## Titanium mesh reconstruction of a dog's cranium after multilobular osteochondrosarcoma resection

Reconstructie van de schedel van een hond met een titanium mesh na het verwijderen van een multilobulair osteochondrosarcoom

### A. Dierckx de Casterlé, <sup>1</sup>B. Van Goethem, <sup>1</sup>A. Kitshoff, <sup>1</sup>S.F.M. Bhatti, <sup>2</sup>I. Gielen, <sup>1</sup>T. Bosmans, <sup>3</sup>H. De Cock, <sup>1</sup>H. de Rooster

<sup>1</sup>Small Animal Medicine, Faculty of Veterinary Medicine, Ghent University, Salisburylaan 133, 9820 Merelbeke, Belgium

<sup>2</sup>Medical Imaging and Small Animal Orthopedics, Faculty of Veterinary Medicine, Ghent University, Salisburylaan 133, 9820 Merelbeke, Belgium

<sup>3</sup>AML/Veterinary Pathology Services, Emiel Vloorsstraat 9, 2020 Antwerpen, Belgium

bart.vangoethem@Ugent.be



An eleven-year-old cavalier King Charles spaniel was presented with a large mass arising from the sagittal crest of the skull. Computed tomography also revealed an intracranial component. A histological diagnosis of multilobular osteochondrosarcoma grade 1 was made from surgical biopsies. Since this tumor type has a moderate aggressive biological behavior characterized by a slow growth, compression of adjacent structures, and only a 30% metastatic rate, surgical resection was performed. A wide partial craniectomy was performed, the skull defect was reconstructed with a designated custom designed titanium mesh and the skin defect closed with a local subdermal plexus flap technique. Histologic evaluation indicated clean surgical margins, which may lead to a long-term survival in this low-grade tumor. Approximately seventeen months after surgical resection, the dog showed no signs of local tumor recurrence or metastasis.

#### **SAMENVATTING**

Een elf jaar oude cavalier-king-charles-spaniël werd aangeboden met een grote massa centraal op het schedeldak. Met behulp van computertomografie werd de intracraniale uitgebreidheid van het letsel vastgesteld. Bioptname en histologisch onderzoek leidden tot de diagnose van een graad-1 multilobulair osteochondrosarcoom. Vermits dit een biologisch matig agressieve tumor is die traag groeit, de nabijgelegen weefsels wegdrukt in plaats van te invaderen en deze laaggradige tumor slechts bij 30% van de patiënten metastaseert, werd besloten tot chirurgische resectie. Een ruime partiële craniëctomie werd uitgevoerd, waarna het defect in het schedelbot werd gereconstrueerd met een specifiek daartoe bestemde titanium mesh. Het huiddefect werd gesloten met een lokale huidflap. De marges werden histologisch beoordeeld en waren vrij van tumorcellen. Volledige chirurgische resectie van deze tumor kan resulteren in een lange overlevingstijd. De beschreven hond was ruim zeventien maanden na de operatie nog steeds in leven, zonder aanwijzingen voor een lokaal recidief of metastasering.

#### INTRODUCTION

Multilobular osteochondrosarcoma (MLO) is an uncommon malignant tumor occurring most often in the flat bones of the canine skull (McGavin and Zachary, 2007). Similar names used for this tumor type include chondroma rodens, multilobular tumor of bone, multilobular osteosarcoma, multilobular osteoma, calcifying aponeurotic fibroma, multilobular chondroma, cartilage analogue of fibromatosis, and juve-

nile aponeurotic fibroma (Dernell et al., 1998). The majority of dogs with MLO are middle-aged to older dogs (median age at presentation eight years), usually belonging to medium or large breeds (median weight 29 kg) (Straw et al., 1989; Groff et al., 1992; Popvitch et al., 1994; Dernell et al., 1998; Loukopoulos et al., 2003; Jubb et al., 2007).

Clinically, these tumors are slow-growing, locally invasive, firm immobile masses on the surface of the skull (Hathcock and Newton, 2000; Pakhrin et al., 2006). Although the bone may be invaded by the tumor, the brain is often compressed rather than infiltrated by the mass, resulting in late-onset neurological signs (Pool, 1990; Dernell et al., 1998). MLO has a characteristic 'popcorn ball' appearance on radiographic imaging, CT and MRI with sharply demarcated, limited lysis of adjacent bone and a course granular mineral density throughout the tumor (Selcer, 1981; Dernell et al., 1998; Webb et al., 2009; Boston, 2010; Forrest, 2013).

The biological behavior makes complete excision the treatment of choice (Straw et al., 1989; Gallegos et al., 2008). Following surgical resection, local tumor recurrence has been reported in 47-58% of dogs with a median time to local recurrence of 420-542 days (Straw et al., 1989; Dernell et al., 1998). The key surgical challenge is therefore to obtain adequate surgical margins. This is complicated by the direct presence of vital structures, of which damage may lead to severe or fatal outcomes, and by the necessity to reconstruct the created skull defect to protect these structures.

A metastatic rate of 56-58% in treated dogs has been reported with a median time to metastasis between 420-797 days (Straw et al., 1989; Dernell et al., 1998). The metastatic sites include mostly the lungs, but also the cerebral cortex, kidneys, pancreas, rib and long bones (Mc Lain et al., 1983; Straw et al., 1989; Dernell et al., 1998).

A histologic grading system has been described in an attempt to predict the biological behavior of this tumor (Banks and Straw, 2004) (Table 1). Local tumor recurrence, metastasis and survival time appear to be correlated to histologic grade (Dernell et al., 1998) (Table 2).

In this case report, the occurrence of MLO in a previously unreported breed, the surgical removal and the use of a designated contourable titanium mesh for the reconstruction of the bone defect in the skull com-

Table 1. Histological grading criteria for MLO (Dernell et al., 1998).

Criteria	Score
Borders	1
Pushing Pushing and invasive	_
Invasive	2 3
Size of lobules	
Small and medium	1
Large	2
Organization	
Well organized	1
Moderately well organized Poorly organized	2 3
1 doily diguinized	
Mitotic figures/10 HPFs*	1
1 to 5 6 to 10	1
>10	2 3
Pleomorphism of cells	
Monomorphic	0
Mild	1
Moderate	2
Marked	3
Necrosis	
None	0
Present	1
Grade	Total
Grade I	7 or less
Grade II Grade III	8 to 12
Grade III	13 or greater

<sup>\*</sup>HPFs = high-powered fields

Table 2. Survival outcome for MLO based on tumor grade (Dernell et al., 1998).

Grade	I	П	Ш
Number of cases	13	17	9
Time to local recurrence			
(days)			
Median	>1.332	782	288
Range	192-1.332	30-782	82-534
Time to metastasis (days) Median Range	>820 720-820	405 28-1.225	321 150-542
Survival time (days)			
Median	>897	520	405
Range	66-797	28-1.487	82-1.670

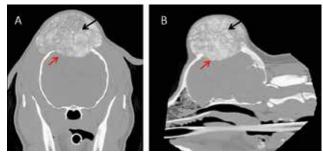


Figure 1. A. Transverse and B. sagittal CT images of the skull. A well-circumscribed, space-occupying mass is present on the skull with a granular, hyperdense aspect (black arrows). Intracranial expansion with dorsal compression of the brain is present (red arrows).

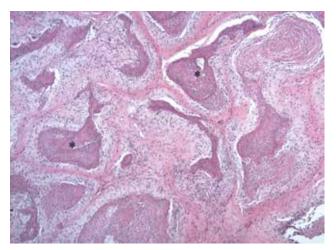


Figure 2. Microscopy of the MLO demonstrating multilobular patterns, consisting of a core of bone matrix (asterisk) surrounded by multiple connective tissue cells (Hematoxylin and eosin stain, 40x).

bined with a single-pedicle advancement flap to close the skin defect are described.

#### CASE DESCRIPTION

An eleven-year-old female castrated cavalier King Charles spaniel with a body weight of 8.6 kg was examined for a large mass on the skull of an elevenmonths duration. The mass was approximately 5 cm in diameter and centered on the top of the skull. The dog was alert, responsive and able to ambulate well on all limbs. Although no neurologic dysfunction was noted, the owners mentioned the recent appearance of behavioral changes, i.e. anxiety and hiding in the corner of the room. Physical examination was unremarkable other than bilateral breed-related exophthalmia and a systolic heart murmur (3/6). The results of complete blood count were within the reference limits; so were the results of serum biochemical analyses except for the elevated activity of alkaline phosphatase (571 U/L; reference range, 23-212 U/L).

A CT scan of the head was performed to evaluate the extent of bone involvement and to determine

the intracranial extension of the mass. The CT scan revealed a 5.5 x 5.0 x 3.7 cm lobulated, mineralized mass arising from the caudal portions of the frontal bones, both parietal bones, the external sagittal crest and the cranial part of the supraoccipital bone (Figure 1). The mass had a heterogeneous aspect with multiple granular, sharply demarcated hyperdense regions. Transcranial extension exerted a mass effect on the brain parenchyma. A CT scan of the thoracic cavity was performed and no pulmonary metastases could be detected. However, partial, positional atelectasis prevented the complete evaluation of all lung lobes.

A core biopsy confirmed the presumed diagnosis of MLO by finding multiple lobules, centered on a core of cartilaginous or bony matrix surrounded by a thin layer of spindle cells (Figure 2). The tumor was classified as grade I based on tumor borders, size of lobules, organization, mitotic figures, cellular pleomorphism and necrosis (Dernell et al., 1998).

Due to the increasing severity of tumor-related behavioral changes and the potential long survival time after surgical resection, the owners opted for surgery. The dog was premedicated intravenously with 0.5 mg/ kg midazolam (Dormicum, Roche B.V., the Netherlands) combined with 0.5 mg/kg methadone (Comfortan, Eurovet Animal Health B.V., the Netherlands). Induction occurred with 4 mg/kg propofol (PropoFlo Plus, Abbott Logistics B.V., the Netherlands) IV, after which a 6-mm endotracheal tube was placed. Maintenance of anesthesia was done with isoflurane (Iso-Flo, Abbott Logistics B.V., the Netherlands) in 100% oxygen through a circle rebreathing system (Cicero, Dräger, Germany). Intraoperative analgesia consisted of a continuous rate infusion of fentanyl (Fentadon, Dechra, Belgium) at 5 μg/kg/h. At the time of the induction of anesthesia, 20 mg/kg cefazolin (Cefazoline, Sandoz N.V., the Netherlands) was administered and repeated every two hours until the end of surgery.

The dog was positioned in sternal recumbency with the head elevated and fixed in a 45-degree angle between head and neck, to avoid intracranial venous congestion that might occur from both jugular compression or positional hypostatic congestion (Otto, 2015) (Figure 3). In order to decrease the risk for tumor recurrence, the overlying skin and a 1-cm margin of surrounding soft tissues and bone, ensuring to include the previous biopsy tract, was resected (Figure 4). Hemostasis was achieved with bipolar electrocoagulation. The temporalis muscles were elevated from the parietal bones and the skull around the mass was cut with a 3-mm high-speed pneumatic burr. The endostium was removed with 2-mm Kerrison rongeurs, after which the entire mass was carefully elevated and dissected from the underlying dura by use of blunt probes (Figure 5). Hemorrhage from the right transverse sinus resulted in a subdural hematoma. To avoid an increase of the intracranial pressure, the dura was incised, the hematoma was evacuated, and the dura was closed with a continuous suture pattern using 5-0 polyglecaprone (Monocryl, Ethicon, Belgium). Gross inspection of the removed mass revealed margins varying from 5 to 10 mm (Figure 6). The excised mass was submitted for histologic evaluation. The results confirmed the diagnosis of a grade-1 MLO and considered the surgical margins free of tumor tissue.

The bone defect was reconstructed by means of a 0.4 mm titanium mesh (MatrixNEURO, Contourable Mesh, DePuy Synthes, Belgium). Briefly, the mesh was molded as a dome overlapping the burred bone edges for 1 cm to allow fixation with 1.5 mm diameter low-profile, self-tapping cortical screws (Figure 7). Since the overlying temporalis muscle was removed, no muscle layer was available to cover the mesh. From the abundant dorsal cervical skin, a single-pedicle advancement flap was elevated and used to cover the mesh and close the skin defect (Figure 8). The underlying cutaneous muscle and deep subcutaneous tissues were closed with 3-0 polyglecaprone in a simple continuous pattern, and the skin was apposed with a subcuticular pattern of 3-0 polyglecaprone.

Postoperative radiographs showed the mesh to be contoured anatomically correct, without any apparent compression of the brain parenchyma (Figure 9). The dog recovered from anesthesia without complications. Postoperative IV analgesics consisted of a continuous rate infusion of lidocaine (Xylocaine, Eurovet Animal Health B.V., the Netherlands) for 2 days at a rate of 30 µg/kg/minute preceded by a single IV loading dose of 2 mg/kg, 10 µg/kg buprenorphine (Vetergesic Multidosis, Patheon UK Limited, United Kingdom) q8h for 11 days, 4 mg/kg carprofen (Rimadyl, Pfizer, Belgium) q24h for 7 days. In addition, 20 mg/kg cefazolin (Cefazolin, Sandoz N.V., the Netherlands) was continued q12h for 11 days. Twenty-four hours after surgery, the dog was quiet and alert but unable to walk due to left-sided hemiparesis. During blinking, the upper eyelids incompletely covered the cornea, for which vitamin A eye ointment (Opticorn A, Ecuphar, Belgium) was instilled q4h. Wound healing occurred without complication, and from the fifth day on, the general condition improved and a progressive improvement of ambulation was noted. On the sev-

Figure 3. Surgical positioning of the dog in sternal recumbency with the head elevated, and the head and neck in a 45-degree angle not to obstruct venous head flow.

enth day, only slight delayed proprioceptive reflexes remained.

Despite the preventive eye ointment, the right eye developed an acute 'melting' corneal ulcer with hypopyon and conus cornea. Initially, atropine sulphate 5 mg/ml eye drops (Atropine, Alfasan B.V., the Netherlands) were instilled q12h and 5 mg/g oxytetracycline + 10.000 I.E./g polymyxin B eye ointment (Terramycin, McNeil Manufacturing, France) q8h. Due to disease progression, there was a risk of corneal perforation on the eight postoperative day, a 360-degrees conjunctival flap was used as a salvage procedure (Pumphrey et al., 2011). Both anesthesia and surgical procedure were without complications. Postoperatively, ophthalmic medication consisted of 3 mg/g ofloxacine (Trafloxal collyre, Baush & Lomb Pharma nv, Belgium) q8h, acetyl cysteine (Lysomucil 10%, Zambon n.v., Belgium) q12h, ketorolac trometamol (Aculare 0.5%, Allergan Pharmaceuticals, Ireland) q6h, and 0.02 mg/kg atropine sulphate (Atropine, Alfasan B.V., the Netherlands) q12h.

On the ninth postoperative day, the dog was bright, alert and ambulating with only mild ataxia. The dog was discharged two days later with continued antibiotics (clavulanic potentiated amoxicillin for 7 days), analgesics (carprofen for 14 days), and local eye medication (ofloxacine, acetyl cysteine, and ketorolac trometamol for 3 weeks).

Eight weeks after surgery, the conjunctival flap dehisced spontaneously, with some remaining mucosa reinforcing the weakened cornea at the ulcer site. The eye medication was discontinued. Six months after surgery, the dog was neurologically normal (ambulation and behavior), the eyelids closed during blinking, and all that remained from the corneal ulcer was some pigmented scar tissue (Figure 10). Lung auscultation was unremarkable, but unfortunately skull and thoracic radiographs were declined by the owner because of financial reasons.

Seven and nine months after surgery, the dog developed a single seizure, which started during rest and was characterized by tonic-clonic convulsions, loss of



Figure 4. A circular incision was made around the base of the tumor, the underlying muscles were incised and the bone was prepared for cranioplasty.

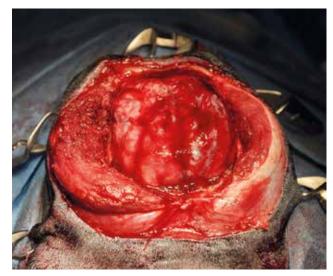


Figure 5. After tumor removal, a large partial craniectomy defect remained that needed reconstruction. A subdural hematoma is seen in the right hemisphere (purple discoloration).



Figure 7. The 0.4-mm titanium mesh (MatrixNEURO, Contourable Mesh, DePuy Synthes, Belgium) was shaped conforming the removed skull piece and fixed with 1.5-mm screws.



Figure 9. Postoperative lateral radiograph showing the titanium mesh fixed on the calvarium after partial craniectomy.

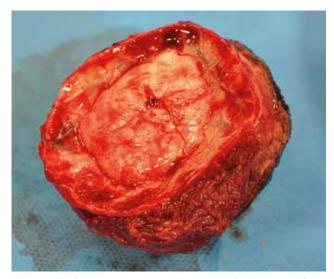


Figure 6. View at the ventral (intracranial) aspect of the tumor. The rim of normal bone present at the craniectomy location determines the macroscopic tumor margins.



Figure 8. From the dorsal neck, a single-pedicle advancement flap consisting of skin and sphincter collimuscle, was mobilized and directed cranially to cover the implant.



Figure 10. Six months after tumor resection, the dog is fully functional. There is some remaining scar tissue visible on the right cornea, but no evidence of local tumor recurrence or metastasis.

consciousness and abnormal autonomous activity. Interictally, the dog behaved normal. On general clinical and neurological examination, no abnormalities were detected. Eleven and thirteen months after surgery, new seizures developed. Complete blood count and serum biochemistry were normal and cardiac ultrasound revealed stable stage B1 mitral endocardiosis. Further diagnostics (magnetic resonance imaging) to search for underlying structural cerebral disorders were offered but unfortunately declined by the owner. The dog was started on 2.5 mg/kg phenobarbital (Phenoleptil, Le Vet B.V., the Netherlands) q12h. At the time of writing, seventeen months after surgery, no further seizures had occurred.

#### **DISCUSSION**

Because of its location, complete surgical removal of MLO is often difficult resulting in local recurrence in about 57% of the cases (McLain et al., 1983; McCalla et al., 1989; Straw et al., 1989; Dernell et al., 1998; Jubb et al., 2007). In a retrospective study on 39 dogs with MLO, complete excision resulted in a disease free interval of 1.332 days versus only 320 days for incomplete removal (Dernell et al., 1998). In addition, the development of metastases is also correlated to the completeness of surgical resection, with the metastatic rate for complete excision being 25% versus 75% for incomplete removal (Dernell et al., 1998). Therefore, aggressive surgical excision with wide margins is the treatment of choice.

In earlier studies on canine cranioplasty, the use of cranial bone allograft (Moissonnier et al., 1997), polymer prostheses (including polytetrafluoroethylene, high-density polyethylene, polyester, polyamide mesh), and polymethylmethacrylate (Moissonnier et al., 1997; Mouatt, 2002; Bryant et al., 2003), and metal plates including platinum, tantalum, cobalt-chromium alloys, aluminum, titanium and stainless steel (Bordelon and Rochat, 2007) have been described.

Polymethylmethacrylate is the most widely used implant material because it is radiolucent, can be easily molded, has acceptable tensile and flexural strength, is widely available, and relatively inexpensive in relation to other implant materials (Moissonnier et al., 1997; Park et al., 2001; Mouatt, 2002; Bryant et al., 2003). Preforming the PMMA using a mold avoids potential for tissue necrosis and creates a custom, lightweight and durable implant (Rosselli et al., 2017). However, potential disadvantages include exothermic reaction associated with polymerization, the brittle nature of the material once it has polymerized, variations in porosity depending on mixing technique, the potential to provide for bacterial colonization (Arens et al., 1996), and the extended duration of surgery caused by the multiple steps for molding (Rosselli et al., 2017). To provide sufficient protection, a thickness of 5-10 mm is recommended (Moissonnier et al., 1997). Thus, in smaller patients and in

patients with defects of the dorsal aspect of the skull, where excessive tension of the overlying soft tissues may lead to necrosis or dehiscence, PMMA may not be the best choice (van Gool, 1985; Moissonnier et al., 1997; Mouatt, 2002; Bryant et al., 2003).

Titanium has been reported to be the most promising of metallic implants (Bordelon and Rochat, 2007). Potential disadvantages include poor malleability, lack of availability and high cost (Moissonnier et al., 1997). Advertised advantages include radiolucency, lower density, corrosion resistance and potentially lower susceptibility to infection (Arens et al., 1996; Rosselli et al., 2017). Bordelon and Rochat (2007) used a 1.5-mm-thick sheet of rigid titanium mesh for a cranioplasty and concluded the benefits in terms of fit, strength, low profile and secure fixation to the remaining calvarium outweighed the costs. The contourable 0.4-mm titanium mesh used in this case is a technologically improved version of the previously described titanium implant, which is even more malleable, yet maintains substantial rigidity due to its dome shape. In a case series of five dogs, Rosselli et al. (2017) demonstrated the titanium mesh did not interfere with postoperative CT or MR images, thereby allowing postoperative imaging for ongoing assessment.

Postoperative radiographs confirmed the correct placement and anatomical contouring of the titanium mesh over the skull defect. The titanium mesh in this case report was secured to the skull with screws. No complications were recorded during or after the procedure attributable to the titanium implant. In previous case series, the use of sutures to secure the mesh to the skull has been documented, claiming that the use of screws could prolong and increase the cost of surgery, create iatrogenic damage to underlying brain parenchyma, and might be associated with postoperative loosening or migration (Rosselli et al., 2017). The screws used in this case report were low-profile, self-tapping screws that were placed monocortically, thereby eliminating the aforementioned potential disadvantages. Follow-up radiographs at seventeen months after surgery showed no signs of implant loosening or migration. Screw fixation to the skull offers a more rigid fixation of the implant than fixing it to overlying soft tissues. Furthermore, suturing the mesh to the overlying temporalis fascia might work for smaller reconstructions, but may not be applicable when there is insufficient overlying muscle or fascia.

Because MLO's are considered to be locally invasive tumors, the overlying skin, biopsy tract and surrounding muscles were removed together with the tumor. A wide deep resection was not necessary since this tumor does not usually invade across the meninges. Other authors have removed MLO's with a linear skin incision and the conservation of superficial muscle layers (Bordelon and Rochat, 2007; Gallegos et al., 2008). Unfortunately, due to the size of the tumor, the location on the sagittal crest and the presence of a biopsy tract, conservation of either skin or muscles would endanger complete surgical resection.

A single pedicle skin flap that advances skin from the dorsal cervical region was used to close the skin defect. The incorporated sphincter colli and cutaneous muscle allowed to cover the mesh with a muscular layer. However, positional differences between the head in the surgical field and in the awake dog may result in undue traction when using this type of skin flap. A ninety-degree rotation flap would have increased surgical morbidity, but could have provided skin for closure without any traction on the upper eyelids (Smith et al., 1991).

Both CT and MR imaging were offered postoperatively for ongoing assessment. Rosselli et al. (2017) have shown there is no interference of image quality using a titanium mesh. Since elemental titanium has a low molecular number, it is minimally attenuating and since it is non-ferrous, it does not create magnetic field inhomogeneities.

The description of the dog of the present study was different from that of previous reviews and case reports (Straw et al., 1989; Groff et al., 1992; Dernell et al., 1998; Loukopoulos et al., 2003; Banks and Straw, 2004; Jubb et al., 2007; Webb et al., 2009; Leonardi et al., 2014). Although multilobular osteochondrosarcoma appears to be a disease of middle-aged to older, medium- to large-breed dogs, scarce reports of young and small dogs do exist (Jacobson, 1971; Diamond et al., 1980; Fukui and Takamori, 1986; Pakhrin et al., 2006; Bordelon and Rochat, 2007).

The dog's exophthalmos was originally attributed to breed-related conformation. But the combination of neurologic changes (behavioral) and ocular changes (exophthalmia) with intracranial tumor growth has been described before (Pletcher et al., 1979; Straw et al., 1989; Pakhrin et al., 2006; Psychas et al., 2009; Leonardi et al., 2014). In a review study on 39 dogs, 5% had neurologic symptoms and 15% exophthalmia (Dernell et al., 1998). It is therefore possible that the presurgical exophthalmia was, at least partially, tumor-induced.

According to some studies, elevated levels of alkaline phosphatase are a negative prognostic factor for osteosarcoma (Ehrhart et al., 1998; Garzotto et al., 2000). For MLO, such a correlation has not been established. Earlier reports have also found elevated levels of alkaline phosphatase and have linked this to tumor-induced increased bone metabolism (Gallegos et al., 2008).

Skull radiographs can easily differentiate between osteoma and MLO (Ling et al., 1974). However, some MLO's don't display typical radiographic characteristics and in these patients, a preoperative histological biopsy is necessary for accurate diagnosis (Stoll et al., 2000). In the current case, the diagnosis was made based on the radiographs, but the biopsy was performed to establish a tumor grade. Dernell et al. (1998) found that the tumor grade is not only correlated

to local recurrence (30% for grade I MLO, 47% for grade II MLO, and 78% for grade III MLO), but also to the presence of metastasis (30% for grade I MLO, 60% for grade II MLO, and 78% for grade II MLO). The identification of a low-grade MLO in the present case offered the owner a more informed decision on the expected course after surgical therapy.

Risk factors for postoperative wound infection are the presence of a metallic implant and a surgical duration longer than two hours (Brown et al., 1997). Prolonged antibiotic therapy is, however, essential when there is a communication between the frontal sinus and mesh implant (Moissonnier et al., 1997). In the present case, where no communication with the frontal sinus existed and where a metallic implant was used that is less susceptible to harboring bacteria, antibiotic therapy was pursued for the postoperative corneal ulcer development and the pending risk for eye perforation.

The role of chemotherapy and radiation therapy in the management of MLO is not well defined. No increase in survival time, time to local recurrence and metastasis have been found between the surgical excision alone and the surgical excision followed by adjuvant therapy (Straw et al., 1989; Dernell et al., 1998; Koch et al., 2000). However, adjuvant therapy may provide some benefit for local tumor control or palliation of unresectable or recurrent lesions (Boston, 2010).

Idiopathic epilepsy was considered very unlikely in this case as the dog developed seizures at the age of eleven years, whereas in most of the dogs with idiopathic epilepsy, the seizure onset is between the age of six months and six years (De Risio et al., 2015). Metabolic and toxic causes of the seizures were also considered unlikely as the blood examination was normal, and no history of toxin exposure was mentioned in the anamnesis. In this case, the possibility of a structural cause for the seizures seemed most likely (De Risio et al., 2015). Taking into account the previous medical issues in this dog, regrowth of the skull tumor or the development of scar tissue at the surgery site seem the two most likely possibilities (De Risio et al., 2015). However, other structural causes, such as neoplasm, inflammation and vascular disorders, should be taken into account as well (De Risio et al., 2015).

#### **CONCLUSION**

Despite the extensiveness of the process in the skull and the compression of intracranial structures, a dog with MLO may reach a prolonged survival time if the tumor has been removed with complete margins. The titanium mesh used in this case offers a smooth, fast and uncomplicated method to reconstruct the cranium of a dog.

#### **REFERENCES**

- Arens S., Schlegel U., Printzen G., Ziegler W.J., Perren S.M., Hansis M. (1996). Influence of materials for fixation implants on local infection. An experimental study of steel versus titanium DCP in rabbits. *Journal of Bone and Joint Surgery 78 B*, 647-651.
- Banks T.A., Straw R.C. (2004). Multilobular osteochondrosarcoma of the hard palate in a dog. *Australian Veterinary Journal* 82, 409-412.
- Bordelon J.T., Rochat M.C. (2007). Use of a titanium mesh for cranioplasty following radical rostrotentorial craniectomy to remove an ossifying fibroma in a dog. *Journal of the American Veterinary Medical Association 231*, 1692-1695
- Boston S.E. (2010). Craniectomy and orbitectomy in dogs and cats. *Canadian Veterinary Journal* 51, 537-540.
- Bhatti SF, De Risio L, Muñana K, Penderis J, Stein VM, Tipold A, Berendt M, Farquhar RG, Fischer A, Long S, Löscher W, Mandigers PJ, Matiasek K, Pakozdy A, Patterson EE, Platt S, Podell M, Potschka H, Rusbridge C, Volk HA. (2015). International veterinary epilepsy task force consensus proposal: medical treatment of canine epilepsy in Europe. *BMC Veterinary Research* 28, 176-192.
- Brown D.C., Conzemius M.G., Shofer F., Swann H. (1997). Epidemiologic evaluation of postoperative wound infections in dogs and cats. *Journal of the American Veterinary Medical Association* 210, 1302-1306.
- Bryant K.J., Steinberg H., McAnulty J.F. (2003). Cranioplasty by means of molded polymethylmethacrylate prosthetic reconstruction after radical excision of neoplasms of the skull in two dogs. *Journal of the American Veterinary Medical Association 223*, 67-72.
- Dernell W.S., Straw R.C., Cooper M.F., Powers B.E., LaRue S.M., Withrow S.J. (1998). Multilobular osteo-chondrosarcoma in 39 dogs: 1979-1993. *Journal of the American Animal Hospital Association 34*, 11-18.
- DePuy Synthes Companies (2015). MatrixNEURO. The next generation cranial plating system. Internetreferentie: <a href="http://emea.depuysynthes.com/hcp/cmf/products/qs/matrixneuro-system">http://emea.depuysynthes.com/hcp/cmf/products/qs/matrixneuro-system</a>
- De Risio L, Bhatti S, Muñana K, Penderis J, Stein V, Tipold A, Berendt M, Farqhuar R, Fischer, A, Long S, Mandigers PJ, Matiasek K, Packer RM, Pakozdy A, Patterson N, Platt S, Podell M, Potschka H, Batlle MP, Rusbridge C, Volk HA. (2015). International veterinary epilepsy task force consensus proposal: diagnostic approach to epilepsy in dogs. *BMC Veterinary Research* 28, 148-159.
- Diamond S.S., Raflo C.P., Anderson M.P. (1980). Multilobular osteosarcoma in the dog. *Veterinary Pathology* 17, 759-780.
- Ehrhart N., Dernell W.S., Hoffman W.E., Weigel R.M., Powers B.E., Withrow S.J. (1998). Prognostic importance of alkaline phosphatase in serum from dogs with appendicular osteosarcoma: 75 cases (1990-1996). *Journal of the American Veterinary Medical Association 213*, 1002-1006.
- Forrest L.J. (2013). The Cranial and Nasal Cavities: Canine and Feline. Multilobular Osteochondrosarcoma. In: Thrall D.E. (editor). *Veterinary Diagnostic Radiology*. Elsevier Saunders, Missouri, p. 121.
- Fukui K., Takamori Y. (1986). Multilobular osteoma (chondroma rodens) in a Pekingnese. *Veterinary Record* 118, 483.

- Gallegos J., Schwarz T., McAnulty J.F. (2008). Massive midline occipitotemporal resection of the skull for treatment of multilobular osteochondrosarcoma in two dogs. *Journal of the American Veterinary Medical Association* 233, 752-757.
- Garzotto C.K., Berg J., Hoffmann W.E., Rand W.M. (2000) Prognostic significance of serum alkaline phosphatase activity in canine appendicular osteosarcoma. *Journal of Veterinary Internal Medicine* 14, 587-92.
- Groff J.M., Murphy C.J., Pool R.R., Koblik P., Bellhorn R. (1992). Orbital multilobular tumour of bone in a dog. *Journal of Small Animal Practice* 33, 597-600.
- Jubb K.V.F., Kennedy P.C., Palmer N. (2007). Bone and joints. In: *Pathology of Domestic Animals*. Fifth edition, Volume 1, Elsevier, St. Louis, p. 119-120.
- Koch B.B., Karnell L.H., Hoffman H.T., Apostolakis L.W., Robinson R.A., Zhen W., Menck H.R. (2000). National cancer database report on chondrosarcoma of the head and neck. *Head Neck* 22, 408-425.
- Leonardi L., Carrano A., Stoppini L., Floris M. (2014). Multilobular tumor of the zygomatic bone in a dog. *Open Veterinary Journal* 4, 9-11.
- Ling G.V., Morgan J.P., Pool R.R. (1974). Primary bone tumors in the dog: a combined clinical, radiographic, and histologic approach to early diagnosis. *Journal of the American Veterinary Medical Association* 165, 55-67.
- Loukopoulos P., Thornton J.R., Robinson W.F. (2003). Clinical and pathologic relevance of p53 index in canine osseus tumors. *Veterinary Pathology* 40, 237-248.
- McCalla T.L., Moore C.P., Turk J., Collier L.L., Pope E.R. (1989). Multilobular osteosarcoma of mandible and orbit in a dog. *Veterinary Pathology* 26, 92-94.
- McGavin M.D., Zachary J.F. (2007). *Pathologic Basis of Veterinary Disease*. Fourth edition, Elsevier, Mosby, United States, p. 1091.
- McLain D.L., Hill J.R., Pulley L.T. (1983). Multilobular osteoma and chondroma (chondroma rodens) with pulmonary metastasis in a dog. *Journal of the American Animal Hospital Association* 19, 359-362.
- Moissonnier P., Devauchelle P., Delisle F. (1997). Cranioplasty after en bloc resection of calvarial chondroma rodens in two dogs. *Journal of Small Animal Practice* 38, 358-363.
- Mouatt J.G. (2002). Acrylic cranioplasty and axial pattern flap following calvarial and cerebral mass excision in a dog. *Australian Veterinary Journal* 80, 211-215.
- Otto K.A. (2015). Physiology, Pathophysiology and Anesthetic Management of Patients with Neurological Disease. In: Grimm K.A., Lamont L.A., Tranquilli W.J., Greene S.A. en Robertson S.A. (editors). *Veterinary Anesthesia and Analgesia*, Fifth edition of Lumb and Jones, Wiley Blackwell, Iowa, p. 574-582.
- Pakhrin B., Bae I.H., Jee H., Kang M.S., Kim D.Y. (2006). Multilobular tumor of the mandible in a Pekingese dog. *Journal of Veterinary Science* 7, 297-298.
- Park H.K., Dujovny M., Agner C., Diaz F.G. (2001). Biomechanical properties of calvarium prosthesis. *Neurological Research* 23, 267-276.
- Pletcher J.M., Koch S.A., Stedhem M.A. (1979). Orbital chondroma rodens in a dog. *Journal of the American Veterinary Medical Association* 175, 187-190.
- Pool R.R. (1990). Tumors of bone and cartilage. In: Moulton J.E. (editor). *Tumors of Domestic Animals*. Third edition, University of California Press, Berkeley, p. 159-213.
- Popvitch C.A., Weinstein M.J., Goldschmidt M.H., Shofer

- F.S. (1994). Chondrosarcoma: a retrospective study of 97 dogs (1987-1990). *Journal of the American Animal Hospital Association 30*, 81-85.
- Psychas V., Loukopoulos P., Polizopoulou Z.S., Sofianidis G. (2009). Multilobular tumour of the caudal cranium causing severe cerebral and cerebellar compression in a dog. *Journal of Veterinary Science* 10, 81-83.
- Pumphrey S.A., Pizzirani S., Pirie C.G. (2011) 360-degree conjunctival grafting for management of diffuse keratomalacia in a dog. *Veterinary Ophthalmology* 14, 209-213.
- Rosselli D.D., Platt S.R., Freeman C., O'Neill J., Kent M., Holmes S.P. (2017). Cranioplasty using titanium mesh after skull tumor resection in five dogs. *Veterinary Surgery* 46, 67-74.
- Selcer B.A., McCracken M.D. (1981). Chondroma rodens in dogs: a report of two case histories and a review of the veterinary literature. *Journal of Veterinary Orthopedics* 2, 7-11.

- Smith M.M., Payne J.T., Moon M.L., Freeman L.E. (1991). Axial pattern flap based on the caudal auricular artery in dogs. *American Journal of Veterinary Research* 52, 922-925.
- Stoll M.R., Roush J.K., Moisan P.G. (2001). Multilobular tumour of bone with no abnormalities on plain radiography in a dog. *Journal of Small Animal Practice* 42, 453-455
- Straw R.C., LeCouteur R.A., Powers B.E., Withrow S.J. (1989). Multilobular osteochondrosarcoma of the canine skull: 16 cases (1978-1988). *American Veterinary Medical Association* 195, 1764-1769.
- van Gool A.V. (1985). Preformed polymethylmethacrylate cranioplasties: report of 45 cases. *Journal of Maxillofacial Surgery 13*, 2-8.
- Webb J.A., Liptak J.M., Hewitt S.A., Vince A.R. (2009). Multilobular osteochondrosarcoma of the os penis in a dog. *The Canadian Veterinary Journal* 50, 81-84.



# World Veterinary Poultry Association WVPA - Belgian Branch

WVPA België organiseert jaarlijks verschillende studienamiddagen omtrent recente problematiek bij industrieel gehouden pluimvee. De thema's van de studienamiddagen spelen in op de huidige problematiek in de pluimveesector en proberen tegemoet te komen aan de verwachtingen van de pluimveedierenarts.

Thema's die tijdens het academiejaar 2017-2018 aan bod zullen komen, zijn: persisterende Salmonella infecties, drinkwaterkwaliteit en –veiligheid, pluimveewelzijn en de registratiewetgeving omtrent het gebruik van antimicrobiële producten in de pluimveesector.

Deelname aan het programma is gratis voor leden. Graag nodigen wij u uit om uw lidmaatschap te verlengen. Alle informatie over de geplande studienamiddagen, het lidgeld alsook het rekeningnummer waarop het lidgeld kan gestort worden, kan u vinden op de website **www.wvpa.be**.

Met vriendelijke groeten, WVPA België