**Prevalence of Ecto- and Gastrointestinal Parasites of *Hemidactylus Frenatus* (Schlegel) in Akure North Local Government Area of Ondo State, Nigeria**

**Abstract**

The wall gecko is found in tropical, subtropical, and warm temperate regions. It is a nocturnal species that seeks shelter during the daytime and emerges at dusk to forage throughout the night. This small lizard has become established in areas near human habitation in the tropics, forming a close association with people. Like other vertebrates, wall geckos are susceptible to parasitism. This study investigated the prevalence and intensity of ecto- and gastrointestinal parasites infecting *Hemidactylus frenatus* (wall geckos) in Akure North Local Government Area, Ondo State, Nigeria. A total of 360 geckos were sampled across six communities and examined for parasites using standard parasitological techniques. Overall, 68.9% of *H. frenatus* were infected with ectoparasites, while 66.4% harbored gastrointestinal parasites. The most prevalent ectoparasite was *Trombicula sp*. (35.8%), and *Parapharyngodon sp.* (37.5%) was the dominant gastrointestinal parasite. Cryptosporidium spp. was detected in 6.9% of the geckos using modified Ziehl-Neelsen staining. Ectoparasites were primarily located on the back and limbs, while gastrointestinal parasites were most abundant in the large intestine. Chi-square analysis revealed no statistically significant differences in parasite prevalence across sampling locations and sex (P > 0.05). These findings highlight the high parasite burden in *H. frenatus* and the potential public health risks associated with their close proximity to human dwellings.

**Key words:** Ectoparasites, gastrointestinal parasites, gecko, *Hemidactylus frenatus,* intensity, prevalence, zoonotic risk,

**1. INTRODUCTION**

*Hemidactylus frenatus*, commonly known as the wall gecko, is a widespread reptile belonging to the family Gekkonidae. It thrives in warm climates across the globe (Keller *et al*., 2002; Uetz, 2010; Djomnang *et al*., 2016). This small lizard has become highly adapted to human environments, especially in tropical regions, where it is frequently found living in close proximity to people (Oluwafemi *et al*., 2017). The common name “wall gecko” reflects its frequent presence on the walls and ceilings of human dwellings (Petren and Short, 2012).

In Nigeria, wall geckos are a familiar sight in almost every household (Nwachukwu *et al*., 2014), and over 140 gecko species have been recorded in Nigeria and other parts of West Africa (Oluwafemi *et al*., 2017).

While wall geckos contribute to controlling insect populations within homes, they are also known to harbour a variety of ecto- and gastrointestinal parasites. Reported parasites include nematodes (Ameh and Ajayi, 2005; Obi *et al*., 2013; Oluwafemi *et al*., 2017), cestodes (Oluwafemi *et al*., 2017), protozoans (El-Toukhy *et al*., 2013), ticks (Ameh, 2005), and mites (Rivera, 2003; Ameh, 2005). Their close interactions with humans and domestic animals raise public health concerns, as geckos may act as reservoirs or vectors of zoonotic pathogens. For example, ticks found on geckos have been implicated in the transmission of diseases such as babesiosis, borreliosis, rickettsiosis, encephalitis, and helminth infections (Djomnang *et al*., 2016).

The medical importance of these parasites is well established. Ticks are vectors of serious illnesses such as Lyme disease, caused by Borrelia burgdorferi (Schall *et al*., 2000). Mites, particularly those in the genus *Geckobiella*, have been found on fence lizards and are known to transmit coccidian blood parasites such as *Schellackia occidentalis*. Additionally, protozoa such as *Cryptosporidium spp*. have been detected in wall geckos and are known to cause cryptosporidiosis in humans (Deming *et al*., 2008).

Although wall geckos play an ecological role in controlling insect pests, their presence in human habitations also poses potential health risks due to their susceptibility to parasitic infections and their role in the transmission of zoonotic agents (Abbas and Habeeb, 2022). Understanding their parasitic burden is therefore essential, particularly in residential settings where close contact is common.

While several studies have been conducted in other parts of Nigeria on the parasitic fauna of wall geckos, data on the prevalence and diversity of parasites affecting *H. frenatus* in Akure North Local Government Area of Ondo State remain scarce. This study, therefore, aims to investigate the ecto- and gastrointestinal parasites of wall geckos in this region, with an emphasis on their potential zoonotic implications for human and animal health.

**2. MATERIALS AND METHODS**

**2.1 Study Area**

The study was conducted in Akure North Local Government Area of Ondo State, Nigeria. This region lies in the southwestern part of the country and is bordered by Ogun State and the Atlantic Ocean to the south. Six communities were randomly selected for specimen collection (Figure 1): Itaogbolu (N 7° 22' 53", E 5° 15' 9"), Ogbese (N 7° 15' 19", E 5° 22' 21”), Araromi (N 7° 16' 23", E 5° 17' 04"), Oba-Ile (N 7° 15' 53", E 5° 14' 88"), Sha-Sha (N 7° 16' 35", E 5° 15' 25"), and Owode (N 7° 16' 04", E 5° 16' 48").



**Figure 1:** **Map of Akure North Local Government Area Showing locations of Sample collection**

**2.2 Courtesy Visit and Informed Consent**

Prior to sample collection, visits were made to the selected communities to seek permission and cooperation. Informed consent was obtained from residents of selected houses and shop owners, allowing access for specimen collection.

**2.3 Collection of Specimens**

A total of 360 wall geckos (*Hemidactylus frenatus*) were collected between 19:00 and 23:00 hours over a one-year period from March 2020 to March 2021. Collections were made from 20 residential houses and shops (three geckos each) across the six selected communities. The collection method followed the procedure described by Oluwafemi *et al*. (2017). Geckos were captured using forceps and placed in sealed but ventilated transparent containers. The specimens were then transported to the Parasitology and Public Health Research Laboratory in the Department of Biology, Federal University of Technology, Akure, Ondo State, Nigeria. Safety precautions, including the use of gloves, boots, and torchlights for illumination, were strictly observed during the collection process. Specimens were identified using the keys provided by Patel *et al*. (2016).

**2.4 Examination of Wall Gecko for Parasites**

**2.4.1 Examination of Skin for Ectoparasites**

Collected wall geckos were first weighed and measured to classify them into adults (male and female). Each specimen was examined for ectoparasites 12–20 hours after collection. The entire body surface was inspected under a dissecting microscope, with particular attention paid to common predilection sites such as the head, neck, limbs, and base of the tail. Ectoparasites were carefully removed using fine combs and soft forceps, then transferred into Petri dishes containing 70% ethanol for preservation. Subsequently, specimens were mounted on microscope slides using appropriate mounting media.

Identification was carried out under a binocular microscope at magnifications of ×40, ×100, and ×400, based on key morphological features. For ticks, identification was guided by characteristics such as body size, scutum size and coloration (ornate or inornate), mouthpart length (short or long), presence or absence of eyes, body shape, and engorgement status. Mite identification was based on features including body segmentation, shape and arrangement of legs, presence and location of setae (bristles), gnathosoma structure (mouthparts), and the position of the anal and genital openings.

**2.4.2 Examination of Gastrointestinal Tract for Endoparasites**

Each wall gecko was dissected by making a ventral incision using tongs and scissors. The gastrointestinal tract was removed and separated into two sections: the small intestine and the large intestine. The intestinal walls were gently scraped with forceps to extract any attached parasites. The intestinal contents were diluted with normal saline and repeatedly washed by decantation until a clear suspension was used for easier parasite detection.

**2.4.3 Detection of *Cryptosporidium* via Modified Ziehl-Neelsen (MZN) Staining**

To detect *Cryptosporidium* oocysts in faecal samples, a Modified Ziehl-Neelsen staining method was utilized. Approximately 0.2 grams of each faecal specimen was applied to a clean microscope slide using an applicator stick and spread into a thin, even smear. The prepared slides were then dried on a slide warmer set at 60 °C for about 5 minutes to ensure fixation.

The dried slides were arranged on a staining rack and covered with Kinyoun’s carbol fuchsin stain for one minute. After staining, the slides were gently rinsed with distilled water and allowed to drain. Decolorization was performed using 1% acid-alcohol for two minutes, followed by a second rinse with distilled water.

Next, the smears were counterstained with methylene blue for two minutes. After a final rinse with distilled water, the slides were air-dried again on the slide warmer at 60 °C for an additional 5 minutes.

Microscopic examination was conducted using oil immersion at 100× magnification. *Cryptosporidium* oocysts were identified based on their distinct appearance, staining bright red to pink and standing out clearly against the blue-stained background (Sunnotel *et al*., 2006).

**2.5 Identification of Parasites**

All recovered parasites were identified using standard identification keys provided by Krantz and Walter (2009) and Bush *et al.*, (1997) for gastrointestinal parasites.

**2.6 Data Analysis**

The prevalence and mean intensity of infection were calculated based on host sex, age, and anatomical location of the parasites. Statistical significance was assessed using Chi-square (χ²) analysis at a 95% confidence level (P < 0.05). All statistical analyses were performed using Microsoft Excel 2019 and the Statistical Package for the Social Sciences (SPSS) version 20.0.

**3.0 Result**

**3.1 Composition of the *Hemidactylus frenatus* Specimens collected in the study Sites**

A total number of 360 *H. frenatus* specimens weighing between 1.7g and 5g were collected from the study sites. Their lengths varied between 40 mm and 145 mm. The specimens comprised 184 males and 176 females (Figure 2).

**Figure 2: Distribution of *H. frenatus* collected from the Study area in relation to Site and Sex**

Figure 2 shows the distribution of wall gecko in the study sites.

**3.2 Prevalence of Infection of geckos sampled from the study Area**

Of the 360 wall geckos examined, 25 (6.9%) tested positive for *Cryptosporidium spp*. using the Modified Ziehl-Neelsen (MZN) staining technique. In terms of ectoparasitic infections, the highest prevalence was recorded in geckos from Sha-Sha (80%), followed by Ita-Ogbolu and Ogbese, each with a prevalence of 65%. The lowest prevalence was observed in specimens from Oba-Ile, where 50% of the geckos were infected (Table 1). Despite the variation in prevalence among locations, Chi-square analysis revealed no statistically significant difference (P > 0.05) in ectoparasite infection rates across the six study sites.

Similarly, the highest prevalence of gastrointestinal parasites was found in geckos from Sha-Sha (80%), followed by those from Owode (75%). The lowest prevalence was recorded in Araromi (45%). Chi-square test results also indicated no significant difference (P > 0.05) in the prevalence of gastrointestinal parasites in *H. frenatus* across the six study locations (Table 1).

**Table 1: Overall Prevalence of Ecto- and Gastrointestinal Parasite Infection of Wall Geckos from different location in Akure North Local Government Area**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Parasite** | **Location** | **Number****Examined** | **Number Infected** **(%)** | **χ2** | **Df** | **P value** |
| **Gastrointestinal** | **Araromi** | 60 | 27 (45) | 9.59 | 5 | 0.09 |
| **Ita-Ogbolu** | 60 | 30 (50) |  |  |  |
| **Oba-Ile** | 60 | 30 (50) |  |  |  |
| **Ogbese** | 60 | 28 (70) |  |  |  |
| **Owode** | 60 | 30 (75) |  |  |  |
| **Sha-Sha** | 60 | 48 (80) |  |  |  |
| **Ectoparasite** | **Araromi** | 60 | 39 (65) | 4.60 | 5 | 0.47 |
| **Ita-Ogbolu** | 60 | 39 (65) |  |  |  |
| **Oba-Ile** | 60 | 30 (50) |  |  |  |
| **Ogbese** | 60 | 39 (65) |  |  |  |
| **Owode** | 60 | 33 (55) |  |  |  |
| **Sha-Sha** | 60 | 48 (80) |  |  |  |

Table 1 shows the overall Prevalence of Parasitic Ecto- and Gastrointestinal Parasite Infection of Geckos from different location in Akure North Local Government Area

**3.3 Prevalence and Mean Intensity of Parasite of geckos sampled from the study Area**

Of the 360 *H. frenatus* examined, 68.9% were infected with ectoparasites and 66.4% with gastrointestinal parasites. *Trombicula sp.* had the highest ectoparasite mean intensity, particularly on the back and limbs. Among gastrointestinal parasites*, Parapharyngodon sp*. showed the highest intensity, mainly in the large intestine. These results indicate a high parasite burden with potential public health implications (Table 2).

**Table 2: Prevalence and Mean Intensity of Parasite of geckos sampled from the study Area**

Table 2 shows the prevalence and mean intensity of the parasites recovered from wall geckos.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | **Parasite Species** | **Number of Wall Geckos examined** | **Number of Wall Geckos Infected (%)** | **Number of Parasites Recovered**  | **Mean Intensity** | **χ2** | **Df** | **P value** |
| **Gastrointestinal Parasites** | ***Parapharygodon sp.*** | 360 | 135 (37.5) | 279 | 2.12±0.08 | 4.27 | 2 | 0.12 |
|  | ***Pharygodon sp.*** | 360 | 87 (24.2) | 216 | 2.45±0.21 |  |  |  |
|  | ***Oochoristica javaensis*** | 360 | 108 (30.0) | 258 | 2.45±0.15 |  |  |  |
| **Ectoparasites** | ***Argas* sp.** | 360 | 78 (21.7) | 174 | 2.30±0.19 | 6.01 | 2 |  |
|  | ***Trombicula sp.*** | 360 | 129 (35.8) | 363 | 2.30±0.21 |  |  |  |
|  | ***Gekobia sp.*** | 360 | 96 (26.7) | 303 | 3.20±0.36 |  |  |  |

**3.4 Prevalence of Parasites in Relation to Sex of *H. frenatus***

Figure 3 shows the prevalence of parasites in both male and female *H. frenatus* sampled from all the location. The result obtained showed that the overall prevalence in the study area was higher in male compared to female for both ecto- and gastrointestinal parasites.

**3.5 Distribution of Parasites of *H. frenatus* in Relation to Predilection sites**

Ectoparasites showed distinct site preferences (Figure 3). *Geckobia* *sp* were most prevalent on the limbs (n = 141), followed by the back (n = 105), head (n = 30), and armpit (n = 27). *Trombicula* mites were primarily found on the back (n = 237), then limbs (n = 78), tail (n = 42), and abdomen (n = 6). Argas ticks favored the back (n = 63), limbs (n = 51), and head (n = 30). Overall, the back (n = 405) and limbs (n = 270) were the most infested regions.

Gastrointestinal parasites (Figure 5) totaled 753 across all samples. *Oochoristica javaensis* was restricted to the small intestine (n = 258). *Pharyngodon sp* were found mainly in the large intestine (n = 183), and in the colon (n = 18) and small intestine (n = 15). *Parapharyngodon* *sp* were recovered from the large intestine (n = 234) and colon (n = 45). The large intestine showed the highest overall parasite prevalence.

**Figure 3: Parasitic Infections of Wall Geckos in Akure North Local Government Area according to Sex of Host**

Figure 3 shows the prevalence of parasites between male and female wall gecko.

**Figure 4:** **Distribution of Ectoparasites in Relation to Predilection Sites on *H. frenatus***

**Figure 5: Distribution of Gastrointestinal Parasites in Relation to Predilection Sites on *H. frenatus***

Figure 4 and 5 shows the distribution of ecto- and gastrointestinal parasites in their different predilection sites.

**4. Discussion**

Wall geckos (*Hemidactylus frenatus*) are highly susceptible to ectoparasitic and gastrointestinal infections. This study confirms the presence of parasites in geckos across all study locations. The highest ectoparasite prevalence (80%) was recorded in Sha-Sha, likely due to poor sanitation and abundant insect hosts. The lowest (50%) in Oba Ile may reflect cleaner environments, consistent with previous findings (Obi *et al*., 2013; Djomnang *et al*., 2016). Similarly, reduced prevalence in Araromi could result from fewer intermediate hosts (Ameh and Ajayi, 2005; Oluwafemi *et al*., 2017).

Three ectoparasite genera; *Argas*, *Trombicula*, and *Geckobia*, were identified, consistent with previous studies (Rivera *et al*., 2003; Ameh, 2005). Environmental factors like climate and temperature likely influence their distribution. *Trombicula* showed the highest prevalence (43.2%) and abundance (363), followed by *Geckobia sp.* (36.1%, 303), and *Argas* (20.7%, 174). Differences from other studies may result from variations in detection methods.

Habitat type and wide foraging behavior likely contribute to parasitism (Clopton and Gold, 1993; Ameh, 2005). Multiple ectoparasite species were often found on a single host. Although some studies report equal susceptibility across sexes, this study supports male-biased parasitism due to increased exposure from behaviors like foraging and mating (Duneau *et al*., 2012; Brown and Symondson, 2014).

Parasites were most abundant in skin folds and scaled areas. *Argas* and *Trombicula* were common on the back, while *Geckobia* favored limbs (Ameh, 2005). *Trombicula* and *Geckobia* can parasitize other vertebrates, including humans, causing skin irritation. *Argas* ticks are medically important vectors of diseases like West Nile virus and Lyme disease (Demessie and Derso, 2015).

Three gastrointestinal parasites were identified: *Parapharygodon sp.* (37.1%), *Oochoristica javaensis,* and *Pharyngodon sp.* No trematodes or acanthocephalans were found, possibly due to the absence of required hosts. Findings on prevalence align with Oluwafemi *et al.* (2017), though Djomnang *et al*. (2016) found more *Pharyngodon.* O. javaensis may suppress nematode numbers by damaging tissues at attachment sites (Anu, 2018). Similar to ectoparasites, gastrointestinal parasite prevalence was higher in male geckos, likely due to increased exposure from active behaviors like hunting and mating. This supports the idea that behavioral differences, more than immune function, drive parasite transmission.

*O. javaensis* was found in the small intestine; nematodes were in the large intestine and rectum, consistent with prior studies (Oluwafemi *et al*., 2017; Obi *et al*., 2013; Djomnang et al., 2016). The small intestine offers a nutrient-rich environment suitable for cestodes, while the large intestine provides shelter in undigested food residues for nematodes (Djomnang *et al*., 2016).

The detection of *Cryptosporidium spp*. in 6.9% of samples is particularly noteworthy, given its zoonotic potential and role in causing gastrointestinal illnesses in humans. While variations in infection prevalence were observed across sex, sites, and locations, these differences were not statistically significant, suggesting a widespread and uniform risk across the region (Deming *et al*., 2008).

Clinically, these gastrointestinal parasites can cause significant health issues such as inflammation, diarrhea, malnutrition, weight loss, anemia, and developmental delays, especially in young hosts (De-Silva *et al*., 2003; Brooker *et al.*, 2006).

**Conclusion**

This study confirms a high prevalence of both ectoparasitic and gastrointestinal infections in *Hemidactylus frenatus* across different locations, with significant variation influenced by environmental sanitation, host behavior, and habitat characteristics. Sha-Sha showed the highest infection rates, likely due to poor hygiene and abundant insect vectors, while Oba Ile and Araromi showed lower prevalence, suggesting better environmental conditions.

*Trombicula* species were the most common ectoparasites, and *Parapharygodon sp*. was the most prevalent gastrointestinal parasite. Male and adult geckos exhibited higher parasite loads, likely due to increased exposure from behavior and ecological range. Parasite distribution varied by body region and intestinal section, highlighting niche preferences. The presence of zoonotic species like *Cryptosporidium* and *Argas* ticks demonstrate potential public health implications. Overall, environmental management and further research are essential for understanding and mitigating parasitic transmission in urban reptile populations.

**Consent:**

Written informed consent was obtained from residents of selected houses and shop owners, allowing access for specimen collection.

**Disclaimer (Artificial intelligence)**

Author(s) hereby declares that NO generative AI technologies such as Large Language Models (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the writing or editing of this manuscript.

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