A Call for a Rational Polypharmacy Policy: International Insights From Psychiatrists

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Objective Recently, rational polypharmacy approaches have been proposed, regardless of the lower risk and cost of monotherapy. Considering monotherapy as first-line treatment and polypharmacy as rational treatment, a balanced attitude toward polypharmacy is recommended. However, the high prevalence of polypharmacy led the Japanese government to establish a polypharmacy reduction policy. Based on this, the association between the policy and psychiatrists’ attitude toward polypharmacy has been under debate.

Methods We developed an original questionnaire about Psychiatrists’ attitudes toward polypharmacy (PAP). We compared the PAP scores with the treatment decision-making in clinical case vignettes. Multiple regression analyses were performed to quantify associations of explanatory variables including policy factors and PAP scores. The anonymous questionnaires were administered to psychiatrists worldwide.
INTRODUCTION

In the field of psychiatry, polypharmacy refers to the prescription of multiple psychotropic drugs to one patient. Polypharmacy is of concern because it can increase the risk of adverse effects and mortality rate and reduce medication adherence and patients’ quality of life.\(^1\) In addition, polypharmacy imposes a large economic burden. Although most practice guidelines recommend monotherapy as a first-line treatment, polypharmacy has become common practice. The widespread implementation of polypharmacy prompted collaborative research to evaluate the prescription patterns of psychotropic drugs in Asia (Research on Asian Psychotropic Prescription Pattern, REAP) in 2001.\(^2\) The study revealed a high prevalence of polypharmacy in Japan. For example, \(>50\%\) of patients with schizophrenia used \(\geq 2\) psychotropic drugs.\(^3\) This high prevalence of polypharmacy led the Japanese government to establish a polypharmacy reduction policy, and polypharmacy limitations have been adopted in a stepwise manner since the medical fee revision in 2012.\(^4\) Initially, the reduction of the national fee schedule was based on prescriptions of \(\geq 3\) anxiolytics or \(\geq 3\) hypnotics, and the list of prescriptions which apply to this reduction policy has been updated and extended every two years until 2018. Since the introduction of the polypharmacy reduction policy, polypharmacy is less frequently observed.\(^4\)

Considering the decreased polypharmacy rate in Japan, some psychiatrists doubt whether the decrease in polypharmacy rate improved the clinical practice of psychiatry, especially when “rational polypharmacy” is taken into consideration. “Rational polypharmacy” has recently been proposed by several reports, regardless of the low risk and lower cost of monotherapy.\(^5\) One example of “rational polypharmacy” pertains to cross-titration: when switching from one medication to another, the overlap of the two medications is inevitable during a certain period. Another example of “rational polypharmacy” arises when the benefits of polypharmacy are supported by evidence. For example, in a nationwide cohort study in Finland that collected clinical data from \(>62,000\) patients with schizophrenia, within-individual analyses minimizing selection bias were conducted.\(^4\) The analyses revealed that using clozapine plus aripiprazole combination therapy yielded the lowest risk of psychiatric readmission and was associated with a 14%–23% lower risk of rehospitalization relative to the use of clozapine monotherapy. The specific polypharmacy patterns supported by this report would be considered “rational polypharmacy.”

Even when a specific polypharmacy pattern is not supported by previous evidence, polypharmacy can be regarded as rational. In terms of clinical practice, psychiatrists tend to choose the most appropriate treatment strategy for each patient based on their individual circumstances, not big data. When evidence-based monotherapies with optimal dosages and enough duration are ineffective, non-evidence-based polypharmacy prescription can be attempted as a last resort. In this case, careful studies on individual patients (i.e., \(n=1\)) should be performed. Preskorn and Lacey\(^6\) proposed criteria for rational polypharmacy in psychiatry and guidelines for using a combination of medication based on an \(n=1\) trial in clinical practice. If the criteria and guidelines are followed, administering polypharmacy to the patient would be considered rational. In this way, polypharmacy in psychiatry is rational in some settings. However, in the context of “rational polypharmacy,” the negative attitudes of psychiatrists toward polypharmacy might interfere with selecting the best treatment for an individual patient. Considering both monotherapies as a first-line treatment and polypharmacy as a rational treatment, a balanced attitude toward polypharmacy should be recommended.

Aims of the study

To test the hypothesis that public policies, such as laws, rules, and regulations could distort the balanced attitude toward polypharmacy among psychiatrists, we first developed an original questionnaire to evaluate Psychiatrists’ attitudes toward polypharmacy (PAP). Then, the correlation between the results of the original questionnaire and treatment decision-making toward polypharmacy was assessed using the case vignette-based questionnaire to evaluate Psychiatrists’ attitudes toward polypharmacy (PAP). The analyses revealed that using clozapine plus aripiprazole combination therapy yielded the lowest risk of psychiatric readmission and was associated with a 14%–23% lower risk of rehospitalization relative to the use of clozapine monotherapy. The specific polypharmacy patterns supported by this report would be considered “rational polypharmacy.”

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ed to be associated with polypharmacy across countries. To the best of our knowledge, this is the first international survey about both psychiatrists’ attitudes toward polypharmacy and treatment decision-making using case vignettes.

METHODS

In this study, we created a self-report questionnaire for psychiatrists practicing in different countries. The contents of the questionnaire are described in the following paragraph. The survey was developed in Japanese before being translated into English using forward-backward translation. The Japanese or English questionnaire was distributed both online and offline using the psychiatrists’ networks, such as the consortium of REAP, Early Career Section of the World Psychiatric Association, and the Japan Young Psychiatrists Organization. All participants were requested to complete the questionnaire within the survey period between June 1, 2018 and October 31, 2019.

Questionnaire

The questionnaire was composed of the following three sections: a PAP questionnaire, a case vignette-based questionnaire, and a questionnaire covering factors related to polypharmacy. The details of each part are described below.

Psychiatrists’ attitudes toward polypharmacy questionnaire

The questionnaire items were developed based on published research on polypharmacy and expert opinion from several psychiatrists who have been engaged in clinical research on polypharmacy. The questionnaire asking “What do you think about polypharmacy?” was composed of the following four items: 1) depending on the situation, it can be an important treatment option; 2) absolutely needs to be avoided; 3) inevitable in severe cases; and 4) are you careful to avoid polypharmacy in your usual clinical practice? The responses were recorded on a 5-point Likert scale, where a score of 1 indicated “do not agree at all” and a score of 5 indicated “fully agree.” The sum of the scores was obtained by adding the scores of each item after reverse scoring for items 2 and 4. A higher score indicated a more positive attitude of the psychiatrist toward polypharmacy.

Case vignette

“Typical” anonymized cases of patients with schizophrenia were described by professional psychiatrists with extensive experience and used to draft a fictitious case vignette. After confirmation and modification by mental health professionals, including psychiatrists and psychologists, a case vignette questionnaire based on two patients with schizophrenia was developed in Japanese. The Japanese case vignette was translated into English using forward-backward translation (Table 1). Using the case vignette, treatment decision-making, including polypharmacy and non-polypharmacy (e.g., electroconvulsive therapy), were evaluated.

Questionnaire covering factors related to polypharmacy

Ghaemi suggested that five factors were associated with the rise of polypharmacy: scientific, clinical, economic, political, and cultural factors. Based on these five factors, we designed our questions in this fashion:

1) With respect to the scientific factor, clinical evidence on psychotropic medications is known to contribute to polypharmacy. Thus, we added a question on the personal importance of prescription guidelines on psychotropic drugs and recorded responses on a 6-point Likert scale with 1=“no use,” 2=“useful, but not important,” and 6=“useful, very important.”

2) As for the clinical factor, the association between standardized diagnostic criteria and polypharmacy has been suggested as the criteria sometimes result in overlapping diagnoses, which can lead to polypharmacy. In our study, we asked to what extent responders were committed or devoted to the diagnosis of a psychiatric disease using the ICD-10 or DSM-5 and recorded the responses on a 6-point Likert scale with 1=“small extent” and 6=“large extent.”

3) Considering our study’s hypothesis, the policy factor indicating “public policy,” such as laws, rules, and regulations, was evaluated by a question asking about the importance of a limitation on the use of psychotropic drugs using a 6-point Likert scale: 1=“there is no limitation,” 2=“there is a limitation, but it is not important,” and 6=“there is a limitation, and it is very important.”

4) The economic factor was assessed by asking a question about the extent of prescription of psychotropic drugs covered financially by the government or national insurance coverage using a 3-point Likert scale to record responses with 1=“many restrictions,” 2=“a few exceptions,” and 3=“100% covered.”

5) The cultural factor related to the hypothesis that a preference for pharmacological treatments tends to differ across countries. Thus, a question on nationality was added.

Statistical analyses

Normality tests (skewness, kurtosis, and boxplots) were performed to determine if the PAP scores had a normal distribution. When the PAP scores in each country were not normally distributed, the Kruskal-Wallis H test and post hoc Dunn’s pairwise comparison with Bonferroni correction were used to evaluate differences in PAP scores between major countries. Correlations between the PAP scores and treatment decision-making in the case vignette were evaluated using Spearman’s correla-
Table 1. Case vignettes

Case A: 19-year-old male university student
Mr. A is a 19-year-old male university student with a quiet character. He has just entered a university this spring and started to live on his own leaving his parents’ home in a nearby neighborhood. He had difficulty adapting to his new life and developing new friendships. He started missing school. Six months after admission, he stopped going to school and withdrew into his room. Gradually, Mr. A became convinced that he was being watched from the outside and began to feel that he was being monitored at all times and criticized in a scary voice by someone, making it impossible for him to go outside. His concerned family members visited him, but were told, “Why would you come here?” “Did you wiretap me?” by Mr. A with hostile and suspicious eyes. They struggled to persuade him and took him to your hospital, where Mr. A kept murmuring with fear and was diagnosed with schizophrenia. Risperidone was prescribed (up to 6 mg/day) for one month after the admission, and he no longer felt like he was being monitored, which enabled him to go outside. He was discharged from your hospital two-months after the admission. He resumed living on his own and returned to the university after two-months of recuperation at his parents’ home. However, when he actually returned to school, he felt attacked with words, such as “He is weird” by other students. He withdrew into his room again and started sitting around all day long. Although his doctor additionally prescribed aripiprazole (24 mg/day), it seemed not to be sufficiently effective. He remains to be withdrawn, but is desperate to return to school at any cost. His family members also hope for a fast road to recovery.

Case B: 46-year-old male office worker
Mr. B is a 46-year-old man and worked as an office worker after graduating from university. When he was 21 years old, he experienced auditory hallucinations and persecutory delusions. He was diagnosed with schizophrenia and underwent therapeutic treatments in an outpatient department. His psychotic symptoms were ameliorated by treatment with risperidone (4 mg/day). However, he gradually stopped going to the hospital and finally stopped treatment 3 years ago when he was 43 years old. One day, he suddenly started screaming meaninglessly near an elementary school. A neighbor found him coming at a student with a hammer and reported him to the police. Then, Mr. B was admitted to your hospital. After admission, Mr. B tirelessly wandered in his cell severely agitated while mumbling incoherently. His speaking and behavior were disorganized. Although olanzapine (20 mg/day) was prescribed for him, he remained excited all day and was severely aggressive toward others all night. Therefore, risperidone (4 mg/day) was added for agitation, and chlorpromazine (50 mg/day) was added for insomnia. After the addition of drugs, he calmed down and his behavior and speech were gradually organized.

Our study (Table 2). The six countries with the highest number of participants were Japan, Korea, Nepal, Iran, Thailand, and India (n=65, 54, 49, 40, 33, and 19, respectively). The average age of participants was 36.7 years (standard deviation [SD]=8.6), and the average amount of clinical experience was 8.5 years (SD=7.8). Sixty-one percent of participants were male (n=212). The average PAP score was 12.6 (SD=3.0).

Psychiatrists’ attitudes toward polypharmacy scores and treatment decision-making toward polypharmacy across countries
The PAP scores and treatment decision-making toward polypharmacy in case vignettes are plotted across countries in Figures 1 and 2. Although a normal distribution was confirmed for the overall PAP scores, a normal distribution was not confirmed for the PAP scores in each country. A significant difference in PAP scores between major countries was revealed by the Kruskal-Wallis H test ($\chi^2$-value=71, degrees of freedom=6, and p=0.0001). The post hoc Dunn’s pairwise comparison with Bonferroni correction revealed significant differences between Korea and all other countries, as well as between Iran and the other major countries except for Thailand (Table 3).
Factors related to polypharmacy

| Clinical factor                        | Total (N=347) | Japan (N=65) | Korea (N=54) | Nepal (N=49) | Iran (N=40) | Thailand (N=33) | India (N=19) | Others (N=87) |
|----------------------------------------|---------------|--------------|--------------|--------------|-------------|-----------------|--------------|---------------|
| Age, years                             | 36.7±8.6      | 37.3±9.4     | 34.3±5.9     | 32.3±4.9     | 37.7±6.2    | 32.2±3.2        | 34.3±5       | 41.7±10.8     |
| Male, N (%)                            | 212 (61)      | 53 (82)      | 44 (81)      | 32 (65)      | 12 (18)     | 6 (18)          | 14 (74)      | 51 (59)       |
| Years of clinical experience           | 8.5±7.8       | 8.8±8.9      | 6.5±5.7      | 4.9±4.0      | 8.0±6.0     | 5.8±2.7         | 7.3±4.3      | 12.7±9.8      |
| PAP score (a total sum score after reverse scoring of *)| 12.6±3.0      | 11.7±2.7     | 15.3±2.1     | 11.3±2.9     | 13.4±2.6    | 12.5±2.6        | 10.6±2.7     | 12.4±3.0      |
| Depending on the situation, it can be an important treatment option (not at all=1, extremely=5) | 3.5±1.2       | 3.2±1.1      | 4.4±0.6      | 2.9±1.1      | 3.6±1.0     | 3.5±1.1         | 3.1±1.4      | 3.6±1.1       |
| Absolutely needs to be avoided (not at all=1, extremely=5)* | 2.5±1.2       | 2.6±1.1      | 1.7±0.7      | 3.1±2.2      | 2.1±1.1     | 2.8±1.0         | 2.9±1.4      | 2.7±1.3       |
| Inevitable in severe cases (not at all=1, extremely=5) | 3.7±1.0       | 3.4±1.1      | 4.1±0.9      | 3.7±1.0      | 3.9±0.7     | 3.8±0.9         | 2.8±1.3      | 3.7±1.0       |
| Are you careful to avoid polypharmacy in your usual clinical practice? (not at all=1, extremely=5)* | 4±1.0         | 4.3±0.7      | 3.2±0.9      | 4.2±1.0      | 4±0.9       | 4.1±0.8         | 4.5±1.0      | 4.2±1.0       |

Factors related to polypharmacy

| Policy factor          | 2.9±2.1       | 4.8±1.0      | 2.4±1.9      | 1.9±1.8      | 1.4±1.3     | 4.5±1.6         | 2.2±2.1      | 2.6±2.0       |
| Scientific factor      | 4.1±1.9       | 3.4±2.0      | 3.1±1.8      | 5.1±1.3      | 2.5±2.1     | 4.3±1.5         | 5.3±1.2      | 4.9±1.4       |
| Economic factor        | 1.8±0.6       | 2.0±0.3      | 1.6±0.5      | 1.5±0.6      | 2.0±0.6     | 1.5±0.6         | 1.2±0.4      | 1.9±0.7       |
| Clinical factor        | 5.1±1.0       | 4.7±0.9      | 4.7±1.2      | 5.5±0.7      | 4.7±1.3     | 5.5±0.7         | 5.5±1.0      | 5.2±1.1       |

PAP: Psychiatrists’ attitudes toward polypharmacy

**Correlation between Psychiatrists’ attitudes toward polypharmacy scores and treatment decision-making in the case vignettes**

The PAP scores reflected the positive decision-making toward polypharmacy in the case vignettes; item A-4 (How likely are you to add a new antipsychotic drug to his present medication?), item B-1 (How likely are you to agree with the additional risperidone at 4 mg/day?), and item B-3 (How likely are you to think that this three-type-combined medication is too much?) were correlated with the PAP scores (Spearman rho = 0.28, 0.21, and -0.29; p<0.0001, p=0.00001, and p<0.0001; respectively). The other treatment decision-making parameters (e.g., electroconvulsive therapy, dosage adjustment, or replacement of already taken psychotics) were not correlated with the PAP scores (Table 4). The PAP scores and treatment decision-making toward polypharmacy in the case vignettes are plotted in Figure 3.

**Results of multiple linear regression analyses on predicting the Psychiatrists’ attitudes toward polypharmacy scores**

The results of the multiple regression analyses performed to predict the PAP scores are presented in Table 5. A significant association was found (F[10, 277]=8.88, p<0.001), with an R2 of 0.243. The policy factor was significantly negatively predicted by the PAP scores (β=-0.20, p=0.004). Regarding the cultural factor among the six major countries, culture in Korea was associated with higher PAP scores (β=-0.17, p=0.006), whereas culture in India and Nepal was associated with lower scores (β=-0.15, p=0.01, and β=-0.17, p=0.006, respectively). When a dummy variable for sex (male=1, female=2) and the years of clinical experience were also introduced as independent variables, similar results were obtained (Table 6).

**DISCUSSION**

We developed an original questionnaire to evaluate attitudes toward polypharmacy among psychiatrists. The PAP scores re-
flected decision-making toward polypharmacy in clinical case vignettes. The PAP scores differed among psychiatrists from different countries: in Korea, high PAP scores were found among psychiatrists. Beyond these differences between countries (cultural factor), the policy factor also predicted PAP scores.

In our study, a policy factor was significantly associated with relatively lower PAP scores. This association between the limitation on the use of psychotropic drugs and lower PAP scores is compatible with the hypothesis that the Japanese polypharmacy reduction policy has influenced psychiatrists’ attitudes and reduced the prevalence of polypharmacy. To the best of our knowledge, this is the first report indicating the effect of policy on PAP, although the effects of policy on attitudes toward mental disorders and smoking behavior have been reported previously.

In the context of the effects of a policy on psychiatrists’ attitudes and decision-making toward polypharmacy, the content of the policy is important. In Japan, polypharmacy limitations have been placed as follows: 1) in 2012, a reduction of the medical fee reimbursement was introduced for prescriptions of three or more anxiolytics or three or more hypnotics; 2) in 2014, the prescription of four or more antidepressants or four or more antipsychotics was added as a condition for this reduction; 3) in 2016, the rule was strengthened, and prescription of three or more anxiolytics, hypnotics, antidepressants, or antipsychotics was targeted for reimbursement reduction; and 4) in 2018, the prescription of four or more anxiolytics and hypnotics was added as a condition to this reduction scheme. Evidently, this Japanese policy has focused on the mere number of medications, lacking consideration for medical conditions or the severity of mental disorders. This sweeping policy can hamper the rational polypharmacy that is required for specific settings. Furthermore, it can lead to psychiatrists refusing the medical examination of patients with severe mental disorders because they

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Table 3. Comparison of Psychiatrists’ attitudes toward polypharmacy scores in six major countries

|               | Japan | Korea | Nepal | Iran | Thailand | India |
|---------------|-------|-------|-------|------|----------|-------|
| Korea         | -6.6436*** | 0.0000 |       |       |          |       |
| Nepal         | 0.7456  | 6.9160*** | 1.0000 | 0.0000 |          |       |
| Iran          | -2.8719* | 3.0255* | -3.3657** | 0.0428 | 0.0261 | 0.0080 |
| Thailand      | -1.2879 | 4.3035*** | -1.8489 | 1.3077 | 0.0002  | 0.6770 | 1.0000 |
| India         | 1.1698  | 5.6492*** | 0.6187  | 3.1385* | 2.0028  |       |
| Others        | -1.3643 | 5.8188*** | -2.0454 | 1.8835 | 0.2615  | -2.0668 | 1.0000 | 0.4069 |

Significance levels of p<0.05, p<0.01, and p<0.001 are indicated by *, **, and *** respectively. Dunn’s pairwise comparison with Bonferroni correction.
Religious beliefs possibly influence attitudes toward polypharmacy, as indicated in our study. The time trend of polypharmacy rates also indicated that the prevalence rate of polypharmacy is continuously increasing only in Japan across the four time points examined in our previous study. This sequential increase in polypharmacy would be related to the positive attitudes toward polypharmacy among Korean psychiatrists. Considering the similarity of PAP scores to the time trend of polypharmacy rate, the PAP scores can be used as a marker to predict the time course of polypharmacy prevalence, although further studies are required.

In our study, PAP were assessed, although we focused on the prevalence rate of polypharmacy in collaborative research on the prescription patterns of psychotropic drugs in Asia (REAP). Compared to the prevalence rate, it would be easier to interpret the results about psychiatrists’ attitudes across countries as the polypharmacy rate is more susceptible to local factors, such as drug approval, over-the-counter drugs (non-prescription drugs), and the barriers to psychiatric treatments. Regarding drug approval, for example, in Japan, brexpiprazole is approved at 2 mg/day during the maintenance period for patients with schizophrenia, whereas 2–4 mg is approved by the United States Food and Drug Administration. This difference in the approved drug dosage may lead to polypharmacy. For a patient whose appropriate dosage of brexpiprazole is 4 mg, 2 mg brexpiprazole is not effective; thus, more than one type of psychotic medication needs to be prescribed in Japan. Regarding over-the-counter drugs, the types and dosages of non-prescription drugs differ among countries. Drugs without prescriptions are not usually included in the prescription survey, which makes it difficult to compare the prevalence of polypharmacy across countries. Furthermore, barriers to psychiatric treatments can delay seeking care and treatment, potentially leading to the deterioration of mental conditions. If only severe cases are treated because of barriers in a country, the prevalence of polypharmacy will increase. In this way, the prevalence rate of polyphar-
macy in psychiatry is influenced by both local medical factors and PAP.

PAP have been assessed in several reports, although these are much fewer than studies on polypharmacy rates. Previous surveys on psychiatrists’ attitudes were conducted in America, Japan, and Nigeria and have shown similarities in the PAP among both high and low polypharmacy prescribers.15-17 Contrary to these previous results, our results indicated a significant association between attitude toward polypharmacy and treatment decision-making toward polypharmacy using case vignettes. This discrepancy between our results and previous findings could be attributed to differences in evaluating the following two aspects: 1) attitudes toward polypharmacy and 2) treatment decision-making toward polypharmacy. In previous studies, psychiatrists’ attitudes were evaluated based on the levels of concerns toward polypharmacy and the reasons for justifying or being against polypharmacy, whereas treatment decision-making toward polypharmacy was evaluated by the self-reported percentage of patients on polypharmacy. In our present study, psychiatrists’ attitudes were evaluated using original questions based on expert opinion, which does not target justification or concerns about polypharmacy because we thought that common knowledge about the advantages and disadvantages of polypharmacy could distort the answers focusing on concerns and justification. In addition, in our study, treatment decision-making toward polypharmacy was evaluated using clinical case

Figure 3. Psychiatrists’ attitudes toward polypharmacy (PAP) scores and treatment decision-making toward polypharmacy. PAP scores and treatment decision-making toward polypharmacy in case vignettes are plotted.
vignettes, which are useful tools for measuring health professionals’ attitudes.18

This study has several limitations. First, there may be an inconsistency between the real attitudes and attitudes recorded in our survey. PAP were evaluated using self-reported questionnaires, and their scores were validated using the results of the attitudes in response to case vignettes. It is possible that the self-reported answers were distorted by some factors, such as social desirability bias. In addition, the attitudes in response to a case vignette cannot completely and accurately reflect actual attitudes in clinical settings. Second, the study results cannot be generalized to psychiatrists in each country. Due to the process of distributing questionnaires and the voluntariness of participation, participants in our study may have had more interest in polypharmacy than the average psychiatrist in the country (self-selection bias). Finally, only five factors (i.e., scientific, clinical, economic, policy, and cultural factors), already known to be related to polypharmacy, were introduced as one question per factor in our study. Whether other factors, such as socioeconomic status, religion, race, hospital type, and the professional field, have an impact on the attitudes toward polypharmacy cannot be evaluated, although we have discussed the possible impact of religion.

In conclusion, we developed a questionnaire that reflected treatment decision-making toward polypharmacy among psychiatrists. Based on this questionnaire, international comparison and multiple regression analyses were conducted. Our results suggested that psychiatrists’ attitudes may be affected by cultural differences in each country. Beyond these cultural differences, the policy factor significantly influenced psychiatrists’ attitudes. Considering its effect on treatment decision-making, thoughtful policies on polypharmacy considering “rational polypharmacy” should be established. We also discussed the possibility that the PAP scores may predict the future prevalence rate of polypharmacy. If psychiatrists’ attitudes are useful for predicting future prevalence rates, they may be useful for better policymaking. Further research is warranted to assess whether predictions can be made using PAP scores.

### Table 5. Multiple regression analysis results

|                  | Coefficients | Std. err. | t-value | p>|t|  | Beta | VIF |
|------------------|--------------|-----------|---------|------|------|-----|
| (Constant)       | 13.233       | 0.946     | 13.99   | <0.001 |      |     |
| Policy factor    | -0.297       | 0.102     | -2.91   | 0.004 | -0.195 | 1.65 |
| Scientific factor| -0.072       | 0.102     | -0.71   | 0.479 | -0.044 | 1.42 |
| Economic factor  | 0.055        | 0.308     | 0.18    | 0.858 | 0.011 | 1.29 |
| Clinical factor  | 0.066        | 0.169     | 0.39    | 0.699 | 0.021 | 1.12 |
| Cultural factor  | Japan        | -0.221    | 0.573   | -0.39 | 0.700 | -0.028 | 1.93 |
|                  | Korea        | 2.864     | 0.560   | 5.12  | <0.001 | 0.336 | 1.58 |
|                  | Nepal        | -1.699    | 0.608   | -2.79 | 0.006 | -0.173 | 1.41 |
|                  | Iran         | 0.572     | 0.649   | 0.88  | 0.379 | 0.058 | 1.56 |
|                  | Thailand     | 0.550     | 0.627   | 0.88  | 0.381 | 0.055 | 1.46 |
|                  | India        | -2.271    | 0.879   | -2.58 | 0.010 | -0.151 | 1.25 |

VIF, variance inflation factor

### Table 6. Multiple regression analysis results including sex and years of clinical experience

|                  | Coefficients | Std. err. | t-value | p>|t|  | Beta | VIF |
|------------------|--------------|-----------|---------|------|------|-----|
| (Constant)       | 13.592       | 1.207     | 11.26   | <0.001 |      |     |
| Policy factor    | -0.263       | 0.103     | -2.54   | 0.012 | -0.173 | 1.62 |
| Scientific factor| -0.054       | 0.105     | -0.52   | 0.606 | -0.033 | 1.42 |
| Economic factor  | -0.042       | 0.319     | -0.13   | 0.895 | -0.008 | 1.32 |
| Clinical factor  | -0.054       | 0.178     | -0.30   | 0.763 | -0.017 | 1.15 |
| Cultural factor  | Japan        | -0.226    | 0.601   | -0.38 | 0.707 | -0.029 | 2.05 |
|                  | Korea        | 2.856     | 0.612   | 4.66  | <0.001 | 0.338 | 1.84 |
|                  | Nepal        | -1.669    | 0.673   | -2.48 | 0.014 | -0.167 | 1.58 |
|                  | Iran         | 0.711     | 0.701   | 1.02  | 0.311 | 0.069 | 1.61 |
|                  | Thailand     | 0.639     | 0.667   | 0.96  | 0.339 | 0.066 | 1.64 |
|                  | India        | -2.209    | 0.894   | -2.47 | 0.014 | -0.167 | 1.29 |
|                  | Sex          | -0.133    | 0.383   | -0.35 | 0.073 | -0.208 | 1.27 |
|                  | Years of clinical experience | 0.026 | 0.023 | 1.14 | 0.257 | 0.068 | 1.25 |

VIF, variance inflation factor
Availability of Data and Material
The datasets generated or analyzed during the study are available from the corresponding author on reasonable request.

Conflicts of Interest
Seon-Cheol Park, a contributing editor of the Psychiatry Investigation, was not involved in the editorial evaluation or decision to publish this article. All remaining authors have declared no conflicts of interest.

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