Factors Associated with Postoperative Diabetes Insipidus after Pituitary Surgery

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Background: Determining risk factors for diabetes insipidus (DI) after pituitary surgery is important in improving patient care. Our objective is to determine the factors associated with DI after pituitary surgery.

Methods: We reviewed records of patients who underwent pituitary surgery from 2011 to 2015 at Philippine General Hospital. Patients with preoperative DI were excluded. Multiple logistic regression analysis was performed and a predictive model was generated. The discrimination abilities of the predictive model and individual variables were assessed using the receiving operator characteristic curve.

Results: A total of 230 patients were included. The rate of postoperative DI was 27.8%. Percent change in serum Na (odds ratio [OR], 1.39; 95% confidence interval [CI], 1.15 to 1.69); preoperative serum Na (OR, 1.19; 95% CI, 1.02 to 1.40); and performance of craniotomy (OR, 5.48; 95% CI, 1.60 to 18.80) remained significantly associated with an increased incidence of postoperative DI, while percent change in urine specific gravity (USG) (OR, 0.53; 95% CI, 0.33 to 0.87) and meningioma on histopathology (OR, 0.05; 95% CI, 0.04 to 0.70) were significantly associated with a decreased incidence. The predictive model generated has good diagnostic accuracy in predicting postoperative DI with an area under curve of 0.83.

Conclusion: Greater percent change in serum Na, preoperative serum Na, and performance of craniotomy significantly increased the likelihood of postoperative DI while percent change in USG and meningioma on histopathology were significantly associated with a decreased incidence. The predictive model can be used to generate a scoring system in estimating the risk of postoperative DI.

Keywords: Diabetes insipidus; Vasopressins; Neurosurgery; Postoperative complications; Neuroendocrinology

INTRODUCTION

Antidiuretic hormone (ADH) is a nonapeptide produced in the magnocellular neurons of the hypothalamic supraoptic and paraventricular nuclei. It is transported via the hypothalamo-hypophyseal tract, terminating in the neurohypophysis where it matures and is stored for future release [1]. Disturbances in ADH secretion leading to disorders of water balance are among the most common perioperative complications in patients undergoing transsphenoidal surgery [2]. Untreated diabetes insipidus (DI) patients can develop hypernatremia and increased serum osmolality which can lead to dehydration and impaired mental status in severe cases [3].

Usually a complication of the postsurgical course after pitu-
Diabetes Insipidus after Pituitary Surgery

Methods

Study design and setting
This is a retrospective cohort study conducted at the Philippine General Hospital (PGH), a tertiary university hospital and referral center for pituitary cases. This study was conducted after approval by the University of the Philippines Manila Research Ethics Board.

Study population and sampling
All patients who underwent pituitary surgery in PGH from January 2011 to December 2015 were included. Those who had preoperative DI: urine output (UO) (>2.5 mL/kg body weight per hour or at least 4 L per day), urine specific gravity (USG) ≤ 1.005 or urine osmolality < 200 mOsm/kg H2O with or without hypernatremia or hyperosmolality present prior to surgery were excluded as well as those given medications that can potentially affect serum Na levels and UO. Patients with missing charts were also excluded.

Definition of terms
We defined postoperative DI as the presence of documented voluminous UO (>2.5 mL/kg body weight per hour or at least 4 L per day), with a USG of ≤ 1.005 or urine osmolality < 200 mOsm/kg H2O with or without hypernatremia or hyperosmolality or as clinically diagnosed by the attending endocrinologist, developing anytime during hospital admission following pituitary surgery. Other causes of polyuria such as postoperative water diuresis, glycosuria from diabetes and steroid induced hyperglycemia were differentiated from DI through the findings of normal or increased USG (> 1.005) in these cases. If hypotonic polyuria was observed, other potential causes besides DI were first considered.

Data collection
Patients included in the study were identified through the computerized databases of the Section of Endocrinology and records in the Department of Pathology. Data were collected using a standardized data collection form. Study subjects were anonymized and were given code numbers as identifiers. Information gathered included age, sex, size of mass, type of mass (macroadenoma vs. microadenoma), histopathology, functionality of tumor, type of surgery, status of surgery, type of surgeon, and other postoperative complications. Laboratory parameters such as preoperative serum Na and USG and first postoperative serum Na and USG were also collected. At our institution, the first postoperative serum Na and USG are usually collected upon arrival at the neurosurgical critical care unit or at least within 6 hours postoperatively. Percent change in serum Na was computed as the difference between postoperative and preoperative serum Na divided by the preoperative serum Na. Percent change in USG was also computed similarly.

Statistical analysis
Demographic data were presented using descriptive statistics. Quantitative variables were expressed using means and standard deviations for normally distributed data and median and interquartile range (IQR) for non-normally distributed data, while qualitative variables were expressed using frequencies and percentages. Normality of distribution for continuous variables was tested using the Shapiro-Wilks test. Student t test and Wilcoxon rank sum test were used to compare continuous variables between two groups with normal and non-normal distribution, respectively. Chi-square test was used to compare categorical variables. Prior to predictive modeling, tests for collinearity, confounding, and interaction were performed. Multicollinearity
was assumed if the variable inflation factor was >10. Collinear variables that were identified included change in serum Na and USG, and postoperative serum Na and USG; hence, these were excluded in the logistic model. A variable was tested as a possible confounder if the independent variable was related to both the dependent variable (postoperative DI) and the percent change in serum Na. Confounding was considered if the change in odds ratio (OR) in the crude and adjusted logistic model was >10%. Histopathology, functionality, and complications were identified as significant confounders. After checking for interaction, no variables were identified to significantly modify the relationship of postoperative DI and percent change in serum Na. During the univariate logistic regression analysis, variables found to be significant at $P<0.2$ were entered into an exploratory multivariate logistic regression model for postoperative DI. After univariate analysis, status of surgery (initial vs. repeat), sex, size of mass, and type of mass (macroadenoma vs. microadenoma) were excluded from the full model.

Multivariate logistic regression was then performed to examine whether percent change in serum sodium was independently associated with postoperative DI after adjustment for age, surgeon (consultant vs. trainee), percent change in USG, preoperative serum Na, preoperative USG, type of surgery, histopathology, functionality, and surgical complications. Independent predictors of postoperative DI were identified using the hierarchical method of variable selection with multivariate logistic regression analysis. The significant variables ($P<0.05$) after hierarchical stepwise elimination formed the final predictive model: percent change in serum Na, percent change in USG, preoperative serum Na, preoperative USG, type of surgery, and histopathology. The Hosmer-Lemeshow test was used to determine goodness-of-fit of our model. The discrimination abilities (diagnostic accuracy) of the predictive model as well as individual variables—preoperative serum Na and USG, percent change in serum Na and USG, were assessed with the area under the receiver operator characteristic (ROC) curve. All the statistical analyses were performed using STATA 13 (StataCorp, College Station, TX, USA).

**RESULTS**

A total of 242 patient records were initially retrieved. Of these, 10 patients were excluded due to the presence of preoperative DI, while two patients were excluded because they were given mannitol postoperatively which can affect sodium levels and UO. There were 230 patients included in the final analysis, with 124 females (53.9%) and 106 males (46.1%) and a mean age of 44 years. The demographic and clinical data are summarized in Table 1. Majority of the pituitary masses were greater than 1 cm (96.9%) with a mean size of 3.3 cm. The most common histopathology was that of pituitary adenoma (82.2%), followed by meningioma (8.3%) and craniopharyngioma (8.3%). Most of the tumors were non-functioning (82%). Among the functional pituitary tumors, prolactinoma was the most common (8.5%).

Transsphenoidal surgery (both nasal and labial approaches) was performed in 170 patients (73.9%). Majority of the cases underwent pituitary surgery for the first time (88.2%). Two thirds of the cases were performed by a neurosurgery consultant. The most common postoperative complication other than DI was infection (3.5%) followed by postsurgical hemorrhage (2.6%).

In this 5-year period, 64 patients developed postoperative DI with an overall rate of 27.8%. The mean onset of postoperative DI in this study is 27.38 hours after surgery. All laboratory parameters were non-normally distributed and presented as median (IQR) as seen in Table 2. Notably, the absolute and percent change in serum Na and USG (preoperative to 6 hours postoperative) were all statistically greater among patients who developed DI.

In the univariate analysis (Table 3), age (OR, 0.97; 95% confidence interval [CI], 0.95 to 0.99), craniopharyngioma (OR, 6; 95% CI, 2.17 to 16.7), and craniotomy (OR, 2.5; 95% CI, 1.20 to 4.86) were associated with postoperative DI, without adjustment for other factors. All laboratory parameters (except for preoperative and postoperative serum Na) met the level of statistical significance to be included in the exploratory multivariate logistic regression modeling (cut-off $P<0.2$). However, since the preoperative and postoperative serum Na are some of our major variable of interest, we decided to include it in the multivariate logistic regression analysis.

In the final logistic regression model (Table 4), laboratory parameters that were independently associated with postoperative DI include percent change in serum Na (OR, 1.39; 95% CI, 1.15 to 1.69), percent change in USG (OR, 0.53; 95% CI, 0.33 to 0.87), and preoperative serum Na (OR, 1.19; 95% CI, 1.02 to 1.4). The higher the percent change in serum Na and preoperative serum Na, the higher the probability of postoperative DI. Patients with meningioma are less likely to have postoperative DI compared to pituitary adenoma (OR, 0.05; 95% CI, 0.004 to 0.70). On the other hand, patients who underwent craniotomy are five times more likely to develop postoperative DI compared to the transnasal approach. The presence of craniopharyn-
### Table 1. Demographic Data and Clinical Characteristics of Patients

| Characteristic                        | No DI (n=166) | DI (n=64) | Overall | P value |
|---------------------------------------|---------------|-----------|---------|---------|
| Age, yr                               | 46±14.1       | 38.6±16.4 | 44±15.1 | <0.001  |
| Sex                                   |               |           |         | 0.460   |
| Female                                | 92 (55.4)     | 32 (50.0) | 124 (53.9)|
| Male                                  | 74 (44.6)     | 32 (50.0) | 106 (46.1)|
| Mass size, cm                         | 3.3±1.1       | 3.4±1.2   | 3.3±1.1 | 0.135   |
| <1                                    | 6 (4.2)       | 0         | 6 (3.1) |         |
| ≥1                                    | 138 (95.8)    | 52 (100.0)| 190 (96.9)|
| Histopathology                        |               |           |         | 0.003   |
| Pituitary adenoma                     | 139 (84.2)    | 50 (78.1) | 189 (82.2)|
| Meningioma                            | 18 (10.9)     | 1 (1.6)   | 19 (8.3) |         |
| Craniopharyngioma                     | 6 (3.6)       | 13 (20.3) | 19 (8.3) |         |
| Adenocarcinoma                        | 1 (0.6)       | 0         | 1 (0.4) |         |
| Glioma                                | 1 (0.6)       | 0         | 1 (0.4) |         |
| No data                               | 1 (0.6)       | 0         | 1 (0.4) |         |
| Pituitary adenoma functionality       |               |           |         | 0.770   |
| Nonfunctioning                        | 113 (81.3)    | 42 (84.0) | 155 (82.0)|
| TSHoma                                | 1 (0.7)       | 0         | 1 (0.5) |         |
| Prolactinoma                          | 12 (8.6)      | 4 (8.0)   | 16 (8.5) |         |
| GH-secreting                          | 6 (4.3)       | 1 (2.0)   | 7 (3.7) |         |
| Cushing’s disease                     | 5 (3.6)       | 0         | 5 (2.7) |         |
| GH and prolactin secreting            | 0             | 1 (2.0)   | 1 (0.5) |         |
| No data                               | 2 (1.4)       | 2 (4.0)   | 4 (2.1) |         |
| Type of surgery                       |               |           |         | 0.007   |
| Nasal approach                        | 96 (57.8)     | 30 (46.9) | 126 (54.8)|
| Labial approach                       | 36 (21.7)     | 8 (12.5)  | 44 (19.1) |         |
| Craniotomy                            | 32 (19.3)     | 25 (39.1) | 57 (24.8)|         |
| Craniectomy                           | 0             | 1 (1.6)   | 1 (0.4) |         |
| Biopsy                                | 2 (1.2)       | 0         | 2 (0.9) |         |
| Status of surgery                     |               |           |         | 0.130   |
| Initial (primary)                     | 148 (90.2)    | 53 (82.8) | 201 (88.2)|
| Repeat                                | 16 (9.8)      | 11 (17.2) | 27 (11.8)|         |
| Surgeon                               |               |           |         | 0.110   |
| Consultant                            | 117 (70.5)    | 38 (59.4) | 155 (67.4)|
| Resident                              | 49 (29.5)     | 26 (40.6) | 75 (32.6)|         |
| Other postoperative complications     |               |           |         | 0.530   |
| None                                  | 151 (91.0)    | 57 (89.0) | 208 (90.4)|
| Death                                 | 1 (0.6)       | 0         | 1 (0.4) |         |
| Hemorrhage                            | 4 (2.4)       | 2 (3.1)   | 6 (2.6) |         |
| Infection                             | 4 (2.4)       | 4 (6.3)   | 8 (3.5) |         |
| CSF leak                              | 3 (1.8)       | 0         | 3 (1.3) |         |
| Hydrocephalus                         | 2 (1.2)       | 0         | 2 (0.9) |         |
| Pneumocephalus                        | 1 (0.6)       | 1 (1.6)   | 2 (0.9) |         |

Values are expressed as mean±SD or number (%).
DI, diabetes insipidus; TSHoma, thyroid stimulating hormone secreting adenoma; GH, growth hormone; CSF, cerebrospinal fluid.
alignment (OR, 0.5; 95% CI, 0.09 to 2.66) and the use of labial approach (OR, 0.64; 95% CI, 0.20 to 1.99) also seem to have less likelihood of postoperative DI; however, these associations did not reach statistical significance.

Evaluation of goodness of fit indicates a good overall fit of the model with our data (Pearson chi² = 130.3, P = 0.25; Hosmer-Lemeshow chi² = 7.9, P = 0.45). The sensitivity and specificity of the model was at 64% and 89%, respectively. In addition, the positive and negative predictive value was at 78% and 80%, respectively. The predictive model was able to correctly predict postoperative DI in 80% of cases.

**Table 2. Laboratory Characteristics of Patients with and without Postoperative Diabetes Insipidus**

| Characteristic                   | No DI (n=166) | DI (n=64) | P value  | 95% CI            |
|----------------------------------|---------------|-----------|----------|-------------------|
| Preoperative serum Na, mmol/L    | 139.2 (5.0)*  | 139.5 (4.0) | 0.772    | 138.6–139.7       |
| Postoperative serum Na, mmol/L   | 141.0 (4.0)   | 144.9 (6.5) | <0.001   | 141.7–143.0       |
| Preoperative USG                 | 1.015 (0.01)  | 1.015 (0.015) | 0.890    | 1.015–1.017       |
| Postoperative USG                | 1.016 (0.014) | 1.005 (0.01) | <0.001   | 1.013–1.020       |
| Percent change in serum Na, %    | 1.40 (4.32)   | 3.60 (4.70) | <0.001   | 1.84–2.97         |
| Absolute change in serum Na, mmol/L | 2 (6.0)   | 5 (6.5)    | <0.001   | 2.48–4.01         |
| Percent change in USG, %         | 0.2 (1.76)    | –0.5 (1.07) | <0.001   | 0.14–1.79         |
| Absolute change in USG           | 0.002 (0.017) | –0.005 (0.011) | <0.001 | –0.003 to 0.005  |

Values are expressed as median (interquartile range). Median (interquartile range) was used for non-normally distributed data. DI, diabetes insipidus; CI, confidence interval; Na, sodium; USG, urine specific gravity.

**Table 3. Univariate Analysis**

| Variable                           | OR    | 95% CI         | P value |
|------------------------------------|-------|----------------|---------|
| Age, mean                          | 0.97  | 0.95–0.99      | 0.001   |
| Female sex                         | 1.24  | 0.70–2.22      | 0.460   |
| Mass size, cm                      | 1.00  | 0.99–1.00      | 0.867   |
| Mass type, microadenoma            | 1.00  | 0.99–1.01      | 0.291   |
| Histopathology                     |       |                |         |
| Meningioma                         | 0.15  | 0.02–1.19      | 0.073   |
| Craniopharyngioma                  | 6.00  | 2.17–16.70     | 0.001   |
| Pituitary adenoma functionality    |       |                |         |
| Prolactinoma                       | 0.90  | 0.27–2.94      | 0.857   |
| GH-secreting                       | 0.45  | 0.05–3.83      | 0.464   |
| Type of surgery                    |       |                |         |
| Labial approach                    | 0.71  | 0.30–1.70      | 0.442   |
| Craniotomy                         | 2.50  | 1.29–4.86      | 0.007   |
| Status of surgery                  |       |                |         |
| Repeat                             | 0.97  | 0.87–1.08      | 0.555   |
| Surgeon                            | 1.60  | 0.90–2.98      | 0.109   |
| Other postoperative complications  |       |                |         |
| Hemorrhage                         | 1.32  | 0.24–7.43      | 0.749   |
| Infection                          | 2.65  | 0.64–10.9      | 0.178   |
| Pneumocephalus                     | 2.65  | 0.16–43.07     | 0.493   |
| Preoperative serum Na              | 0.99  | 0.99–1.00      | 0.350   |
| Postoperative serum Na             | 0.99  | 0.99–1.00      | 0.350   |
| Preoperative USG                   | 0.99  | 0.99–1.00      | 0.100   |
| Postoperative USG                  | 0.99  | 0.99–0.99      | 0.002   |
| Percent change in serum Na         | 1.22  | 1.11–1.35      | <0.001  |
| Absolute change in serum Na        | 1.17  | 1.09–1.26      | <0.001  |
| Percent change in USG              | 0.65  | 0.48–0.87      | 0.004   |
| Absolute change in USG             | 3.10e 10⁻⁶ | 1.22e ¹⁰⁻⁷–7.92e⁻⁷ | 0.003 |

OR, odds ratio; CI, confidence interval; GH, growth hormone; Na, sodium; USG, urine specific gravity.
*The ‘e’ values in exponential form (1.22×10⁻⁷).*

**Receiver operator characteristic curves**

Fig. 1 shows the ROC curves for the different laboratory parameters that were included in the final model. Individually, the preoperative serum Na (area under curve [AUC], 0.47) and USG (AUC, 0.44) as well as USG (AUC, 0.32) were poor tests in discriminating postoperative DI. The percent change in serum Na (AUC, 0.70) can only fairly discriminate postoperative DI based on the AUC of 0.70 and on the appearance of the ROC curve.

On the other hand, our predictive model incorporating all four laboratory parameters as well as histopathology and type of surgery has good diagnostic accuracy in predicting postoperative DI with an AUC of 0.83 with an ROC curve shown below (Fig. 2).

**DISCUSSION**

This is the first study in the Philippines to investigate factors associated with postoperative DI after pituitary surgery. The DI
incidence in our study is 27.3% higher than the reported 20% by Alinsonorin et al. [8] in 2003. This may be due to the lower number of patients who underwent pituitary surgery reported in the earlier study (87 vs. 230). However, the incidence in our

| Variable                                      | OR    | 95% CI        | P value |
|-----------------------------------------------|-------|---------------|---------|
| Percent change in serum Na                    | 1.39  | 1.15–1.69     | 0.001   |
| Percent change in USG                         | 0.53  | 0.33–0.87     | 0.012   |
| Preoperative serum Na                         | 1.19  | 1.02–1.40     | 0.026   |
| Preoperative USG                              | 2.84e-27 | 4.54e-06–177.68 | 0.071   |
| Histopathology                                |       |               |         |
| Meningioma                                    | 0.05  | 0.004–0.70    | 0.025   |
| Craniopharyngioma                             | 0.50  | 0.09–2.66     | 0.415   |
| Type of surgery                                |       |               |         |
| Labial approach                               | 0.64  | 0.20–1.99     | 0.437   |
| Craniotomy                                    | 5.48  | 1.60–18.80    | 0.007   |

OR, odds ratio; CI, confidence interval; Na, sodium; USG, urine specific gravity.
*The ‘e’ values in exponential form (2.84×10^{-27}).

Fig. 1. Receiver operator characteristic (ROC) curves of various laboratory parameters. (A) Preoperative sodium. (B) Preoperative urine specific gravity. (C) Percent change in sodium. (D) Percent change in urine specific gravity.

Fig. 2. Receiver operator characteristic (ROC) curve of the predictive model.
study is still higher than that reported by Nemergut et al. [9] (18.3%), and Schreckinger et al. [10] (16.3%). The mean time of the onset of postoperative DI in our study was 27.38 hours. While most of the postoperative DI resolved prior to discharge, 18 out of the 64 or 28.13% of those who developed postoperative DI were sent home on desmopressin. However, we cannot ascertain if they progressed to permanent DI since we were not able to follow the patients after discharge.

Unlike previous studies, age, sex, tumor size, and CSF leak were not associated with an increased incidence of postoperative DI in this study. Moreover, in contrast to previous reports [9,12], craniopharyngioma did not appear to be associated with postoperative DI in our study. This may have been due to the relatively small number of craniopharyngioma cases in this present study. Earlier reports found conflicting results with regards to association of mass size with postoperative DI. In some studies, patients with macroadenoma were more likely to have postoperative DI [5,13], whereas other studies found microadenoma to be more associated with DI [6,9]. In our study, tumor size did not correlate with the incidence of postoperative DI. Although microadenoma could have been underrepresented in the study. Similar to the results of Nemergut et al. [9], repeated surgery in this study did not increase the incidence of postoperative DI.

It has been proposed in one study that surgeries done by more experienced surgeons tend to have lower likelihood of developing DI postoperatively [14]; however, our study showed no statistically significant difference with regards to who performed the operation. This can be attributed to a higher volume of surgeries done by experienced surgeons in this study compared to those done by trainees (67% vs. 33%).

After multivariate logistic regression analysis, only percent change in serum Na, preoperative serum Na, and performance of craniotomy were found to be associated with increased risk of postoperative DI, while percent change in USG and meningioma on histopathology were associated with decreased incidence of DI. The reason why meningiomas are associated with a lower risk of postoperative DI in this study is still uncertain. Performance of craniotomy in the study was found to have significantly increased likelihood of developing postoperative DI. This may be due to a larger tumor size and the extent of tissue involvement when craniotomy is done.

While increased UO is a hallmark of the diagnosis of postoperative DI, there are several factors that affect urine volume postoperatively such as postoperative mobilization of fluids and medications. Hence, serum Na and USG are used to differentiate these causes from DI. UO postoperatively is difficult to interpret; thus, we only included more objective measures to determine the presence of DI.

Although the study of Schreckinger et al. [12] found percent change in serum Na to be good predictor of DI with a 96% sensitivity and 50% specificity, our study showed percent change in serum Na to be poor in discriminating postoperative DI based on ROC. Furthermore, preoperative serum Na, preoperative USG, and percent change USG were also not shown to be good predictors of postoperative DI. Hence, specific cut-offs for each variable were not obtained. However, incorporating all these variables into our predictive model showed good diagnostic accuracy in predicting postoperative DI.

The predictive model from our study can be used to generate a scoring system that will aid physicians in estimating point-of-care risk of postoperative DI, which may improve postoperative monitoring and management.

There are limitations related to this study. Although there is a correlation of tumor location with postoperative DI in a previous study [11], our study was not able to analyze tumor location because it was not recorded in most of the patients’ charts. Moreover, a separate analysis for transient and permanent DI was also not done in this study because we were not able to follow patients after discharge. Another limitation of this present study was that other supporting biochemical data in diagnosing DI such as urine osmolality and serum osmolality were not commonly done in our hospital and were not part of the analysis therefore we relied only on the USG, urine volume and serum Na.

In conclusion, our study found that greater percent change in serum Na, preoperative serum Na, and performance of craniotomy significantly increased the likelihood of postoperative DI while percent change in USG and meningioma on histopathology were significantly associated with a decreased incidence of postoperative DI. The predictive model from this study has a good diagnostic accuracy in predicting postoperative DI and can be used to generate a scoring system that will aid physicians in estimating the risk of postoperative DI.

CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported.

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