

## Bovine Gastrointestinal Trematodosis In Nigeria: A Review

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**Abstract:** In this review, the economic importance of bovine gastrointestinal trematodosis in Nigeria was reviewed, bearing in mind that infections with these parasites affect cattle productivity either directly or indirectly. The epidemiological factors influencing the occurrence of the parasites were reviewed and three species (*Dicrocoelium*, *Fasciola* and *Paramphistomum*) that commonly occur in Nigeria were discussed. The prevalence of infections with these parasites was also reviewed both globally and in Nigeria. Various diagnostic techniques for each of the parasites were discussed as well as control option applicable to the Nigerian situation. Based on existing literatures and data on the epidemiological features of trematode infections in cattle, control and prevention methods are reviewed and modified grazing practices are recommended as this will help reduce host-parasite contact.

**Key words:** Gastrointestinal, Trematodosis, *Dicrocoelium*, *Fasciola*, *Paramphistomum*, Nigeria.

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### I. Introduction

Trematode parasitism is one of the major problems lowering cattle productivity around the world (Vercruysse and Claerebout, 2001). Among the prevailing parasitic diseases in Nigeria, trematode infections remain a major problem in cattle and other ruminants. They occur in vast water lodged and marshy grazing fields, a condition known to be ideal for propagation and maintenance of the snail intermediate hosts and hence high prevalence of infection (Solomon and Abebe, 2007). Based on the habitat they usually parasitize, trematodes may also be classified into blood, hepatic, lung and intestinal trematodes. This study focused on the hepatic and intestinal trematodes, comprising mainly *Fasciola* spp, *Dicrocoelium* spp and *Paramphistomum* spp.

Adult flukes lay operculated eggs (they are oviparous), the embryo in the eggs develops into pyriform (pear-shaped), ciliated larva (miracidium). Under the stimulus of light and moisture, the miracidium releases an enzyme which attacks the proteinaceous cement holding the operculum in place. This later springs open and the miracidium emerges within few minutes during hatching then penetrates the soft tissues of the snails, being aided by a cytolytic enzyme. The penetration lasts for about 30 minutes after which the cilia are lost and miracidia develop into elongated sacs called sporocysts containing germinal cells. The cells develop into rediae which migrate to the hepato-pancreas of the snail. From the germinal cells of the rediae arise the final stages, the cercariae. The cercariae (young flukes with tail) emerge actively from the snail, usually in considerable numbers. Unlike nematodes where one egg develops to an adult, single trematode egg may eventually develop into hundreds of adults due to the phenomenon of paedogenesis (Urquhart et al., 1996). Stimulus for emergence depends on species but most commonly a change in temperature or light intensity. Infected snails continue to produce cercariae indefinitely, though most infected snails die prematurely due to gross hepato-pancreas destruction (Urquhart et al., 1996).

The cercariae swim for some time and attach themselves to vegetations, shed their tails and encyst. This stage is called metacercariae. The encysted metacercariae have great potential for survival extending to months. Once ingested, the outer cyst wall is removed mechanically during mastication. Rupture of the inner cysts occurs in the intestine leading to hatching of the metacercariae. The juvenile flukes then penetrate the intestinal mucosa and migrate to the predilection sites where they become adults after several weeks. There in the predilection site, they reproduce and exert effect causing damage and disease in the affected host.

Gastrointestinal trematode infections are a major part of the health problem affecting cattle. Several workers have reported economic losses in cattle production due to these infections in the aspects of reduction in meat quantity and quality, milk production, organ condemnation (liver), loss of draught power, reproductive failure, mortality as well as risk of contracting zoonotic species (Hossain et al., 2011; Odigie and Odigie, 2013). However, their focus was mainly on *Fasciola* and did not cover other equally important trematode species that

cause substantial economic losses as well. Therefore, this review was done in order to provide relevant information that will help cover gap for intended researchers.

### **Economic importance of gastrointestinal trematodes**

Infections due to gastrointestinal trematodes are of great veterinary importance in cattle, accounting for considerable economic losses. The losses encountered are seen in the areas of mortality, morbidity, organ condemnation and also decreased meat quantity and quality, milk production, production efficiency and loss of draught power (Malone et al., 1998; Chaudhri, 2000; Bianchin et al., 2007; Javed, 2008; Addis et al., 2014). The economic impact of these trematodes in cattle is enormous. Great losses are evident especially where farmers have little or no knowledge on the disease (Ozung et al., 2011). Losses are more common during the rainy season when more cattle are exposed to the flukes.

Significant economic losses to global agriculture have been estimated to be more than \$3.2 billion annually due to infection with trematodes (Olsen et al., 2015). In Europe, Ayaz et al., (2014) reported an incidence of infections up to 77%. Losses due to these parasites in the UK and Ireland alone are greater than £70.67 million a year (Bekele et al., 2010) and a Swiss study by Morgan et al., (2013) reported £37.2 million loss in cattle due to subclinical infection.

In Africa, gastrointestinal trematodosis is considered as one of the most important helminth infection in cattle with reported prevalence of 30-90% (Nwosu and Srivastava, 1993; Bunza et al., 2008; Ayalew et al., 2016). Losses due to this infection have been reported by Kithuka et al., (2002) who reported up to \$0.26 million annual loss due in cattle slaughtered in Kenya, Abebe et al., (2010) also reported the loss of \$8,312 annually in Ethiopia and about \$45,271.07 was lost in South Africa between 2010 and 2012 (Ishmael et al., 2017).

Nigerian livestock industry contributes about 5% of the gross domestic product (GDP), and is about a quarter of total agricultural output (BBI, 2004). Losses in the livestock industry due to infection by these parasites consequently affect the overall economy of the country negatively. In Maiduguri Borno State for example, Biu et al., (2006) reported the loss of \$1,415.85 due to the infections among cattle over a 6-year period. Similarly, Uduak (2014) reported the loss of \$1,683.09 in cattle as a result of similar infections in Port Harcourt. If reports from the 36 States and the FCT were available, the enormity of the situation will be clearer.

## **II. Epidemiology**

### **Epidemiology of dicrocoeliosis**

The parasite *dicrocoelium* has a worldwide distribution and this may be due to several factors supporting the distribution and survivability of the parasite. These include environmental factors, intermediate host and definitive host factors. Various species of the parasite have been reported in areas where they are endemic. *Dicrocoelium dendriticum* was reported by Rudolphi (1819) as the specie with widest distribution commonly found Europe, Asia and also North America. Looss (1907) was among the earliest to have reported the endemicity of *D. hospes* in Western Central and East Africa. *D. chinensis* was also reported by Tang et al., (1983) to be distributed in China, Japan and East Siberia. Dispersal and contamination of pastures by eggs of *dicrocoelium* can be by domestic and wild ruminants and also by hares and rabbits (Boray, 1985). The eggs passed through faeces are highly resistant to temperature variations and can persist on pasture for up to 20 months (Urquhart et al., 1996). This may serve as reservoir for infection. Though the eggs can persist for long time in the environment, favourable condition of temperature and moisture supports the hatchability of the egg.

*Dicrocoelium* have wide host-range involving several mollusk species (Manga-Gonzalez et al., 2001). The role played by molluscs in epidemiology of *dicrocoeliosis* is very important as *D. hospes* egg hatching and miracidium liberating only occurs in the intestine of the molluscs intermediate hosts. Moreover, the parasites multiply enormously by asexual reproduction inside the mollusk (Urquhart et al., 1996). Therefore, this increases the likelihood of parasite transmission. The mollusk intermediate hosts are then of paramount importance as both the ingestion of *D. hospes* eggs and survivability of the parasite within depends solely on the mollusk activity. Manga-Gonzalez et al., (2001) reported that development of larval stages in the 1<sup>st</sup> intermediate host can be influenced by specie, age, nutritional state of molluscs, infective dose, ambient temperature and relative humidity.

Schuster (1993) reported on measuring the infection rate for *D. dendriticum* in the snail *Helicella obvia* in Germany for four grazing seasons. He observed fluctuation in the population structure of the snail with small snails predominating from April to June medium sized snails from July to September and largest snails in the spring of the following year. He further asserted that young snails were less involved in the epidemiology than medium-shell diameter snails and largest snails were more susceptible to *Dicrocoelium*. This may be due to active metabolism and good nutritional condition for developing sporocysts in the largest snails (Alunda and Rojo-Vazquez, 1983). Conversely, decrease infection with *Dicrocoelium* may occur after high infection rate in the largest snails. This could be as a result of death of the heavily infected snails in which sporocysts cause

disruption of hepatopancrease (Urquhart et al., 1996), then impairment of reproductive activity and shortening of life expectancy (Schuster, 1992). Maturation of sporocysts follows the snails' life cycle thus becoming active more often in spring (Otranto and Traversa, 2002).

Ants serve as the second intermediate host and their relevance in the epidemiology of dicrocoeliosis is mainly due to their abundance, wide distribution and alteration to their behavior caused by the parasite in their brain which facilitates their ingestion by definitive hosts as they settle on tips of grasses. Manga-Gonzalez et al., (2001) report about 21 ant species from the genus *formica* have been described as receptive to this parasite in different countries. The number and size of metacercariae is directly related to the ant species and sex (Schuster, 1991). Variations in the number of *Dicrocoelium* metacercariae exist among different ant species and even among the same species. This variability could be due to season, as it appears higher in summer (Parachivescu et al., 1976), difference in affinity for slime-balls by the ant species (Loos-Frank., 1978). Plant topping by ants is caused by encystment of the metacercariae matures in the stomach of the ant. Tetany like behavior by infected ants occurs due to decrease in high intensity and temperature which alter the ants behavior and favour ingestion of the parasite by definitive host (Urquhart et al., 1996).

The main definitive hosts of *Dicrocoelium* parasite are herbivores. However, infection has been reported in rodents and primates and even humans (Schuster, 1993). Susceptibility to infection among various animal species differs. Jithendran and Bhat, (1996) reported that sheep are more susceptible to dicrocoeliosis than goat. Animal age and relative susceptibility to the parasite have not been fully elucidated. Manga-Gonzalez et al., (1991) reported higher mean parasitic burden in lambs than in adults while a contrast result was reported by Ducommun and Pfister (1991).

Stress inducing factors such as animal transportation and confinement enhance *Dicrocoelium* egg production probably and inducing immune depression in animals (Sotiraki et al., 1999). Gender is considered as an important epidemiological factor. Susceptibility to infection by the parasite varied greatly among males and females (Phiri et al., 2005). Investigation on the effect of sex on infection with *D. hospes* by Asanji and Williams (1984) who reported that female had more infection than males. The higher rate in females may be because dairy heifers and cows grazed for several seasons, acquiring infections while steers and Oxen spend considerable period of their life in fattening units (Ducommun and Pfister, 1991). Seasonality of infection is favoured by movement of animals from low land to mountain pastures where they become infected by ants and then bring back the infection to low land during winter.

Human infection could be related to consumption of fresh vegetables where infected ants attached or particular Asian food habits involving the consumption of ants (Azizova et al., 1988). Gastrointestinal symptoms due to dicrocoeliosis were reported in a patient in Germany by Rack et al., (2004). Also, Karadag et al., (2005) reported a case of biliary obstruction by *D. dendriticum* in a 65yr old patient.

### **Epidemiology of fasciolosis**

Fasciolosis is caused by *Fasciola hepatica* which commonly predominate in Europe and South America and *Fasciola gigantica* which predominates in the tropics of Africa and Asia (Boray, 1985). There is an overlap in their distribution in the central Asia and East African regions, where hybrid forms of the parasites have been isolated (Mascoma, 2005). The occurrence of the disease is fundamentally linked to the mollusc of the genus *Lymnea* spp which acts as the intermediate host shading the infective stage called metacercaria (Mascoma, 2005). There are three main factors influencing the production of large number of metacercaria necessary for breakout of fasciolosis, these include temperature, moisture and availability of suitable snail habitats.

Temperature plays a vital role in the development of *fasciola* egg. Temperature of about 9.5<sup>0</sup>c favours the development of *F. hepatica* and higher temperature limit above 30<sup>0</sup>c inhibit the development (Rowcliffe and Ollerenshaw, 1960). The variation in the rate of development of the eggs with temperature may be influenced by topography. Dinmik and Dinmik (1962) reported on longer development time of *F. gigantica* egg in the Kenyan high lands due to temperature variation between 10<sup>0</sup>c and 22<sup>0</sup>c. Kendall (1965) also reported that a minimum temperature 10<sup>0</sup>c is needed for the development of *Lymnea truncatula* and maximum growth occurs at 18-27<sup>0</sup>c. At 15<sup>0</sup>c, *F. hepatica* cercaria will complete development in the snail in 80 days, decreasing to less than 20days at 30<sup>0</sup>c (Gettinby and Byrom, 1991). There is no development below 10<sup>0</sup>c and above 20<sup>0</sup>c mortality of the cercaria increases, one infected the snail appears to remain infected for life (Urquhart et al., 1996). Temperature limits of 12-14<sup>0</sup>c is suitable for survival of metacercariae but this is reduced in hot condition. Boray and Enigk (1964) reported that metacercaria can remain viable only 3days at 20<sup>0</sup>c whereas at 10<sup>0</sup>c they can survive longer period of 107days. For prolonged survival, the relative humidity needs to be above 70%. Metacercariae can survive on moist hay for 8 months but few days on silage.

Increased water in the environmental favour both the intermediate host and intermediate stages. Fluke eggs will not develop in fecal mass, moisture is necessary to break up the mass and even the presence of soil may reduce the rate of development of the egg. (Rowcliffe and Ollerenshaw, 1960). Eggs in moist faeces can survive up to 10 weeks, but if the fecal material dries out rapid mortality of the eggs may occur (Ollerenshaw,

1966). Free swimming cercariae released from the snails encyst on vegetation surface by secreting an encyst wall and losing their tail. The longevity of metacercariae depends on moisture and temperature in the environments. Environmental colonization by the snail intermediate host is greatly influenced by moisture when water dries up snail like *Lymnea truncatula* undergo aestivation during which the transmission of the parasite is suspended (Malone and Craig, 1990). In some areas like Australia and Spain, the prevalence of fasciolosis is high due to increase moisture Uriarte et al., (1985). The total fluke counts and fecal egg counts are highest at the beginning of the wet season in West Africa (Schilhorn Van Veen, 1980) and this coincides with the time when snail populations are rising. Large numbers of infected snails are present at the end of the wet season and beginning of the dry season and this support ruminant hosts infection. Chaudhri et al., (1993) gave a similar report on the developmental cycle of snails and infection in cattle and buffalo in India which is related to the rainy season.

The snail intermediate host *Lymnea natalensis* was reported by Ndifon and Ukoli (1989) to be widely spread throughout Africa and also distributed in Nigeria usually confined to permanent water bodies. The snail is commonly found in areas with rainfall over 1000mm (Schilhorn Van Veen, 1980) and tolerates relative high temperatures (Njoku-Tony, 2011) consistent with tropical climates. Andrews (1999) reported that temperature between 10-25<sup>o</sup>c, low elevation areas, presence of flooded and wetlands areas, irrigated rice cultivations and extensive livestock breeding are factors that contribute to the maintenance of the mollusc and spread of fasciolosis. These are important factors for the development of large number of metacercariae which would increase the infection rates (Taylor et al., 2007). In Uganda, Howell et al., (2005) reported on the abundance of *L. natalensis* at lower altitudes below 1800m while *Galba truncatula* (intermediate host of *F. hepatica*) was only found on high altitudes above 3000m. The presence of the two snail intermediate hosts indicates the possibility of finding the two fasciola parasites in Uganda.

In Nigeria, Schilhorn Van Veen (1980) reported that developing *Lymnea* snails are washed away in torrential streams after heavy rain and this may be important in the spread of fasciolosis. He also reported that abundance of snails at the beginning of dry season with peak in the middle of dry season and decreased towards the end of the dry season when ponds and streams dry up. This was supported by Ndifon and Ukoli (1989) who reported that dry season conditions favour snails and this was said to be due to low turbidity reduced currents and substantial growths of algae and macrophytes. According to Taylor et al., (2007), optimal development of fluke eggs to miracidia occurs at the start of the wet season and development within the snail is completed by the end of rainy season. Therefore, shedding of the cercariae coincides with dry season and more animals graze around streams and pond by that time, thereby predisposing them to infections. Ikeme and Obioha (1993) reported that herdsmen migrate in search of water and grazing during the dry season and thousands of cattle often converge on the few ponds, which fail to dry up.

Cattle and other herbage feeding ruminants are the common definitive hosts to the fasciola parasite and also humans (Funatsu and Bargnes, 2001). Elelu et al., (2016) reported the influence of age in the distribution of fasciolosis in cattle where he obtained adult cattle to be more susceptible than weaners Cattle. A similar report was given by Pfukenyi et al., (2006) in Zimbabwe and Nzalawahe et al., (2014) in Tanzania. This was attributed to longer exposure period to infection by the adults (Pfukenyi et al., 2006). Older cattle serve as constant source of *F. gigantica* infection for susceptible young ones (Pfukenyi et al., 2005).

Epidemiology of fasciolosis is significantly influenced by management system (Keyyu et al., 2005). A work conducted in the Lake Chad area by Jean-Richard et al., (2014) reported high prevalence rate in cattle managed extensively in South Western Nigeria. Cattle usually come in contact with snail infected habitat during extensive communal grazing. However, under intensive can be supplied to the cattle thereby minimizing the likelihood of outbreak of fasciolosis.

There are varying reports on the seasonal infection rates of *F. gigantica* across globe. Umar et al., (2009) reported high infection rates in the beginning of the dry season in Nigeria. A similar report was obtained by Phiri et al., (2005) in Zambia who reported on bovine fasciolosis with higher fluke abundance in post rainy season. Also Pfukenyi et al., (2005) reported seasonal variation with increase during the end of the dry season. However, all year occurrence of infection was reported in Southern Nigeria by Gboeloh, (2012) and this was due to favourable climatic condition which favours the development of parasites and the increase in the density of snail intermediate hosts (Nzalawahe et al., 2014).

Human infection with *Fasciola* has been reported in various countries and this infection occurs mainly in rural areas where consumption of aquatic plants such as water-cress is high. A recent case-control study conducted in Pen, found fasciolosis to be positively associated with familiarity with aquatic plants, drinking Alfa juice and dog ownership (Marcos et al., 2006).

### **Epidemiology of Paramphistomosis**

*Paramphistomum* is one of the common parasites in the rumen and reticulum and of sheep, goats, cattle and water buffaloes. Light infection with the parasites doesn't cause serious damage to the animals but massive number of immature *Paramphistomum* can migrate through intestinal tract causing parasitic gastroenteritis with

high morbidity and mortality rates in young animals (Urguhart et al., 1996). Paramphistomosis is widely distributed, but highest prevalence has been reported in tropical and subtropical regions particularly in Africa, Asia, Australia, Eastern Europe and Russia (Rolfe et al., 1991). The major epidemiological variable influencing worm burdens of animals is the infection rate from pastures and it is influenced by the climatic requirement for egg hatching, development and survival of the larvae in pasture (Melaku et al., 2012).

In Nigeria, rumen flukes are commonly seen in the abattoir among slaughtered cattle (Bunza et al., 2008). The snail vector of the disease (*Lymnea*, *Planorbis* and *Bulinus*) has also been reported in Nigeria (Ndifon and Ukoli, 1989, Brown and Kristensen, 1993). The species reported to occur in Nigerian domestic livestock include *Paramphistomum microbothrium*, *Carymerius gregarious*, *Carymerius spatiosus*, *Cotylophorum cotylophorum* (Schillhorn Van Veen et al., 1975) and *Paramphistomum cervi* (Bogatko, 1975). Other species recovered in Nigeria as reported by Dube et al., (2013) include *Cylopylicranocoeilium microbothriodes*. Conditions favourable for natural increases in the population of snail intermediate hosts usually occur at the beginning of and end of rainy season (Mukaratirwa et al., 1996). Shiff (1964) also reported that egg production and abundance of the juvenile snails was high during the beginning and end of the summer rainy season. Therefore, the snail intermediate host populations undergo marked seasonal variations in density with generally low densities during the rainy period and high densities in the post rainy periods (Pfunkenyi et al., 2005). The numerical size of the population is greatly influenced by several factors like flooding, desiccation and temperature (Woolhouse and Chandiwana, 1989).

Outbreaks of disease generally occur in the dry months of the year when the receding water uncovers herbage contaminated with encysted metacercariae. Foster et al., (2008) reported that dispersal of snails by flooding events and changes in farm management practices may be responsible for the apparent emergence of the parasite in the U.K. Similarly, Pfunkenyi et al., (2005) reported that fecal egg count is seen towards the end of the dry season and during the wet months of the year (October to March) in Zimbabwe. Similar report on seasonal pattern of paramphistomosis was also obtained by Pfunkenyi et al., (2005) who reported that the timing of the season pattern may vary based on location, length of the rainy season and grazing habits of the cattle.

Infections with *Paramphistomum* vary between altitudes (high and low). In Zimbabwe Pfunkenyi et al., (2005), reported the prevalence of patent amphistome infections with spatial heterogeneity, varying from 0% to 19.5% on high veld and 0% to 20.5% in the low veld. This difference in patent infected snails in space is attributed to recent river conditions (flood and draughts) and patchy contamination of the water by excreta (Woolhouse and Chandiwana, 1989).

Cattle infection by *Paramphistomum* has been reported in Nigeria, with a highest prevalence of 56%. This may be associated with grazing of animals around river valley which provide suitable breeding sites for snail intermediate hosts off the parasites (Bunza et al., 2008). Overcrowding of animals at grazing sites due to scarcity of feed and watering sites may favour the establishment and spread of paramphistomosis (Ayalwa et al., 2016). Susceptibility to paramphistomosis may vary among cattle of different ages. This was reported to be more in older cattle than young ones according to Ayalwa et al., (2016) in Ethiopia. However, other workers reported no significant difference between age groups (Titi et al., 2010; Khederi et al., 2015). Peak infection of cattle with *Paramphistomum* is seen during October to November (post rainy season) (Ayalwa et al., 2016). During the dry periods, breeding of snails and development of the larval flukes slow down or stop completely and snails undergo a state of aestivation (Soulsby, 1982, Urguhart et al., 1996).

### III. Prevalence of gastrointestinal trematodosis

#### Global prevalence of gastrointestinal trematodosis

The prevalence of trematode infections in cattle in some selected countries of the world is presented in Table 1.

**Table 1: The global prevalence of gastro-intestinal trematodosis in cattle**

| Country   | Species of parasites            | Prevalence (%) | References                      |
|-----------|---------------------------------|----------------|---------------------------------|
| Turkey    | <i>Dicrocoelium dentriticum</i> | 23.6           | Gargili et al., 1999            |
| Sudan     | <i>F. gigantic</i>              | 30.0           | El-manan et al., 2001           |
| Italy     | <i>F. hepatica</i>              | 11.1           | Cringoli et al., 2002           |
| Algeria   | <i>F. hepatica</i>              | 27.0           | Mekround et al., 2004           |
| Zimbabwe  | <i>F. gigantic</i>              | 37.1           | Pfunkenyi and Mukaratirwa, 2004 |
| Zambia    | <i>F. gigantic</i>              | 48.9           | Phiri et al., 2005              |
| Argentina | <i>F. hepatica</i>              | 40.0           | Kleiman et al., 2007            |
| Spain     | <i>F. hepatica</i>              | 71.0           | Paz-silva et al., 2007          |
| Iran      | <i>Dicrocoelium dentriticum</i> | 66.0           | Ahmadi et al., 2010             |
| Brazil    | <i>F. hepatica</i>              | 21.3           | Alves et al., 2011              |
| Ethiopia  | <i>F. gigantic</i>              | 45.3           | Abebe et al., 2011              |
| Australia | <i>F. hepatica</i>              | 17.8           | Duscher et al., 2011            |
| Portugal  | <i>Paramphistome</i>            | 12.0           | Arias et al., 2011              |
| Tanzania  | <i>F. gigantic</i>              | 23.3           | Njoku-Tomy, 2011                |
| Algeria   | <i>Paramphistome</i>            | 45.7           | Titi et al., 2014               |
| Ethiopia  | <i>Paramphistome</i>            | 51.8           | Ayelew et al., 2016             |

|               |               |      |                        |
|---------------|---------------|------|------------------------|
| Spain         | Paramphistome | 6.0  | Ferraras et al., 2014  |
| Belgium       | Paramphistome | 28.0 | Malrait et al., 2015   |
| Netherland    | Paramphistome | 15.8 | Ankum, 2015            |
| India         | Paramphistome | 75.6 | Swarnakar et al., 2014 |
| New Caledonia | Paramphistome | 70.0 | Cauquil et al., 2016   |

It can be seen that trematodosis is endemic in various countries of the world and has been cause of great economic losses in the livestock industry across the globe. India appeared to have the highest prevalence of 75.6% while the least prevalence of 6% was reported in Spain. Reports obtained from other countries are between this range.

### Prevalence of trematodosis in Nigeria

The incidence of trematode infection in Nigeria is presented in Table 2.

**Table 2: Prevalence of gastrointestinal Trematodosis in Nigeria**

| City/State   | Spp of parasite | Prevalence (%) | References                 |
|--------------|-----------------|----------------|----------------------------|
| Borno        | Dicrocoelium    | 18.3           | Nwosu and srivastava, 1993 |
| Plateau      | Fasciola        | 71.6           | Fabiyi and Adeleye, 1982   |
| Sokoto       | Paramphistome   | 56.0           | Bunza et al., 2008         |
| Zaria        | Fasciola spp    | 23.4           | Raji et al., 2010          |
| Abia         | Dicrocoelium    | 3.1            | Amadi et al., 2012         |
| Adamawa      | Fasciola        | 21.8           | Ardo et al., 2012          |
| Maiduguri    | Fasciola        | 14.8           | Biu et al., 2013           |
| Edo          | Fasciola        | 11.5           | Odigie and Odigie, 2013    |
| Ebonyi       | Fasciola        | 37.9           | Ngele and Ibe, 2013        |
| Ibadan       | Dicrocoelium    | 2.5            | Olubukola et al., 2014     |
| Portharcourt | Fasciola        | 1.7            | Uduak, 2014                |
| Kogi         | Dicrocoelium    | 39.0           | Iyaji et al., 2018         |
| Kwara        | Dicrocoelium    | 7.3            | Elelu et al., 2016         |
| Plateau      | Dicrocoelium    | 22.3           | Omowaye et al., 2012       |
| Benin city   | Paramphistome   | 2.2            | Edosomwan & Shoyeni 2012   |
| Ebonyi city  | Paramphistome   | 18.8           | Nwigwe et al., 2013        |
| Kwara        | Paramphistome   | 16.1           | Elelu et al., 2016         |
| Kaduna       | Paramphistome   | 41.7           | Nnabuife et al., 2013      |

Apparently, trematode infections are widespread in the country because there has been report from all the geopolitical zones. The highest prevalence of 71.6% was recorded in Plateau northcentral Nigeria and least prevalence of 1.7% was reported from Portharcourt, southsouth Nigeria. All the other areas recorded values that are between these two extremes

## IV. Diagnosis of trematode infections

### Diagnosis of dicrocoeliosis

This is commonly achieved based on coprological examination for eggs and necropsy findings which is similar to that of fasciolosis since they have common predilection site. Sotiraki et al., (1999) examined the effect of stress on the intensity of egg excretion in sheep where they reported that stressed animals excreted more eggs. Campo et al., (1999) also reported higher excretion of Dicrocoelium dendriticum eggs in faecal samples collected in the afternoon than those collected in the morning from dame animals. Presence of fluke eggs in liver, faeces and bile were compared by Braun et al., (1995) and they found significant differences among them with highest intensity reported in the bile. Thienpont et al., (1979) reported of a modified technique for analysis of bile sample. Postmortem diagnosis is carried out on the basis of postmortem findings at the point of slaughter. Thickening and extension of the biliary ducts as well as hepatic indurations are the most significant macroscopically observable changes in the liver in dicrocoeliosis. In the thickened biliary ducts there are numerous flukes of various sizes, which are sometimes located in the gall bladder (Duchacek and Lamka, 2003). However, Klimas et al., (1994) reported that not all cases of dicrocoeliosis can be recognized at slaughter inspection of meat.

Serological tests for the detection of dicrocoeliosis in sheep and goat was estimated by Jithendran et al., (1996). Counter-current immunoelectrophoresis was reported by Duchacek and Lamka (2003) as most sensitive in the diagnosis of Dicrocoelium dendriticum. This specific and rapid test of epidemiological examination of sheep and goats makes it possible to diagnose dicrocoeliosis already in the prepatent period. The antibodies are detectable with the help of ELISA test (Haralabidis, 1987). High antibodies titer against Dicrocoelium parasite was reported by Ambrosi et al., (1980) to be detected in 4-8 weeks before the appearance of eggs in faeces. Otrando and Traversa (2002) reported that the use of ELISA in detecting dicrocoeliosis is important for

epidemiological studies and also help in early diagnosis leading to treatment and decreasing economic losses. Sandoval et al., (1999) reported on the use of restriction fragment length polymorphism (PCR-RFLP) of mitochondrial genes using common restriction enzymes to study genetic variability of *Dicrocoelium* species.

### **Diagnosis of fasciolosis**

The most common method of diagnosis is by faecal egg counts and pathological lesions in the liver during post mortem examination at the abattoir. Thienpoint et al., (1979) reported a modified technique for diagnosing fasciolosis using bile sample. A survey of cattle fasciolosis in Kwara state as reported by Elelu et al., (2016) showed that fecal analysis had higher prevalence rate than liver examination and this revealed lack of sensitivity of the abattoir method of diagnosis as positive samples are likely to be lost. Therefore, multiple stool sampling or the combination of different diagnostic tests should be considered to enhance diagnostic accuracy (Johansen et al., 2010). Indeed, a single FLOTAC (new multivalent flotation method) can show a considerable higher sensitivity for *Fasciola hepatica* diagnosis multiple Kato-Katzes thick smears, McMaster or sedimentation slides (Duthaler et al., 2010). In the sub-tropical and tropical countries with distinct wet and dry seasons, optimal development of fluke eggs to miracidia occurs at the start of the wet season and development within the snail is completed by the end of rains (Taylor, 2007). The dry season therefore coincides with the snail shedding of cercariae and more animals grazing closer to streams and ponds thereby predisposing them to infection.

Serological diagnostic method to detect antibodies, such as indirect ELISA (Ardo 2013; Aliyu et al., 2014) and direct ELISA (Fagbemi et al., 1997) has been reported in Nigeria. Testing of precipitating antibodies by the use of Agar Gel precipitation test (AGPT) was also reported by Adedokun et al., (2008), this method has been shown to detect more positive cases than faecal and bile egg counts. A polymerase chain reaction technique (PCR) has also been used to detect *Fasciola gigantica* infection status of snail intermediate host (Velusamy et al., 2004, Kaset et al., 2010). Amplification of specific fragment of mitochondrial DNA in fecal samples in sheep by the use of PCR technique as reported by Martinez-Perez et al., (2012) can be of importance in the early detection of *Fasciola hepatica* infection.

### **Diagnosis of Paramphistomosis**

Provisional diagnosis is usually made on history and clinical signs of the disease (anorexia and projectile diarrhea) and the presence of immature Paramphistomes in fluid faeces or at postmortem examination. Faecal examination for eggs at this stage is usually unrewarding as the disease is in the prepatent phase (Urquhart et al., 1996). The immature flukes are conical, pink in color and 1-5 mm long (Soulsby, 1982). The faecal sedimentation technique is the most suitable for identifying the eggs in faeces. The eggs are oval and operculated, resembling that of *Fasciola gigantica* but they are larger and transparent (Zojac and Conboy, 2006). The eggs of the two species are easily distinguished by the addition of a contrast stain like methylene blue. At postmortem there is marked hemorrhagic enteritis with large numbers of immature flukes embedded in duodenal mucosa. Marked fall in total plasma proteins due to increased leakage of plasma albumin as a result of pluck feeding habits of the immature flukes (Kusiluka et al., 1996). Adult flukes are found in the rumen and reticulum during postmortem, they are pear-shaped and red in colour, approximately 1cm long with a sucker at the tip of the cone and another ventrally at the posterior end (Waal, 2011).

The diagnosis of paramphistomosis could be influenced by season as snail intermediate hosts predominate in the monsoon and post monsoon season and this coincides with the season where majority of cases are reported (Swarnakar et al., 2014, Titi et al., 2014). Similarly, Rangel-Ruiz et al., (2003) reported that livestock infected with *Paramphistomum cervi* occurred more frequently during the rainy season. Few cases may be encountered during the dry periods due to interference of the snails breeding and development which slow down or even stop completely (Urquhart et al., 1996).

Enzyme linked immunosorbent assay (ELISA) is being practiced as the most effective diagnostic technique for detection of anti-parasitic antibodies (Shabih et al., 2006). Indirect plate ELISA was also able to demonstrate the antibody titer at different week post infection in experimental cattle as reported by Estuningsih et al., (2004). A similar report was obtained by Kaur et al., (2009) who reported variation in immune response at weekly intervals in all groups of experimental cattle. The use of molecular technique in the identification of paramphistome species has been reported in Asia and some African countries (Lofty et al., 2010). Ankhum (2015) reported that modified Dorsman technique is effective in faecal diagnosis of paramphistomosis.

### **Control and Prevention**

Instituting control against trematodes in ruminant involve the use of various techniques which could be independent or integrated approach (Santos, 2012). These techniques aimed at reduction of the number of the snail intermediate hosts through biological or chemical means, environmental manipulation and grazing management practices (Asrat, 2004). A number of chemotherapeutic agents are in use for the control of trematode infections.

The control of human infection may be achieved by strict control of water cress and other metacercariae carrying aquatic plants for human consumption especially in endemic areas (Mas-coma et al., 2005). Drinking water must be boiled or purified. Intergrated control approaches and inter-sectoral collaboration between public health and veterinary medicine has also been suggested for control (Keiser and Utzinger, 2009).

## V. Conclusion

Gastrointestinal trematode infections are a significant limiting factor in cattle and other ruminant production, therefore development of reliable diagnostic techniques and sustainable strategies for their control is a priority. In order to develop sustainable control, gap in knowledge must be identified to guide researchers. The outcome of such research would provide useful information that will aid in the disease control and this will optimize the production efficiency in order to meet the growing population of Nigeria. Therefore, integrated approach towards the successful control of these infections is required. Thus, it is recommended that farmers should adopt modified grazing practices and government should develop national control policies that will curb this menace.

## References

- [1]. Abebe F., Behabloom M., and Berhanu M. (2011): Major trematode infection of cattle slaughtered at Jimma municipality abattoir and the occurrence of the intermediate hosts in selected water bodies of the zone. *Journal of Animal and Veterinary Advances* **10**(12): 1592-1597.
- [2]. Abebe, R., Abunna, F., Berhane, M., Mekuria, S., Megersa, B. & Regassa, A. (2010). Fasciolosis: Prevalence, financial losses due to liver condemnation and evaluation of a simple sedimentation diagnostic technique in cattle slaughtered at Hawassa Municipal abattoir, Southern Ethiopia. *Ethiopia Veterinary Journal*, **14**, 39-51.
- [3]. Addis M. and Fetene A. (2014): An abattoir survey on the prevalence and Monetary Loss of Fasciolosis Among Cattle Slaughtered at Dangila Municipal Abattoir, Ethiopia. *Journal of Veterinary Medicine and Animal Health*. **6**(12): 309-316.
- [4]. Adedokun O.A., Ayinmode A.B. and Fagbemi B.O. (2008). Seasonal prevalence of Fasciola gigantica infection among the sexes in Nigerian cattle. *Vet. Res.* **2**(1): 12-14.
- [5]. Ahmadi N.A. and Meshkekar M. (2010). Prevalence and long term trend of liver fluke infections in sheep, goats and cattle slaughtered in Khuzastern, Southeastern Iran. *Journal of Paramedical Sciences*, **1**(2): 25-31.
- [6]. Aliyu A.A., Ajogi I.A., Ajanusi O.J. and Reuben R.C. (2014). Epidemiological studies of Fasciola gigantica in cattle in Zaria, Nigeria using coprology and serology. *Journal of Public health and Epidemiology*. **6**(2); 85-91.
- [7]. Alunda J.M. and Rojo F.A. (1983). Effect of infection rate and host age on the intramulluscan development of *Dicrocoelium dendriticum*. *Helminthologia* **20**: 251-258.
- [8]. Alves D.P., Carneiro Junior O.S., Almeida B.R., Avelar B.R. and Leao A.G.C. (2011). Distribution and factors associated with Fasciola hepatica infection in cattle in the south of Espitito Santo States, Brazil. *Journal of Venomous Animals and Toxins Including Tropical Diseases*. **17**, 271-276.
- [9]. Amadi A.N.C., Avoaja D.A. and Essien E.J. (2012): Epidemiology of helminth parasites of West African Dwarf goat (*Capra Hircus*) in Umuariaga in Ikwuano L.G.A., Abia State. *J. Appl. Sci. Environ. Manage.* **16**(4): 359-362.
- [10]. Ambrosi M., Baldelli B., Piergili Fioretti D., Polidori G.A., Grelloni V., Moretti A. and Principato M. (1980). Dicrocoeliosi ovina: insorgenza e decorso della infezione da *Dicrocoelium dendriticum* studiati con metodi parassitologici e sierologici (ELISA) in quattro gruppi di ovine di traccia. *Rivista di Parassitologia*. **41**: 299-307.
- [11]. Andrews S.J. (1999). The life cycle of *Fasciola hepatica*. In: Dalton J. (Ed.). *Fasciolosis*. New York: CAB international, Chapter 1. Pp 1-20.
- [12]. Ankum L. (2015). Presence and pathogenicity of Paramphistomidae in ruminants in the Netherlands. Final report, Utrecht University Repository. <https://dspace.library.uu.nl/handle/1874/306065>.
- [13]. Ardo M.B., Aliyara Y.H. and Lawal H. (2013). Prevalence of Bovine Fasciolosis in Major abattoir of Adamawa State Nigeria. *Bayero Journal of pure and Applied Sciences*, **6**(1): 12-16.
- [14]. Arias M., Lomba C., Dacal G., Vasquez L., Pedreira J., Francisco I., Pineiro P., et al., (2011). Prevalence of mixed trematode infections in an abattoir receiving cattle from northern Portugal and north-west Spain. *Vet. Rec.*, **168**, 408.
- [15]. Asanji M.F. and Williams M.O. (1984) the effect of sex on seasonal variation in single and double infection of cattle in Sierra Leone by *Dicrocoelium hospes* and *Fasciola gigantica*. *Vet. Parasitol.* **15**: 247-255.
- [16]. Asrat, M. (2004): Infection prevalence of ovine in irrigation schemes along the Upper Awash River Basin and effects of strategic anthelmintic treatment in selected upstream.
- [17]. Ayalew G., Tilahun A., Aylate A., Teshale A. and Getachew A. (2016): A Study on Prevalence of Paramphistomum in cattle Slaughtered in Gondar Elfora Abattoir, Ethiopia. *J. Vet. Med. Anim. Health* **8**(8): 107-114.
- [18]. Ayaz S., Ullah R., AbdEl-Salam N.M., Shams S. and Niaz S. (2014). *Fasciola hepatica* in some Buffaloes and Cattle by PCR and Microscopy. *The Scientific World Journal*, 2014.
- [19]. Azizova O.M., Sagieva A.T., Irailova S., Sadykov V.M., Shirinov N.Sh., Mukhitdinov Sh.M., Muuminov A., Ismatov L., Adilova N.B. and Saidaliev T.S. (1988). *Dicrocoelium lanceolatum* infection in man (on autopsy data). *Meditsinskaya Parazitologiya i Parazitarnye Bolezni*, **2**: 26-28.
- [20]. Bekele, M., Tesfey, H. and Getachew, Y. (2010). Bovine Fasciolosis: prevalence and its economic loss due to liver condemnation at Adwa Municipal Abattoir, North Ethiopia. *Ethiopian Journal of Applied Sciences and Technology*, **1**: 39-47.
- [21]. Better Business initiative (BBI) Nigerian Economics Summit Group (NESG) Consultative Form with livestock farmers in northern Nigeria. 2004 [http://www.pak-nigeria.org/pdfs/livestock\\_forum\\_summary.pdf](http://www.pak-nigeria.org/pdfs/livestock_forum_summary.pdf).
- [22]. Bianchin C., Kichel T. and Honer R. (2007). The effect of the control of endo and ecto parasites on weight gains in cross breed cattle in the central region of Brazil. *Trop. Anim. Hith. Prod.* **39**(4): 287-296.
- [23]. Biu A.A., Paul B.T., Konto M and Ya'uba A.M (2013). Cross sectional and phenotypic studies on fasciolosis in slaughter cattle in Maiduguri, Nigeria. *Journal of Agriculture and veterinary sciences volume* **5**(2): 155 – 162.
- [24]. Biu A.A., Ahmed M.I. and Mshelia S.S., (2006): Economic assessment of losses due to parasitic disease in common in Maiduguri abattoir, Nigeria. *African Scientist*; **7**: 143-145.



- [25]. Bogatko W. (1975). Mass mixed infection with *Paramphistomum cervi* and *Fasciola gigantica* in cattle in northern Nigeria. *Medycyna Weterynaryjna*, **31**, 469-470.
- [26]. Boray J.C. (1985). Flukes of domestic animals. In *Parasites, Pests and Predators*. (World Animal Science, B2) (ed. Gaafar, S. M., Howard, W. E. and Marsh, R.E.), pp179-218. Amsterdam, Elsevier Science Publishers B. V.
- [27]. Boray J.C. and Enigk K. (1964). Laboratory studies on the survival and infectivity of *Fasciola hepatica* and *Fasciola gigantica* metacercariae. *Z. Tropenmed. Parasit.* **15**, 324-331.
- [28]. Braun U., Wolfensberger R. and Hertzberg H. (1995). Diagnosis of liver flukes in cows – a comparison of the findings in the liver, in the faeces, and in the bile. *Schweiz Arch Tierheilkde*, **137**: 438-444.
- [29]. Brown D.S. and Kristensen T.K. (1993). A field guide to African fresh water snails: West African species, Danish Bilharziasis Laboratory.
- [30]. Bunza M.D.A., Ahmad A. and Fana S.A. (2008). Prevalence and fluke burden of *Paramphistomiasis* in ruminants slaughtered at Sokoto central abattoir, Sokoto. *Nigerian Journal of Basic and Applied Sciences*, **2**: 287-292.
- [31]. Campo R., Manga-Gonzalez M.Y., Gonzalez-Lanza C., Rollinson D. and Sandoval H. (1999). Characterization of adult *Dicrocoelium dendriticum* by isoelectric focusing. *Journal of Helminthology*, **72**: 109-116.
- [32]. Cauquil L., Hue T., Hurlin J.C., Seale K., Skuce P. and Zadoks R. (2016). Prevalence and sequence-based identity of rumen fluke in cattle and deer in New Caledonia. *PLoS ONE*, **11**(4): e0152603.
- [33]. Chaudhri S.S. (2000). Studies on the incidence and epidemiology of *Paramphistomiasis* in camels and sheep. *Information Bulletin*, **23**: 269-274.
- [34]. Chaudhri S.S., Gupta R.P. and Sangwan A.K. (1993). Helminth diseases in ruminants of Haryana and their control a review. *Agric. Rev. Karnal*, **14**: 121-132.
- [35]. Cringoli G., Rinaldi L., Veneciono V., Capelli G. and Malone J.B. (2002). A cross-sectional coprological survey of liver flukes in cattle and sheep from an area of the Southern Italian Apennines. *Vet. Parasitol.*, **108**, 137-143.
- [36]. Dinmick J.A. and Dinmick N. (1962). The growth of *Paramphistomum microbothrium fiscoeder* to maturity and its longevity in cattle. *Bulletin of Epizootic Diseases of Africa*, **10**: 27-31.
- [37]. Dube S., Onyedineke N.E. and Aisien M.S.O. (2013). Ceylonocotyle, Bothriophoron and Callicophoron species parasitic in some Nigerian cattle. *Advances in BioResearch*, **4**, 38-43.
- [38]. Duchacek L. and Lamka J. (2003). *Dicrocoeliosis* – the present stage of knowledge with respect to Wildlife species. *Acta Veterinaria Brno*, **72**: 613-626.
- [39]. Ducommun D. and Pfister K. (1991). Prevalence and distribution of *Dicrocoelium dendriticum* and *Fasciola hepatica* infections in cattle in Switzerland. *Parasitol. Res.*, **77**, 364-366.
- [40]. Duscher R., Duscher G., Hofer J., Tichy A., Prost H. and Joachim A. (2011). *Fasciola hepatica*- Monitoring the milky way? The use of tank milk for liver fluke monitoring in dairy herds as base for treatment strategies. *Vet. Parasitol.*, **178**, 273-278.
- [41]. Duthaler U., Rinaldi L. and Maurelli M.P. (2010). *Fasciola hepatica*: comparison of the sedimentation and FLOTAC technique for the detection and quantification of fecal egg counts in rats. *Exp. Parasitol.* **126**: 161-166.
- [42]. Edosomwan E.U. and Shoyemi O.O. (2012). Prevalence of gastrointestinal helminth parasites of cattle and goats slaughtered at abattoirs in Benin City, Nigeria. *African Scientist*, **13**(2): 109-114.
- [43]. Elelu, N., Ambali, A., Coles, G. C. & Eisler, M. C. (2016). Cross-sectional study of *Fasciola gigantica* and other trematode infections of cattle in Edu Local Government Area, Kwara State, North-central Nigeria. *Parasites & Vectors*, **9**, 47.
- [44]. Elmanan A.M., Bushara H.O. and Majid A.M. (2001). Some aspect of bovine fasciolosis in northern Gazira and Khartoum State. *The Sud. J. Vet. Res.*, **17**: 35-40.
- [45]. Estuningsih E.S., Widjajanti S., Diwinata G.A. and Piedrafta D. (2004). Detection of Coproantigens by Sandwich ELISA in sheep experimentally infected with *Fasciola gigantica*. *Tropical Biomedicine*, **21**: 51-56.
- [46]. Fabiyi, J.P. and Adeleye, G.A. (1982). Bovine Fascioliasis on the Jos Plateau, Northern Nigeria with particular preference to economic importance. *Bulletin of Animal Health and Production in Africa*, **30**: 41-43.
- [47]. Fagbemi B.O., Aderibigbe O.A. and Guobadia E.E. (1997). The use of monoclonal antibody for the immunodiagnosis of *Fasciola gigantica* infection in cattle. *Veterinary Parasitology*, **69**: 231-240.
- [48]. Ferreras M.C., Gonzalez-Lanza C., Perez V., Fuentes M., Banavides J., Mezo M., Gonzalez-Warleta M., et al., (2014). *Callicophoron daubneyi* (Paramphistomidae) in slaughtered cattle in Castilla Y Leon (Spain). *Vet. Parasitol.*, **191**, 252-263.
- [49]. Foster A.P., Otter A., O'sullivan T., Cranwell M.P., Twomey D.F., Miller M.F. and Taylor M.A. (2008). Rumen fluke (paramphistomosis) in British cattle. *Vet. Record*, **162**: 528.
- [50]. Gargili A., Tuzer E. and Gulamber A. (1999). Prevalence of liver fluke infections in slaughtered animals in Trakya (Thrace), Turkey. *Turkey Journal of Veterinary Animal Science*, **23**, 115-116.
- [51]. Gboeloh L.B. (2012). Seasonal prevalence of *Fasciola gigantica* in slaughter cattle in major abattoirs in Port Harcourt. *Advances in Agriculture, Scientific and Engineering Research*, **2**(9): 336-340.
- [52]. Gettinby G. and Byrom W. (1991). Weather based computer experiment on parasites. *Prev. Vet. Med.*, **11**, 293-308.
- [53]. Haralabidis S. (1987). An ELISA-CELISA double test for multifaceted immunodiagnosis of parasitic diseases. *Helvetic Armed Forces Medical Review*, **21**(5): 137.
- [54]. Hossain M.M., Paul S., Rahman M.M., Hossain F.M.A., Hossain M.T. and Islam M.R. (2011) Prevalence and economic significance of caprine fasciolosis at sylhet district of Bangladesh. *Pak. Vet. J.* **3**(12): 113 – 116.
- [55]. Howell A., Baylis M., Smith R., Pinchbeck G. and Williams D. (2015). Epidemiology and impacts of *Fasciola hepatica* exposure in high yielding dairy herds. *Prev. Vet. Med.* **121**(1-2): 41-48.
- [56]. Ikeme M.M. and Obioha F. (1993). *Fasciola gigantica* infestations in cattle trade in Eastern Nigeria. *Bulletin on Epizootiology and Distribution in Africa*, **21**: 259-264.
- [57]. Ishmael F.J., Borden M., Ezekiel G. and Voster M. (2017). A quantitative assessment of causes of Bovine liver condemnation and its implication for food security in the Eastern Cape Province South Africa. *Sustainability*, **9**, 736, doi:10.3390/su9050736.
- [58]. Iyaji F.O., Yaro C.A., Peter M.F. and Abutu A.E. (2018). *Fasciola hepatica* and associated parasite, *Dicrocoelium dendriticum* in slaughter house in Ayigba, Kogi State, Nigeria. *Advances in Infectious Diseases*, **8**: 1-9.
- [59]. Javed U.K. (2008). Epidemiology, economic importance and therapy of paramphistomosis in cattle and buffaloes in Pakistan.
- [60]. Jean-Richard V., Crump L., Abicho A.A., Nare N.B., Greter-Steinmann H., Hattendorf J., Schelling E. and Zinsstag J. (2014). Prevalence of *Fasciola gigantica* infection in slaughtered animals in Southeastern Lake Chad area in relation to husbandry practices and seasonal water levels. *BMC Veterinary Research*, **10**, 81.
- [61]. Jithendra K. and Bhat T.K. (1996). Prevalence of *Dicrocoeliosis* in sheep and goats in Himachal Pradesh, India. *Veterinary Parasitol.*, **61**: 265-271.

- [62]. Jithendra K.P., Vaid J. and Krishna L. (1996). Comparative evaluation of agar gel precipitation, counterimmunoelectrophoresis and passive haemagglutination test for the diagnosis dendriticum infection in sheep and goats. *Vet. Parasitol.* **61**: 151-156.
- [63]. Johansen M.V., Sithithaworn P. and Bergquist R. (2010). Towards improved diagnosis of zoonotic trematode infections in southeast Asia. *Adv. Parasitol.* **73**: 171-195.
- [64]. Karadag B., Bilici A., Doventas A., Kantarci F., Selcuk D., Dincer N., Oner Y.A. and Erdinler D.S. (2005). An unusual case of biliary obstruction caused by *Dicrocoelium dendriticum*. *Scand. J. Infect. Dis.*, **7(5)**: 385-388.
- [65]. Kasset C., Eursitthichai V., Vichasri-Grams S., Viyanant V. and Grams R. (2010). Rapid identification of Lymnaeid snails and their infection with *Fasciola gigantica* in Thailand. *Experimental Parasitology*, **126**, 482-488.
- [66]. Kaur S., Singla L.D., Hassan S.S. and Juyal P.D. (2009). Standardize application of indirect plate ELISA for immunodiagnosis of paramphistomosis in ruminants. *J. Parasit. Dis.* **33**:70-76.
- [67]. Keiser J. and Utzinger J. (2009). Foot-borne Trematodiasis. *Clinical Microbiology Reviews*, **22**, 466-483.
- [68]. Kendall S.B. (1965). Relationship between the species of *Fasciola* and their moluscan hosts. In "Advances in Parasitology" (ed. Ben Dawes) **Vol. 3**, pp 59-95. Academic Press, London and New York.
- [69]. Keyyu J, Kassuku A., Msalilwa L., Monrad J. and Kyugaard N. (2006): Cross sectional prevalence of helminth infection in cattle on traditional, small scale dairy farms in Iringa district. *Tanz. Vet. Res. Commun.* **30**: 45-55.
- [70]. Khederi J., Radfar M., Borji H. and Mirzaci M. (2015). Prevalence and intensity of *Paramphistomum* spp. in cattle from Southeastern Iran. *Iran Journal of Parasitology*, **10**, 268-272.
- [71]. Kithuka, J.M., Maingi, N., Njeruh, F.M. and Ombui, J.N. (2002). The prevalence and economic importance of bovine fasciolosis in Kenya- an analysis for abattoir data. *Onderstepoort Journal of Veterinary Research*, **69**: 255-262.
- [72]. Kleiman F., Pietrokovsky S., Prepelitchi L., CArbajo A.E. and Wisnivesky-Colli C. (2007). Dynamics of *Fasciola hepatica* transmission in Andean Patagonian Valleys, Argentina. *Veterinary Parasitology*, **145**, 274-286.
- [73]. Klimas M., Schuster R. and Hirschmann R.U. (1994). Vorkommen und Verbreitung von *Dicrocoelium dendriticum* in Nord-West-Thuringen. *Mh Vet-Med.* **49**: 317-322.
- [74]. Kusiluka L.J.M., Kambarage D.M., Harrison L.J.S., Matthewman R.W. and Daborn C.J. (1996). Gastrointestinal helminths of goats and sheep in Tanzania. *Vet. J.* **16**: 53-58.
- [75]. Lofty W.M., Brant S.V., Ashmawy K.I., Devkota R., Mkoji G.M. and Loker E.S. (2010). A molecular approach for identification of *Paramphistomes* from Africa and Asia. *Veterinary Parasitology*. **174**: 234-240.
- [76]. Loos-Frank B. (1978). Zum Verhalten von Ameisen der Gattung *Formica* (Hyemenoptera: Formicidae) gegenüber Schleimballen des Kleinen Leberegels *Dicrocoelium dendriticum* (Digenea: Dicrocoeliidae) und über infektionsbedingte Veränderungen ihrer Hamolymphe. *Entomologica Germanica*. **4**: 12-23.
- [77]. Looss D. (1907). Notitez zur Helminthologie Aegyptiens VII. Über einige neue Trematoden der aegyptischen Fauna Zentral-blatt für Bakteriologie, Parasitkunde. Infektions Krankheiten und Hygiene. Abteilung 1 Originale, **43**: 478.
- [78]. Malone J.B. and Craig T.M. (1990). Cattle liver flukes, risk assessment and control. *Compendium on Continuing Education for Practising Veterinarians*, **12**, 747-754.
- [79]. Malone JB, R. Gommès, J. Hansen, J.M. Yilma, J Slingenberg, F. Snijders F, Nachtet Of, Ataman EA (1998). Geographic information system on the potential distribution and abundance of *fasciola hepatica* and *F. gigantica* in East Africa based on Food and Agriculture organization databases, *Veterinary Parasitology*. **78**, 87-101.
- [80]. Malrait K., Verchave S., Skuce P., Van Loo H., Vercruysee J. and Charlier J. (2015). Novel insight into the pathogenic importance, diagnosis and treatment of the rumen fluke (*Calicophoron daubneyi*) in cattle. *Vet. Parasitol.*, **207**, 134-139.
- [81]. Manga-Gonzalez M.Y., Gonzalez-Lanza C., Cabanase E. and Campo R. (2001). Contributions to and Review of *Dicrocoeliosis*, with special reference to the intermediate hosts of *Dicrocoelium dendriticum*. *Parasitology*. **123**: 91-114.
- [82]. Manga-Gonzalez M.Y., Gonzalez-Lanza C. and Del-Pozo P. (1991). Dynamics of the elimination of *Dicrocoelium dendriticum* (Trematode; Digenea) eggs in the faeces of lambs and ewes in Porma Basin (Leon, NW Spain). *Annales de Parasitologie Humaine et Camparee*, **66**, 57-61.
- [83]. Marcos L., Maco V. and Samalvides F. (2006). Risk factors for *Fasciola hepatica* infection in children: A case control study. *Trans. R. Soc. Trop. Med. Hyg.*, **100**: 158-166.
- [84]. Mas-coma S., Bargues M.D. and Valero M.A. (2005). Fascioliasis and other plant borne trematode zoonoses. *Int. J. Parasitol.*; **35**: 1255-1278.
- [85]. Mekroud, A., Benakhla, A., Vignoles, P., Bondelaud, D., and Dreyfuss, G. (2004). Preliminary studies on the prevalences of natural fasciolosis in cattle, sheep and the host snail (*Galba truncatula*) in northeastern Algeria. *Parasitology Research*, **92**: 502-505.
- [86]. Melaku S. and Addis M. (2012). Prevalence and intensity of *Paramphistomum* in ruminants slaughtered at Debre Zeit industrial abattoir, Ethiopia. *Globe Vet.* **8**:315-319.
- [87]. Morgan E.R., Charlier J., Hendrickx G., Briggeri A., Catalan D., Von Samson-Himmelstjerne G. and Vercruysee J. (2013). Global change and helminth infections in grazing ruminants in Europe: impacts, trends and sustainable solutions. *Agriculture*, **3**(3), 484-502.
- [88]. Mukaratirwa S., Siegismund H.R., Kristensen T.K. and Chandiwana S.K. (1996). Genetic structure and parasite compatibility of *Bulinus globosus* (Gastropoda: Planorbidae) from two areas of different endemicity of *Schistosoma haematobium* in Zimbabwe. *International Journal for Parasitology*. **26**: 269-280.
- [89]. Ndifon G.T. and Ukoli F.M.A. (1989). Ecology of fresh water snails in Southwestern Nigeria, I: distribution and habitat preferences. *Hydrobiologia*. **171**: 231-253.
- [90]. Ngele K.K. and Ibe E. (2014). Prevalence of fasciolosis in cattle slaughtered at Eke market abattoir, Afikpo, Ebonyi state Nigeria. *Animal Research International*, **11**(2): 1958 – 1963.
- [91]. Njoku-Tony R.F. (2011). Bovine Fasciolosis among slaughtered cattle in selected abattoirs in Imo State, Nigeria. *World Rural Observations*. **3**(1):59-63.
- [92]. Nnabuiwe, H. E., Dakul, A. D., Dogo, G. I., Egwu, O. K., Weka, P.R., Ogo, I. N., Onovoh, E. O. & Obaloto, B. O. (2013). A study on helminthiasis of cattle herds in Kachia grazing reserve of Kaduna State, Nigeria. *Veterinary World*, **6**, 936-940.
- [93]. Nwigwe J.O., Njoku O., Odikamnor O. and Uhuo A.C. (2013): Comparative study of intestinal helminth and protozoa of cattle and goats in Abakaliki metropolis of Ebonyi State, Nigeria. *Adva. Appl. Sci. Res.* **4**: 223-227.
- [94]. Nwosu, C.O. and Srivastava, G.C. (1993) Liver flukes infection in Borno State, Nigeria. *Veterinary Quarterly* **15**:182-183.
- [95]. Nzalawahe J., Kassuku A., Stothard J., Coles G. and Eisler M. (2014). Trematode infections in cattle in Arumeru District, Tanzania are associated with irrigation. *Parasite Vectors*. **7**(1): 107.
- [96]. Odigie B.E and Odigie I.O. (2013). Fasciolosis in cattle: A survey of abattoir in Egor, Ikpoba-okha and Oredo LGAs of Edo State using historochemical techniques. *IJB AIR*; **2**(1): 1-9.

- [97]. Ollerenshaw C.B. (1966). The approach to forecasting the incidence of fascioliasis over England and Wales 1958-1962. *Agric. Met.* **3**, 35-53.
- [98]. Olsen A., Frankena K., Rene BØdker N.T., Thamsborg S.M., Enemark H.L. and Halasa T. (2015). Prevalence, risk factors and spatial analysis of liver fluke infections in Danish cattle herds. *Infection*, **6**, 7.
- [99]. Olubokola D.A., Emmanuel C.U., Victor O.A., Oyetunde A.A. and Simeon I.B.C. (2014). Gastrointestinal helminths in slaughtered cattle in Ibadan, South-Western Nigeria. *Journal of Veterinary Medicine*, Volume 2014, Article (II) 923561, 6 pages.
- [100]. Omowaye S.O., Idachaba O.S. and Falola O.O. (2012). The prevalence of parasitic infection in bile of cattle slaughtered in Jos abattoir, Plateau State, Nigeria. *Global Journal of Bio-science and Biotechnology*, **1**, 121-123.
- [101]. Otranto D. and Traversa D. (2002). A review of dicrocoeliosis of ruminants including recent advances in the diagnosis and treatment. *Veterinary Parasitology*, **107**: 317-335.
- [102]. Ozung P.O., Owa P.U. and Oni K.O. (2011). An assessment of the prevalence of fascioliasis of ruminants in Ikom abattoir of Cross River State, Nigeria. *Continental Journal of Veterinary Sciences*, **5**: 1-5.
- [103]. Parachivescu D., Hurginsiu I. and Popescu S. (1976). Bioecologic and biochemical research upon formicidae complementary hosts of *Dicrocoelium lanceolatum* fluke (Stiles and Hassal, 1989). *Archiva Veterinaria*, **11**(12), 159-178.
- [104]. Paz-Silva A., Hillyer G.V., Arias M.S., Sanchez-Andrade R., Pedreira J., Suarez J.L., et al., (2007). A cross-sectional study of fasciolosis in autochthonous cattle from NW Spain by using a 2.9-kDa recombinant protein. *Intern. J. Appl. Res. Vet. Med.*, **5**(2): 52-56.
- [105]. Pfukenyi D.M., Mukaratirwa S., Willingham A.N. and Monrad J. (2006a). Epidemiological studies of *Fasciola gigantica* infections in cattle in the highveld and lowveld communal grazing areas of Zimbabwe. *Onderstepoort Journal of Veterinary Research*, **73**, 37-51.
- [106]. Pfukenyi D.M., Mukaratirwa S., Willingham A.N. and Monrad J. (2006b). Epidemiological studies of *Fasciola gigantica* infections in cattle in the highveld and lowveld communal grazing areas of Zimbabwe. *Onderstepoort Journal of Veterinary Research*, **73**, 179-191.
- [107]. Pfukenyi D.M., Mukaratirwa S., Willingham A.L. and Monrad J. (2005). Epidemiological studies of Amphistome infections in cattle in the high veld and low veld communal grazing areas of Zimbabwe. *Onderstepoort Journal of Veterinary Research*. **72**: 67-86.
- [108]. Pfukenyi, D. and Mukaratirwa, S. (2004): A retrospective study of the prevalence and seasonal variation of *F. gigantica* in cattle slaughtered in the major abattoirs of Zimbabwe between 1990 and 1999. *Onderstepoort Journal of Veterinary Research* **71**(3): 181-187.
- [109]. Phiri A.M., Phiri I.K., Sikazunge C.S. and Monrad J. (2005). Prevalence of fasciolosis in Zambian cattle observed at selected abattoirs with emphasis on Age, sex and origin. *Journal of veterinary Medicine, series B* and **52**(9): 414-416.
- [110]. Rack J., Adusu E. and Jelinek T. (2004). Human infection with *Dicrocoelium dendriticum*. *Dtsch. Med. Wochensch.* **129**(47): 2538-2540.
- [111]. Raji M. A., Salami S.O and Ameh J.A. (2010) Pathological conditions and lesions observed in slaughtered cattle in Zaria abattoir. *Journal of clinical pathology and forensic medicine* Vol. **1**(2). Pp. 9-12.
- [112]. Rangel-Ruiz L.G., Albores-Brahms S.T. and Gamboa-Aguilar J. (2003). Seasonal trends of *Paramphistomum cervi* in Tabasco, Mexico. *Vet. Parasitol.* **116**:217-215.
- [113]. Rolfe P.F., Boray J.C., Nichols P. and Collins G.H. (1991). Epidemiology of paramphistomiasis in cattle. *International J. Parasitol.* **21**: 813-819.
- [114]. Rowcliffe, S.A. and Ollerenshaw, C.B. (1960). Observations on the bionomics of the egg of *Fasciola hepatica*. *Ann. Trop. Med. Parasitol.*, **54**: 172-181.
- [115]. Sandoval H., Manga -Gonzalez Y.M., Campo R., Garcia P., Castro J.M.A. and De La Vega M.P. (1999). Preliminary study on genetic variability of *Dicrocoelium dendriticum* determined by random amplified polymorphic DNA. *Parasitology International*, **48**, 21-26.
- [116]. Santos, M. (2012). Genetic characterization of Portuguese *Fasciola hepatica* isolates instituto de Hygiene e Medicina Tropical, Universidade Nova de Lisboa. MSc thesis (unpublished).
- [117]. Schiff C.J. (1964). Studies on *Bulinus* (*Physopsis*) *globosus* in Rhodesia. III: Bionomics of a natural population existing in a temporary habitat. *Annals of Tropical Medicine and Parasitology*. **58**: 240-255.
- [118]. Schillhorn Van Veen T.W. (1980). Dynamics of *Lymnaea natalensis* populations in the Zaria area (Nigeria) and the relation to *Fasciola gigantica* infections. *Acta Tropica* **37**: 183-194.
- [119]. Schillhorn Van Veen T.W., Shonekan R.A.O. and Fabiyi J.P. (1975). A host-parasite checklist of helminth parasites of domestic animals in Northern Nigeria. *Bulletin of Animal Health and Production in Africa*, **23**, 269-288.
- [120]. Schuster R. (1991). Factors influencing the metacercarial intensity in ants and the size of *Dicrocoelium dendriticum* metacercarial cyst. *Journal of Helminthologie*. **65**: 275-279.
- [121]. Schuster R. (1992). Zur Beeinflussung von *Helicella obvia* durch *Dicrocoelium* - Parthenitae. *Angewandte Parasitologie*. **33**: 61-64.
- [122]. Schuster R.K. (1993). Infection patterns in the first intermediate host of *Dicrocoelium dendriticum*. *Veterinary Parasitology*, **47**(3-4): 235-243.
- [123]. Shabih H.S. and Juyal P. (2006). Diagnosis of *Paramphistomum* in Domestic Ruminants in Punjab (India). *Indian Journal of Animal Sciences*, **42**(4): 272-282.
- [124]. Solomon W. and Abebe W. (2007): Prevalence study of ruminant fasciolosis in areas adjoining upper Blue Nile Basin, North Western Ethiopia. *Ethiop. Vet. J.* **11**: 68-83.
- [125]. Sotiraki S.T., Leontides L.S. and Himonas C.A. (1999). The effect of transportation and confinement stress on egg production by *Dicrocoelium dendriticum* in sheep. *J. Helminthol.* **73**: 337-339.
- [126]. Soulsby E.J.L. (1982): *Helminthes, Arthropods and Protozoa's of domestic Animals* 7<sup>th</sup> ed. Bailliere Tindall, London. P 836.
- [127]. Swarnakar G., Kumawat A., Sanger B., Roat K. and Goswami H. (2014): Prevalence of amphistome parasites (*Trematoda: Digenea*) in Udaipur of Southern Rajasthan, India. *Int. J. Curr. Microbial. App. Sci.* **3**(4): 32-37.
- [128]. Tang C. Tang Z., Tang L., Cui Q., Lu H. and Qian Y. (1983). Studies on the biology and epizootics of *Dicrocoelium chinensis* in Eastern inner Mongol Autonomous region. *Acta Zoologica Sinica*, **29**: 340-349.
- [129]. Taylor M.A., Coop R.L. and Wall R.L. (2007). *Veterinary Parasitology*. 3<sup>rd</sup> ed. Oxford: Blackwell Publishers. Pp 113-139.
- [130]. Thienpont D., Rochette F. and Vanparijs O.F.J. (1979). Diagnosing helminthiasis through coprological examination. Janssen Research Foundation, Beerse Belgium. 112.
- [131]. Titi A., Mekroad A., Sedraoui S., Vignoles P. and Rondelaud D. (2010). Prevalence and intensity of *Paramphistomum daubunyi* infections in cattle from Northeastern Algeria. *Journal of Haematology*. **84**: 177-181.

- [132]. Titi A., Mekroud A., Chibat M., Boucheikhchoukh M., Zein-Eddine R., Djuikwo-Teukeng F.F., Vignoles P., Rondelaud D. and Dreyfuss G. Ruminal (2014). Paramphistomosis in cattle from Northeastern Algeria: Prevalence, Parasite burden and Species Identification. *Parasite*, **21**, 50.
- [133]. Uduak, A. (2014). Incidence of Bovine Fasciolosis and its Economic implications at Trans-Amadi Abattoir Port Harcourt, Nigeria. *Acta parasitological Globalis*, **5**(3): 206-209.
- [134]. Umar A.G., Nwosu C.O. and Philip H.R. (2009). Seasonal prevalence and economic importance of Bovine Fascioliasis in Jalingo abattoir, Taraba State Nigeria. *Nig. Vet. J.* **30**: 44-50.
- [135]. Uriarte J., Cabaret J. and Tanco J.A. (1985). The distribution and abundance of parasitic infections in sheep grazing on irrigated or non-irrigated pastures in North-Eastern Spain. *Ann. Rech. Vet.* **16**(4): 321-325.
- [136]. Urquhart GM Amous J. Duncan JL Dunn AM. And Jennings FW (1996): *Veterinary Parasitology 2<sup>nd</sup> Edn.*, Black Well Science Ltd, London, UK., pp. 307.
- [137]. Velusamy R., Singh B.P. and Raina O.K. (2004). Detection of Fasciolagigantica infection in snails by polymerase chain reaction. *VeterinaryParasitology*, **120**, 85-90.
- [138]. Vercruyysse J. and Claerebout E. (2001). Treatment vs non-treatment of helminth infections in cattle: Defining the threshold. *Vet. Parasitol.*, **98**: 195-214.
- [139]. Waal D.T. (2011). Paramphistomum- a brief review. *Irish Vet. J.* **63**: pp; 5.
- [140]. Woolhouse M.E.J. and Chandiwana S.K. (1989). Spatial and temporal heterogeneity in the population dynamics of *Bulinus globosus* and *Biomphalaria pfeifferi* and in the epidemiology of the infection with Schistosomes. *Parasitology*. **98**: 21-34.
- [141]. Zajac A.M. and Conboy G.A. (2006). *Veterinary clinical parasitology (7<sup>th</sup> edition)* Blackwell Publishing, Oxford. Pp. 320.

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