

PREVALENCE OF INTESTINAL HELMINTH PARASITES OF CHICKEN (*Gallus gallus domesticus* Linnaeus, 1758) IN LALITPUR DISTRICT, NEPAL

*Janak Raj Subedi**
Tasneem Mujahid
Bijay Cheetri

ABSTRACT

This study was aimed to determine the prevalence of intestinal helminth parasites of local chicken (*Gallus gallus domesticus*) from Lalitpur district, Nepal. It was carried out from February 2014 to March 2015. A total of 125 samples (93 alimentary canals of freshly killed chicken and 32 stool samples from ground and fowl runs) were collected. Out of 93 alimentary canals 17 were collected from free range chicken and 76 were from slaughtered house i.e. from poultry farm chicken. The faecal samples were collected in the sterile vials containing 2.5 percent potassium dichromate. Depending upon the convenience, post mortem examination method, differential floatation method and direct smear method were used for the examination of samples. The present study showed only 40 percent of all the poultry examined as infected. Overall 5 species of nematodes, 1 species of cestode and 4 unidentified species were found to be prevalent in chicken from Lalitpur district. The highest prevalence rate was found with *Heterakis gallinarum* (22.4%) followed by *Capillaria* species (16%), *Ascaridia galli* (10.4%), unidentified species (4.8%) and *Raillietina tetragona* (4%). Statistically there was a significant difference in the prevalence of helminth species ($\chi^2=33.83$; $p< 0.05$; $\alpha=1$). Likewise, there was significant difference in the prevalence rate of helminth parasites in free range chicken and poultry chicken ($\chi^2=22.055$; $p< 0.05$; $\alpha=1$).

Keywords: Chicken, Nematode, Cestode, Lalitpur, Helminth

INTRODUCTION

A domestic fowl or chicken (*Gallus gallus domesticus*), belonging to the family Phasianidae, is a sub species of Red Jungle Fowl. It is one of the most common and widespread domestic animals, with a total population

* Mr. Subedi is a Lecturer, Central Department of Zoology, Kirtipur, TU. Mujahid and Chhetri are associated with Central Department of Zoology, Kirtipur. TU.

of more than 19 billion as of 2011 (UNFAO, 2011). It is one of the most common and domesticated birds than any other species in the world. Human keep chickens primarily as a source of food, consuming both their meat and their eggs (UNFAO, 2011). Parasitic infection in chicken is the major problem in Nepal which leads to economic loss of the country. Domestic fowls are more often infected due to unhygienic management practices, malnutrition, lack of veterinary supervision and also the complicated life cycle of the parasites (Nair and Nadakal, 1981). Chicken infected with parasites show retarded growth, decreased egg production, reduced weight gain, significant haemoglobin depression (Nair and Nadakal, 1981), villous atrophy, catarrhal enteritis, granuloma formation in duodenum, desquamation of villi and submucosal glands congestion, inflammatory reaction and vacuolation of epithelial cells (Kurkure and Ganorkar 1998).

Parasitism is an association in which the parasite is metabolically dependent to a greater or lesser extent to the host. Gastro- intestinal parasites are however the most prevalent and most devastating parasites affecting chicken productivity (Swaton *et al.*, 2003). According to and village chicken are raised mainly under the free range (scavenging) product system, with partial or no housing and this predisposes the chicken to disease and parasites especially helminths (Muchadeyi *et al.*, 2004; Mwale and Masika, 2009 and Swaton *et al.*, 2003). In village, chicken are raised mainly under the free range (scavenging) with partial or no housing and this predisposes the chicken to disease and parasites especially helminths (Muchadeyi *et al.*, 2004; Mwale and Masika, 2009 and Swaton *et al.*, 2003). Different types of helminth parasites infect the chicken flocks. Worms find cozy places to stay in the crop, gizzard, intestine, caecum, windpipe and even the eyelids (Gauthier and Ludlow, 2013). On the basis of their site of location helminths are of different types, the worm which are found in caecum of large intestine are called caecal worms (*Heterakis* SPP.), worms which are found in eye are called eye worm (*Oxyuris mansoni*), Gape worms are found in trachea (*Syngamus trachea*) (Gauthier and Ludlow, 2013). These worms are also called “red- worm” or “forked-worm” and birds infected with gape worm show “open mouth breathing characteristics”. Round worm (*Ascaridia*) and tape worms (*Raillietina*) are found in intestine while thread worm (*Capillaria*) is found in crop or oesophagus (Janquera, 2017). The eggs and immature stages of many parasitic worms can live outside of the chicken host for a long time, possibly several years, whereas some parasitic worms spend part of their life cycle in other creatures such as

from Lalitpur District. Out of 93 alimentary canals 17 were collected from free range chicken and 76 were from slaughtered house i.e. from poultry farm chicken. These samples were collected from different places of Lalitpur district from February 2014 to March 2015. The faecal samples were collected in the sterile vials containing 2.5% potassium dichromate with the help of wooden stick. Potassium dichromate was used as preservative that helped to maintain the morphology of eggs and also prevented further development of helminth eggs. The alimentary canals were collected in polythene bags and were immediately brought to the laboratory for the examination. (Gurung, 2016). Depending upon the convenience, post mortem examination method, differential floatation method and direct smear method were used for the examination of samples. For post mortem examination method the alimentary canal of chicken was cut longitudinally from oesophagus to rectum including both caecal tubes. All worms visible to the naked eye were removed using thumb forceps and brush (Fowler, 1990). Cestodes were whole mounted for identification while nematodes were fixed in glycerol jelly and observed under the microscope. The freshly collected helminth parasites were kept in normal saline before fixation. Permanent slides (whole mount slides) of parasites were prepared for their identification according to the method described by Cable (1957).

For differential floatation method the saturated salt solution of specific gravity 1.2 was prepared by allowing an excess of common salt to boil in a basin until a scum was formed on the surface. It was cooled and stored in a bottle leaving an excess of undissolved salt at the bottom. Four grams of fecal material was taken in a test tube and a few drops of salt solution were added (Hansen and Perry, 1994). It was then stirred with a glass rod or a small piece of stick so as to make an even emulsion. After that more salt solution (15 to 20 ml according to the capacity of the test tube used) was added till the test tube was nearly full, stirring was continued through the process. Any coarse matter, which floats up, were removed without fear of removing any egg, as an egg takes a long time (20 to 30 minutes) to come to the surface of the fluid. At this stage the test tube was placed on the level surface and the final filling of the test tube was done by means of a dropper until a convex meniscus was formed. A glass slide was carefully laid on the top of the test tube so that its center is in contact with the fluid. The preparation was allowed to stand for 20 to 30 minutes, after which the glass slide was quickly lifted, turned over smoothly so as to avoid spilling of the liquid and was examined under the microscope (Chatterji, 2009).

For direct smear method method small amount of feces was placed on a slide . A drop of normal saline or Lugol’s solution was added to the feces and mixed thoroughly. Since we were looking for the helminth eggs , larva and cysts Lugol’s solution or normal saline was used. Then the fecal materials were covered with cover slip. The cover slip was moved around until it laid flat. The smear film was made thin so that the light from the microscope was able to pass through the sample in order for us to examine it. The slides were examined at 10X and 40X objective lens. (Chatterji, 2009).

Statistical Analysis

Prevalence of infection of identified species in each area was calculated as the number of individual chicken infected by a specific helminth species at the time of study was divided by the total number of chicken examined and multiplied by 100. Variation in the prevalence of gastrointestinal helminths in relation to helminth species and chicken type were analysed by using Chi- square statistics. In all cases $p < 0.05$ was considered indicative of statistically significant difference or association.

RESULTS AND DISCUSSION

The present study revealed that 40% among all poultry examined were infected by one or more species of helminth parasites. 36 intestine of chicken out of 93 were infected with helminth parasites while 14 stool samples or droppings were found positive for helminth eggs.

Prevalence Rate on the Basis of Species

The highest prevalence of *Heterakis gallinarum* (22.4%) was seen followed by *Capillaria* species (16%), *Ascaridia galli* (10.4%), and *Raillietina tetragona* (4%). Similarly the prevalence of unidentified species was found 4.8%. Statistically, there was a significant difference in the prevalence of helminth species ($\chi^2=33.832$, $p < 0.05$) with the highest prevalence of *H. gallinarum*.

Table 1: Species- wise Prevalence Rate (%), n= 125

Name of Parasite	No. of Chicken Infected by		T	Prevalence Rate
<i>Ascaridia galli</i>	Adult	Egg	13	10.40%
	9	4		
<i>Heterakis gallinarum</i>	23	5	28	22.40%
<i>Capillaria</i> SPP.	17	3	20	16%
<i>Raillietina tetragona</i>	5	0	5	4%
Unidentified	4	2	6	4.80%

where, t = total no. of infected chicken, Prevalence Rate= $(t/n) \times 100\%$, n= 125.

Prevalence Rate on the Basis of Class

Table 2: Class- wise Prevalence Rate (%), n= 125

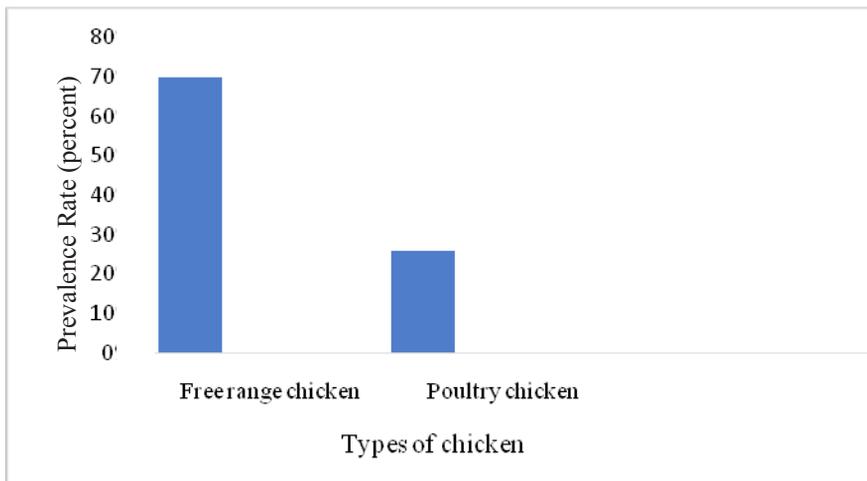
Name of class	t	Prevalence rate
Nematoda	61	48.80%
Trematoda	5	4%
Cestoda	0	0%
Unidentified	6	4.80%

where, t=total no. of infected chicken, Prevalence rate= $(t/n) \times 100\%$, n= 125.

Prevalence Rate on the Basis of Chicken Type

Out of 40 samples examined from free range chicken, 28 were found to be infected with the prevalence rate of 70% while from 85 samples of poultry chicken (from slaughtered house), 22 were infected with the prevalence rate of 25.88%. Statistically there was significant difference in the prevalence rate of helminth in free range chicken and poultry chicken ($\chi^2=22.055$, $p < 0.05$, $\alpha=1$).

Figure 2: Prevalence on the Basis of Chicken Type.



The domestic fowl or chicken harbours many intestinal parasites due to its feeding habit. According to Muchadeyi *et al.* (2004) and Mwale and Masike (2009) village chicken which are raised mainly under the free range (scavenging) product system, with partial or no housing have higher rate of infection of disease and parasites especially helminthes (Swaton

et al., 2003). Likewise, Mwale and Masike (2011) reported that village chickens increase rural farmers' nutritional and income status. Nonetheless, chicken productivity is chiefly affected by gastro-intestinal parasites and there is dearth of information on the prevalence of these parasites in village chickens in South Africa.

The study showed 40% among all poultry examined were infected by one or more species of helminth parasites which agrees with the work of Sudhir (2013) who found 51.67% infection in free range chicken in India. The present study is also more or less similar to the report of other worker who reported the prevalence rate of 41.4% (Tesfaheyawet *et al.*, 2010), 53.00% (Matur *et al.*, 2010) and 37.9% (Dawet *et al.*, 2012). In the present study six species of helminth were identified comprising five nematodes and one cestode compared to seven species of helminth identified by Adang *et al.*, (2014) comprising of six cestodes and one nematode, five species by Kose *et al.* (2009) comprising four nematodes and one cestodes, five species by Rayyan *et al.*, (2010) comprising of three nematodes and two cestodes and three species by Matur (2002) comprising two cestodes and one nematode

Out of 40 samples examined from free range chicken, 28 were found to be infected with the prevalence rate of 70% while 85 samples of poultry farm chicken (from slaughtered house) were found to have prevalence rate of 25.88% which agrees with the work of Hamad (2013) and Yoriyo *et al.* (2005). Similarly, Mikail and Adamu (2008) reported high infection rate in free range chicken (92.66%).

Statistically there was a significant difference in the prevalence rate of helminth in free range chicken and poultry chicken ($p < 0.05$). This may be because the free range chicken or backyard poultry are more susceptible to parasitic infection. The main food of backyard chicken consists of different types of seeds, kitchen wastage, insects, slugs, earthworm, etc. insects slugs worms act as intermediate host of many bioparasites (Soulsby, 1982).

CONCLUSION

The study was carried out to find the prevalence of intestine helminth parasites of local chicken in Lalitpur district. A total of 93 intestine and 32 stool sample of local chicken, *Gallus gallus domesticus*, were collected from different places of Lalitpur district. The collected samples were examined thoroughly for the presence of helminth parasites in the form of adult or egg. The present study showed that only 40% of all the poultry examined were

infected. The highest prevalence rate was found with *Heterakis gallinarum* (22.4%) followed by *Capillaria* species (16%), *Ascaridia galli* (10.4%) and *Raillietina tetragona* (4%). Similarly the prevalence of unidentified species was (4.8%).

The infection was more in free range chicken (70%) as they are reared in unhygienic environment and are more susceptible to parasitic infection due to their feeding habit. The intestine which were collected from slaughter house were with least infection (25.88%) as they were bought from poultry farm where chicken are reared in hygienic environment with medication.

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