INTRODUCTION

An association between depression and altered immunity has been suggested by many research groups (1, 2), although immunological findings have not been consistently demonstrated. Immune changes associated with depression include decreased mitogen-induced lymphocyte proliferation and natural killer (NK) cell activity, and enumerative changes like increases in the total number of white blood cells and neutrophils, and a reduction in the number of lymphocytes (2, 3). Recently, depression is also thought to be associated with activation of some aspects of cellular immunity increasing secretion of proinflammatory cytokines such as interleukin-1, interleukin-6, and tumor necrosis factor-alpha that can elicit depressive or anxiety symptoms, as contrasted with the initial opinion that depression can compromise immune system (4, 5).

Patients experiencing a major depressive episode along with a comorbid panic disorder have a greater number of T cells, and more mitogen-induced lymphocyte proliferation than depressed patients without panic disorder (6). Panic disorder comorbidity seems to contribute to immune variances in patients with major depressive disorder, and it seems reasonable to postulate that some immunological alterations may be attributed to panic disorder. However, relatively few studies have been carried out to observe immune changes in panic disorder, and these studies do not reveal any consistent immunological differences for panic disorder patients versus normal subjects. Marazziti et al. (7) showed that panic disorder patients do not differ from healthy controls in immune cell number, except for T helper (Th) cells, which were significantly lowered. However, many other studies reported no differences in the numbers and proportions of immunocytes in panic disorder without concurrent depression (8-10). In addition, some studies have indicated that B-cell numbers are altered in panic disorder patients, although not all in the same direction (10-12).

We postulated that panic disorder patients have altered immunity versus normal healthy subjects, and thus we examined differences in lymphocyte subset counts between panic disorder patients and control subjects. We also tried to identify the relationship between clinical variables including mood states, and lymphocyte subsets in panic disorder patients.

MATERIALS AND METHODS

All subjects were medically healthy ones, and they included 20 panic disorder patients and 20 age- and gender-matched normal control subjects (10 males and 10 females in each group). The panic disorder patients who had other major psychiatric disorders including other anxiety disorders, major depressive disorder, and substance abuse were excluded from this study. The patients were new outpatients who had visited the Samsung Medical Center in Seoul, and all met the DSM-IV criteria for panic disorder using the Anxiety Disorder Interview Schedule for DSM-IV (13). Patients were
not receiving any regular anti-panic medication at the time of study, although some of the patients took alprazolam intermittently at the time of study, but had never received any treatment for panic disorder previously. They had a relatively recent onset of illness (38.5±50.3 weeks), and 17 among 20 panic patients showed less than 1 yr of illness duration.

Physical examinations were performed with comprehensive history taking and some clinical laboratory tests (EKG, and liver function and thyroid function testing) so as to exclude panic symptoms secondary to other medical conditions. Healthy age- and gender-matched controls were recruited through an internet advertisement in Seoul. All subjects were required to be in good health, and this was defined as having no acute or chronic medical or psychiatric illness that could affect immunity (2, 14-17). Control subjects did not take any medication known to affect the immune system.

The Institutional Review Board of Samsung Medical Center approved the study protocol, and a written informed consent was obtained from all subjects after the procedures had been fully explained. Background data were collected, including demographic data and medical and psychiatric histories. Body Mass Index (BMI), which might affect immune function, was calculated as weight in kilograms divided by height in meters squared; BMIs ranged from 18 to 30 among all 40 subjects. Subjects were instructed to fast for 12 hr prior to the study and to refrain from caffeinated beverages, alcohol, and smoking for 24 hr prior to the study. We began the study at 8 a.m., and upon arrival, a 20-gauge catheter was inserted into a forearm vein. After a physical examination and blood pressure monitoring, subjects were quietly rested for 30 min. At the end of this period, blood samples were collected. Four milliliters of blood was drawn initially and discarded and then at least 3 milliliters was drawn into an EDTA tube. The total white blood cell count and the differential count were obtained for panic disorder patients showed a significant positive correlation with T cell number (r=0.498, p=0.026), and Th cell number (r=0.444, p=0.050), but HAMA scores correlated negatively with NK cell proportion (r=-0.529, p=0.016). STAIS scores for panic disorder patients showed a significant positive cor-

**RESULTS**

The characteristics of panic disorder patients and control subjects are shown in Table 1. No significant differences were evident with respect to age, sex, or BMI between the two groups. Panic disorder patients had significantly higher HAMA, HAMD, STAIS and STAIT scores than control subjects.

The patients with panic disorder had a lower T suppressor (Ts) cell proportion (t=-2.63, p=0.012) and a higher Th/Ts ratio (t=-2.57, p=0.014), than controls. Total WBC number (t=-1.85, p=0.072) was slightly higher in panic disorder patients, but this did not reach statistical significance (Table 2).

Ten panic patients with a HAMD score of >17 had a lower BMI (p=0.051) than 10 panic disorder patients with a HAMD score of <17. More depressive panic patients had lower numbers (p=0.036) and proportions (p=0.013) of NK cell, but showed no differences with respect to other cumulative immune variables versus less depressive panic disorder patients.

Age, gender, and BMI showed no association with immune variables. For panic disorder patients, HAMA scores were positively correlated with T cell number (r=0.498, p=0.025), T cell proportion (r=0.495, p=0.026), and Th cell number (r=0.444, p=0.050), but HAMA scores correlated negatively with NK cell proportion (r=-0.529, p=0.016). STAIS scores for panic disorder patients showed a significant positive cor-

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**Table 1. Characteristics of panic patients and control subjects**

| Variable | Control subjects | Panic patients | t    | p Value |
|----------|------------------|----------------|------|---------|
|          | n=20 (Male=10)   | n=20 (Male=10) |      |         |
| Mean     | SD               | Mean           | SD   |         |
| Age (yr) | 32.4±4.5         | 34.9±7.0       | -1.37| 0.180   |
| BMI (kg/m²) | 22.1±1.8         | 21.8±3.1       | 0.38 | 0.704   |
| HAMD     | 3.3±2.8          | 6.8±6.3        | -3.36| <0.01*  |
| HAMA     | 4.5±3.0          | 10.1±7.26      | -3.26| <0.01*  |
| STAIS    | 37.3±8.9         | 48.7±9.5       | -3.81| <0.01*  |
| STAIT    | 41.2±11.0        | 51.7±8.3       | -3.31| <0.01*  |

BMI, Body Mass Index; HAMD, Hamilton Depression Rating Scale; HAMA, Hamilton Anxiety Rating Scale; STAIS, State Trait Anxiety Inventory-State form; STAIT, State Trait Anxiety Inventory-Trait form. *: p<0.01.
Lymphocyte Subsets in Panic Disorder

Table 2. Immune measures in panic patients and control subjects

| Variable               | Control subjects | Panic patients | t    | p Value |
|------------------------|------------------|----------------|------|---------|
|                       | N=20 (Male=10)   | N=20 (Male=10) |      |         |
| Mean ± SD              | Mean ± SD        |                |      |         |
| Total WBC              | 5400.0 ± 1245.6  | 6475.0 ± 2284.9| -1.85| 0.072   |
| Lymphocyte No.         | 2017.8 ± 504.0   | 2062.5 ± 544.8 | -0.39| 0.699   |
| Lymphocyte %           | 37.9 ± 7.1       | 33.6 ± 7.5     | 1.88 | 0.068   |
| CD3 T cell No.         | 1444.2 ± 464.2   | 1469.5 ± 436.1 | -0.18| 0.860   |
| CD3 T cell %           | 70.8 ± 7.4       | 70.2 ± 8.5     | 0.22 | 0.824   |
| CD4 Th cell No.        | 778.0 ± 303.2    | 861.8 ± 217.5  | -1.09| 0.283   |
| CD4 Th cell %          | 37.9 ± 8.4       | 42.3 ± 7.8     | -1.74| 0.090   |
| CD8 Ts cell No.        | 626.1 ± 228.0    | 553.5 ± 225.8  | 1.01 | 0.318   |
| CD8 Ts cell %          | 30.8 ± 6.7       | 25.7 ± 5.6     | 2.63 | 0.012   |
| CD4/CD8 ratio          | 1.3 ± 0.5        | 1.8 ± 0.6      | -2.57| 0.014   |
| CD16 NK cell No.       | 274.6 ± 123.6    | 332.6 ± 184.1  | -1.17| 0.250   |
| CD16 NK cell %         | 14.2 ± 6.6       | 16.1 ± 8.0     | -0.81| 0.422   |
| CD19 B cell No.        | 299.0 ± 92.0     | 280.4 ± 102.5  | 0.60 | 0.550   |
| CD19 B cell %          | 15.0 ± 3.2       | 13.7 ± 4.0     | 1.13 | 0.264   |

WBC, White blood cell; Th cell, T helper cell; Ts cell, T suppressor cell; NK cell, Natural killer cell.

* p<0.05.

Table 3. Predictive variables for immune changes in panic patients by stepwise multiple regression analyses

| Outcome factor | Good predictors | β  | t    | p    | R²  |
|----------------|-----------------|----|------|------|-----|
| CD3 No.        | HAMA            | 0.544 | 2.594 | 0.020 | 0.296 |
| CD3 %          | STAI            | 0.533 | 2.517 | 0.023 | 0.284 |
| CD4 No.        | HAMA            | 0.480 | 2.188 | 0.044 | 0.230 |
| CD16 %         | HAMA            | -0.493 | -2.266 | 0.038 | 0.243 |

response, macrophage activation, and the enhanced activity of NK cells and cytotoxic T cells. Thus, a reduction in the Ts cell proportion and an elevation of the Th/Ts ratio may reflect altered immunity, which suggests a possibility of T cell activation in panic disorder. Thus, examining the functional measures of T cell activation will be needed in a future study.

Some panic patients in this study had more depressive features (HAMD score >17), even though they did not meet the major depressive disorder DSM-IV criteria. Indeed, both panic disorder and major depressive disorders are highly comorbid and are probably affected by the same serotonin and norepinephrine neurotransmitter systems. However, patients with panic disorder are known to have immune findings that differ from those with major depression. Moreover, the depressive symptoms of panic disorder patients were not found to affect any enumerative immune variables, except for NK cell proportion in this study. Thus, a depressed mood per se does not seem to play an important role in altered measures of immunity in panic disorder patients.

With regard to stress, a reduction in the number of lymphocytes and a suppression of lymphocyte proliferative response to mitogen were demonstrated by a psychiatric fellowship examination. Speech stress in healthy men was found to result in significant increases in Th cells and NK cells and a reduction in the Th/Ts ratio versus baseline. Gerritsen and colleagues also found that a laboratory induced fear situation (public speaking) produced changes in the experimental subjects’ immune systems, and specifically reduced the numbers of Th cells and T lymphocytes. Heightened stress as well as heritable factors has been suggested to contribute to the onset of panic disorder. Although a panic attack itself can also be a severe stressor, previous findings described the above seem to be at odds with ours, even though we did not directly measure stress levels in this study. In the present study, we tried as much as possible to reduce the effects of stress per se does not play an important role in altered measures of immunity in panic disorder patients.

DISCUSSION

We found that panic disorder patients exhibited no differences with respect to the absolute number of immune cells, except for a reduced proportion of Ts cells and an increased Th/Ts ratio, as compared with normal healthy subjects. Marazziti et al. (7) reported significantly fewer Th cells and reduced Th/Ts ratios in panic disorder patients, and others have found increases (11, 12) or decreases (21) in B cell populations in panic disorder. However, our findings support a lack of a relation between this disorder and the absolute numbers of Th cells (9-12, 21), the B cells (7, 9), and NK cell (9, 10, 12, 21).

Suppressor T cells are known to suppress the action of other immune cells, most notably B-cells and T-cells, and thereby prevent the establishment of an immune response. Helper T cells act on other cells in the immune system to promote various aspects of immune response, including immunoglobulin isotype switching and affinity maturation of antibody

response, macrophage activation, and the enhanced activity of NK cells and cytotoxic T cells. Thus, a reduction in the Ts cell proportion and an elevation of the Th/Ts ratio may reflect altered immunity, which suggests a possibility of T cell activation in panic disorder. Thus, examining the functional measures of T cell activation will be needed in a future study.

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With regard to stress, a reduction in the number of lymphocytes and a suppression of lymphocyte proliferative response to mitogen were demonstrated by a psychiatric fellowship examination. Speech stress in healthy men was found to result in significant increases in Th cells and NK cells and a reduction in the Th/Ts ratio versus baseline (25). Gerritsen and colleagues also found that a laboratory induced fear situation (public speaking) produced changes in the experimental subjects’ immune systems, and specifically reduced the numbers of Th cells and T lymphocytes. Heightened stress as well as heritable factors has been suggested to contribute to the onset of panic disorder (27). Although a panic attack itself can also be a severe stressor, previous findings described the above seem to be at odds with ours, even though we did not directly measure stress levels in this study. In the present study, we tried as much as possible to reduce the effects of stress per se does not play an important role in altered measures of immunity in panic disorder patients.

Multiple regression analyses revealed that anxiety levels in panic disorder patients correlated positively with T cell number and proportion, and Th cell number, but anxiety levels were found to be negatively correlated with NK cell proportion in the present study. Weizman (28) reported that interleukin 3 (IL3) production is negatively correlated with

response, macrophage activation, and the enhanced activity of NK cells and cytotoxic T cells. Thus, a reduction in the Ts cell proportion and an elevation of the Th/Ts ratio may reflect altered immunity, which suggests a possibility of T cell activation in panic disorder. Thus, examining the functional measures of T cell activation will be needed in a future study.

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severity of anxiety, and that IL3 might be sensitive to the presence of anxiety. Hence, our findings also support that anxiety levels in panic disorder patients could affect immunity, although the mechanism and meaning of altered immunity in panic disorder remain unknown.

Several limitations of our study should be noted. First, the sample size in our study was relatively small, although there were very few immune studies with a large sample size in panic disorder. Second, there is a possibility that panic disorder patients with severe symptoms might have been excluded because the patients in this study had a relatively recent illness onset and did not take any regular medication at the time of study commencement. Third, our study was limited to enumerative immune measures, although recent immunological studies have included other functional immune measures, such as mitogen induced lymphocyte proliferation and NK cell cytotoxicity, as well as enumerative measures. In addition, the result of increased Th/Ts ratio in this study differed from that of our recent study (29), which showed no significant difference in Th/Ts ratio between panic disorder patients and normal control subjects. It is difficult to explain why this difference occurred, but one possible explanation would be that panic patients with different anxiety levels were involved in each study. Thus, in the future, a prospective study with a larger sample size using functional immune measures, as well as enumerative ones, will be needed to confirm the contradictory immune finding in panic disorder.

In conclusion, panic disorder patients showed a reduced Ts cell proportion and an increased Th/Ts ratio. Moreover, higher levels of anxiety were found to be related to an altered T cell population in panic disorder patients. Quantitative immune differences may reflect altered immunity in panic disorder.

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