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# First Report of *Dictyocaulus arnfieldi* Infestation in a Horse in Mashhad, Iran

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#### Abstract

This article describes a case of *Dictyocaulus arnfieldi* infestation in a horse with signs of mild pneumonia in Mashhad zoo. In hematological examinations leukocytosis, neutrophilia with degenerative left shift; and in parasitological examination the first stage larvae of *Dictyocaulus arnfieldi* were detected in the feces. According to the authors' knowledge, this is the first report of *D. arnfieldi* infestation in Iran.

Keywords: Dictyocaulus arnfieldi, pneumonia, horse

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#### Introduction

Pneumonia can be caused by parasites in the horse. Dictyocaulus arnfieldi is the true lungworm found in the horses, belonging to the superfamily of Trichostrongyloidea. It has been reported from various parts of the world. (Bakirci and et al., 2004; Feinstein, 1979; Fikru and et al., 2005; Gothe and Heil, 1984) The natural host of the parasite is donkey, and comparably, large numbers of parasites can accumulate in the lungs of this host without clinical signs. Donkey can act as a reservoir for horses.(Beelitz and et al., 1996). The parasite has also been isolated from 2 zebras (Ashizawa and et al., 1980). It has a direct life cycle, and a prepatent period of approximately 12 weeks (Round, 1976a). Migrating *D*. arnfieldi (Cobbold 1884) is usually the etiological agent of parasitic pneumonitis in the adult horse, whereas Parascaric equorum is the most common etiological agent of parasitic pneumonitis in weanlings (Clayton, 1986). The present report describes a case of D. arnfieldi infestation in a 7-year-old stallion.

### **Case Presentation**

The stallion was kept in a zoo in Mashhad metropolice, northeast of Iran as a recreational horse with minimal exercise, almost entirely used by people for riding and taking photos. There was a history of housing the stallion with five donkeys together for a limited period of time recently. The horse ration was consisted of alfalfa hay and barley grain. The complained about chronic owner a nonproductive cough that caused annoyance of the visiting people. The horse was examined in the zoo on April 2008.

A thorough clinical examination was performed. Additionally, blood and fecal samples were collected for hematological and parasitological examinations, respectively. The fecal samples were Baermannized for the detection of larvae and also, egg per gram (EPG) count was performed by floatation method. Fecal culture was also performed for 7 days at 25 °C for harvesting and diagnosis of larvae. The affected horses were treated with ivermectin ( $200\mu g/kg$ , P.O., Ivermctin 1%, Nasr Co., Fariman, Iran). Two weeks later, fecal samples were collected again to evaluate the response to treatment.

# Results

On physical examination, the horse was in good body condition (fat), weighing around 400 kilograms, with a nonproductive cough, without nasal discharge. Rectal temperature was 38°C, heart rate 40 beats/min, and respiratory rate 40 breaths/min. The other clinical examination parameters were normal, except some crackles were auscultable over the topographic area of the lungs. On a clinical basis, bacterial pneumonia was considered as the first tentative diagnosis. Other probable tentative diagnoses were considered such as chronic hypersensitivity pneumonitis, chronic obstructive pulmonary disease. fungal pneumonia, immune mediated interstitial or vascular disease and unusual drug reactions, (Burks, 1998), as well as foreign body in the trachea.

The hematological results were as follows: PCV 44 (normal range: 32-47%), Hemoglobin 15.9 (normal range: 11-17g/dl), RBC 8.96 (Normal range: 6.4-10 × 106), MCV 49 (normal range: 43-54 fl) MCH 17.7 (normal range: 15-19 pg) MCHC 36.1 (normal range: 34-37 g/dl), total white blood cells 17100 (normal range: 5200-13900/µl), adult neutrophils 12825 (normal range: 2200-7400/µl), Band neutrophils 171 (normal range: 0-100/µl), eosinophils 171 (normal range: 0-600/µl), lymphocytes 3591 (1100- 5300/µl), monocytes 342 (normal range: 0-900/µl), platelets  $5.84 \times 105/\mu$ l, total protein 6.7 (normal range: 5.5-7.3 g/dl), fibrinogen 300 (normal range: 100-500 mg/dl). The EPG was 130 eggs per gram of faeces. No Parascaris equorum eggs were observed on fecal examination. The first stage larvae of D. arnfieldi were diagnosed according to the observation of a terminal small spike (Figure 1). Larvae per gram (LPG) were 5. Other larvae which were harvested after fecal culture belonged to small strongyles. The larvae of *D*. *arnfieldi* were also isolated from another 9 years old horse in the same premise in the zoo, with neither an apparent history of exposure to donkeys nor overt clinical signs.

## Discussion

In several studies, 50-80% of donkeys have been found infected with D. arnfieldi (Clayton and Duncan 1981; Klei, 1986). Before the advent of the anthelmintic, ivermectin, the prevalence of D. arnfieldi infection in horses in Kentucky at necropsy was approximately 11%, while it was 2% in live horses at the same time and at the same region, based on fecal examination (Lyons and et al., 1985a; Lyons and et al., 1985b). This difference may suggest the difficulty in antemortem diagnosis of D.arnfieldi in horses. However, patent infections have been found in horses, resulting in disease occurrence in closed herds with no exposure to donkeys or mules (Lyons and et al., 1985b; Clayton and Duncan, 1981). Experimental reproduction of the infestation has been established in pony foals (Clayton and Duncan, 1981). These findings support previous reports of horse-to-horse transmission (Lyons and et al., 1985a; Lyons and et al., 1985b; Round, 1976b). The presence of migrating larvae and adult worms in the lung parenchyma and bronchi activate both the humoral and cellular immune systems and immune-mediated results in an bronchopneumonia, subsequently with release of IgE, infiltration of mast cells, and basophils release of leading to histamine and eosinophilic chemotactic factor (Mackay and Urquhart, 1979), with deleterious effects on both parasite and the surrounding respiratory cells, causing accumulation of mucopurulent exudate in the lumen. (Mackay and Urguhart, 1979; Nicholls and et al., 1979) Direct damage by the adult parasites results in chronic catarrhal bronchitis with hyperplasia and thickened epithelium as well as focalz edema

hemorrhage. Secondary and immunosuppression and the colonization of secondary bacterial infections may occur (Burks, 1998). The lack of nasal discharge, normal temperature, good appetite and good body condition in both horses in our observation might be due to low parasite burden, and/or short course of the disease after exposure to donkeys. The diagnosis of D. arnfieldi infestation in horses may be somewhat difficult. The animals are not typically inappetent or dull in the case of D. arnfieldi (Boyle and Houston, 2006) However, it has been reported that cough and dyspnea were always accompanied infected horses (Andersen and Fogh, 1981a). Hematological and serum biochemical results may be normal in horses with parasitic pneumonitis. However, secondary bacterial infection if occurs. inflammatory changes, such as a mature neutrophilia and hyperfibrinogenemia may be evident following parasitis pneumonitis. This feature may prompt the clinician to follow an antimicrobial therapy. From this viewpoint, some horses with parasitic pneumonitis may show some clinical improvement following antimicrobial treatment (Boyle and Houston, 2006), that may be misleading and be followed by the relapse of clinical signs. In this case, the horse may show leukcytosis and neutrophilia with regenerative left shift implicating a bacterial pneumonia. It seems that we should consider the possibility of concurrent bacterial pneumonia; however, secondary bacterial infections may occur by the presence of the adult parasites in the bronchi (Burks, 1998). Diagnostic confirmation of parasitic pneumonitis due to D. arnfieldi depends on several considerations. Bronchoalveolar lavage and transtracheal wash in order to harvest some mucus may reveal eosinophilic reaction that is most commonly found in cases of D. arnfieldi infestation in horses and may strongly implicate D. arnfieldi infestation. Normal eosinophil percentages of the total nucleated cell count in bronchoalveolar lavage fluid are 1.0±1.4% (Mair et al., 1987). In the

apparently absence of the evidence of the parasite, outbreak of the eosinophilic bronchitis in horses, and also a pony possibly associated with D. arnfieldi have been reported (Mackay and Urquhart, 1979, Church, 1983). The response to appropriate anthelmintic treatment is an alternative procedure for diagnosis. А history of commingling of horses with donkeys, and/or grazing in common pastures should be considered in the diagnosis of parasitic pneumonitis caused by D. arnfieldi in horses that show signs of respiratory disease (Boyle and Houston, 2006; Burks, 1998).

Finding *D. arnfieldi* larvae in the feces is not common, as it was a discrepancy between estimated prevalence rate of infestation at necropsy or antemortem diagnosis in Kentucky (11 vs. 2%) (Lyons and *et al.*, 1985a; Lyons and *et al.*, 1985b). However, patent infections have been found in horses, resulting in disease occurrence in closed equine herds with no exposure to donkeys (Andersen and Fogh, 1981).

The affected horses showed a good response to antihilmintic treatment. In the subsequent fecal examination two weeks later, both horses showed no evidence of *D*. *arnfieldi* larvae in their feces, as well as a high drop in EPG of other parasitic nematodes.

Up to the authors' knowledge, this is the first report of D. arnfieldi infestation in Iran. The *D. arnfieldi* infection in horses should be considered when horses show clinical signs of respiratory disease when commingled with donkey, or shared parstures with donkeys when they are not commingled apparently. The relapse of clinical signs in horses that was treated for a bacterial pneumonitis should arouse suspicion of parasitic pneumonitis. With the difficulty in D. arnfieldi Larvae capture in horse feces, it is highly advisable to transfer fecal samples in the vicinity of ice, because the loss of the parasite larvae has been reported insignificant in 4 ° C, but not at 16 and 20 ° C (Rode and Jorgensen, 1989) that might increase the chance of larval recovery in feces. With the difficulty of larval isolation in feces, other methods of diagnosis, including bronchoalveolar lavage and/ or a good response of horses with signs of respiratory disease to antihilmintic treatment shouldn't be overlooked.

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Figure 1. Dictyocaulus arnfieldi larva. The terminal spike is evident and the larva is about 480 $\mu$ m. (Lugol 10% staining, 400X)

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# اولین گزارش آلودگی اسب با دیکتیوکولوس آرنفیلدی در مشهد

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#### چکیدہ

در اردیبهشت ماه ۱۳۸۷، یک راس نریان، که قبلاً با چند راس الاغ در یک جایگاه قرار داشت، با علائم سرفه های خودبه خودی و متناوب در باغ وحش مشهد معاینه شد. برای آزمایش های خون شناختی و انگل شناختی نمونه های خون و مدفوع تهیه شد. نتایج آزمایشگاهی لکوسیتوز، نوتروفیلی همراه با انحراف به چپ جبرانی، و همچنین لارو مرحله ی اول دیکتیوکولوس آرنفیلدی در مدفوع را نشان داد. تاکنون آلودگی به این انگل در ایران گزارش نشده است.

واژگان کلیدی: دیکتیوکولوس آرنفیلدی، پنومونی، اسب