in Turkish children and adolescents.

**Social Responsiveness Scale (SRS)**

The SRS is a parent-completed questionnaire exploring emotionally appropriate reciprocal social interaction and communication in children. It consists of 65 items rated on a four-point scale from “Not true” to “Almost always true” by the parent based on the child’s behavior over the past six months and generating one total score. Higher total scores indicate a greater severity of social impairment. The reliability and validity of the Turkish version were conducted and presented in the 19th National Child and Adolescent Psychiatry Congress in 2009. The scale was shown to be valid and reliable in Turkish children and adolescents.

**Aberrant behavior checklist (ABC)**

The ABC has 58 items which are rated on a four-point scale ranging from 0 to 3. The items are scored into five subscales as follows: (i) Irritability, Agitation, Crying, (ii) Lethargy, Social Withdrawal, (iii) Stereotypic Behavior, (iv) Hyperactivity, Noncompliance, and (v) Inappropriate Speech. The Turkish version was developed and the congruent and criterion validity of the ABC was investigated in a Turkish clinical sample.

**Childhood Autism Rating Scale (CARS)**

The CARS has 15 items that are rated on a seven-point scale of behavioral observation ranging from 1 to 4 including intermediate values. The scale was developed as a tool to aid the diagnosis of Autism Spectrum Disorder (ASD) by Schopler. It was adapted to Turkish language and the cut-off of the CARS was calculated as 29.5 for Turkish children.

**Statistical analysis**

The sample size was calculated based on a previous study (the effect size [Cohen’s d value] was found to be 1.25 for MI score). We calculated the sample size using the G*Power version 3.1 software (Heinrich-Heine-Universität Düsseldorf, Düsseldorf, Germany) and, accordingly, at least 25 patients per group were needed with an alpha coefficient of 0.05 and a power of 99% based on the t-test. Statistical analysis was performed using the PASW for Windows version 17.1 software (SPSS Inc., Chicago, IL, USA). The groups were compared in terms of demographic characteristics and psychiatric diagnosis using the chi-square tests for categorical variables and Mann-Whitney U test for continuous variables. Continuous variables were expressed in mean ± standard deviation (SD) or median (min-max), while categorical variables were expressed in number and frequency. A p value of <0.05 was considered statistically significant.

**RESULTS**

Baseline demographic and clinical characteristics of the children with SB and typically developing children are shown in Table 1. Both groups were similar with respect to age and sex distribution.

Table 2 shows the distribution of psychiatric diagnoses. Accordingly, 51.6% of the children with SB had at least one psychiatric diagnosis and 6.5% of them had two or more diagnosis. The most common psychiatric disorders were internalizing disorders and attention deficit hyperactivity disorder (ADHD). The rate of psychiatric disorders was significantly higher in
### TABLE 1
Demographic and clinical characteristics of children with SB and TD children

|                     | SB (n=31) | TD (n=36) | p   |
|---------------------|-----------|-----------|-----|
| **Age (year)**      | 9.4±2.8   | 9.8±2.4   | 0.49|
| **Sex**             |           |           |     |
| Male                | 11 (35.5%)| 16 (44.4%)| 0.46|
| Female              | 20 (64.5%)| 20 (55.6%)|     |
| **Lesion level**    |           |           |     |
| Sacral              | 10 (14.9%)|           |     |
| Lumbosacral         | 7 (10.4%) |           |     |
| Midlumbar           | 11 (16.4%)|           |     |
| Thoracic            | 3 (4.5%)  |           |     |
| **Hoffer ambulatory criteria** | | | |
| Community walker    | 12 (17.9%)|           |     |
| Household walker    | 8 (11.9%) |           |     |
| Exercise walker     | 7 (10.4%) |           |     |
| Nonwalker (nonambulator) | 4 (6%)  |           |     |
| **Psychiatric diagnosis** | 16 (51.6%)| 5 (13.9%) | 0.001|

**WISC-R**

|                     | Mean±SD  | Median | Min-Max | Mean±SD  | Median | Min-Max | p   |
|---------------------|----------|--------|---------|----------|--------|---------|-----|
| **Verbal IQ**       | 7.5±3.1  | 7.5    | 2-16    | 10.3±3.1 | 10.5   | 1-16    | 0.001|
| **Performance IQ**  | 10.4±2.8 | 10     | 6-19    | 11.9±2.0 | 12     | 7-16    | 0.015|
| Picture completion  | 9.5±3.1  | 10     | 1-15    | 9.0±3.2  | 11     | 6-16    | 0.042|
| Picture arrangement | 10.3±3.0 | 9      | 2-17    | 10.3±3.0 | 11     | 5-18    | 0.11  |
| **ABC total**       | 15.5±15.7| 10     | 0-67    | 14±13.7  | 10     | 0-55    | 0.87  |
| Hyperactivity       | 5.0±5.8  | 3.5    | 0-24    | 4.6±5.4  | 2.5    | 0-21    | 0.84  |
| Irritability        | 5.6±6.3  | 3      | 0-28    | 5.3±6.0  | 3      | 0-23    | 0.91  |
| Lethargy            | 2.7±2.9  | 1      | 0-9     | 2.5±2.9  | 2      | 0-13    | 0.82  |
| Stereotypic behavior| 2.3±3.0  | 1.5    | 0-11    | 1.3±2.2  | 0      | 0-9     | 0.13  |
| **BRIEF**           | 59.8±13.9| 58     | 37-94   | 57.3±12.7| 56     | 36-91   | 0.44  |
| Inhibit             | 22.0±6.2 | 21     | 15-39   | 20.8±5.8 | 19.5   | 15-43   | 0.41  |
| Shift               | 19.2±4.8 | 19     | 11-27   | 17.9±4.3 | 18     | 11-27   | 0.23  |
| Emotional control   | 18.6±4.6 | 18     | 11-29   | 18.7±4.2 | 19     | 10-28   | 0.98  |
| Metacognition Index | 84.7±18.5| 86     | 52-127  | 77.9±14.8| 75.5   | 53-123  | 0.11  |
| Initiate            | 14.5±3.5 | 15     | 8-21    | 13.6±2.7 | 13.5   | 8-19    | 0.26  |
| Working memory      | 19.3±4.8 | 18     | 12-32   | 17.3±3.6 | 17     | 11-29   | 0.06  |
| Plan/organize       | 25.3±6.7 | 25     | 14-39   | 21.4±4.8 | 21     | 14-37   | 0.007 |
| Organization of materials | 11.9±3.7 | 11  | 8-20 | 12.7±3.0 | 12 | 8-22 | 0.32 |
| Monitor             | 13.7±3.3 | 14     | 8-21    | 12.9±3.2 | 12     | 8-20    | 0.3   |
| Total               | 144.5±30.5| 142 | 92-216 | 135.2±24.6| 130 | 88-200 | 0.17 |
| **SRS**             | 45.8±29.6| 45     | 5-124   | 30.2±24.3| 22     | 6-93    | 0.02  |

SB: Spina bifida; TD: Typically developing; SD: Standard deviation; Min: Minimum; Max: Maximum; WISC-R: Wechsler Intelligence Scale for Children-Revised; ABC: Aberrant Behavior Checklist; BRIEF: Child Behavioral Rating Inventory of Executive Functions; SRS: Social Responsiveness Scale; P-values in bold indicate statistical significance.

### Table 2
Psychiatric diagnosis according to K-SADS-PL

|                     | SB  | TD  | p   |
|---------------------|-----|-----|-----|
| Attention-deficit/hyperactivity disorder | 8  | 25.8| 3  | 8.3 | 0.054 |
| Internalizing disorders | 9  | 29  | 1  | 2.8 | 0.004 |
| Generalized anxiety disorder | 3  | 9.7 | 1  | 2.8 | 0.23  |
| Separation anxiety disorder | 3  | 9.7 | 0  | 0   | 0.056 |
| Social anxiety disorder | 2  | 6.5 | 0  | 0   | 0.12  |
| Specific phobia | 2  | 6.5 | 1  | 2.8 | 0.46  |
| Tic disorders | 1  | 3.2 | 0  | 0   | 0.27  |

K-SADS-PL: Kiddie Schedule for Affective Disorders and Schizophrenia Present and Lifetime Version; SB: Spina bifida; TD: Typically developing children; P-values in bold indicate statistical significance.
Psychosocial problems in children with SB

the children with SB (p=0.001). The percentage of internalizing disorders rate was also significantly higher in SB group (p=0.004). The ADHD rate was higher in the SB group (25.8%; p=0.054). Information, vocabulary, picture completion subtests of WISC-R were lower in the SB group than the control group (p=0.001, p=0.015, and p=0.042, respectively). The WISC-R subtests did not significantly differ according to lesion level or ambulatory status (p>0.05) (Tables 3 and 4). There were no statistically significant differences between the children with SB and typically developing children in terms of the ABC scores (p>0.05). Stereotypic behavior subtest scores of the ABC did not significantly differ between the groups (p>0.05). Irritability and lethargy subtest scores of the ABC were significantly lower in the children with high level (p=0.02 and p=0.01, respectively), while hyperactivity, irritability, and lethargy subtest scores of the ABC were significantly lower in the non-community walkers, compared to the community walkers (p=0.03, p=0.003, and p=0.001, respectively) (Tables 3 and 4).

In addition, planning and organization subscale scores of the BRIEF were significantly higher in the SB group than the control group (p=0.007) (Table 1). The initiate and organization of materials subtest scores were significantly lower in children who were community walkers than the non-community walkers (p=0.02, p=0.002, and p=0.03, respectively) (Table 4).

All children scored below the CARS cut-off of 30 for ASD. The SRS total score was significantly higher in the children with SB than typically developing children (p=0.02) (Table 1). However, there was no statistically significant difference between either lower lesion level group and upper lesion level group, or the community walkers and non-community walkers (p>0.05) (Tables 3 and 4).

### Table 3

|                | Low level (n=17) | High level (n=14) |          |          |          |          |          |
|----------------|-----------------|-------------------|----------|----------|----------|----------|----------|
|                | Mean±SD         | Median            | Min-Max  | Mean±SD  | Median   | Min-Max  | p        |
| WISC-R         |                 |                   |          |          |          |          |          |
| Verbal IQ      |                 |                   |          |          |          |          |          |
| Information    | 7.5±3.0         | 7                 | 2-16     | 7.6±3.4  | 8        | 3-14     | 0.72     |
| Vocabulary     | 10.3±2.6        | 10                | 7-15     | 10.6±3.1 | 10       | 6-19     | 0.8      |
| Performance IQ |                 |                   |          |          |          |          |          |
| Picture completion | 10.4±2.9  | 10.5              | 5-15     | 8.6±3.3  | 10       | 1-13     | 0.24     |
| Picture arrangement | 9.8±2.8  | 9.5               | 6-17     | 8.2±3.6  | 8.5      | 2-15     | 0.24     |
| ABC total      | 21.7±17.5       | 15.5              | 0-67     | 9.5±11.4 | 5        | 0-40     | 0.02     |
| Hyperactivity  | 6.5±6.0         | 5                 | 0-24     | 3.6±5.4  | 2        | 0-20     | 0.07     |
| Irritability   | 8.5±7.6         | 5.5               | 0-28     | 2.7±2.9  | 2.5      | 0-9      | 0.02     |
| Lethargy       | 4±3.4           | 3.5               | 0-9      | 1.2±1.5  | 1        | 0-4      | 0.01     |
| Stereotypic behavior | 2.7±3.3  | 2                 | 0-11     | 2±2.9    | 1        | 0-9      | 0.42     |
| BRIEF          |                 |                   |          |          |          |          |          |
| Behavioral Regulation Index | 63.3±12.9 | 62                | 39-94    | 57.6±15.1| 53.5     | 37-89    | 0.26     |
| Inhibit        | 23.3±5.8        | 22                | 15-39    | 21.1±6.8 | 19.5     | 15-37    | 0.24     |
| Shift          | 21.4±6.6        | 21                | 13-27    | 17.6±4.7 | 17.5     | 11-26    | 0.09     |
| Emotional control | 19.1±4.6  | 18                | 11-29    | 18.9±4.6 | 18.5     | 11-28    | 0.71     |
| Metacognition Index | 88±17.5  | 92                | 57-121   | 79.9±20.3| 76.5     | 52-127   | 0.13     |
| Initiate       | 15.6±3.3        | 16                | 10-21    | 12.9±3.4 | 12       | 8-21     | 0.02     |
| Working memory | 18.9±4.2        | 19                | 13-28    | 19.4±5.8 | 18       | 12-32    | 0.83     |
| Plan/organize  | 26.7±7          | 26                | 17-39    | 23.9±6.8 | 23.5     | 14-37    | 0.19     |
| Organization of materials | 13.1±3.9 | 13                | 8-20     | 10.4±3.1 | 9        | 8-16     | 0.02     |
| Monitor        | 13.7±3.6        | 15                | 8-19     | 13.4±3.2 | 13.5     | 8-21     | 0.5      |
| Total          | 151.3±27.8      | 151               | 96-193   | 137.6±34.8| 131.5    | 92-216   | 0.14     |
| SRS            | 50.3±31.6       | 50                | 11-124   | 39.6±9.1 | 30.5     | 5-103    | 0.21     |

SD: Standard deviation; Min: Minimum; Max: Maximum; WISC-R: Wechsler Intelligence Scale for Children-Revised; ABC: Aberrant Behavior Checklist; BRIEF: Child Behavioral Rating Inventory of Executive Functions; SRS: Social Responsiveness Scale; P-values in bold indicate statistical significance.
DISCUSSION

In the present study, children with SB and typically developing children were compared in terms of psychiatric symptomatology, cognitive functioning, behavioral problems, executive functions, and social responsiveness. Also, we investigated whether psychiatric symptomatology, cognitive functioning, behavioral problems, executive functions, and social responsiveness varied as a function of two variables in children with myelomeningocele: lesion level or community ambulatory status. Our study results showed that psychiatric pathology was more common among children with SB than typically developing children.

Similar to the results of the present study, children with SB were found to be at risk for exhibiting higher levels of internalizing symptoms (e.g., anxiety and depression), compared to their peers in previous studies. Ammerman et al. detected ADHD and internalizing disorders as the two most prevalent diagnostic categories using the Child Symptom Inventory (CSI). Burmeister et al. also reported the ADHD prevalence as 31% in children with SB using the Swanson Nolan Achenbach Pelham-IV (SNAP IV). Fletcher et al. reported impaired behavioral adjustment in children with SB using the Child Behavior Checklist (CBCL). However, the CSI, SNAP IV, and CBCL are all parent report screening tools with a lower diagnostic value. Different from the previous studies, in the present study, we examined the children using a semi-structured interview to reach a final diagnosis. Although the literature reports that children with SB have an elevated risk of depression, interestingly none of the participants in the present study reached the diagnostic cut-off for major depressive disorder (MDD). A possible explanation can be that reaching the diagnostic cut-off for MDD is more unlikely using a semi-structured interview compared to a self- or parent-report rating scale. Therefore, it is likely that over-diagnosis is an issue with rating scales, in general.

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**TABLE 4**

Comparison of outcomes according to ambulatory status

|                      | Non-community walker (n=19) | Community walker (n=12) | p    |
|----------------------|----------------------------|-------------------------|------|
|                      | Mean±SD | Median | Min-Max | Mean±SD | Median | Min-Max |      |
| WISC-R               |          |        |        |          |        |        |      |
| Verbal IQ            |          |        |        |          |        |        |      |
| Information          | 8±3.5    | 8      | 3-16   | 6.7±2.3  | 7      | 2-10   | 0.36 |
| Vocabulary           | 10.7±2.9 | 10     | 6-19   | 10.1±2.8 | 9      | 7-15   | 0.53 |
| Performance IQ       |          |        |        |          |        |        |      |
| Picture completion   | 9.1±3.4  | 10     | 1-15   | 10.5±2.4 | 11     | 7-13   | 0.37 |
| Picture arrangement  | 8.6±3.4  | 10     | 2-15   | 9.8±3.0  | 9      | 7-17   | 0.56 |
| ABC total            | 9.6±10.8 | 6      | 0-40   | 27.6±7.7 | 25     | 7-67   | **0.003** |
| Hyperactivity        | 3.6±5.0  | 2      | 0-20   | 8±6.5    | 6      | 0-24   | **0.03** |
| Irritability         | 2.8±2.9  | 2      | 0-9    | 11.1±7.9 | 10     | 2-28   | **0.003** |
| Lethargy             | 1.3±1.5  | 1      | 0-4    | 5.3±3.3  | 6      | 1-9    | **0.001** |
| Stereotypic behavior| 2.0±3.8  | 1      | 0-9    | 3.2±3.8  | 2      | 0-11   | 0.41 |

**BRIEF**

|                      | Mean±SD | Median | Min-Max | Mean±SD | Median | Min-Max | p    |
|----------------------|---------|--------|---------|---------|--------|---------|------|
| Behavioral Regulation Index | 57.7±1  | 55     | 37-89   | 66±13.1 | 62.5   | 45-94   | 0.25 |
| Inhibit              | 21.5±6.4 | 21     | 15-37   | 23.6±6.2 | 22     | 15-39   | 0.45 |
| Shift                | 18.4±9.4 | 18     | 11-27   | 21.9±3.9 | 21     | 14-27   | 0.08 |
| Emotional control    | 18.2±4.4 | 18     | 11-28   | 20.5±4.7 | 19     | 13-29   | 0.56 |
| Metacognition Index  | 78.9±19.0 | 80    | 52-127  | 93.9±15.6 | 93   | 66-121  | **0.02** |
| Initiate             | 12.9±3.2 | 12     | 8-21    | 16.8±2.7 | 16.5  | 12-21   | **0.002** |
| Working memory       | 18.4±5.4 | 18     | 12-32   | 20.7±3.7 | 21     | 16-28   | 0.07 |
| Plan/organize        | 23.7±6.3 | 24     | 14-37   | 28.4±7.3 | 26.5  | 19-39   | 0.07 |
| Organization of materials | 11.3±7.3 | 10    | 8-19    | 13.3±3.7 | 12.5  | 9-20    | **0.03** |
| Monitor              | 12.9±3.2 | 13     | 8-21    | 14.7±3.4 | 16    | 9-19    | 0.06 |
| Total                | 139.6±32.0 | 134 | 92-216  | 159.9±25.7 | 155 | 111-193 | **0.05** |
| SRS                  | 41.7±28.9 | 32     | 5-103   | 51.7±33.6 | 49   | 12-124  | 0.32 |

SD: Standard deviation; Min: Minimum; Max: Maximum; WISC-R: Wechsler Intelligence Scale for Children-Revised; ABC: Aberrant Behavior Checklist; BRIEF: Child Behavioral Rating Inventory of Executive Functions; SRS: Social Responsiveness Scale; P-values in bold indicate statistical significance.
Psychiatric evaluation is necessary for identifying psychopathology in children with SB.

In the current study, we also evaluated the intelligence levels and behavioral parameters related to executive functions. The three subtests of WISC-R including information, vocabulary, and picture completion were within the average range, but lower than the control group. Although two groups had no significant differences in most subtests of BRIEF, planning and organizing components of executive functions were scored higher in children with SB. These findings are in line with the previous literature.\(^4,7,30\) Children with SB demonstrated typical patterns of cognitive deficits, such as difficulties in perceptual, visuo-motor, mnemonic areas and impairments in executive functioning, including working memory and ability to inhibit response.\(^7,31-33\) On the other hand, most of BRIEF subtest scores, MI, BRI and the total scores did not significantly differ between the two groups. This finding can be explained by the sample characteristics, as the children with SB were clinically followed by a multidisciplinary team in our study and, therefore, their care was most likely managed better than the general SB population. Also, it is important to note that presence of hydrocephalus is likely to have influenced the intellectual and neuropsychological functioning of children. Previously, it has been concluded that hydrocephalus rather than SB itself causes cognitive deficits and lower IQ in children with SB.\(^30\) Similarly, children with hydrocephalus showed a poorer performance IQ than verbal IQ. However, this discrepancy was not observed in children with only SB (without hydrocephalus). Following the shunt operations, the prevalence of mental retardation in SB population has decreased.\(^30\) In our study, we included only children with no intellectual disabilities without any history or presence of shunt dysfunction to compare the children with SB to their peers.

Furthermore, we assessed social responsiveness which involves elements of social cognition in this study. Previous reports have suggested that children with SB tend to have fewer social contacts and difficulties in unstructured social situations, more problems with social relationship and are more socially immature.\(^7,8\) Attention and executive function are necessary for social interactions in children with SB.\(^7,8\) In our study, the SRS total score was significantly higher in children with SB, compared to typically developing children, supporting the previous findings on social functioning of children with SB. Neurocognitive deficits may be associated with the social cognition deficits. In addition, posterior attention system plays a critical role in joint attention development, which is vital for social competence;\(^34\) and it has been hypothesized that executive dysfunction in children with SB is related to posterior attention system;\(^35,36\) therefore, problems related to posterior attention system may lead to social cognitive impairments in children with SB. Although none of the children were diagnosed as having ASD according to the CARS or clinical evaluation, as previously reported, social cognition abilities may be impaired. Even if children do not have autism, they should be supported in terms of social skills.

In the present study, outcomes in relation to lesion level and ambulatory status were also investigated. The results showed that the lesion level had no significant association with psychopathology or intellectual capacity as measured by the WISC-R. Similar to the findings, Holmbeck et al.\(^6\) demonstrated that child psychosocial adjustment did not significantly differ in relation to lesion level. Similarly, Ammerman et al.\(^3\) found no statistically significant difference between the lower and higher lesion groups and related to the ambulation status (i.e., no assistance, assistance, wheelchair) in terms of psychiatric symptomatology. Different from the previous studies, in the present study, the children were categorized according to their ambulatory status with the Hoffer criteria which evaluates ambulatory status in the context of the International Classification of Functioning. Children with a lower lesion level and children who are community ambulators demonstrated statistically more important impairments in the initiation and organization of materials subtests of BRIEF and showed more maladaptive behaviors in ABC.

In the study by Rose and Holmbeck,\(^7\) the spinal lesion level was not found to be a predictor of BRIEF performance. Ammerman et al.\(^3\) found that ambulation status or lesion level had no relationship with the presence of ADHD. Supporting the study of Ammerman et al.\(^3\) Tarazi et al.\(^23\) reported that the BRIEF MI and BRI scores were not significant related to the lesion level. In contrast, Burmeister et al.\(^25\) noted significant differences in the lesion level and ambulation among the ADHD subtypes (I= inattentive type; hyperactive/impulsive type [ADHD-HI]; C= combined type). They detected that children with SB and ADHD-C were more likely
to have spinal lesions below the thoracic level. As a possible explanation of this result, the researchers suggested that, since children with sub-thoracic spinal lesions were more independent and ambulatory than others, combined type ADHD was more prevalent in this group. In the present study, the initiate and organization of materials subtests of BRIEF and hyperactivity, irritability and lethargy subtests of ABC were found to be lower in the non-community walkers. The common aspect of the aforementioned subtests is being related to movement. Despite being used in many of the previous studies, none of the psychiatric measurement tools have been validated in children with SB and they are not specifically developed for this population. From this point of view, they might have been not very useful in reflecting the exact status of children with SB. Since children with SB are at a high risk for poor neuropsychological and psychological outcomes, development of new SB specific tools is needed for appropriate treatment and diagnosis.

Nonetheless, the current study has several limitations. First, the cross-sectional nature of the study limits to learn how neuropsychological and behavioral problems may change over time. A longitudinal study would provide more useful information about psychosocial functioning. Second, the participants of the present study were selected from a group of children who were long followed by a multidisciplinary team. Therefore, it limits the generalization of results. Third, environmental factors such as socioeconomic status, family environment, and resilience factors such as coping skills could impact mental health; however, such factors were unable to be assessed in the present study.

On the other hand, to the best of our knowledge, this study is the first to assess psychiatric symptomatology using a gold-standard semi-structural interview, which appears as the main strength of the present study.

In conclusion, psychiatric disorders are more prevalent in children with SB and impairment in certain aspects of executive functioning appears to be associated with the lesion level and ambulatory status of children with SB. However, there is still a need for developing assessment tools which are sensitive for assessing neuropsychological functioning in this patient population.

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REFERENCES
1. Holmbeck GN, Devine KA. Psychosocial and family functioning in spina bifida. Dev Disabil Res Rev 2010;16:40-6.
2. Oakeshott P, Hunt GM, Poulton A, Reid F. Expectation of life and unexpected death in open spina bifida: a 40-year complete, non-selective, longitudinal cohort study. Dev Med Child Neurol 2010;52:749-53.
3. Ammerman RT, Kane VR, Slomka GT, Reifisch MD, Gadow KD. Psychiatric Symptomatology and Family Functioning in Children and Adolescents with Spina Bifida. J Clin Psychol Med Settings 1998;5:449-65.
4. Kelly NC, Ammerman RT, Rausch JR, Ris MD, Yeates KO, Oppenheimer SG, et al. Executive functioning and psychological adjustment in children and youth with spina bifida. Child Neuropsychol 2012;18:417-31.
5. Snow JH. Executive Processes For Children With Spina Bifida. Children's Health Care 1999;28:241-53.
6. Holmbeck GN, Faier-Routman J. Spinal lesion level, shunt status, family relationships, and psychosocial adjustment in children and adolescents with spina bifida myelomeningocele. J Pediatr Psychol 1995;20:817-32.
7. Rose BM, Holmbeck GN. Attention and executive functions in adolescents with spina bifida. J Pediatr Psychol 2007;32:983-94.
8. Holbein CE, Lennon JM, Kolbuck VD, Zebracki K, Roache CR, Holmbeck GN. Observed differences in social behaviors exhibited in peer interactions between youth with spina bifida and their peers: neuropsychological correlates. J Pediatr Psychol 2015;40:320-35.
9. Sirzai H, Dogu B, Demir S, Yilmaz F, Kuran B. Assessment on self-care, mobility and social function of children with spina bifida in Turkey. Neural Regen Res 2014;9:1234-40.
10. Pauly M, Cremer R. Levels of mobility in children and adolescents with spina bifida-clinical parameters predicting mobility and maintenance of these skills. Eur J Pediatr Surg 2013;23:110-4.
11. Gioia GA, Isquith PK, Guy SC, Kenworthy L. BRIEF: Behavior rating inventory of executive function. Lutz, FL: Psychological Assessment Resources; 2000.
12. Constantino JN, Gruber C. The Social Responsiveness Scale. Los Angeles, CA: Western Psychological Services; 2005.
13. Aman MG, Singh NN, Stewart AW, Field CJ. The aberrant behavior checklist: a behavior rating scale for the assessment of treatment effects. Am J Ment Defic 1985;89:485-91.
14. Kaufman J, Birmaher B, Brent D, Rao U, Flynn C, Moreci P, et al. Schedule for Affective Disorders and Schizophrenia for School-Age Children-Present and Lifetime Version (K-SADS-PL): initial reliability and validity data. J Am Acad Child Adolesc Psychiatry 1997;36:980-8.
15. Göklер B, Ünal F, Pehlivantürk B, Pehlivantürk ÇE, Akdemir D, Taner Y. Okul çağı çocukları için duygulanım bozuklukları ve şizofreni görüşme çizelgesi -şimdi ve yaşam boyu şekli-Türkçe uyarlamasının geçerlik ve güvenirliği. Çocuk ve Gençlik Ruh Sağlığı Dergisi 2004;11:109-16.

16. Savaşır I, Şahin N. Wechsler Çocuklar İçin Zeka Ölçeği (WISCR) El Kitabı. Ankara: Türk Psikologlar Derneği Yayınları; 1995.

17. Mahone EM, Zabel TA, Levey E, Verda M, Kinsman S. Parent and self-report ratings of executive function in adolescents with myelomeningocele and hydrocephalus. Child Neuropsychol 2002;8:258-70.

18. Batan SN, Öktem-Tanör Ö, Kalem E. Reliability and validity studies of Behavioral Rating Inventory Of Executive Function (BRIEF) in a Turkish normative sample. Elementary Education Online 2001;10:894-904.

19. Ünal S, Güler AS, Dedeoğlu C, Taşkın B, Yazgan Y. Social Responsiveness in a clinical sample of ADHD: Comparison with a community control group, in 19th National Child and Adolescent Psychiatry Congress. April 14-18, 2009, Antakya, Turkey.

20. Karabekiroğlu K, Aman MG. Validity of the aberrant behavior checklist in a clinical sample of toddlers. Child Psychiatry Hum Dev 2009;40:99-110.

21. Schopler E, Reichler RJ, DeVellis RF, Daly K. Toward objective classification of childhood autism: Childhood Autism Rating Scale (CARS). J Autism Dev Disord 1980;10:91-103.

22. Sucuoğlu B, Öktem F, Akkök F. Otistik çocukların değerlendirilmesinde kullanılan ölçekler ilişkisi ve çalışmanın bir gelişmesi. 3P Derg 1996;4:116-21.

23. Tarazi RA, Zabel TA, Mahone EM. Age-related differences in executive function among children with spina bifida/hydrocephalus based on parent behavior ratings. Clin Neuropsychol 2008;22:585-602.

24. Fletcher JM, Brookshire BL, Landry SH, Bohan TP, Davidson KC, Francis DJ, et al. Behavioral adjustment of children with hydrocephalus: relationships with etiology, neurological, and family status. J Pediatr Psychol 1995;20:109-25.

25. Burmeister R, Hannay HJ, Copeland K, Fletcher JM, Boudousquie A, Dennis M. Attention problems and executive functions in children with spina bifida and hydrocephalus. Child Neuropsychol 2005;11:265-83.

26. Appleton PL, Ellis NC, Minchom PE, Lawson V, Böll V, Jones P. Depressive symptoms and self-concept in young people with spina bifida. J Pediatr Psychol 1997;22:707-22.

27. Holmbeck GN, DeLucia C, Essner B, Kelly L, Zebracki K, Friedman D, et al. Trajectories of psychosocial adjustment in adolescents with spina bifida: a 6-year, four-wave longitudinal follow-up. J Consult Clin Psychol 2010;78:511-25.

28. Nicholls EG, Arango-Lasprilla JC, Olivera Plaza SL, Mendez N, Quintero L, Velasco Trujillo DM, et al. Psychological Functioning in Youth With Spina Bifida Living in Colombia, South America. J Pediatr Psychol 2015;40:602-8.

29. Pinquart M, Shen Y. Depressive symptoms in children and adolescents with chronic physical illness: an updated meta-analysis. J Pediatr Psychol 2011;36:375-84.

30. Wills KW. Neuropsychological Functioning in Children With Spina Bifida and/or Hydrocephalus. Journal of Clinical Child Psychology 1993;22:247-65.

31. Brown TM, Ris MD, Beebe D, Ammerman RT, Oppenheimner SG, Yeates KO, et al. Factors of biological risk and reserve associated with executive behaviors in children and adolescents with spina bifida myelomenigocele. Child Neuropsychol 2008;14:118-34.

32. Mahone EM, Hagelthorn KM, Cutting LE, Schuerholz LJ, Pelletier SF, Rawlins C, et al. Effects of IQ on executive function measures in children with ADHD. Child Neuropsychol 2002;8:52-65.

33. Burro F, Cama A, Lertora V, Veneselli E, Rossetti S, Pezzuti L. Intellectual efficiency in children and adolescents with spina bifida myelomenigocele and shunted hydrocephalus. Dev Neuropsychol 2018;43:198-206.

34. Mundy P, Newell L. Attention, Joint Attention, and Social Cognition. Curr Dir Psychol Sci 2007;16:269-74.

35. Brewer VR, Fletcher JM, Hiscock M, Davidson KC. Attention processes in children with shunted hydrocephalus versus attention deficit-hyperactivity disorder. Neuropsychology 2001;15:185-98.

36. Dennis M, Barnes MA. The cognitive phenotype of spina bifida meningomyelocele. Dev Disabil Res Rev 2010;16:31-9.