

Syndactyly in a litter of cats

In this case report, we describe the clinical and radiographic features of a litter of kittens affected with complex syndactyly. We also provide guidelines for the diagnosis, possible treatment and prevention of propagation of this condition. This is the first report of syndactyly in a litter of kittens and syndactyly affecting both the pectoral and pelvic limbs.

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INTRODUCTION

The kittens described in this case report were diagnosed with syndactyly, a dysostosis characterised by a congenital lack of differentiation between two or more digits (Swanson 1976, Ger 1998, Kozin 2003). Syndactyly results when the mesenchyme between radial swellings of the hand or footplate of the developing extremity fails to break down. As a result, there is incomplete separation of skin, fibrous tissue and/or bone in utero (Noden and de Lahunta 1985, Ogden and Grogen 1987, Gilbert 2003).

In human beings, syndactyly is classified as simple or complex, as complete or incomplete, and as complicated or uncomplicated (Ger 1998, Kozin 2003, Dao and others 2004). In simple syndactyly, the abnormal interconnection between adjacent digits consists only of skin and fibrous tissue, whereas in complex syndactyly there is a lack of separation of adjacent bone(s), in addition to skin and fibrous tissue deformities. Syndactyly is called complete when adjacent digits (phalanges 1 to 3) are connected throughout their entire length with soft tissue and/or bone and incomplete when they are not connected throughout their entire length (Ger 1998, Kozin 2003, Dao and others 2004). All cases in the veterinary literature with complex syndactyly have a complete simple syndactyly. Complex syndactyly is further classified as complicated or uncomplicated. Complicated complex syndactyly is associated with other anomalies whereas uncomplicated complex syndactyly is not associated with other anomalies (Kozin 2003). This classification has also been used for syndactyly feline and canine patients (Tables 1 and 2) (Towle and Breur 2004).

The incidence of syndactyly in human beings is approximately two or three per 10,000 live births (Kozin 2003). The mode of transmission of familial syndactyly is considered to be autosomal dominant with variable expressivity and incomplete penetrance. This means that syndactyly in human beings is propagated in family lines, may not be present in full form in each patient (phenotypic variation) and may skip a generation (Kozin 2003). It is also associated with defects of the second chromosome (Kozin 2003). Syndactyly in human beings has been associated with a mutation of the homeobox D cluster-13 (*Hoxd-13*) gene, fibroblast growth factor receptors, ligand Wnt-1 and transcription factor Msx-2 (Daluisi and others 2001, Arias and Stewart 2002, Gilbert 2003). In human beings, complicated complex syndactyly is seen in association with symbrachydactyly (Poland syndrome), constriction bands and acrocephalosyndactyly (Apert syndrome) (Kozin 2003).

Information about syndactyly in the veterinary literature is sparse. To the authors' knowledge, all canine (Baum 1889, Gehring and Schröder 1961, Riser 1964, Leipold and Guffy 1973, Dallman and Brown 1980, Carrig and others 1981, Renoy and Balligand 1991, Richardson and others 1994, Schultz and Watson 1995) and feline (Howe 1902, Hays 1917, Boehringer 1975) reports of syndactyly are limited to isolated cases with the exception of one case report describing complicated complex syndactyly in a cat (Searle 1953) and one case report describing complicated complex syndactyly in a family of Australian shepherd dogs (Freeman and others 1988; Table 2). The incidence, aetiology and prognosis of feline syndactyly are unknown. In this case report, we will describe a litter of kittens with syndactyly and review clinical signs, treatment, prognosis, aetiology and possible mode of inheritance of feline syndactyly.

CASE HISTORY

Six kittens from a single litter were presented to the Purdue University Veterinary Teaching Hospital (PUVTH) for evaluation of paw deformities on multiple limbs.

Table 1. Nomenclature of syndactyly

Simple syndactyly	Abnormal interconnection between adjacent digits consisting only of skin and fibrous tissue
Complex syndactyly	Lack of separation of adjacent bone(s), in addition to skin and fibrous tissue deformities
Incomplete syndactyly	Adjacent digits (phalanges 1 to 3) are not connected throughout their entire length with soft tissue and/or bone
Complete syndactyly	Adjacent digits (phalanges 1 to 3) are connected throughout their entire length with soft tissue and/or bone
Uncomplicated syndactyly	Syndactyly not associated with other abnormalities
Complicated syndactyly	Syndactyly associated with other abnormalities

The queen was obtained from the humane society by its foster owner (L. R. T.). The queen's gestation and vaccination history were unknown. The queen, a domestic shorthair, did not have any gross skeletal deformities. The foster owner (L. R. T.) immediately noticed the paw deformities after the kittens were born.

The kittens were approximately five weeks old upon initial presentation to PUVTH. There were two female and four male kittens. Four of the six kittens (three males and one female) in the litter were affected. Physical examination was unre-

markable and no other abnormalities besides the manus and pes deformities were detected. Physical examination findings consisted of abnormal or absent digits, abnormal shape, fusion or absence of digital pads and abnormal shape or fusion of nails. No lameness was detected in any of the kittens during the initial visit. Radiographs of affected limbs showed varying degrees of complex syndactyly. Kitten 1 had syndactyly of all four feet (Table 3 and Fig 1). Kittens 2, 3 and 6 had pelvic limb syndactyly (Table 3 and Figs 2 to 4).

Table 2. Classification of syndactyly

	Feline references	Canine references
Simple		
Complete		● Richardson and others (1994)
Incomplete		
Complex		
Complete		
Complicated		● Schultz and Watson (1995) (partial terminal transverse, hemimelia and an anomalous eighth lumbar vertebra)
Uncomplicated	● Hays (1917) ● Boehringer (1975) ● Present study	● Riser (1964) ● Leipold and Guffy (1973) ● Dallman and Brown (1980) ● Renoy and Balligand (1991)
Incomplete		
Complicated	● Howe (1902)* (polydactyly) ● Searle (1953) (ectrodactyly)	● Gehring and Schröder (1961) (oligodontia, cheiloschisis) ● Carrig and others (1981) (ectrodactyly) ● Freeman and others (1988) (cleft palate, polydactyly, shortened tibia-fibula, brachygnathism, scoliosis)
Uncomplicated	● Hays (1917) ● Present study	● Baum (1889)

*Not reported in detail

One kitten died before the eight-week re-evaluation appointment. Gross post-mortem findings included mesenteric lymphadenopathy and pneumonia. Virology was negative for feline panleucopenia, coronavirus and feline herpes via the fluorescent antibody test. Calici virus was isolated from the spleen and tongue via virus isolation assay. The ultimate cause of death was not determined. Physical examination for the remaining kittens was normal during the second appointment and no lameness was noted during observation.

Kitten 1 had complex incomplete uncomplicated syndactyly of both the pectoral limbs and pelvic limbs. Orthopaedic examination and radiographs determined that kitten 1 (Fig 1) had incomplete separation of the third phalanx of the second and third digits of the right pectoral limb, incomplete separation of the third and second phalanges of the second and third digits of the left pectoral limb, and incomplete separation of the first to the third phalanges of the second to fourth digits of the right and left pelvic limbs. Both pectoral limbs and the right pelvic limb had only three digital pads; the remaining pads were normal. The digital pads of the left pelvic limb were fused into one pad. In addition, two nails were present in the pectoral limbs and right pelvic limb of the affected digits, whereas in the left pelvic limb, there was one large nail for the digits 2 to 4.

Kitten 2 had complex incomplete uncomplicated syndactyly of both pelvic limbs. Orthopaedic examination and radiographs confirmed that kitten 2 had normal pectoral limbs and demonstrated incomplete separation of the third and second phalanges of digits 3 and 4 (Fig 2). The right pelvic limb had three digital pads; the remaining pads were normal. The left pelvic limb had two digital pads; the remaining pads were normal. In addition, two nails were present in each of the affected digits.

Kitten 3 had complex incomplete uncomplicated syndactyly of the right pelvic limb and complex complete uncomplicated syndactyly of the left pelvic limb. Orthopaedic examination and radiographs confirmed that kitten 3 had normal pectoral limbs, incomplete separation of

Table 3. Syndactyly deformities in the litter of kittens

	Classification and description of affected digits			
	Right pectoral limb	Left pectoral limb	Right pelvic limb	Left pelvic limb
Kitten 1	Complex/Incomplete P3 of digits 2 to 3	Complex/Incomplete P2 and P3 of digits 2 to 3	Complex/Incomplete P1 to P3 of digits 2 to 4	Complex/Incomplete P1 to P3 of digits 2 to 4
Kitten 2	Normal	Normal	Complex/Incomplete P2 and P3 of digits 3 to 4	Complex/Incomplete P2 and P3 of digits 3 to 4
Kitten 3	Normal	Normal	Complex/Incomplete P1 to P3 of digits 2 to 3	Complex/Complete P1 to P3 of digits 2 to 3
Kitten 4	Normal	Normal	Normal	Normal
Kitten 5	Normal	Normal	Normal	Normal
Kitten 6 (died prematurely)	Normal	Normal	Complex/Incomplete P2 to P3 of digits 3 to 4	Complex/Incomplete P1 to P3 of digits 3 to 4

the first to third phalanges of digits 2 and 3 of the right pelvic limb, and complete fusion of the first to third phalanges of digits 2 and 3 of the left pelvic limb (Fig 3). Each pelvic limb had three digital pads;

the remaining pads were normal. In addition, two nails were present in the right pectoral limb for the fused digits and one large nail was present in the left pelvic limb for the affected digits.

Kittens 4 and 5 had normal pectoral and pelvic limbs upon physical and radiographic examination.

Kitten 6 had complex incomplete uncomplicated syndactyly of the right and left pelvic limbs. Orthopaedic and radiographic examination revealed that the third and second phalanges of digits 3 and 4 were completely joined in the right pelvic limb and the first to third phalanges of digits 3 and 4 were partially joined (Fig 4). The right pelvic limb had three digital pads, whereas the left pelvic limb had four digital pads. One large nail was present in the right pelvic limb for the affected digits, while two nails were present in the left pelvic limb for the affected digits.

It has now been a year since the kittens were diagnosed with syndactyly. To date, all kittens have been successfully adopted and have been free of lameness.

DISCUSSION

In this case report we have presented a litter of kittens with various forms of complex uncomplicated syndactyly involving both pectoral and pelvic limbs. Four of six kittens in the litter were affected. In one kitten, all four feet were affected, whereas in the other three only the pelvic limbs were affected. None of the kittens was lame, required medical or surgical treatment or had other gross skeletal anomalies.

Feline syndactyly is a rare condition, and to our knowledge, there are only four previous reports of this condition. Thus, information on the clinical presentation and concurrent anomalies, clinical and

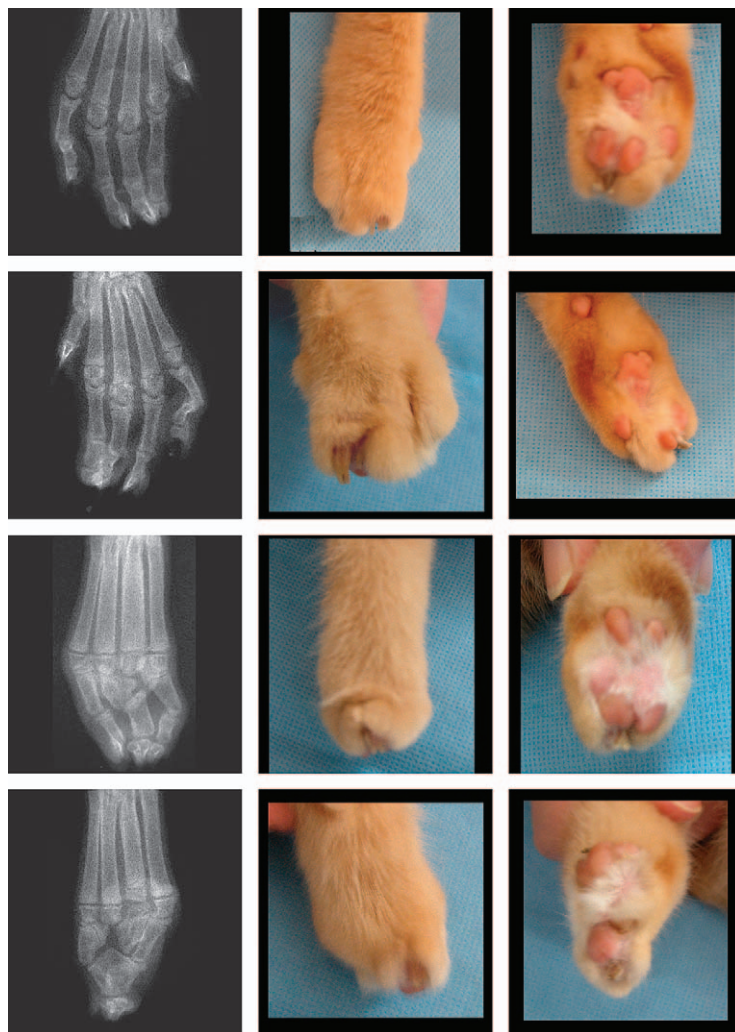


FIG 1. Foot deformities in kitten 1. Row 1: right pectoral limb; Row 2: left pectoral limb; Row 3: right pelvic limb; Row 4: left pelvic limb

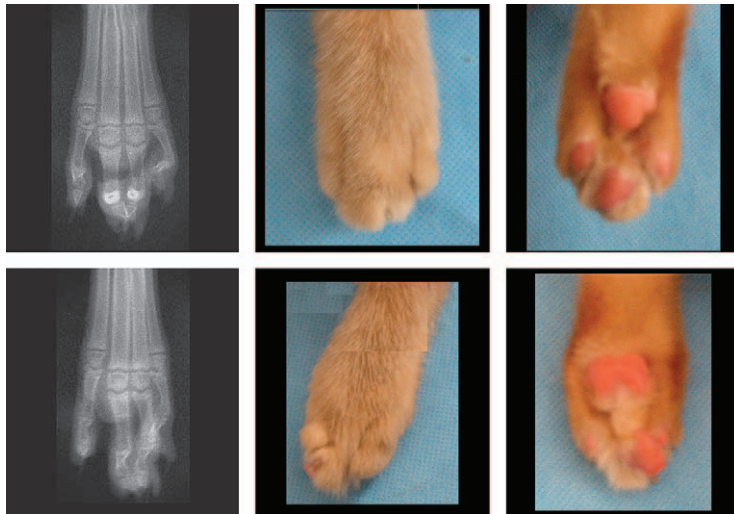


FIG 2. Foot deformities in kitten 2. Row 1: right pelvic limb; Row 2: left pelvic limb

radiographic findings and possible links to heredity and methods of prevention is limited. All reported cases were affected with complex syndactyly. In two of the previous reports, the affected cats also had ectrodactyly (Searle 1953) or a family history of polydactyly (Howe 1902). Thus, although the kittens in the present report did not have concurrent anomalies, feline syndactyly may be associated with polydactyly or ectrodactyly (complicated syndactyly). This parallels canine syndactyly, in which approximately half of the reported cases are complicated (Table 2). All affected cats in the previous four and in

this report were presented without lameness and the kittens in the present report had not developed lameness at the time of writing. This is similar to canine syndactyly, where all reported cases with complex syndactyly were without lameness and only cases with simple syndactyly were presented with lameness.

Paw changes in syndactyle cats include abnormal or absent digits, abnormal shape, fusion or absence of digital pads and abnormal shape or fusion of nails. Dissection of feline syndactyle legs demonstrated that deep digital flexor muscles to fused digits were often absent (Howe

1902, Hays 1917). In general, the musculature corresponded to the skeletal anomalies, for example, fused digits corresponded to single-muscle units and some muscles were lacking in entirety. Radiographic findings consisted of complete or incomplete joining of phalanges 1 to 3 and metacarpal or metatarsal bones (Howe 1902, Hays 1917, Searle 1953, Boehringer 1975). Both the external and radiographic features of feline syndactyly are similar to those of canine complex syndactyly. However, a lack of cutaneous separation between digits without osseous abnormalities, as seen in canine simple syndactyly (Richardson and others 1994), has not been observed (Howe 1902, Hays 1917, Searle 1953, Boehringer 1975).

An inheritable form of feline syndactyly has been reported (Searle 1953). This report was from a cattery where the female was allowed to breed freely. In addition, affected kittens also had ectrodactyly deformities (Searle 1953). The proportion of kittens with either defect, in successive litters, was 0 of 5, 2 of 6, 2 of 7, 3 of 4 and 0 of 4. Both males and females were affected. It was concluded that ectrodactyly combined with syndactyly was due to the action of a heterozygous gene with a variable expression. In another report, syndactyly was reported in a colony of polydactyle cats (Howe 1902). Although the incidence of syndactyly in this colony was not reported, it may be that syndactyly was an inherited trait, just like polydactyly (Howe 1902). In the present report, environmental and family histories were not available. However, because four of the six kittens in the litter were affected, a genetic background for the syndactyly is not unlikely (Thrusfield 1988, Patterson and others 1989). Thus, it may be that almost all reported feline syndactyly cases have a genetic aetiology.

The presented litter and previous reports yield guidelines for the diagnosis, treatment and prevention of feline syndactyly. Because syndactyly may be a hereditary trait, a detailed history with questions about syndactyly in littermates and previous litters (including male:female ratio) and other related cats should be obtained. Information on drug exposure, maternal disease or radiation exposure during the first and second trimester of gestation

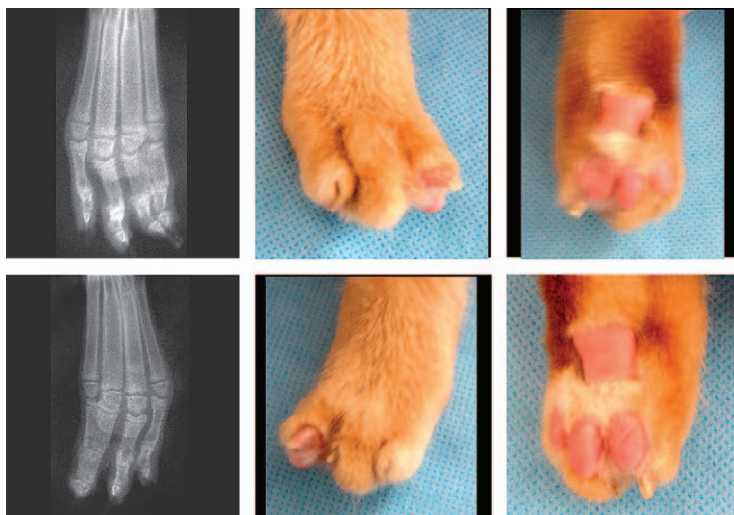


FIG 3. Foot deformities in kitten 3. Row 1: right pelvic limb; Row 2: left pelvic limb

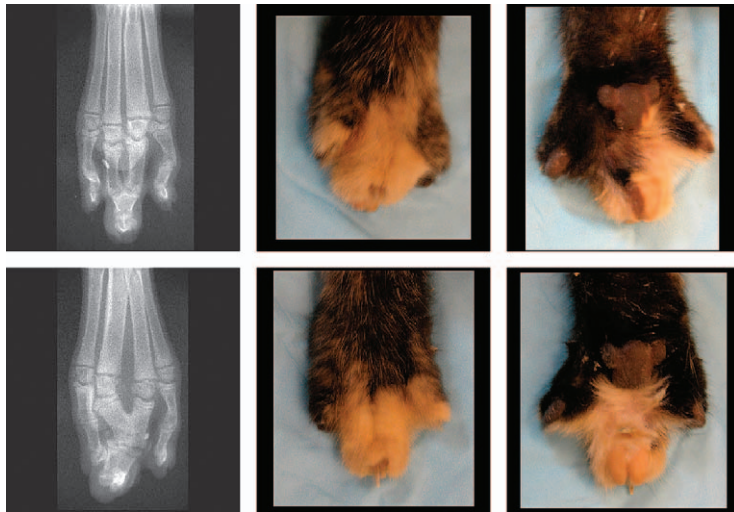


FIG 4. Foot deformities in kitten 6. Row 1: right pelvic limb; Row 2: left pelvic limb

should be collected to determine possible causative environmental factors (Towle and Breur 2004). The clinician should perform a thorough general physical examination to detect other congenital anomalies. The paws should be carefully examined for abnormal or absent digits, abnormal shape, fusion or absence of digital pads and abnormal shape or fusion of nails. Radiographic changes may be dramatic and can include complete or incomplete fusion of phalanges and metatarsal or metacarpal bones. The diagnosis of complex syndactyly is based on clinical and radiographic findings. It appears that feline complex syndactyly causes minimal to no discomfort and reported patients affected with this condition have presented without lameness. Therefore medical or surgical treatment of this condition is not recommended. Because the condition may be hereditary, breeding of affected animals should be discouraged and it is recommended that affected animals be neutered or spayed.

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References

ARIAS, A. M. & STEWART, A. (2002) Patterns in three dimensions. In: *Molecular Principles of Animal Development*. Eds A. M. Arias and A. Stewart, Oxford University Press, Oxford. pp 363-400

BAUM, H. (1889) Ein Fall von Syndaktylie beim Hunde. *Deutsche Zeitschrift für Tiermedizin* **15**, 81-90 (In German)

BOEHRINGER, B. T. (1975) Foot deformity in a cat. *Feline Practice* **5**, 50

CARRIG, C. B., WORTMAN, J. A., MORRIS, E. L., BLEVINS, W. E., ROOT, C. R., HANLON, G. F. & SUTER, P. F. (1981) Ectrodactyly (split-hand deformity) in the dog. *Veterinary Radiology* **22**, 123-144

DALLMAN, M. J. & BROWN, R. E. (1980) Syndactyly in the dog. *Canine Practice* **1**, 21-24

DALUISKI, A., YI, S. E. & LYONS, K. M. (2001) Original communications: the molecular control of upper extremity development: implications for congenital hand anomalies. *Journal of Hand Surgery* **26A**, 8-22

DAO, K. D., SHIN, A. Y., BILLINGS, A., OBERG, K. C. & WOOD, V. E. (2004) Surgical treatment of congen-

ital syndactyly of the hand. *Journal American Academic Orthopedic Surgery* **12**, 39-48

FREEMAN, L. E., SPONENBERG, D. P. & SCHABDACH, D. G. (1988) Morphologic characterization of a heritable syndrome of cleft lip/palate, polydactyly, and tibial/fibular dysgenesis in Australian Shepherd dogs. *Anatomia Histologia Embryologia* **17**, 81

GEHRING, H. & SCHRÖDER, U. (1961) Syndaktylie und Oligodontie bei einem Papillon-Rüden. *Deutsche Tierärztliche Wochenschrift* **68**, 352-353

GER, E. (1998) Syndactyly. In: *Congenital Malformation of the Hand and Forearm*. Ed D. Buck-Gramcko. Churchill Livingstone, London. pp 131-141

GILBERT, S. F. (2003) Development of the tetrapod limb. In: *Developmental biology*. 7th edn. Sinauer Associates Inc, Sunderland. pp 523-546

HAYS, G. P. (1917) A case of syndactylous cat. *Journal of Morphology* **30**, 65-82

HOWE, F. (1902) A case of abnormality of cats' paws. *American Naturalist* **36**, 511-526

KOZIN, S. H. (2003) Current concepts review: upper extremity congenital anomalies. *Journal of Bone and Joint Surgery* **85A**, 1564-1576

LEIPOLD, H. W. & GUFFY, M. M. (1973) Syndactyly in a German shepherd dog. *Veterinary Medicine/Small Animal Clinician* **68**, 910-911

NODEN, D. & DE LAHUNTA, A. (1985) Limb development. In: *The Embryology of Domestic Animals: Developmental Mechanisms and Malformations*. Eds D. M. Noden and A. de Lahunta, Williams & Williams Co, Baltimore, MD, USA. pp 196-210

OGDEN, J. A. & GROGON, P. (1987) Prenatal skeletal development and growth of the musculoskeletal system. In: *The Scientific Basis of Orthopaedics*. Eds J. A. Albright and R. A. Brandt. Appleton & Lange, Norwalk. pp 47-89

PATTERSON, D. F., AGUIRRE, G. A., FYFE, J. C., GIGER, U., GREEN, P. L., HASKINS, M. E., JEZYK, P. F., NODEN, D., DE LAHUNTA, A. & MEYERS-WALLEN, V. N. (1989) Is this a genetic disease. *Journal of Small Animal Practice* **30**, 127-139

RENOY, B. P. & BALLIGAND, M. (1991) Un cas de syndactylie chez le chien. *Annales de Médecine Veterinaire* **135**, 43-44 (In French)

RICHARDSON, E. F., WEY, P. D. & HOFFMAN, L. A. (1994) Surgical management of syndactyly in a dog. *Journal of the American Veterinary Medical Association* **205**, 1149-1151

RISER, W. H. (1964) What is your diagnosis? *Journal of the American Veterinary Medical Association* **145**, 169-170

SCHULTZ, V. A. & WATSON, A. G. (1995) Lumbosacral transitional vertebra and thoracic limb malformations in a Chihuahua puppy. *Journal of the American Animal Hospital Association* **31**, 101-106

SEARLE, A. G. (1953) Hereditary "split-hand" in the domestic cat. *Annals of Eugenics* **17**, 279-282

SWANSON, A. B. (1976) A classification for congenital limb malformation. *Journal of Hand Surgery* **1**, 8-22

THRUSFIELD, M. (1988) Is it hereditary?: the cause of disease. *Journal of Small Animal Practice* **29**, 603-609

TOWLE, H. A. & BREUR, G. J. (2004) Dysostoses of the canine and feline appendicular skeleton. *Journal of the American Veterinary Medical Association* **225**, 1685-1692