Antebrachial chondrodysplasia in New Zealand white rabbits (Oryctolagus cuniculus)

T Pulker\(^a\), A Carstens\(^b\) and J Williams\(^c\)

**ABSTRACT**

Rabbits obtained from a South African rabbit breeder exhibited deformities of the distal forelimbs. The clinical, radiological and histological presentation of mid-antebrachial valgus formation (a.k.a distal foreleg curvature) in these rabbits was consistent with an autosomal recessive trait associated with a chondrodystrophic lesion of the distal ulna epiphysis 1st described in the 1960s. The impact this might have on South African farming enterprises and laboratory facilities has not been established, but the heritability and welfare implications of this condition make it a concern. Mildly affected animals can adapt to the deformity with some housing adjustments, but severely affected animals may require humane euthanasia.

**Keywords:** antebrachium, chondrodysplasia, hereditary conditions, Oryctolagus cuniculus, rabbit, valgus.


**INTRODUCTION**

The use of rabbits in South Africa is confined mainly to farming enterprises and laboratory animal centres. Low-cost protein sources are constantly being sought to guarantee food security and rabbit meat is 1 of the avenues being explored. Diseases in a rabitry can have profound effects on meat quality and economic viability. Pathological lesions observed during meat inspections in northern Italy abattoir rabbits mainly affected the integument, digestive and urinary systems\(^a\). Non-traumatic developmental musculoskeletal disorders are not commonly detected in abattoir rabbits during the slaughter process\(^b\). This may in part be due to the fact that many of the dysplastic conditions in rabbits are lethal mutations, associated with low survival rates. Production losses can be expected due to the decrease in progeny reaching slaughter age\(^c\). However, 1 form of chondrodysplasia in rabbits, known as distal foreleg curvature, is not associated with a change in growth rate or survival\(^d\), but these animals should be removed from the breeding stock to prevent dissemination of this mutation.

In a laboratory setting, the presence of ‘silent’ conditions can have varied or even unknown effects on research results. Although non-traumatic developmental musculoskeletal disorders in rabbits are not often diagnosed, these disorders have the potential to compromise the animal’s well-being. Once recognised, various measures can be implemented to counteract this negative impact such as non-slippery flooring, improved access to food, water and shelter and regular monitoring.

**CASE HISTORY**

Nine of 18 approximately 3-month-old female New Zealand white rabbits, obtained from a local rabbit breeder and housed at the University of Pretoria Biomedical Research Centre (UPBRC), had varying degrees of lateral deviation of the forelimbs from the mid ulna–radius diaphyses distally, with the manus directed laterally away from the body. Three of the 9 animals were unilaterally affected on visual inspection. Physical examination of these animals revealed no clinical abnormalities apart from the deviated forelimbs associated with difficulty in adducting these limbs.

Radiographic findings of the forelimbs of 2 of these animals were suggestive of some form of thoracic limb chondrodysplasia. Additional radiographs were recommended, including an unaffected rabbit to serve as a control.

Four of the bilaterally affected rabbits (including the initial 2 animals sent for radiographs) and 1 unaffected rabbit were radiographed at the Onderstepoort Veterinary Academic Hospital, Department of Diagnostic Imaging, after intramuscular sedation with 35 mg/kg ketamine (Anaket-V, Bayer) and 5 mg/kg xylazine (Chanazin 2 %, Bayer). Both cranio-caudal and mediolateral views of the antebrachiae, elbows and manus were taken. Cranio-caudal views of the tibia, and distal and mediolateral views of the femur and distal views of both hind limbs were taken of the 2 affected animals and the unaffected animal.

Owing to severe locomotor deficits 8 months later 1 of the bilaterally affected rabbits was euthanased on humane grounds with 200 mg/kg sodium pentobarbitone intravenously (Euthapent, Kyron Laboratories) and necropsied.

**RESULTS**

Radiographic findings of the 4 affected animals were similar. Mild cranial (143–160°) and moderate medial (131–148°) bowing of the mid-radial and, to a similar extent, mid-ulnar diaphyses was noted in all affected animals with no indication of metabolic bone disease (Fig. 1A,B). These deformities were not evident on the unaffected animal’s radiographs (Fig. 2). The changes were restricted to the radius and ulna, as no radiographic changes indicative of chondrodysplasia were discovered on hind-limb and humeral views. Mild to moderate elbow degenerative joint disease was observed in all 5 radiographed animals. As similar changes were noted in the unaffected animal, this was considered an incidental finding. Bilateral stifle degenerative joint disease was evident in all 3 rabbits that had hind-limb radiographs taken. Two of these animals had bowing of the antebrachium and the 3rd had a normal antebrachium.

The only abnormality found macroscopically during necropsy examination of 1 of the affected rabbits was the bilaterally symmetrical bowing and shortening of the radii/ulnae. These were fixed in
formalin in toto after which they were placed in 8% formic acid, which was replaced every 4–5 days, to decalcify the bones. Only 0.5 cm of the distal and proximal radii/ulnae could be sectioned after 2 weeks of decalcification, suggesting normal mineralisation of these bones. Histologically these sections showed healthy bone and periosteum with medullary cavities containing healthy bone marrow.

At the time of diagnosis, the mean weight of the affected rabbits was 4.3 ± 0.33 kg while the unaffected animals had a mean weight of 4.69 ± 0.43 kg. Upon implementation of a restricted diet, for weight control reasons, 3 months later the mean weight for the affected animals was 4.88 ± 0.4 kg and 5.31 ± 0.39 kg for the unaffected animals. Over the rest of the year, the mean weights for the animals fluctuated between 4.6–4.8 kg (affected) and 4.7–4.9 kg (unaffected).

**DISCUSSION**

Skeletal dysplasias are a group of defects associated with short limbs, abnormal bone shape and/or increased bone fragility, which primarily affect bone formation or remodelling. Despite the formulation of an extensive classification system for human skeletal dysplasias, a complete system for the corresponding animal conditions has not been reported. Skeletal dysplasias are a complex and diverse group of abnormalities. A brief overview of the recognised conditions in rabbits is summarised in Table 1.

Animal skeletal dysplasias are often propagated by inbreeding or excessive use of an animal carrying the relevant gene. It is, however, important to exclude non-genetic causes of dysplasia before condemning a stud animal. For the rabbits in question it was not possible to rule out infectious agents, toxin exposure or mineral deficiencies that could result in metabolic bone disease during gestation.

Table 1: Recognised gene mutations resulting in skeletal dysplasia in rabbits.

<table>
<thead>
<tr>
<th>Alleles</th>
<th>Description</th>
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<tr>
<td>Achondroplasia</td>
<td>ac/ac Disproportionate dwarf Lethal mutation</td>
</tr>
<tr>
<td>Chondrodystrophy</td>
<td>cd/cd Disproportionate dwarf Lethal mutation</td>
</tr>
<tr>
<td>Dachs</td>
<td>Da/Da or Da/da Viable chondrodystrophic dwarf Non-lethal mutation</td>
</tr>
<tr>
<td>Distal foreleg curvature</td>
<td>fc/fc Foreleg deformities Non-lethal autosomal recessive mutation</td>
</tr>
<tr>
<td>Dwarf</td>
<td>Dw/Dw Pituitary/proportionate dwarfism Semi dominant lethal mutation</td>
</tr>
<tr>
<td>Dwarf</td>
<td>nan/nan Proportionate dwarf-nansomia May be the same as Dw</td>
</tr>
<tr>
<td>Dwarf</td>
<td>zw/zw Proportionate dwarf (zwergwuchs) May be the same as Dw</td>
</tr>
<tr>
<td>Pelger</td>
<td>Pg/Pg Chondrodystrophic dwarf, leukocytes affected Semi-dominant mutation</td>
</tr>
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and early development, as the animals were only obtained in early adulthood. These causes were considered unlikely, however, owing to a lack of further clinical abnormalities and the localised nature of the lesion.

A relatively common clinical manifestation in rabbits, referred to as ‘splay leg’, presents as the inability to adduct any or all of the limbs. Splay leg can be used to describe the physical posture of mutants with syringomyelia, hypoplasia pelvis, femoral luxation, hereditary distal foreleg curvature or achondroplasia restricted to shoulder and hip joints. It has also been used to describe young animals raised on slippery surfaces that present with the same inability to adduct the limbs. Usually the latter cases can be corrected by placing the animal on non-slippery floors. Two of the affected rabbits were housed together on plastic mesh flooring. It has been described in young animals raised on slippery floors. The latter cases can be corrected by placing the animal on non-slippery floors. Two of the affected rabbits were housed together on plastic mesh flooring. Although this non-slippery flooring improved mobility, the adduction problem was not resolved.

Radiographic findings suggested chondrodysplasia. This is a form of generalised skeletal dysplasia defined as a defect in or disordered development of cartilage. The clinical, radiological and histological findings of these 4 affected rabbits correspond with lesions described in Beveren, Dutch, Belgian and French Silver rabbits in the 1960s. The cause of the deformity in these breeds was found to be a chondrodystrophic lesion in the distal ulna epiphysis. The lesion appeared from 2 weeks of age and progressed until approximately 2 months, from which resolution of the lesion by approximately 2 months, from which time regression occurred rapidly with complete radiographic and histological resolution of the lesion by approximately 4 months of age. The only indication of the condition after this was the permanent deformity of the distal forelegs. Animals displaying this deformity had no other clinical abnormalities; body growth and physical condition remained unchanged. This correlates with the findings that the 4 affected animals had no other clinical abnormalities and were in good physical condition. The radiographs were taken when the rabbits were over 4 months of age once the lesion had healed and only the permanent deformity could be visualised. The rabbit sent for necropsy was over a year of age at the time of necropsy, hence the lack of histological deformities in the radius and ulna. Although no significant difference in weights between the affected and unaffected animals could be detected and the growth of the animals was similar, the affected animals were slightly lighter than the unaffected animals, indicating potentially lower meat production from these animals.

This distal ulna epiphyseal lesion has been described in a comprehensive overview. It was found to be an interruption of normal endochondral ossification, resulting in persistent chondrocytes that continued to multiply. Without degradation of the cartilaginous matrix, an increasingly broad cartilaginous zone developed, delaying calcification. This irregular cartilage plate during the progressive stage of bone growth gave rise to the bow shape of the ulna shaft. As a firm anatomical connection exists between the radius and ulna, the curvature of the radius was secondary to the ulna deformity. Similarly, the positional changes of the paw and carpus were secondary to the ulna lesion. No other skeletal changes or signs of dwarfism were described. Various breeding tests indicated an autosomal recessive mode of inheritance. Serum and bone calcium, phosphorus and haematological examinations were performed as well as a comprehensive study of the lesion progression both radiologically and histologically.

As this is to the authors’ knowledge the 1st report of chondrodysplasia in New Zealand white rabbits in South Africa, the importance and prevalence of this condition are not known. Even though this condition is unlikely to have a major effect on meat production, there are welfare implications that need to be considered in both farming enterprises and laboratory facilities. Heritability is a concern in specified pathogen-free breeding establishments in South Africa owing to the diminishing genetic pool and difficulty in importing new genetic material. Mildly affected animals are generally able to compensate for the deformities with a few housing adjustments and regular welfare monitoring. Severely affected animals may require humane euthanasia.

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REFERENCES