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Referring Veterinarian: Dr David Eaton

Postal Address: 24665 155th Ave, Eldridge,

Clinic Name: Scott County Animal Hospital

Email: scahdave@netins.net

Phone: 5632858624

Patient Information

Client: Williams, Sarah Patient ID: 10848-8235 PennHIP Num: 101103

Sex: Male

Date Submitted: 27 Sep 2016

Breed: STANDARD POODLE

Tattoo Num:

Fax: 5632856798

Reg. Name: Magnificant Marley Microchip Num: 985112004440450

Date of Birth: 30 Apr 2015 Weight: 20 lbs/9.1 kgs Date of Report: 27 Sep 2016 Patient Name: Marley

Registration Num: PR18584106 Species: Canine

Age: 17 months

Date of Study: 27 Sep 2016

Findings

Distraction Index (DI): Right DI = 0.40, Left DI = 0.33.

Osteoarthritis (OA):No radiographic evidence of OA for either hip.

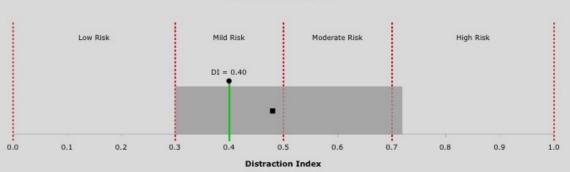
Cavitation/Other Findings: None.

Interpretation

Distraction Index (DI): The laxity ranking is based on the hip with the greater laxity (larger DI). In this case the DI used is 0.40.

OA Risk Category: The DI is between 0.31 and 0.49. This patient is at mild risk for hip OA. Distraction Index Chart:

STANDARD POODLE



Breed Statistics: This interpretation is based on a cross-section of 2721 Canine patients of the STANDARD POODLE breed in the AIS PennHIP database. The gray strip represents the central 90% range of DIs (0.30 - 0.72) for the breed. The breed average DI is 0.48. The patient DI is the solid circle (0.40).

summary: The degree of laxity (DI = 0.40) falls within the central 90% range of DIs for the breed. This amount of hip laxity places the hip at a mild risk to develop hip OA. No radiographic evidence of OA for either hip.

Interpretation and Recommendations: No OA/Mild Risk: Low risk to develop radiographic evidence of hip OA early in life, however OA may manifest after 6 years of age or later. Risk of OA increases as DI, age, body weight, and activity level increase. OA susceptibility is breed specific, larger breeds being more susceptible. Recommendations: Evidence-based strategies to lower the risk of dogs developing hip OA or to treat those having OA fall into 5 modalities.* For detailed information, consult these documents.* Use any or all of these modalities as needed: 1) For acute or chronic pain prescribe NSAID PO short or long term. Amantadine can be added if response is marginal or if a neuropathic component to the pain is suspected. 2) Optimize body weight, keep lean, at BCS = 5/9. 3) Prescribe therapeutic exercise at intensities that do not precipitate lameness. 4) Administer polysulfated glycosaminoglycans IM or SQ, so-called DMOAD. 5) Feed an EPA-rich prescription diet preventatively for dogs at risk for OA or therapeutically for dogs already showing radiographic signs of OA. At the present time there is inadequate evidence to confidently recommend any of the many other remedies to prevent or treat OA. Studies are in progress. Consider repeating radiographs at periodic intervals to determine the rate of OA progression and adjust treatment accordingly. Older dogs may show clinical signs such as chronic pain, reluctance to go stairs or jump onto the bed, and stiffness particularly after resting. It is unlikely that end-stage hip disease will develop for dogs at this risk level so surgical therapy for the pain of hip OA would rarely be indicated.

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