Imaging and surgical outcomes of spinal tumors in 18 dogs and one cat

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Clinical and magnetic resonance imaging (MRI) findings, histological appearances and surgical outcomes of 18 dogs and one cat with spinal tumors are presented. Medical records of the cases admitted for spinal disorders were reviewed, and cases of spinal tumors that were diagnosed by MRI and confirmed by histological examination were included in this study. T1 weighted, T2 weighted and contrast enhanced T1 weighted images were taken and interpreted to evaluate the spinal tumors. The tumors were diagnosed as: meningioma (n = 6), ependymoma (n = 1), nerve sheath tumor (n = 4), metastatic spinal tumor (n = 3), osteosarcoma (n = 2), osteoma (n = 1), rhabdomyosarcoma (n = 1), and nephroblastoma (n = 1). Thirteen cases underwent surgical operation and the remaining six cases were euthanized at the request of the owners. The neurological status of the surgical cases did not deteriorate, except for one dog that showed ependymoma in the early period after the operation. These results indicate the potential for surgical gross total tumor removal of vertebral tumors to provide better quality of life and surgical collection of histological specimens for definitive diagnosis. For effective case management, dedicated MRI examination is important to accurately evaluate the spinal tumors, and surgical treatment is useful for extradural and intradural-extradural spinal tumors.

Keywords: dog, histology, magnetic resonance imaging, spinal tumor

Introduction

Spinal neoplasia involves the spinal cord, dura, exiting peripheral nerves, or paraspinal tissues (e.g., the vertebrae and ligaments) and results in clinical signs of spinal cord dysfunction [2]. Spinal tumors can be classified as primary, originating from spinal meningeal and paraspinal tissues, or as secondary, metastasizing from other locations. They are also classified by anatomic location, in relation to the dura and spinal cord, and based on whether there is spinal involvement [7,13].

Magnetic resonance imaging (MRI) has a high soft-tissue resolution and plays a central role in depiction of spinal tumors, allowing them to be classified as extradural, intradural-extraduillary or intramedullary, which is very useful in tumor characterization. Extradural tumors are the most common spinal tumors in dogs and cats. These tumors include primary and secondary mesenchymal tumors (osteosarcoma, fibrosarcoma, and chondrosarcoma), hemangiosarcoma, multiple myeloma and other plasma cell tumors, liposarcoma, and lymphosarcoma [2,8,20,26]. Intradural but extramedullary tumors include meningiomas and nerve sheath tumors [6,32,33]. Meningiomas are most often found in the cervical area, followed by the lumbar area. Intramedullary tumors, which include astrocytomas, oligodendrogliomas, ependymomas and nephroblastomas, arise from cells within the spinal cord parenchyma [9,13,16,33]. Additionally, there is an unusual intradural but extramedullary tumor that has been referred to by various terms, including neuroepithelioma and spinal cord blastoma [4,13,29].

Treatment options for spinal tumors include surgical removal, radiation therapy and chemotherapy [10,11,12,17]. The prognosis depends on the degree of local resection, degree of spinal infiltration, spinal cord damage before and during surgery, surgeon’s experience with spinal neoplastic conditions, and tumor type [2]. This study was conducted to describe the clinical and MRI findings and histological features of 19 cases of spinal tumor, and to evaluate the surgical outcomes of 12 dogs and one cat.

Materials and Methods

Medical records at the Ankara University Faculty of
Veterinary Medicine Department of Surgery from November 2007 to February 2015 were reviewed for spinal tumors diagnosed by MRI and confirmed histologically. The results of neurological examination, neurological grade, conventional radiography findings, MRI findings, lesion localization, treatment modalities and outcomes were also evaluated [18].

Clinical examination: General physical, clinical and orthopedic examinations, and later neurologic examination scheme for spinal disease [27] were carried out, and paraplegia was graded according to the paraplegia score for dogs reported by Levine et al. [18]. In all dogs, complete blood count, serum biochemical profile, thoracic radiography and abdominal ultrasonography were performed to determine possible metastasis. Clinical outcomes were followed by neurological examination of the changes in neurological status and clinical findings after operation.

MRI acquisitions
All MRI examinations were performed with a 1.5 Tesla MR machine (Vision Plus; Siemens, Germany). T1 (time of repetition [TR]/time of echo [TE], 400/20 ms) and T2 (TR/TE, 2900/96 ms) weighted (W) turbo-spin echo images were obtained with 2 mm slice thicknesses. Gadolinium diethylenetriaminepentaacetic acid (Magnevist; Bayer, Germany) was used as the paramagnetic contrast medium, and was administered (dose, 0.2 mmol/kg) intravenously. All images were evaluated by board certified human neuroradiologists (OA) and veterinary surgeons (OB, PC).

Surgical technique
Laminectomy or hemilaminectomy were conducted to reach the lesion or remove the mass. The tumors were removed under the operation microscope in all cases. In two cases with osteosarcoma and one case with osteoma, spondylectomy was conducted to remove the mass. The created defects were filled with polymethylmethacrylate.

Histological examination
Histologically collected samples of each mass were fixed with 10% buffered formalin embedded in paraffin and cut to 5 µm-thick sections. All sections were stained with hematoxylin and eosin (H&E), phosphotungstic acid and Masson’s trichrome stains. Some sections were stained using pan-cytokeratine, S-100, glial fibrillary acidic protein, α-smooth muscle actin (α-SMA) and vimentin (Dako, USA).

Outcome
The outcomes were provided after neurological examination and/or by telephone conversation with the owner or referring veterinarians.

Results
Eighteen dogs and one cat matched the case selection criteria for this study. The clinical, MRI, histological examination and surgical outcomes are summarized in Table 1. The sex distribution was 12 males and 6 females for dogs, as well as one female cat. Median age at presentation was 7.68 years (range, 2 to 12 years old), and median body weight for dogs was 16 kg (range, 7 to 52 kg). The breed distribution was mixed breed (n = 7), Sharpei (n = 1), Cocker Spaniel (n = 2), Poodle (n = 4), Siberian Husky (n = 1) long haired Collie (n = 1), Rottweiler (n = 1) Golden Retriever (n = 1), and one domestic short hair cat. Meningioma was seen in five dogs and one cat (66.66% of all intradural-extramedullary tumors). Localization of the tumor was in the lumbar area in three cases, in the cervical area in two cases, and in thoracic area in one cat. The clinical signs were progressive and in different degrees of neurological dysfunction. Characteristic MRI findings of all cases with meningioma were seen as hypointense on T1-weighted (T1W), slightly hyperintense on T2-weighted (T2W) MR sequences, and contrast-material enhancement on postcontrast T1W images. Meningeal tail was seen in two cases on postcontrast T1W images. Generally, the location of meningiomas was intradural-extramedullar on MRI. Cases were treated surgically, except when the owner opted for euthanasia. The longest survival time with better outcome was seen for sarcomatous meningioma (> 6 month). Neurological improvement was not observed in one case for one month after surgery, after which it was lost to follow up (case No. 3). The remaining two dogs survived for 3 and 5 months. The cat with meningioma underwent a second surgery because of tumor recurrence after 3 months, but neurological status was not improved after another one month, and it was then euthanized at the request of the owner. Histologically, the most common pattern was laminated whorls of elongated cells. Neoplastic cells had variably distinct borders with moderate cytoplasm, oval to fusiform shape and occasionally vesicular nuclei and nucleolus.

A dog with osteosarcoma in the T12 vertebra showed acute onset paraplegia with intact deep pain perception that disappeared 48 hours later. However another case with osteosarcoma in C2 had progressive ataxia and cervical pain. Characteristic MRI findings of osteosarcoma as sclerotic and lytic areas were seen. Both showed good contrast enhancement on postcontrast T1W images. Partial spondylectomy was conducted to remove the gross tumor completely in both cases, and the created cavity was filled with polymethylmethacrylate (PMMA) (T12) and screw + polymethylmethacrylate (C2). The case with cervical osteosarcoma was euthanized on day 46 and the case with thoracic osteosarcoma euthanized on day 21 because of pulmonary metastasis. Osteoma was diagnosed in a 2 year old dog with a history of progressive paraparesis. The radiopaque and lytic areas were prominent upon direct
Table 1. Clinical, magnetic resonance imaging (MRI), histopathological examination and surgical outcomes in canine and feline patients with spinal tumors

| Number | Signalment | Neurological exam | MRI | Localization | Histological diagnosis | Treatment | Outcomes |
|--------|------------|------------------|-----|--------------|-----------------------|-----------|----------|
| 1      | 7 yr FN dog Mixed breed | HLs ataxia for about 1 year, nonambulatory paraplegia with intact DPP withdrawal, patellar and perianal reflexes intact, spinal pain (G2) | Hypo Hyper ++ | A mass intradural extramedullary located from the caudal border of L1 to the caudal border of L2 at the L1 side with compression to the spinal cord | Sarcomatous meningioma | Hemilaminectomy Durotomy: the border of tumor was not clear from the spinal cord. It was resected under the operation microscope | 2 months after surgery the neurological status improved from (G2) to (G4) |
| 2      | 9 yr MI dog Mixed breed | Progressive paraparesis for 1 month, patellar reflex depressed at the Lt side, spinal pain (G2) | Hypo Hyper ++ | Two intradural extramedullary, dorsally located solid lesions at the L4-5 levels | Meningioma | Laminectomy and the mass was removed gross totally | 6th months after surgery it was still (G4) |
| 3      | 11 yr F1 dog Poodle | Progressive HLs ataxia for 1 month, nonambulatory paraplegia without DPP, patellar and withdrawal reflexes depressed (G0) | Hypo Hyper ++ | Two masses intradural extramedullary located with heterogeneous nature at L4 and L5-L6 | Meningioma | Laminectomy and the masses totally removed | Neurologically improved (G3), CP deficit at the Lt side was persistent for 1 month. The dog was euthanatized after deteriorating neurologic status at the 3rd month post-surgery (G1) |
| 4      | 12 yr FN dog Poodle | Neck pain for 4 months, ataxia in all limbs, ambulatory tetraparesis | Hypo Hyper ++ | The solid mass intradural extramedullary located at the C2-C3 Lt side, with meningeal tail in post contrast T1 W images | Cervical meningioma | None | Neurological status was not improved at 1 month. Lost from follow up |
| 5      | 11 yr MI dog Long haired Collie | Neck pain for 1 month and occasional crying | Hypo Hyper ++ | A solid mass intradural extramedullary located, with the meningeal tail at C2 | Meningioma | Euthanized | None |
| 6      | 2.5 yr F1 cat Domestic short hair | 2 month ataxia and one month inability to use HLs, intact DPP (G2) | Hypo Hyper ++ | A mass intradural extramedullary Lt dorsolaterally located at T2 | Meningioma | Laminectomy and tumor gross removed | 3 months after surgery the cat could walk (G3), at 4th month she deteriorated neurologically (G1) |
| 7      | 13 yr FN dog Rottweiler | HLs ataxia for 4 months and recent 2 weeks progressed to all limbs ataxia, cervical pain | Hyper Hyper ++ | X-ray: radiolucency at the C3 vertebral body. MRI: mass extradural located at the C2 vertebral body invaded to the epidural space, heterogeneous nature and compressing the spinal cord | Osteosarcoma | Hemilaminectomy + corpectomy + Lt side screw and polymethylmethacrylate | Clinical signs improved for the first week, then gradually deteriorated. The dog was euthanized at the 46th postoperative day because of pulmonary metastasis |
| Number | Signalment | Neurological exam | MRI | Localization | Histological diagnosis | Treatment | Outcomes |
|--------|-------------|--------------------|-----|--------------|------------------------|-----------|----------|
| 8      | 2 yr MI dog Mixed breed | HLs ataxia for one month normal perineal, patellar and withdrawal reflexes (G3) | Hypo Hypo | + | X-ray: radiopacity in T7-T8 vertebral body invading to the right side paraspinal muscle. MRI: a mass extradurally located in the T7-T8 vertebral body and invading to the vertebral body, T2 hypointense lesion at the spinal cord | Osteoma | Rt hemilaminectomy + corpectomy. The mass elongated to the paraspinal muscle was removed, it was not bloody and had a granular structure. The defect at the vertebral body was filled with PMM | 1 month postoperative clinical signs were better (G4). The dog was still alive at 11 months after surgery (G5) |
| 9      | 7 yr MI dog Golden Retriever | Acute onset paraplegia with intact DPP, G1, 2 days later DPP lost (G0) | Hypo Hyper | + | X-ray: radio opacity at the T12 vertebral body. A mass extradural located at the vertebral body, invading to the epidural space at T12 | Osteosarcoma | The mass was gross totally removed and the cavity was filled with PMMA and a screw. | 10 days after surgery, deep pain recurred (G1). Day 21 after surgery pulmonary metastasis was observed and the dog was euthanized |
| 10     | 6.5 yr MI dog Mixed breed | Progressive paraparesis for 4 months. Nonambulatory paraplegia (G0), spinal reflexes depressed | Hypo Hyper | + | Multifocal unshaped intramedullary located at the spinal cord paranchyme, T5-L6 | Metastasis Malignant Mesenchymal tumor | Proxicam Euthanized 2 weeks after surgery (G0) |
| 11     | 9 yr Fl dog Cocker Spaniel | Surgery for perineal adenocarcinoma (3 months prior to presentation). Progressive HLs ataxia, non-ambulatory paraplegia without DPP. Normal perineal, patellar and withdrawal reflexes (G1) | Hypo Hyper | + | Multifocal masses extradurally located at T10-L3, lumbar sacral area in both bone and soft tissue, solid and nodular characteristic | Metastasis Adenoarcinoma | Euthanasia requested. Euthanized |
| 12     | 7 yr MI dog Mixed breed | Living in shelter, HLs ataxia for about 1 month, non ambulatory paraparesis for the last 3 days (G2), normal perineal, patellar and withdrawal reflexes | Hypo Hyper | + | Multifocal masses, intradural extramedullary located in cranial thoracic area in paraspinal muscles and lymph node, compressing spinal cord and causing myelomalacia at the T5 level | Metastasis Squamous cell carcinoma | Euthanasia requested Euthanized |
| 13     | 12 yr Fl dog Poodle | HLs ataxia for 2 months, paraplegia without DPP. Mases at the lumbar paraspinal muscles (G0) | Hypo Hyper | + | Multifocal masses located at the paraspinal muscles in all lumbar areas, invading to the epidural space extradurally and L6 vertebral body | Rhabdomyosarcoma invading to the epidural space (especially L2) in multilocation | Euthanasia requested Euthanized |
| Number | Signalment | Neurological exam | MRI | Localization | Histological diagnosis | Treatment | Outcomes |
|--------|-------------|-------------------|-----|--------------|------------------------|-----------|----------|
| 14     | 9 yr MI dog | Progressive spinal ataxia in all limbs for 3 months. Non ambulatory tetraparesis, neck pain, head tilting to the Rt side, muscle atrophy at the cervical muscles | Hypo Hyper ++ | A mass extradurally located at the Rt side of the C4 vertebral body and occupying the intervertebral foramen | Nerve sheath tumor | Rt side hemilaminectomy rhizotomy was conducted | Inability to walk for one month. Ambulatory tetra-paresis, with neck tilting to the right side. Muscle atrophy at the cervical muscles, euthanatized two months post surgery |
| 15     | 7 yr MI dog Sharpei | Progressive HLs ataxia, normal perineal, patellar and withdrawal reflexes, atrophy of paraspinal muscles, intact DPP, (G2) | Hypo Hyper | Mass extradurally located at the Rt paraspinal muscles, invading through the intervertebral foramen to the epidural space at the T9 vertebra level | Nerve sheath tumor | Hemilaminecoty and the mass surrounding the nerve root was removed from the epidural space and elongated between muscles. Second operation: (2 months later because of deterioration, and intradural extramedullary located mass) gross totally removed | 2 month observation: (G3) 3 month observation: neurological status had not progressed and the dog was euthanized 1m later after a second operation |
| 16     | 4 yr MI dog Cocker Spaniel | 2 week kyphosis, paraparesis without DPP at the Lt, normal perineal reflexed, patellar and withdrawal reflexes normal at the Rt and depressed at the Lt (G2) | Hypo Hyper | Mass extradurally located at the L4-L5 level, Lt side intraforaminal location, nodular, lobulated spikula contours | Nerve sheath tumor, myositis, calcification and hemorrhage at the paraspinal muscle | Hemilaminectomy gross totally removed | Neurological status had not improved at 1 m. Loss of follow up |
| 17     | 9 yr MI dog Poodle | HLs ataxia for 2 months, spinal pain at the thoracolumbar area (G2) | Hypo Hyper ++ | Solid mass intradural extramedullary, Lt side located mass T13-L1 | Nerve sheath tumor | Hemilaminectomy surgically removed | Improved 1 month after surgery (G4). Later died from gastroenteritis |
| 18     | 6 yr MI dog Siberian Husky | Progressive para-paresis for 2 months, intact DPP, depressed perineal, patellar and withdrawal reflexes (G2) | Hypo Hyper ++ | Intramedullary solid lesion at the L6 level, Diastematomyelia L5-L6 | Ependymoma | Laminectomy gross totally removed | DPP was lost after the operation (G0) Euthanasia requested |
| 19     | 2 yr MI dog Mixed breed | HLs ataxia for 6 months and unable to use HLs for 1 month. Paraplegia without DPP, normal perineal, patellar and withdrawal reflexes (G0) | Hypo Hyper | A mass intradural extramedullary located at the T12 level and syringomyelia | Nephroblastoma | Euthanasia requested | Euthanized |

G0, no voluntary movement seen when the dog is supported; G1, intact limb protraction with no ground clearance; G2, intact limb protraction with inconsistent ground clearance; G3, intact protraction with ground clearance > 75% of steps; G4, ambulatory with consistent ground clearance and significant paresis ataxia that results in occasional falling; G5, ambulatory with consistent ground clearance and mild paresis ataxia that does not result in falling; G6, normal gait [18]. +, moderate enhancement; ++ , good enhancement. MI, male intact; MN, male neutered; FI, female intact; FN, female neutered; yr, year; Lt, left; Rt, right; HL, hind limbs; DPP, deep pain perception; Hypo, hypointense; Hyper, hyper intense; CME, contrast-material enhancement.
radiography. The tumor appeared in T7–T8 vertebral bodies, extended to the epidural space and pedicles and compressed the spinal cord from the right side. The tumor was hypointense on both T1W and T2W images, and showed ring like heterogeneous contrast-material enhancement on postcontrast T1W images (Fig. 1). The tumor size was 29 × 33 mm on MRI, and it was completely removed by hemilaminectomy and partial spondylectomy, which created a defect that was filled with PMMA. Histologically, it was a well differentiated ovoid bone tumor (Fig. 2) in which the cells had hyperchromatic nuclei that were eccentrically located in dark stained cytoplasm and produced thin spicules of tumor osteoid and matrix.

Clinical manifestation of cases with metastatic tumor consisted of neurologically weighted symptoms, which were the chief complaints proposed by the owner, and clinical signs were progressive. Metastatic spinal tumor was diagnosed in four cases and malign mesenchymal tumor, squamous cell carcinoma (panels A and B in Fig. 3), adenocarcinoma and rhabdomyosarcoma was diagnosed histologically. Tumors were multifocal, unshaped, and hypointense on T1W and hyperintense on T2W images. All tumors were located epidurally except for one case with metastatic mesenchymal tumor (case No. 10), in which it invaded the spinal cord.

Nerve sheath tumor was diagnosed in four dogs, all of which
had progressive proprioceptive ataxia, pain and different degrees of paresis, as well as atrophy of the related muscles. The tumor was located epidurally in all cases except for one with recurrence of clinical signs, and on the second MRI the mass was seen as both extradural and intradural (Case No. 15). Even though the mass was completely removed during the second operation, the neurological status was not improved and the dog was euthanized in response to the owner’s request one month later. Hemilaminectomy with rhizotomy was conducted in all cases with nerve sheath tumor, and they survived for 2 or 3 months, although one case was lost to follow up after 1 month (Case No. 16). Nerve sheath tumor was diagnosed in one dog at L4–L5. After this tumor was removed surgically, clinical improvement was seen; however, the dog died from an unrelated disease 1 month later. Histologically, the tumor was well circumscribed without encapsulation and mainly composed of spindle-shaped cells with thin and eosinophilic cytoplasms and atypical plump, ovoid, and vesicular nuclei.

An intramedullary solid lesion that was confirmed histologically as ependymoma at L6 and accompanied diastematomyelia was removed surgically, but neurological status deteriorated and the animal was euthanized (panels A and B in Fig. 4). Upon histological examination, the ependymoma was composed of pseudorosettes, and immunohistochemical examination revealed vimentin and glial fibrillary acidic protein immunoreactivity in the neoplastic cells (panel C in Fig. 4).

One case with a mass located at the T12 level that was intradural-extradural and mimicked meningioma was histologically diagnosed as nephroblastoma. This dog was euthanized without any treatment. Histologically, the tumor was composed of uniform cells that were cuboidal to ovoid with indistinct cell borders and scant to mildly amphiphilic. Immunohistochemically, neoplastic cells, especially in tubules and glomeruloid structures, stained pan cytokeratin. Conversely, fibrovascular stroma and capsules were stained α-SMA and vimentin sera.

**Discussion**

Characteristic signs and location of spinal tumor in MRI provided ante-mortem prediction of the various histological types of neoplasms. However, histological confirmation should be conducted for definitive diagnosis. Surgical removal of spinal tumors without neurological derioriation after the operation was observed for all except one case with ependymoma in this study, which indicates potential for

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**Fig. 3.** (A) Contrast enhanced mid-sagittal T1 weighted image of thoracolumbar spine of a dog with metastatic squamous cell carcinoma. Metastatic tumors were contrast enhanced in T1W images (arrow heads). (B) Metastatic squamous cell carcinoma. H&E stain. 100× (B).

**Fig. 4.** Mid-sagittal T2W image of lumbar spine (A) transverse T2W image at the level of L6 (B) in a dog with ependymoma hyperintense masses located in the spinal cord parenchyma in T2W images. (C) Pseudorosette with central fibrovascular stroma (ependymoma). H&E stain. 400× (C).
treatment by gross total tumor removal and providing specimens for histological examination.

Older and larger breeds of dogs have been reported to be more susceptible to spinal tumors [5,15]; however, there may be a tendency for younger animals to develop tumors of neural origin [9,19,24]. The mean age of the dogs presented in this study was 7.68 years (range, 2–12 years), which is similar to that reported in the literature. In this case series, the youngest dog (2 years old) had nephroblastoma that occurred between 6 months and 3 years of age, but it has also been reported in dogs as old as 7 years [19,31].

Clinical signs related to spinal tumors can be noticed by the owner earlier before presentation to the veterinarian. This interval was reported to be about 1 month in a recent study [25]. The clinical signs of a spinal tumor represent spinal cord dysfunction and pain, but these are not pathognomonic [6,14,28]. In the present study, the time of determining abnormal clinical signs was not clear, and in some cases the owners could not remember the exact time of onset of neurogenic disorders. The clinical signs were progressive in all cases except for one case with osteosarcoma, in which acute onset of clinical signs was noticed by the owner. The later situation could be related to the growing mass invading to the epidural space and causing damage to the neural structure.

Distribution of spinal tumors according to location relative to the dura mater and histopathology has been reported as intramedullary (approximately 15%) [26], extradural (approximately 50%) and intradural-extradural (approximately 35%), which does not differ greatly from the observations in humans [23,24,26,31]. In the cases reported in this study, intramedullary tumor location was observed in one ependymoma and one metastasis case (2/19 in this study, intramedullary tumor location was observed in one case), indicating that this is not a pathognomonic MRI finding for meningioma. Meningioma revealed similar characteristics to MRI finding for meningioma. Meningioma revealed similar signal on T2w images [3,5,6,30]. In this case series, nerve sheath tumors showed different clinical signs according to anatomical location and usually progressed to proprioceptive impairment and sometimes to paraplegia or tetraplegia caused by spinal cord compression and related spinal cord pathophysiologic alterations. Nerve sheath tumors are isointense on T1W images and have an atypical marked high signal on T2w images [3,5,6,30]. In this case series, nerve sheath tumor was hypointense on T1W image and highly hyperintense on T2W images, and contrast-material was enhanced in all cases. Even though the tumor was located epidurally in the first operation, it had grown intradurally 2 months later and was completely removed after durotomy. Paraspinal muscle atrophy was seen in all cases, but it was more remarkable in two cases (case Nos. 14 and 15), and all cases with nerve sheath tumor showed progressive clinical signs.

Clinical signs of intradural extramedullary ependymomas appeared in the later stage of the disease, and occurred in the conus medullaris and the filum terminale in humans [9,30]. Canine spinal cord nephroblastoma, which is also known as Wilms’ tumor, generally occur between T10-L2 spinal cord segments in large breed dogs [19]. The ependymoma case presented in this study was admitted with hindlimb ataxia for 2 months. In addition to the tumor, there was diastematomyelia at the level of L6 in MRI, which deteriorated after surgical removal of the tumor. The case of nephroblastoma was accompanied by syringomyelia in MRI. Even though this case was not treated surgically, the owner allowed a biopsy, which enabled the case to be histologically confirmed.

Any malignant tumor can metastasize to bone, but the most common metastatic spinal tumors found in women are from the breast and lung, while those in men are from the prostate and lung in humans [30]. Metastases can also affect the spinal cord [8,21]. Both extradural and intramedullary metastasis is possible. Carcinomas are one of the most common types of tumors associated with extradural metastasis. In some instances, clinical signs of the metastasis may be apparent before clinical signs of the primary tumor [2]. In this case series, the first clinical signs noted by the owner were neurological signs that were located extradurally (n = 2), intramedullary (n = 1) and intradural extramedullary (n = 1), and none of them were treated surgically.

Bone tumors may be evident on survey radiographs as osteolytic/osteoproliferative processes. Classically, vertebral tumors do not cross the joint space (intervertebral disk); however, vertebral tumors do on occasion invade adjacent vertebral bodies and therefore appear to “jump” the joint.
Extradural compression of the spinal cord overlying the vertebral body rather than the intervertebral disk space is indicative of neoplasia [2]. Spinal osteosarcoma was reported as 15% [15]. Both cases with osteosarcoma were predicted from survey radiography. However, the tumor margins and exact location with compressing spinal cord was diagnosed by MRI. Interestingly, in one case, acute onset paraplegia occurred. This case was treated with partial spondylectomy, and the created defect was filled with polymethylmethacrylate. Surgical procedures were well tolerated by the dogs, and their neurological status was not deteriorated during the early postoperative period. Partial spondylectomy for bone tumors and reconstruction with polymethylmethacrylate was found to be a versatile and cost effective method. However, wide surgical margins are essential for surgical oncology by radical spondylectomy, which involves more complex spinal instrumentation. Nevertheless, in both cases with osteosarcoma, the tumors were completely resected. This suggest that, if the radical spondylectomy and spinal instrumentation is combined with adjuvant or neoadjuvant therapy, the survival time could have been longer.

Osteoid osteoma and osteoblastoma are commonly seen as benign osteogenic bone neoplasms. Histologically, these tumors resemble each other, and if the lesion is larger than 1.5 cm it is accepted as osteoblastoma, which is most frequently located in the axial skeleton in humans. This tumor has a tendency to affect the posterior part of the vertebra and occurs primarily in the pedicle and the posterior elements, not in the vertebral body. As a result, this tumor often requires surgical resection. [1,30]. The case with osteoma presented in this study showed invasion of the tumor to the lateral part of the vertebra and that it included vertebral bodies larger than 1.5 cm. Even though the lesion was removed completely and the created defect repaired satisfactorily, radical spondylectomy and more sophisticated spinal instrumentation can be considered in humans to minimize recurrence. The case considered in this study was still alive with slight hindlimb ataxia at the time that this manuscript was written. To the best of the author’s knowledge, this is the first case of spinal osteoma in dogs to be reported.

The method of choice for visualising spinal tumors is MRI. The tumor’s involving structures, its relationship with the duramater, degree of spinal cord compression, and changes in spinal cord paranchyme can be visualised by MRI [11,29]. In this case series, all cases were predicted as tumors by MRI when compared with histological results. The characteristic MRI signs of the tumor were found to be reliable for diagnosis of spinal tumors and determination of the the surgical plane, and the multifocal appearance of the metastatic tumors on MRI differentiated them from other primary spinal cord tumors.

Extradural tumors are removed surgically, but both intradural/extradural and intramedullary tumors can be successfully resected [2]. Radiotherapy and chemotherapy are also treatment options that can be applied individually or in combination with surgery. In this case series, only surgical treatments were applied, and the results of surgery were found to be strategic except for intramedullary tumor (case No. 18), in which neurological status had deteriorated.

In conclusion, spinal neoplasia should be considered in cases with progressive neurological signs, and MRI is a useful and effective method for diagnosis of spinal tumors. Operative management is a strategic for epidural and intradural-extradural spinal tumors according to the surgical outcomes. Partial spondylectomy and reconstruction of vertebral bodies with polymethylmethacrylate can be suggested for bone tumors. Further studies are needed to correlate clinical findings, and characteristic signs, as well as the shape of tumor in MRI, histological findings and outcomes after different treatment.

Conflict of Interest

There is no conflict of interest.

References

1. Atesok KI, Alman BA, Schemitsch EH, Peyser A, Mankin H. Osteoid osteoma and osteoblastoma. J Am Acad Orthop Surg 2011, 19, 678-689.
2. Bagley RS. Spinal neoplasms in small animals. Vet Clin North Am Small Anim Pract 2010, 40, 915-927.
3. Bailey CS. Long-term survival after surgical excision of a schwannoma of the sixth cervical spinal nerve in a dog. J Am Vet Med Assoc 1990, 196, 754-765.
4. Blass CE, Kirby BM, Kreeger JM, Gossett KA. Teratomatous medulloblastoma in the spinal cord of a dog. J Am Anim Hosp Assoc 1988, 24, 51-54.
5. Bradley RL, Withrow SJ, Snyder SP. Nerve sheath tumors in the dog. J Am Anim Hosp Assoc 1982, 18, 915-921.
6. Brehm DM, Vite CH, Steinberg HS, Haviland J, van Winkle T. A retrospective evaluation of 51 cases of peripheral nerve sheath tumors in the dog. J Am Anim Hosp Assoc. 1995, 31, 349-359.
7. Byrne TN. Spinal cord compression from epidural metastasis. N Engl J Med 1992, 327, 614-619.
8. Cooley DM, Waters DJ. Skeletal metastasis as the initial clinical manifestation of metastatic carcinoma in 19 dogs. J Vet Intern Med 1998, 12, 288-293.
9. de Vries-Chalmers Huyk van Papendrecht HR, Vos JH, van Nes JH. Spinal cord epedymoma in two young dogs. Vet Q 1988, 10, 205-210.
10. Ferretti A, Scanziani E, Colombo S. Surgical treatment of a spinal cord tumor resembling nephroblastoma in a young dog. Prog Vet Neurol 1993, 4, 84-87.
11. Fingeroth JM, Prata RG, Patnaik AK. Spinal meningiomas in dogs: 13 cases (1972-1987). J Am Vet Med Assoc 1987, 191, 720-726.
12. Gavin PG, Bagley RS. Practical Small Animal MRI. pp. 179-199, Wiley Blackwell, Ames, 2009.
13. Gilmore DR. Intraspinal tumors in the dog. Compend Contin Educ Pract Vet 1983, 5, 55-64.
14. Gilmore DR. Neoplasia of the cervical spinal cord and vertebrae in the dog. J Am Anim Hosp Assoc 1983, 19, 1009-1014.
15. Heyman SJ, Diefenderfer DL, Goldschmidt MH, Newton CD. Canine axial skeletal osteosarcoma: a retrospective study of 116 cases (1986 to 1989). Vet Surg 1992, 21, 304-310.
16. Huisenga M, Henrich M, Frese K, Burkhardt E, Kuchelmeister K, Schmidt M, Reimacher M. Extraventricular neurocytoma of the spinal cord in a dog. Vet Pathol 2008, 45, 63-66.
17. Jeffery ND. Treatment of epidural haemangio-sarcoma in a dog. J Small Anim Pract 1991, 32, 359-362.
18. Levine GJ, Levine JM, Budke CM, Kerwin SC, Au J, Vinayak A, Hettlich BF, Slater MR. Description and repeatability of a newly developed spinal cord injury scale for dogs. Prev Vet Med 2009, 89, 121-127.
19. Liebel FX, Rossmeisl JH Jr, Lanz Ol, Robertson JL. Canine spinal nephroblastoma: long-term outcomes associated with treatment of 10 cases (1996-2009). Vet Surg 2011, 40, 244-252.
20. Luttgen PJ, Braund KG, Brawner WR Jr, Vandeveld M. A retrospective study of epidural haemangio-sarcoma in a cat. J Small Anim Pract 1980, 21, 213-226.
21. Macpherson GC, Chadwick BJ, Robbins PD. Intramedullary spinal cord metastasis of a primary lung tumour in a dog. J Small Anim Pract 1993, 34, 242-246.
22. McDonnell JL, Tidwell AS, Faissler D, Keating J. Magnetic resonance imaging features of cervical spinal cord menigiomas. Vet Radiol Ultrasound 2005, 46, 368-374.
23. Newton HB, Malkin MG. Overview of spinal cord tumor epidemiology. In: Newton HB, Jolesz FA (eds.). Handbook of Neuro-Oncology Neuroimaging. 1st ed. pp. 31-35, Academic Press, New York, 2008.
24. Pancotto TE, Rossmeisl JH Jr, Zimmerman K, Robertson JL, Werre SR. Intramedullary spinal cord neoplasia in 53 dogs (1990-2010): distribution, clinicopathologic characteristics, and clinical behavior. J Vet Intern Med 2013, 27, 1500-1508.
25. Petersen SA, Sturges BK, Dickinson PJ, Pollard RE, Kass PH, Kent M, Vernau KM, Lecouteur RA, Higgins RJ. Canine intraspinal menigiomas: imaging features, histopathologic classification, and long-term outcome in 34 dogs. J Vet Intern Med 2008, 22, 946-953.
26. Prata RG. Diagnosis of spinal cord tumors in the dog. Vet Clin North Am 1977, 7, 165-185.
27. Sharp NJH, Wheeler SJ. Patient examination. In: Sharp NJH, Wheeler SJ (eds.). Small Animal Spinal Disorders: Diagnosis and Surgery. 2nd ed. pp. 19-32, Elsevier Mosby, New York, 2005.
28. Shell L, Dallman M, Sponenberg P. Chondrosarcoma in a cat presenting with forelimb monoparalysis. Compend Contin Educ Pract 1987, 9, 391-397.
29. Summers BA, deLahunta A, McEntee M, Kuhajda FP. A novel intradural extramedullary spinal cord tumor in young dogs. Acta Neuropathol 1988, 75, 402-410.
30. Van Goethem JW, van den Hauwe L, Ozsarlak O, De Schepper AM, Parizel PM. Spinal tumors. Eur J Radiol 2004, 50, 159-176.
31. Wright JA, Bell DA, Clayton-Jones DG. The clinical and radiological features associated with spinal tumours in thirty dogs. J Small Anim Pract 1979, 20, 461-472.
32. Yoshioka MM. Meningioma of the spinal cord in a cat. Compend Contin Educ Pract 1987, 9, 34-38.
33. Zaki FA, Prata RG, Hurvitz AI, Kay WJ. Primary tumors of the spinal cord and meninges in six dogs. J Am Vet Med Assoc 1975, 166, 511-517.