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Prevalence of gastrointestinal parasite, *Paramphistomum* in domestic animals (Cows and Buffaloes) of district Swat and Charsadda, KP, Pakistan

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Abstract

An investigation was carried out to study the prevalence of *Paramphistomum* in domestic cattle's. For this purpose samples were collected from slaughter houses, dairy farms and houses, from two districts (Swat and Charsadda) of KP Pakistan during month of November 2011 to April 2012. Infection rate was 15% in buffaloes and 10% in cows, in district Swat and 10.8% in buffaloes and 7.5% in Cows, in district Charsadda. Overall the highest month wise prevalence in cows 12.5% and in buffaloes 20% was recorded during March. It was also observed that the highest infection rate was recorded in male animals (10%) than female (7.2%) in case of cows and in case of buffaloes the prevalence rate in male was (17%) and in female (10%). Area wise prevalence indicate that infection in buffaloes and cows of district Swat was higher (15% and 10% respectively) than Charsadda (10.8% and 7.5% respectively).

Keywords: *Paramphistomum*, Prevalence, Buffaloes, Cows, Swat, Charsadda

1. Introduction

Paramphistomosis is a disease of domestic and wild ruminants, which is caused by trematode and commonly belong to the family Paramphistomatidae, which in their early stage present in the small intestine and abomasums, from here they pass to rumen and lastly lodge as an adult trematode [16].

Paramphistomum spp. are Platyhelminth (flatworm) parasites (Platyhelminthes: Trematoda: Digenea) responsible for Paramphistomosis i.e. gastrointestinal parasitic disease in domesticated animals, which causes heavy economic losses [3]. *Paramphistomum* are rounded in transverse section, pear-shaped worms and characterized by possessing distinct suckers which are two in numbers, one for attachment purposes and the other is for mouth. The oral sucker contains a muscular pharynx, which is actually the beginning of the diverged caecum (alimentary canal). The posterior sucker or acetabulum is well developed and present at the posterior end of rumen flukes. The flukes are hermaphroditic organisms and the genital pore is situated ventral medially in the anterior third, acting as a conjoint opening for both male and female sex organs [9].

Trematode parasites of ruminant cattle have a world-wide distribution and even valuable zoonotic organisms among the helminths [11]. In some countries these parasites are considered as a major constriction on productivity. Likewise, *paramphistomum* infections (e.g. *Paramphistomum cervi*, *P. ichikawai* and *P. microbothrium*) are widespread in distribution, mainly in Asia. In Asia the prevalence rates (30-60%) are still recorded in some areas [13].

Paramphistomum cervi is a trematode parasite which attacks livestock badly thereby disturbing their productivity. The adult trematode is located in the rumen of ruminant and immature trematode in the snail intestine (duodenum) [5]. In chronic infection, the liver may be turned pale and shows fibrosis. Tissue and organ damages were caused by *Paramphistomum cervi* which causing poor production of milk, meat, skin and retardation in growth [10].

Paramphistomum cervi is particularly disease causing species, which migrating juveniles and causes severe hemorrhage and enteritis, mostly causing death of the host. Secondary infection also complicates the problem [14]. The main epidemiological variable inducing worm problems of animals is the infection rate from pastures. Paramphistomosis is also influenced by the climatic conditions for hatching of egg, development and existence of larvae in pasture [17].

The harm caused by this infection in bovine affects production, since these parasites provoke a lower nutritious conversion, a loss of weight and/or a decrease in milk production, which cause economic losses. In some areas of India, the Republic of South Africa and Australia, the mortality of cattle has reached 80 or 90% in sheep and cattle. This illness is distributed all around the world, but its highest frequency has been registered in tropical and subtropical regions [12].

The filtration technique with sieves and sedimentation is the most accurate method for identification of eggs in feces, producing stronger evidence in the sediment of sample during microscopy and study. Eggs show similarity in shape with *Fasciola hepatica*, but *Paramphistomum* eggs are somewhat larger (160 – 180 μ) [16].

Most licensed drugs for the treatment of *Fasciola* are not active against paramphistomosis. Only few drugs are efficient against either mature and/or immature flukes, namely; *niclosamide*, *oxyclozanide*, *rafoxanide*, and *resorantel* [19].

Gastrointestinal helminthes prevalence in ruminants has been reported in different areas at different times from 25.1 to 92% in Pakistan [1, 2, 6, 8, 13].

Paramphistomosis is thought to be a main constraint on productivity in many areas. Likewise, *paramphistome* infections (e.g *Paramphistomum cervi*) remain widespread, mainly in Asia. In Asia the prevalence rates of 30-60% are still recorded [13, 15].

The present research was designed to report the seasonal and the overall prevalence of the *paramphistomum* in cows and buffaloes. Furthermore to suggest preventive measures and prevalence of *paramphistomum*, taking this background the study was conducted to find out prevalence of *Paramphistomum* in buffaloes, cows, in district Swat and Charsadda of Khyber Pakhtunkhwa, Pakistan.

2. Materials and Methods

2.1 Epidemiological Study

2.1.1 Study area

Fecal samples of cows and buffaloes were collected from two different Districts (Swat and Charsadda) of Khyber Pakhtunkhwa Pakistan, from November, 2011 to April, 2012 shows in Fig 1



Fig 1: Encircled positions in the map showing study areas: (1) Swat, (2) Charsadda.

2.1.2 Study design

Study design was based under laboratory condition.

2.1.3 Instrumentation

Tools used in this study were as follows:

Cotton, Refrigerator, Cover slip, Slide, Beaker, Glass rod, Gloves, Microscope, Motor and Pestle.

2.2 Coprological examination

2.2.1 Fecal sample collection

From each districts a total of 20 fecal samples each for buffaloes and cows were monthly collected in plastic jars which was clearly labeled with gender, species, date and place of collection. For the presence of *Paramphistomum* eggs these samples were examined on the same day by direct microscopic examination [18]. Samples which were not examined on the same day were preserved in formalin 3.0% to prevent the eggs development and hatching.

2.2.2 Direct microscopic examination

A small amount of fecal sample was mixed with water in a beaker. Few drops of it were place on a glass slide, covered with cover glass and *Paramphistomum* eggs were observed under microscope (10x10 and 10x40). From each sample 3-5 slides were prepared following the direct microscopic examination. Eggs were identified on the basis of morphology [20]. Prevalence of infection was monthly recorded. The age, sex, area wise prevalence was noted. The prevalence of the disease was recorded following the modified formula described by Thursfield (1986):

$$\text{Prevalence (\%)} = \frac{\text{No. of infected individuals at particular point in time}}{\text{No. of total individuals at particular point in time}} \times 100$$

2.2.3 Sedimentation method [7]

First we weighted approximately 3 gm. of faces in container 1. Then we poured 40-50 ml of water in container and filtered the fecal suspension through a tea strainer or double-layer of cheese cloth in container 2. Poured the filtered material into a test tube and allow to sediment for 5 minutes. The supernatant were removed very carefully, re-suspend the sediment in 5 ml of water and allow to sediment for 5 minutes. The supernatant discarded very carefully by pipetting, decantation. Stained the sediment by adding one drop of ethylene blue and transferred the sediment to a micro slide. Cover with a cover slip and was observed under microscope.

3. Results

During the study period of November 2011 to April 2012, an epidemiological survey of 480 samples of cows and buffaloes, were carried out in the two districts of KP (Swat and Charsadda). A total of 240 fecal samples from each district were examined out of which shows the following prevalence percentage (%).

Month, Area and Gender Wise Prevalence of *Paramphistomum* in Cows of KP

An overall prevalence (%) of *Paramphistomum* was found 8.75% in Swat and Charsadda from November 2011 to April 2012. To find out the year's specific period associated with highest prevalence rate, when month wise data recorded, it showed an overall highest prevalence (12.5%) was in March, followed by a decline to the lowest prevalence (5%) in April. Area wise prevalence in cows, two districts of KP province indicates the infection was highest at Swat (10%) followed by (7.5%) in Charsadda. Overall sex wise prevalence showed male (10%) were more susceptible than females (7.2%) for *Paramphistomum* infection. (Table 1. Fig. 2).

Table 1: Month, Area and Gender wise prevalence of *Paramphistomum* in cows at two districts (Swat and Charsadda) of KP from Nov 2011 to April 2012

	Factors	Total no. of sample observed	Total no. of sample infected	Prevalence (%)
Time (Months)	Nov	40	4	10%
	Dec	40	3	7.5%
	Jan	40	3	7.5%
	Feb	40	4	10
	March	40	5	12.5%
	April	40	2	5%
Areas	Swat	120	12	10%
	Charsadda	120	9	7.5%
Gender	Female	111	8	7.2%
	Male	129	13	10%
Total		240	21	8.75%

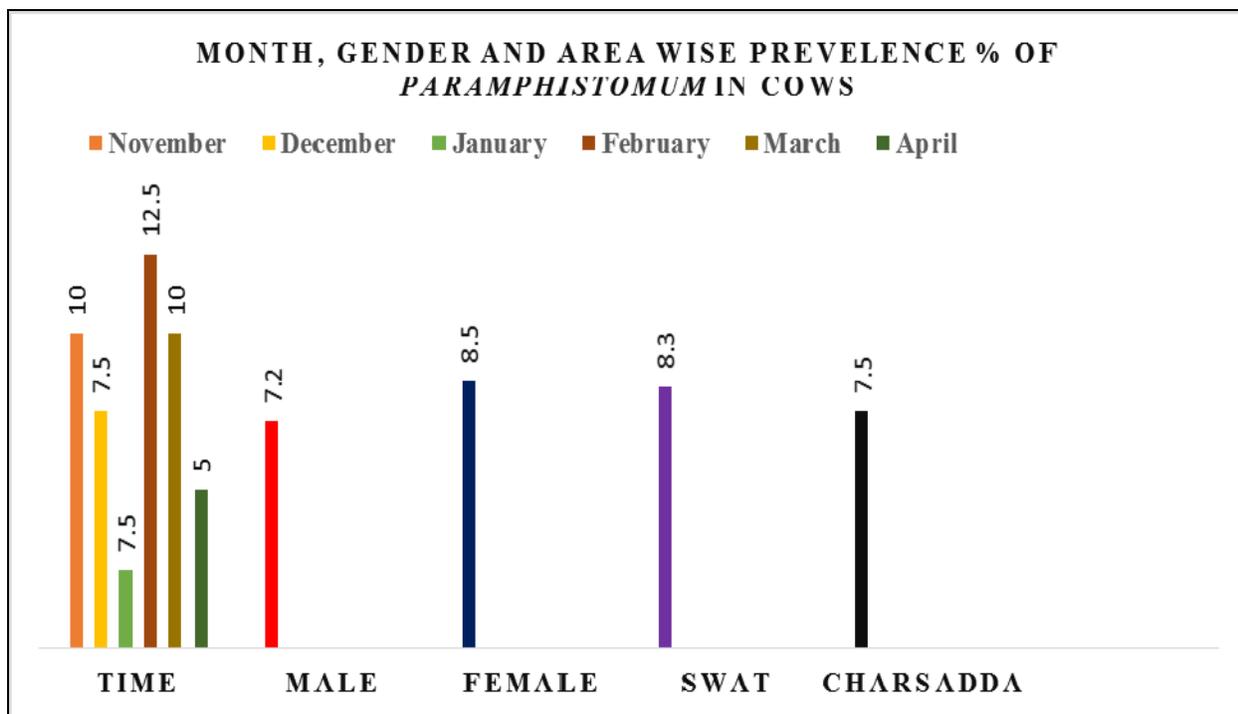


Fig 2: Shows Month wise, Area wise and Gender wise prevalence of *Paramphistomum* in Cows during the study period from November 2011 to April 2012.

3.1 Month, Area and Gender Wise Prevalence of *Paramphistomum* in Buffaloes of KP

Total 240 fecal samples of buffaloes were examined in two districts of KP (Swat and Charsadda). The overall prevalence 12.9% was recorded shown in Table 4.2. To find out the year’s specific period associated with highest prevalence rate,

when month wise data recorded, it showed an overall highest prevalence (20%) was recorded in March, which was decreased in April (10%). The lowest prevalence (7.5%) was recorded in November. Overall sex wise prevalence showed male (17%) were more susceptible than females (10%) in both districts. (Table 2,-; Fig. 3)

Table 2: Month, Area and Gender wise prevalence (%) of *Paramphistomum* in buffaloes at two districts (Swat and Charsadda) of KP from Nov 2011 to April 2012.

	Factors	Total no. of sample observed	Total no. of sample infected	Prevalence (%)
TIME (Months)	Nov	40	3	7.5%
	Dec	40	4	10%
	Jan	40	6	15%
	Feb	40	6	15%
	March	40	8	20%
	April	40	4	10%
Areas	Swat	120	18	15%
	Charsadda	120	13	10.8
Gender	Female	140	14	10%
	Male	100	17	17%
Total		240	31	12.9%

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