Changes of Neurotransmitters in Youth with Internet and Smartphone Addiction: A Comparison with Healthy Controls and Changes after Cognitive Behavioral Therapy

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ABSTRACT

BACKGROUND AND PURPOSE: Neurotransmitter changes in youth addicted to the Internet and smartphone were compared with normal controls and in subjects after cognitive behavioral therapy. In addition, the correlations between neurotransmitters and affective factors were investigated.

MATERIALS AND METHODS: Nineteen young people with Internet and smartphone addiction and 19 sex- and age-matched healthy controls (male/female ratio, 9:10; mean age, 15.47 ± 3.06 years) were included. Twelve teenagers with Internet and smartphone addiction (male/female ratio, 8:4; mean age, 14.99 ± 1.95 years) participated in 9 weeks of cognitive behavioral therapy. Meshcher-Garwood point-resolved spectroscopy was used to measure γ-aminobutyric acid and Glx levels in the anterior cingulate cortex. The γ-aminobutyric acid and Glx levels in the addicted group were compared with those in controls and after cognitive behavioral therapy. The γ-aminobutyric acid and Glx levels correlated with clinical scales of Internet and smartphone addiction, impulsiveness, depression, anxiety, insomnia, and sleep quality.

RESULTS: Brain parenchymal and gray matter volume–adjusted γ-aminobutyric acid-to-creatine ratios were higher in subjects with Internet and smartphone addiction (P = .028 and .016). After therapy, brain parenchymal- and gray matter volume–adjusted γ-aminobutyric acid-to-creatine ratios were decreased (P = .034 and .026). The Glx level was not statistically significant in subjects with Internet and smartphone addiction compared with controls and posttherapy status. Brain parenchymal- and gray matter volume–adjusted γ-aminobutyric acid-to-creatine ratios correlated with clinical scales of Internet and smartphone addictions, depression, and anxiety. Glx/Cr was negatively correlated with insomnia and sleep quality scales.

CONCLUSIONS: The high γ-aminobutyric acid levels and disrupted balance of γ-aminobutyric acid-to-Glx including glutamate in the anterior cingulate cortex may contribute to understanding the pathophysiology and treatment of Internet and smartphone addiction and associated comorbidities.

ABBREVIATIONS: ACC = anterior cingulate cortex; bp- = brain parenchymal volume-adjusted; GABA = γ-aminobutyric acid; gm- = gray matter volume-adjusted; MEGA-PRESS = Meshcher-Garwood point-resolved spectroscopy; wm- = white matter volume-adjusted

Internet addiction is a behavioral addiction characterized by uncontrolled use of the Internet with tolerance, withdrawal symptoms, and compulsiveness. The prevalence of Internet addiction ranges from 1.5% to 8.2% and is much higher in adolescents and young adults in the Far East. Internet gaming addiction was listed as a research criterion for behavioral addiction in the fifth version of the Diagnostic and Statistical Manual of Mental Disorders. In recent years, a preoccupation with smartphones and their worldwide spread has resulted in smartphone addiction.

The mesocorticolimbic system is a dopaminergic projection engaged in common neurobiologic pathways of substance addiction. The anterior cingulate cortex (ACC) is part of the mesocorticolimbic system and is associated primarily with salient networks activated by reward-related stimuli. The ACC has been postulated to play a critical role with the insula in substance...
addiction, PET, SPECT, and electroencephalogram results reportedly show brain regions associated with substance addiction. The ACC has been one of the most frequently implicated regions in Internet addiction.

Substance addiction is associated with the neurotransmitter changes in the mesocorticolimbic system. Nicotine and alcohol indirectly enhance dopamine release via modulation of \( \gamma \)-aminobutyric acid (GABA) and glutamatergic neurons. MR spectroscopy using an editing pulse to create J-coupling can separate GABA and glutamate signals from other stronger overlying metabolite signals and has been used to characterize neurotransmitter changes in dynamic and interactive psychiatric disorders. The use of MR spectroscopy to study the ACC with respect to Internet and smartphone addiction can clarify common neurobiologic mechanisms for behavioral and substance addictions and provide clinical intervention to reduce the prevalence and related functional impairments in young people.

Currently, neurotransmitter changes have not been researched in terms of Internet and smartphone addiction. The purpose of this study was to reveal the associations between neurotransmitter changes and Internet and smartphone addiction and compare them with those in healthy controls and subjects postcognitive behavioral therapy results. In addition, correlations between neurotransmitter changes and affective changes in youth diagnosed with Internet and smartphone addiction were investigated.

**MATERIALS AND METHODS**

**Participants**

The institutional review board Korea University Ansan hospital approved this prospective study, and informed consent was obtained from the adolescents and parents. We included young people between the ages of 10 and 24 years who met the following criteria: 1) \( >50 \) points on the Internet Addiction Test modified from the Young Diagnostic Questionnaire; 2) \( >35 \) points on the Smartphone Addiction Scale-short version for adolescents; and 3) \( >75 \) points on the summed Internet Addiction Test and Smartphone Addiction Scale. Subsequently, the Mini-International Neuropsychiatric Interview was administered to exclude subjects who met the diagnostic criteria of psychotic disorders, such as schizophrenia spectrum and other psychotic disorders, bipolar I disorder, or substance use disorder.

We included sex- and age-matched healthy controls who met the following criteria: 1) \( <30 \) points on the Internet Addiction Test; 2) \( <30 \) points on the Smartphone Addiction Scale; and 3) \( <60 \) points on the summed Internet Addiction Test and Smartphone Addiction Scale.

**Psychology Tests**

The Young Internet Addiction Test measures the severity of Internet addiction and consists of 20 items scored on a 5-point Likert scale, covering the extent to which Internet use affects daily routines, social life, productivity, sleeping patterns, and feelings. The Smartphone Addiction Scale-short version for adolescents measures the severity of smartphone addiction. It consists of 10 items scored on a 6-point Likert scale.

Affective and cognitive characteristics of people with addiction and controls were evaluated using the Barratt Impulsiveness Scale, Hamilton Depression Rating Scale, Spielberger State-Trait Anxiety Inventory, Pittsburgh Sleep Quality Index, Insomnia Severity Index, and Mini-International Neuropsychiatric Interview. In order to the measurement of intelligence, the Korean version of the Wechsler Adult Intelligence Scale-IV for adolescents and adults older than 16 years of age and the Korean version of the Wechsler Intelligence Scale for Children-IV for children from 6 to 16 years of age were used.

**Cognitive Behavioral Therapy**

Cognitive behavioral therapy was modified from the Cognitive-Behavioral Therapy For Internet Gaming Addiction, and the Emotional Identification and Expression Abilities Improvement Program was reinforced. The treatment program consisted of the following 7 areas: recognizing the Internet behavior, modifying the cognitive distortion, finding appropriate alternative activities, promoting self-control, recognizing self-emotions and those of others, expressing emotions, and resolving interpersonal conflicts. The cognitive behavioral therapy consisted of a weekly 75-minute program for 9 weeks. The program was administered to the young subjects with Internet and smartphone addiction who agreed to participate in the therapy. Two or more absences were defined as therapy failure.

**MR Imaging Parameters**

MR imaging was performed within an hour after the psychological tests. In addition, 1 or 2 days after finishing the 9-week cognitive behavioral therapy program, MR imaging rescanning was performed within an hour after the psychological retests. MR imaging data were acquired with a 3T MR imaging scanner with a 32-channel phased array head coil (MAGNETOM Skyra; Siemens Healthcare, Erlangen, Germany). Mesecher-Garwood point-resolved spectroscopy (MEGA-PRESS) was programmed by inserting a dual-band radiofrequency pulse into the manufacturer’s PRESS sequence. A dual-band radiofrequency waveform was obtained using a public domain pulse design tool, MATPULSE software (https://cind.ucsf.edu/education/software/matpulse). GABA measurement was performed using the MEGA-PRESS pulse sequence from a voxel volume of 26.25 mL (voxel dimensions, \( 3.5 \times 3.0 \times 2.5 \text{ cm} \)) in the ACC (Fig 1), with the following acquisition parameters: TR, 2000 ms; TE, 68 ms; spectral width, 2000 Hz; number of oversampled data, 2048; number of signal averages, 256. GABA resonance at 3.01 ppm was detected by application of a refocusing pulse at 1.9 ppm during ON spectra and at 7.5 ppm during OFF spectra. Water signal suppression was achieved with the chemical shift selective imaging technique. First- and second-order shimming was performed for the voxels in the ACC, and the water line widths resulted in \( <16 \text{ Hz} \). Scan time was approximately 9 minutes.

Single-voxel \( ^1 \text{H}-\text{MR} \) spectroscopy with TE = 135 ms was obtained at the same voxels in the ACC (Fig 1), with the following parameters: TR, 2000 ms; number of samples, 64. In addition to MR spectroscopy, T1-weighted structural images were obtained using the MPRAGE sequence with the following...
parameters: TR, 2000 ms; TE, 3.55 ms; flip angle, 8°; section thickness, 1 mm; and acquisition matrix, 256 × 256.

**MR Spectroscopy Data Processing**

The raw GABA MR spectroscopy data were transferred to an offline computer and processed using home-programmed MR spectroscopy analysis software, which was programmed in Python (https://www.python.org/). Raw data were averaged in the time domain and baseline-corrected, and the areas under the GABA and Glx peaks at 3.01 and 3.8 ppm were measured by fitting to double and single Gaussian functions, respectively (Fig 2). The data for 1H-MR spectroscopy at TE = 135 ms were processed using LCModel 6.3 (http://www.lcmodel.com/), and the creatine peak was measured at 3.0 ppm and used as the internal reference. GABA and Glx relative to creatine were quantified, and the GABA-to-creatine ratio (GABA/Cr) and the Glx-to-creatine ratio (Glx/Cr) were obtained. The GABA-to-Glx ratio (GABA/Glx) was also obtained.

The volume fractions of brain parenchyma and gray matter in the MR spectroscopy voxels were obtained from T1 MPRAGE segmentation using the SPM program. Last, GABA/Cr and Glx/Cr are adjusted by multiplying brain parenchymal and gray matter volume fractions for each individual.

**Statistical Analyses**

All statistical analyses were performed using SPSS Statistics 20 (IBM). Statistical significance was defined as P ≤ .05. GABA and Glx differences between the youth with Internet and smartphone addictions and controls were evaluated using the Student t test. Correlations between neurotransmitters and psychological tests were evaluated using the Pearson correlation coefficient. Differences in GABA and Glx levels between pre- and postcognitive behavioral therapy were tested using paired t tests or paired Wilcoxon signed-rank tests based on whether the data met the assumption for a normal distribution.

**RESULTS**

The results of demographics and psychological tests are shown in Table 1. The addiction group consisted of 9 males and 10 females diagnosed with Internet and smartphone addictions. The mean age was 15.47 ± 3.06 years and ranged from 11 to 22 years. The control group consisted of 19 young healthy subjects, sex- and age-matched to the addiction group.

Internet and smartphone addiction scores were significantly higher in the addiction group compared with healthy controls (P < .001) as well as depression (P = .018), state, trait, and total anxiety scores (P < .001, P < .001, and P = .001, respectively); the impulsivity score (P = .001); insomnia severity (P = .006); and poor sleep quality (P = .008). A significant difference in the intelligence quotient was not observed between groups (P = .467).

Eight males and 4 females participated in cognitive behavioral therapy. The mean age was 14.0 ± 1.95 years and ranged from 11 to 17 years. Teenagers with Internet and smartphone addiction significantly improved after 9 weeks of cognitive behavioral therapy based on Internet and smartphone addiction scales (P = .001 and < .001, respectively), though Internet Addiction Test scores and the summed scores of the Internet Addiction Test and Smartphone Addiction Scale still met the criteria of Internet and smartphone addiction. However, psychological and sleep test scores were not significantly changed after therapy.

GABA levels in subjects with Internet and smartphone addictions are summarized in Table 2, and GABA levels after 9 weeks of cognitive behavioral therapy are summarized in Table 3. Brain parenchymal volume-adjusted GABA/Cr (bp-GABA/Cr) was higher in the addiction group compared with the controls (P = .028) and significantly decreased after cognitive behavioral therapy (P = .034). Gray matter volume-adjusted GABA/Cr (gm-GABA/Cr) was also higher in the addiction group (P = .016) and significantly decreased after cognitive behavioral therapy (P = .026; Fig 3A).

Glx levels in subjects with Internet and smartphone addiction are summarized in Table 2, and Glx levels after 9 weeks of cognitive behavioral therapy are summarized in Table 3. A significant difference in Glx levels was not observed in the addiction group compared with the control group. Glx/Cr adjusted brain parenchymal (bp-Glx/Cr) and gray matter volumes (gm-Glx/Cr) were
lower in the addiction group but without statistical significance ($P = .375$ and .587, respectively; Fig 3B). The Glx level was increased after cognitive and behavioral therapy but without statistical significance ($P = .096$ and .131, respectively; Fig 3B).

The results of correlations between neurotransmitters and psychological tests are described in Table 4. The GABA/Cr adjusted with gray-matter volume fraction (gm-GABA/Cr) positively correlated with the Internet Addiction Test and the Smartphone Addiction Scale as well as the sum of both scales. (Fig 4A) The bp-GABA/Cr correlated with the Internet Addiction Test but was not significantly correlated with Smartphone Addiction Scale.

The gm-GABA/Cr significantly correlated with the depression score ($P = .046$), state anxiety ($P = .032$), and total anxiety ($P = \ldots$)
Table 1: Demographics, psychological tests, and pre- and postcognitive behavioral therapy data

<table>
<thead>
<tr>
<th></th>
<th>Internet Addiction (Mean)</th>
<th>Control (Mean)</th>
<th>PreTx (Mean)</th>
<th>PostTx (Mean)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(n = 19)</td>
<td>(n = 12)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sex (M/F)</td>
<td>9:10</td>
<td>1.0</td>
<td>8.4</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (yr)</td>
<td>15.47 ± 3.06 (range, 11–22)</td>
<td>1.0</td>
<td>14.00 ± 1.95 (11–17)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>IAT</td>
<td>63.32 ± 15.15</td>
<td>27.37 ± 4.83</td>
<td>&lt;.001*</td>
<td>68.42 ± 14.74</td>
<td>.001*</td>
</tr>
<tr>
<td>SAS</td>
<td>45.53 ± 7.40</td>
<td>17.89 ± 8.12</td>
<td>&lt;.001*</td>
<td>45.83 ± 6.42</td>
<td>.016</td>
</tr>
<tr>
<td>HRSD</td>
<td>50.16 ± 7.54</td>
<td>39.05 ± 4.93</td>
<td>&lt;.001*</td>
<td>49.27 ± 7.84</td>
<td>.085</td>
</tr>
<tr>
<td>BIS</td>
<td>57.53 ± 7.63</td>
<td>45.58 ± 11.72</td>
<td>&lt;.001*</td>
<td>54.27 ± 6.94</td>
<td>.093</td>
</tr>
<tr>
<td>ISI</td>
<td>6.89 ± 4.977</td>
<td>3.11 ± 2.558</td>
<td>.006*</td>
<td>5.17 ± 3.74</td>
<td>.891</td>
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<tr>
<td>PSQI</td>
<td>5.95 ± 3.24</td>
<td>3.53 ± 1.87</td>
<td>.008*</td>
<td>4.67 ± 2.31</td>
<td>.777</td>
</tr>
<tr>
<td>IQ</td>
<td>95.18 ± 2.88</td>
<td>98 ± 11.12</td>
<td>.467</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note:—PreTx indicates pre-cognitive behavioral therapy; PostTx, post-cognitive behavioral therapy; IAT, Internet Addiction Test; SAS, Smartphone Addiction Scale; HRSD, Hamilton Rating Scale for Depression; STAI, State-Trait Anxiety Inventory; BIS, Barratt Impulsiveness Scale; ISI, Insomnia Severity Index; PSQI, Pittsburgh Sleep Quality Assessment; IQ, intelligence quotient.

* P value ≤ .05

Table 2: Subjects with Internet addiction versus controls

<table>
<thead>
<tr>
<th></th>
<th>bp-GABA/Cr</th>
<th>gm-GABA/Cr</th>
<th>wm-GABA/Cr</th>
<th>bp-GLX/Cr</th>
<th>gm-GLX/Cr</th>
<th>wm-GLX/Cr</th>
</tr>
</thead>
<tbody>
<tr>
<td>IA (Mean)</td>
<td>73.85 ± 22.74</td>
<td>49.10 ± 15.27</td>
<td>24.90 ± 8.25</td>
<td>167.74 ± 43.44</td>
<td>106.87 ± 39.90</td>
<td>56.25 ± 13.74</td>
</tr>
<tr>
<td>Controls (Mean)</td>
<td>56.03 ± 25.25</td>
<td>36.27 ± 16.02</td>
<td>19.77 ± 9.71</td>
<td>179.55 ± 37.30</td>
<td>117.45 ± 25.86</td>
<td>62.67 ± 13.12</td>
</tr>
<tr>
<td>P value</td>
<td>.026*</td>
<td>.016*</td>
<td>.088</td>
<td>.375</td>
<td>587.</td>
<td>.349</td>
</tr>
</tbody>
</table>

Note:—wm-GABA/Cr indicates white matter volume-adjusted GABA-to-creatine ratio; wm-GLX/Cr, white matter volume-adjusted glutamate and glutamine-to-creatine ratio; IA, Internet addiction.

* P value ≤ .05

Table 3: Pre- and postcognitive behavioral therapy data

<table>
<thead>
<tr>
<th></th>
<th>bp-GABA/Cr</th>
<th>gm-GABA/Cr</th>
<th>wm-GABA/Cr</th>
<th>bp-GLX/Cr</th>
<th>gm-GLX/Cr</th>
<th>wm-GLX/Cr</th>
</tr>
</thead>
<tbody>
<tr>
<td>PreTx (mean)</td>
<td>74.20 ± 23.58</td>
<td>46.77 ± 15.95</td>
<td>24.43 ± 8.32</td>
<td>177.19 ± 47.81</td>
<td>119.07 ± 34.37</td>
<td>58.12 ± 14.81</td>
</tr>
<tr>
<td>PostTx (mean)</td>
<td>52.72 ± 14.52</td>
<td>34.60 ± 9.28</td>
<td>18.11 ± 5.67</td>
<td>215.65 ± 48.31</td>
<td>142.46 ± 34.54</td>
<td>73.19 ± 16.40</td>
</tr>
<tr>
<td>P value</td>
<td>.034*</td>
<td>.026*</td>
<td>.071*</td>
<td>.096</td>
<td>.131</td>
<td>.06*</td>
</tr>
</tbody>
</table>

Note:—PreTx indicates pre-cognitive behavioral therapy; PostTx, post-cognitive behavioral therapy.

* P ≤ .05 based on paired t test or Wilcoxon signed rank test.

* wm-GABA/Cr in pre-cognitive behavioral therapy was not in a Gaussian distribution, and the nonparametric Wilcoxon signed rank test was applied in statistical analysis.

FIG 3. The boxplots of GABA/Cr (A) and Glx to Glx/Cr (B) in healthy controls and Internet and smartphone addicted subjects pre- and posttherapy. The horizontal line is the median, and the upper and lower ends of the boxes are the upper and lower quartiles, respectively. The vertical lines represent data ranges. PostTX indicates postcognitive behavioral therapy.
the major excitatory neurotransmitter, and GABA is mostly the main component in Glx based on MR spectroscopy. Glutamate is synthesized from glutamate via decarboxylation. GABA is mostly present in inhibitory local interneurons and is approximately 7-fold more concentrated in gray matter than in white matter. GABA and glutamate are also key opposite modulators of dopamine in mesocorticolimbic pathways, which are closely associated with addiction.

**DISCUSSION**

The results of this study showed that GABA levels were higher in the ACC in young subjects with Internet and smartphone addictions and decreased after 9 weeks of cognitive behavioral therapy. The bp-GABA and gm-GABA correlated with depression and anxiety scores as well as Internet and smartphone addiction scores. The bp-Glx and gm-Glx negatively correlated with insomnia severity and sleep quality.

Recently, the high accessibility of smartphones has led to severe functional impairments compared with conventional Internet addiction. Smartphone addiction is a behavioral or technological addiction based on Internet use and shares core symptoms and risk factors with Internet addiction. Therefore, smartphone addiction could be considered a category of Internet addiction; thus, Internet and smartphone addictions were not separated in this study.

GABA is the main inhibitory neurotransmitter and is present at approximately one-third of all synapses. The GABA concentration in the human brain is approximately 1 mM. GABA is present in inhibitory local interneurons and is approximately 7-fold more concentrated in gray matter than in white matter. Therefore, the gm-GABA level was statistically more significant than wm-GABA in this study (Tables 2–4). Glutamate is the major component in Glx based on MR spectroscopy. Glutamate is the major excitatory neurotransmitter, and GABA is mostly synthesized from glutamate via decarboxylation. GABA and glutamate are also key opposite modulators of dopamine in mesocorticolimbic pathways, which are closely associated with addiction.

In previous in vivo MR spectroscopy studies, the decreased GABA level was associated with depression and autism spectrum disorders. Schizophrenia did not show statistical significance in the meta-analysis but tended to exhibit lower GABA levels. However, the ACC also provides a higher signal-to-noise ratio and a homogeneous magnetic field because the imaging receiver compared with other regions of the mesocorticolimbic system such as the ventral tegmental area, nucleus accumbens, insula, and prefrontal cortex. Therefore, the ACC was thought to satisfy the hypothesis-driven and technical approaches when using MR spectroscopy to study Internet addiction.

In this study, GABA levels were increased in Internet- and smartphone-addicted youth compared with other psychiatric disorders. Two mechanisms can be postulated for the increased GABA levels in Internet and smartphone addiction. One involves the different neurobiology of Internet or...
behavioral addiction, and the other involves tolerance or an anti-reward mechanism. GABA inhibits synaptic signal transmission in the central nervous system. Activation of GABA_A and GABA_B receptors hyperpolarizes neurons and inhibits action potential generation and neurotransmission.24 The inhibition of GABA at the synapse attenuates the function of involved neural networks. Therefore, the increased GABA levels in subjects with Internet and smartphone addiction may be associated with the down-regulation of ACC functions, including impulsiveness control during the decision-making process under conditions of risk.18

The ACC is also important for input integration and regulation of processing in cognitive and emotional neural networks.35 In many neuroimaging and animal studies, the ACC was shown to be associated with affective disorders (Fig 1).35-37 ACC functional loss caused by lesions produces symptoms of emotional instability, inattention, and decreased social interaction.35,37 The emotional and personal traits caused by neurotransmitter derangement include risk factors for Internet and smartphone addictions and may be shared with the pathophysiology of psychiatric comorbidities in Internet and smartphone addiction, such as depression, attention deficit/hyperactivity disorder, and hostility.1,2 In a meta-analysis, Internet addiction was strongly associated with comorbid psychopathology, though the causal interaction could not be defined because of the insufficient longitudinal study.38 In this study, the risk factors and psychiatric comorbidities of Internet and smartphone addiction were possibly associated with decreased ACC function due to increased GABA levels (Table 4 and Fig 4).

Glutamate was shown to enhance the ACC functions in emotion processing and personal traits in previous studies.13-15 The ACC glutamate concentration was positively correlated with impulsivity.14 Conversely, reduced ACC glutamate concentration

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**FIG 4.** The correlations between neurotransmitters and clinical scales of Internet addiction and psychological tests. A, The gm-GABA/Cr is significantly correlated with the Internet Addiction Test (circle and dashed line) and the Smartphone Addiction Scale (x and dotted line) as well as the sum of both scales (square and solid line). B, The gm-GABA/Cr was significantly correlated with the depression scale (circle and dashed line), total anxiety score (x and dotted line), and state anxiety score (square and solid line). C, The gm-Glx/Cr was correlated with insomnia (circle and dashed line) and sleep quality scores (square and solid line). IAT indicates Internet Addiction Test; SAS, Smartphone Addiction Scale; HRSD, Hamilton Rating Scale for Depression; STAI-S, State-Trait Anxiety Inventory to measure state component of anxiety; PSQI, Pittsburgh Sleep Quality Assessment.
increased the risk of exposure to noxious stimulation, concerns regarding potential problems, and the fear associated with uncertainty.13,15 The negative correlations of Glx with insomnia severity and sleep quality in this study may be explained by decreased emotional ACC functions. In this study, the change in GABA levels after cognitive behavioral treatment showed that the GABA change was not the structural change of GABAergic interneurons but a functional change of GABAergic inhibition.29 The neurotransmitters can be reversed and normalized with improvement regarding Internet and smartphone addiction and comorbidities.

This study had several limitations. First is the reproducibility issue of GABA MR spectroscopy. GABA levels can be affected by age, sex, and circadian rhythm. In a previous study, the GABA level was lower in older subjects on the basis of the age-related gray matter loss; however, the tissue-corrected GABA analysis could correct the age effect.28,39 Although the neurotransmitters in youth have been changed by the developing brain volume before and after puberty, the change of neurotransmitters in healthy youth has not yet been reported. Males showed higher specific GABA receptor subunits in tissue analysis,40 and GABA levels were increased at the time of ovulation in females.41 However, in this study, tissue-corrected GABA analysis and exact sex- and age-matched case and control groups were used to reduce the biologic variability. We tried to improve the technical reproducibility of MR spectroscopy as much as possible with the use of large voxel volume and careful positioning in the ACC, as well as preparing very low first- and second-order shimming values.

Another limitation was the relatively small sample size of the study population, particularly in cognitive behavioral therapy. A future study with a larger sample size could clarify the roles of neurotransmitters in subjects with Internet and smartphone addiction. Last, the MR spectroscopy analysis used did not separate the glutamate peak from the glutamine peak. MEGA-PRESS MR spectroscopy with a 1.9-ppm editing pulse can separate the Glx including glutamate in the ACC may contribute to understanding the biochemical and molecular basis of the Internet and smartphone addiction and the associated comorbidities that could be used to devise appropriate treatments.

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REFERENCES


