Cystic intracranial malformations have been reported in animals as a possible cause of seizure, blindness, ataxia, and other neurological signs (6, 7, 16). However, porencephaly could be an incidental finding in asymptomatic dogs. Porencephaly is a rare congenital disorder of the central nervous system characterized by cysts and cavities containing CSF (3, 4, 13), usually connecting the ventricles. Porencephaly in human medicine is considered secondary to environmental factors (toxins, infections), vascular cerebral changes caused by trauma, and hypoxic or congenital defects during the fetal period (16, 17). There are sporadic reports of porencephaly in dogs (3, 8, 13).

Central diabetes insipidus characterized by the inability to concentrate the urine is a well-defined condition in veterinary medicine (18). Etiology of central diabetes insipidus includes obstruction or infiltration of the pituitary gland by tumors, injury, trauma, or infections (2, 18). To the best of the authors’ knowledge, this is the first report in veterinary literature concerning a case of porencephaly complicated by central diabetes insipidus. The aim of this case report was to describe the MRI features of porencephaly lesions and to describe the central diabetes insipidus associated with congenital porencephaly.

A 4-year-old neutered male Terrier dog weighing 3.6 kg was referred to the Veterinary Teaching Hospital of Veterinary Faculty, Ankara University with a history of polyuria and polydipsia for 6 months. Physical observations during hospitalization confirmed polyuria and polydipsia with a water intake of 160 ml/kg/day. The initial blood analyses, including complete blood count and serum profiles, are shown in Table 1. Urinalysis revealed hyposthenuria (urine specific gravity 1.005). Urine osmolality was 245 mOsm/kg (reference interval < 300), and the calculated plasma osmolality was 310 mOsm/kg (reference interval > 300). Urine specific gravity, urine osmolality, and plasma osmolality obtained in a post-water deprivation test were 1.005, 250 mOsm/kg, and 340 mOsm/kg, respectively. No remarkable changes were also observed in abdominal radiography, ultrasonography, or urine culture to differentiate the underlying infectious and metabolic diseases, including adrenal disorders and hypothyroidism. Brain MRI revealed (Fig. 1) right fronto-parietal cystic and cavitary lesions with ex vacuo changes. CSF cytology and culture revealed no remarkable changes. Treatment consisted in applying intranasal desmopressin to the conjunctiva. The dog remains clinically healthy for 2 years. This case report reflects the MRI features of porencephaly and central diabetes insipidus associated with congenital porencephaly.

Keywords: porencephaly, dog, diabetes insipidus
was consistent with central diabetes insipidus. A follow up brain MRI was performed 6 months later, and the lesion was found to be stable. The final diagnosis was porencephaly complicated by central diabetes insipidus. The dog remains clinically healthy 2 years later with desmopressin treatment.

**Discussion**

Porencephaly is a rare cerebral malformation involving cerebral cavities or cysts filled with CSF and connecting the ventricular network and brain surface (7). The condition of focal areas of the cerebrum without residual parenchyma may be described as porencephaly. Neurological signs including seizure have been reported in animals with hydrocephaly, lissencephaly, and porencephaly. However, porencephaly could be an incidental finding in dogs without any symptoms (3, 14). Clinical signs may be absent at birth and may not begin for many years (10). Although epileptic seizure resulting from congenital brain malformations is the most common sign in dogs (15), the case presented here did not show any remarkable clinical signs, including neurological symptoms, for many years. Environmental factors and acquired or perinatal vascular cerebral lesions associated with trauma, viral infections, congenital defects, or inheritance are the possible etiology of porencephaly (5, 12, 17). In this case, no data obtained by anamnesis or diagnostic examination showed a distinct acquired etiology.

**Fig. 1.** MRI images showing cystic and cavitary lesions affecting the dog with porencephaly. A, B – T2 images show a normal size and configuration of the 4th ventricle. No significant signal changes were detected in the brainstem and cerebellum. C, D – No central line deviation is obvious. The cavities are pervaded by thin lines of residual brain tissue. A, B, C, D, E, F – Partial loss of the right-parietal region. Signal changes and cavitations in the right fronto-parietal region connected to the right lateral ventricle, causing enlargement in the lateral ventricle (ex vacuo changes) and the right hemispheric sulcus. Isointense encephalomalacic areas with CSF causing volume loss in all sequences.

**Tab. 1.** Blood analyses in the dog

| CBC                              | Results | References | Serum Profiles          | Results | References |
|----------------------------------|---------|------------|-------------------------|---------|------------|
| Leukocyte (10^9/l)               | 10.32   | 6-17       | Glucose (g/dl)          | 104     | 65-118     |
| Lymphocyte (10^9/l)              | 2.36    | 0.9-5      | Urea (mg/dl)            | 42      | 15-59      |
| Monocyte (10^9/l)                | 0.66    | 0.2-1.5    | Creatinin (mg/dl)       | 1.2     | 0.5-1.5    |
| Neutrophil (10^9/l)              | 7.29    | 3.5-12     | Total Protein (g/dl)    | 4.5     | 5.4-7.8    |
| Eosinophil (10^9/l)              | 0.1     | 0.1-2.5    | Albumin (g/dl)          | 2.7     | 2.8-4      |
| Erythrocyte (10^12/l)            | 9.18    | 5.5-8.5    | Total Bilirubin (mg/dl) | 0.1     | 0.1-0.3    |
| Hemoglobin (g/dl)                | 18.4    | 12-18      | Direct Bilirubin (mg/dl)| 0.06    | –          |
| Hematocrit (%)                   | 53.33   | 37-50      | Alkaline Phosphatase (IU/L) | 15   | < 130     |
| Mean Corpuscular Volume (fl)     | 61      | 60-72      | Alanine Transaminase (IU/L) | 45    | < 50      |
| Mean Corpuscular Hemoglobin (pg) | 19      | 18-125.5   | Aspartate transaminase (IU/L) | 24   | < 40      |
| Mean Corpuscular Hemoglobin Conc (g/dl) | 32.7 | 32-38.5 | Gamma-Glutamyl Transferase (IU/L) | 6 | 6-28 |
| Erythrocyte Distribution Width (%) | 16.3 | 12-17.5 | Chlorine (mg/dl)         | 94      | 102-117    |
| Platelets (10^9/l)               | 471     | 200-500    | Sodium (mmol/l)         | 138     | 147-156    |
| Mean Platelet Volume (fl)        | 9.9     | 5.5-10.5   | Potassium (mmol/l)      | 4.9     | 3.6-5.6    |
|                                  |         |            | Calcium (mg/dl)         | 9.8     | 9-11.3     |
|                                  |         |            | Total Thyroxine (µg/dl) | 2.34    | 0.6-3.6    |
|                                  |         |            | Cortisol (mg/dl)        | 2.68    | 1-4        |
Unlike hydranencephaly, defects in porencephaly are less extensive and limited to the cerebrum (9). Due to the absence of active neoplastic or inflammatory causes on a followup brain MRI, the focal cavitary lesions in the case were defined as porencephaly. It has been reported that central diabetes insipidus may occur as a result of pituitary injury, including trauma, infection, inflammation, and disruption of blood supply (2). It has also been reported that destructive mechanisms causing the development of hydranencephaly may also lead to damage in neurohypophysis, causing central diabetes insipidus (11). The etiology of central diabetes insipidus is described as idiopathic in the absence of structural hypothalamic and pituitary defects. Although it may not be possible to detect structural hypothalamic and pituitary defects on imaging, an underlying unknown pathology may be the cause of central diabetes insipidus (1). The absence of pituitary or hypothalamic lesions on MRI in the dog with porencephaly increased the probability of an unknown pathology between porencephaly and diabetes insipidus.

In conclusion, the association between porencephaly and central diabetes insipidus should prompt clinicians to remain vigilant about the development of central diabetes insipidus in dogs without detectable hypothalamic or pituitary abnormalities on MRI.

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