# Longitudinal Survey of Astrovirus infection in different bat species in Zimbabwe: Evidence of high genetic *Astrovirus* diversity

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**Abstract**

Astroviruses (AstVs) have been discovered in over 80 animal species including diverse bat species and avian species. A study on *Astrovirus* circulation and diversity in different insectivorous and frugivorous chiropteran species roosting in trees, caves and building basements was carried out at 11 different sites across Zimbabwe. Pooled and individual faecal sampling methods were used for this study, collection date ranged from June 2016 to July 2021. In two sites, Magweto and Chirundu, sampling was carried out at monthly intervals from August 2020 to July 2021. Astroviruses and bat mitochondrial genes were amplified using pan-AstVs and CytB /12S RNA PCR systems respectively. Phylogenetic analysis of the *RdRp Astrovirus* sequences revealed a high genetic diversity of astroviruses. All the bat astroviruses tested in this study clustered with the *Mamastrovirus* genus. Two distinct groups of the amplified sequences were identified. One group comprised of sequences isolated from *Hipposideros, Rhinolophus* and *E. helvum spp* clustered with Human Astrovirus strains: *HuAstV* types1-6, *HuAstV*-MLB1-3 and *HuAstV*-VA-1. A second group comprised of the majority of the sequences clustered with different strains of Bat AstVs. Results from the longitudinal study at Magweto and Chirundu showed an overall AstV prevalence of 13.7% and 10.4% respectively. Peaks of AstV infection at Chirundu coincided with the period when juveniles are 4-6 months old. Coinfection of bats with CoVs and AstVs at Magweto and Chirundu sites was 2.6% and 3.5% respectively.

## Introduction

*Astrovirus* are single-stranded positive sense RNA (+ssRNA) viruses. They are non-enveloped with an icosahedral morphology and a genome length of approximately 6.2 to 7.7 Kb [1]. They infect a wide variety of both domestic and wild marine and terrestrial mammals, including humans (*Mamastrovirus*) as well as avian hosts (*Avastrovirus*) [2]. Recently *Astrovirus* infections in fish and insects have been reported [3,4]. Bats and wild birds are considered natural reservoirs of astroviruses [5]. Astroviruses have been found to occur in over 80 non-human species including a diverse number of bat species from Europe, Africa, America and Asia. Human astroviruses (*HuAstVs*) have been identified as causal agents of acute viral gastrointestinal illness worldwide particularly in children, immunocompromised people and the elderly [3–5]. Beyond this well-known clinical manifestation of *Astrovirus* infections, neurovirulent *Astrovirus* infections have also been reported in both humans and domestic animals [6,7]. To note, in humans, the majority of *Astrovirus*-associated encephalitis or meningitis were reported in immunocompromised people [8–10]. According to the International Committee on Taxonomy of Viruses (https://talk.ictvonline.org), *Mamastrovirus* are classified within 19 recognized species, *MAstV*-1 to -19, and two genogroups GI and GII. All classic and novel Human Astrovirus belong to four different species, *MAstV*-1, *MAstV*-6, *MAstV*-8 and *MastV*-9 [2,11,12]. Astroviruses do not seem to have a common reservoir [12]. However, numerous cross-species transmissions of *Astrovirus* have been documented [4]. More particularly, certain *HuAstV* strains are closely related to rodent, feline, mink, ovine and porcine *Astrovirus* [2,4,12].

To date, no human *AstV* strains have been associated with Bat *AstV*. Bat *Astrovirus* belong to *MAstV*-12, and *MastV*-14 to -19 species. Bat-borne viruses represent an extensive research field owing to the plethora of viruses carried out by the Chiropterans. Bats are known to be persistently infected by astroviruses [13]. This is inclusive of many species of insectivorous bats which harbor these viruses [14]. To date, a high diversity of Bat *Astroviruses* from different bat families has been reported worldwide [9,15]. In Europe a variety of bat species belonging to the Yangochiroptera suborder have been discovered to harbor several astroviruses [16]. In Africa, bat astroviruses were reported in Egypt, Gabon, Madagascar, Kenya and Mozambique [5,15,17–19]. Astroviruses are known to occur in bat species with high prevalence and exceedingly high genetic diversity whereby infections in these hosts are minimally pathogenic [20,21]. Despite all these studies, large gaps still exist to map the extent of existence of these viruses in bats' populations [22].

*Astrovirus* prevalence and detection in bat colonies seemed rather correlated to abiotic factor factors such as seasons and year of sampling than biotic factors such as sex and reproductive status or yet viral co-infection [23]. Viral shedding in bats occurs in spatial and temporal pulses that can drive spillover to other animals or humans [24] Shedding pulses in bats determine varying degrees in the

prevalence of viruses where if they do not occur, viruses are rarely detected or at low prevalence and when they occur, higher prevalence is detected [24].

In this study we enlarge the spectrum of bat *Astroviruses* knowledge in Africa by identifying and analyzing Bat *Astroviruses* from different bat species colonies according to season in Zimbabwe.

## Material and Methods

### Sampling approaches and sites

Two different approaches were followed in this study: bat community sampling and individual bat sampling.

#### *Bat community pooled sampling*

Between February 2016 and December 2020, faecal samples from both insectivorous and frugivorous bat species were respectively collected in different sites including caves, an ancient mine and trees in Zimbabwe (Figure 1, Table 1).

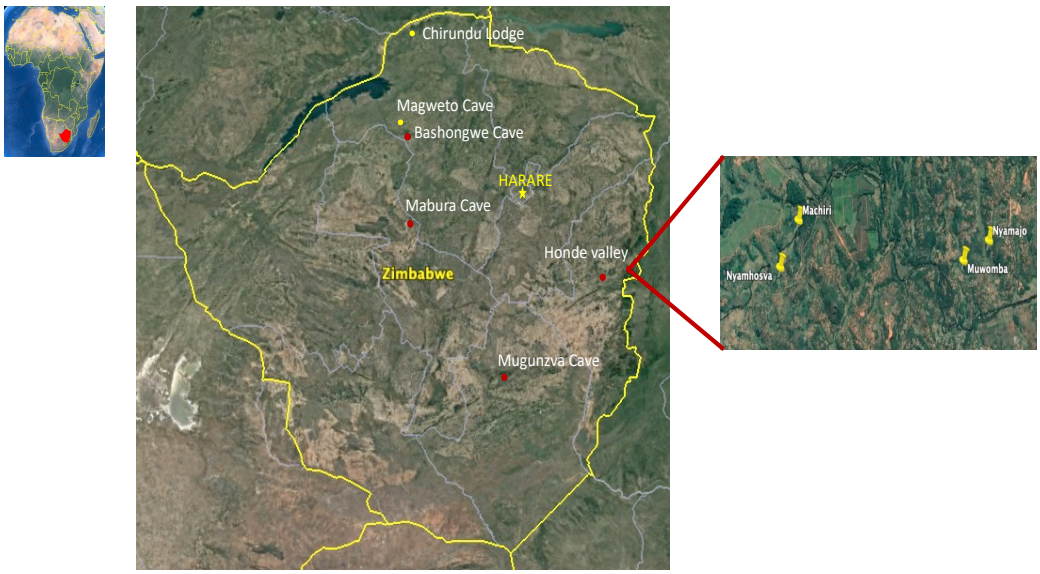



**Figure 1:** The study sites highlighted on the map show the sampling areas for insectivorous and frugivorous bats colonies across Zimbabwe. Red circles represent community approach study sites; yellow circles represent longitudinal survey sites (individual approach).

**Table 1:** Number of Astrovirus positives per each site/sampling year in pooled faecal samples from frugivorous and insectivorous bats at different 14 sites.

| Sampling Sites | type of roost | Date of collect | Bat Species | Number of collected tubes | Number of Tested pools | Number of positive pools |
|---|---|---|---|---|---|---|
| Magweto Cave | Cave | June 2016 | *Hipposideros spp.* | 24 | 8 | 5 |
| Bashongwe Cave | Cave | March 2018 | *Rhinolophus and Hipposideros spp.* | 59 | 7 | 0 |
| | | July 2018 | | 57 | 20 | 0 |
| | | December 2018 | | 135 | 45 | 0 |
| Mabura Cave | Cave | June 2016 | *Hipposideros spp.* | 36 | 12 | 6 |
| | | February 2017 | *Hipposideros spp.* | 22 | 7 | 0 |
| Mugunza Cave | Cave | April 2017 | *Roussetus aegyptus* | 50 | 13 | 0 |
| Honde Valley | | | | | | |
| *Machiri* | Trees | August 2017 | *Eidolon helvum* | 25 | 9 | 0 |
| *Muwomba* | | November 2017 | *Eidolon helvum* | 66 | 22 | 0 |
| *Nyamajo* | | November 2017 | *Eidolon helvum* | 58 | 19 | 0 |
| *Nyamhosva* | | November 2017 | *Eidolon helvum* | 35 | 11 | 0 |
| *Nyamajo* | | January 2018 | *Eidolon helvum* | 17 | 6 | 1 |
| *Machiri* | | January 2018 | *Eidolon helvum* | 7 | 2 | 2 |
| *Muwomba* | | January 2018 | *Eidolon helvum* | 8 | 2 | 0 |
| *Nyamhosva* | | January 2018 | *Eidolon helvum* | 28 | 10 | 1 |
| *Nyamajo* | | February 2020 | *Eidolon helvum* | 92 | 92 | 0 |
| Chirundu | Baobab | December 202 | *Rhinolophus spp.* | 21 | 21 | 1 |
| **Total** | | | | **740** | **306** | **16** |

All these sites were chosen according to the presence of bat colonies and the existence of anthropic activities. For instance, the selected caves and ancient mine selected are regularly visited by local people to collect bat guano which is used as fertilizer, and to hunt bats for consumption. Frugivorous bats roosting sites in trees were close to maize crops or/and fruit tree cultures.

All sites, except for three were visited at different times during the sampling period (Table 1). The same sampling method was used at all sites every session as previously described [25]. Briefly, two square meters of plastic sheets were laid down at each site/per sampling session, underneath the bat colonies and left overnight. Faeces were collected from each plastic sheet at a rate of ≈ 6 grams of pooled faeces in a 15 ml tube with 6 ml of homemade RNA stabilization solution (*https://protocol-online.org/*). Back in the laboratory, samples were stored at -80°C until analysed.

***Bat individual sampling***

Individual bat samples which had already been collected from two study sites for a study on coronaviruses by Chidoti et *al*. were used in this study (Figure 1) [26]. These two sites, one cave (Magweto) and one building basement (Chirundu Farm) had been visited from August 2021 to July 2022 on a monthly basis (Figure 1, Table 2) [26]. Unfortunately, this study was conducted during the COVID 19 crisis and we were not able to access the study sites every month as planned owing to the

121 imposed lock downs. Faecal samples had been collected by placing two square meters plastic sheets
122 underneath the bat colonies. Only one faecal dropping (clean and not contaminated by other faeces
123 or urine) per 20 $cm^2$ was collected, assuming it represented one individual. Faeces were conserved
124 individually in a 1.5 ml tube filled up with 0.5 ml of home-made RNA stabilization solution
125 (*https://protocol-online.org/*) and stored at -80°C before further laboratory analyses.

126 **Table 2:** Prevalence of Astroviruses and Coinfection with Coronaviruses and confidence intervals (CIs) per
127 month at both Magweto and Chirundu sites in faecal samples from individual sampling of insectivorous bat
128 communities

| Site | Reproduction cycle | Month sampled | No of samples tested | No of Astrovirus positives | Prevalence (%) + CI (95%) | No of coinfection CoV and AstV | Prevalence (%) + CI (95%) |
|---|---|---|---|---|---|---|---|
| Chirundu | **Non-gestation** | August, 2020 | 153 | 0 | 0.0 (0.0-0.2) | 0 | 0.0 (0.0-0.2) |
| Chirundu | **Pregnancy** | October, 2020 | 297 | 3 | 1.0 (0.3-2.9) | 0 | 0.0 (0.0-1.3) |
| Chirundu | **Parturition** | November, 2020 | 241 | 3 | 1.2 (0.4-3.6) | 1 | 0.4 (0.05-2.3) |
| Chirundu | **Lactation** | December, 2020 | 159 | 16 | 10.1 (6.3-15.7) | 7 | 4.4 (2.1-8.8) |
| Chirundu | **Weaning** | February, 2021 | 170 | 29 | 17.1 (12.1-23.4) | 18 | 10.6 (6.8-16.1) |
| Chirundu | **4-6 Month juveniles** | March, 2021 | 240 | 51 | 21.3 (16.5-26.8) | 19 | 7.9 (5.1-12.0) |
| Chirundu | **Non-gestation** | May, 2021 | 243 | 47 | 19.3 (14.8-24.7) | 9 | 3.7 (1.9-6.8) |
| Chirundu | **Non-gestation** | July,2021 | 225 | 30 | 13.3 (9.5-18.4) | 9 | 4.0 (2.1-7.4) |
| **Overall prevalence** | | | **1728** | **179** | **10.4 (9.0-11.8)** | **63** | **3.6 (3.8-5.8)** |
| Magweto | **Non-gestation** | September, 2020 | 348 | 18 | 5.2 (3.3-8.0) | 1 | 0.3 (0.05-1.6) |
| Magweto | **Pregnancy** | October, 2020 | 257 | 46 | 17.9 (13.7-23.0) | 7 | 2.7 (1.3-5.5) |
| Magweto | **Parturition** | November, 2020 | 228 | 31 | 13.6 (9.7-18.6) | 3 | 1.3 (0.4-3.8) |
| Magweto | **4-6 Month juveniles** | March, 2021 | 242 | 46 | 19.0 (14.5-24.4) | 20 | 8.3 (5.4-12.4) |
| Magweto | **Non-gestation** | April, 2021 | 242 | 26 | 10.7 (7.4-15.2) | 6 | 2.5 (1.1-5.3) |
| Magweto | **Non-gestation** | June, 2021 | 242 | 47 | 19.4 (14.9-24.9) | 3 | 1.2 (0.4-3.6) |
| **Overall prevalence** | | | **1559** | **214** | **13.7 (12.1-15.5)** | **40** | **2.6 (1.8-3.5)** |

129

130 Periods of gestation (pregnancy), parturition, lactation, weaning and presence of 4-6 months old
131 juveniles had been determined based on observations (from captures and observation of the roosting
132 bats) combined with literature, as already reported in Chidoti et *al.* [26]. The reproductive season at
133 Chirundu and Magweto site had been observed to begin in September to February for the predominant
134 bat species. The different insectivorous bat families observed at both sites had been found to be
135 synchronous regarding their reproductive cycles, and consensus reproduction periods had been
136 determined based on the literature and observations [26, 27].

137 **Nucleic acid extraction and RT-PCR.**

138 ***Community samples***.

139 Nucleic acids extraction was done from all faecal samples as previously described [26]. Briefly,
140 biological material (Faeces) of three or four sample tubes collected from the same plastic sheet were
141 pooled and transferred in a 50 ml tube with 20 ml of PBS 1X then vigorously mixed. Tubes were
142 centrifuged at 4500 rpm for 10 min. The supernatant was first filtered using gauze in order to eliminate
143 faecal matter and transferred in fresh tubes then re-centrifuged at 4500 rpm for 10 min. The

144 supernatant was filtered through a 0.45 µm filter to remove eukaryotic and bacterial sized particles.
145 Seven millilitres of filtered samples were centrifuged at 250,000 g for 2.5 h at 4°C. The pellets were
146 re-suspended in 600 µl $H_20$ molecular grade and 150 µl were used to extract RNA and DNA using
147 NucleoSpin® RNA Kit (Macherey-Nagel, Hoerdt, France) according to the manufacturer's protocol.
148

149 ***Individual samples***

150 Nucleic acids were extracted from 200 µl of faecal samples preserved in 0.5 ml RNA stabilization
151 solution using 5X MagMax Pathogen RNA/DNA Kit (ThermoFisher Scientific, Illkirch-
152 Graffenstaden, France), as already described in [26]. The faeces were vortexed vigorously (30Hz)
153 using Retsch MM400 Tissue lysser for 5 min to fully homogenise and mix the faecal particles,
154 followed by centrifugation at 16000 g for 3 min to fully separate the supernatant from the faecal
155 debris. A volume of 130µl of the supernatant was used for the isolation and purification stage of the
156 nucleic acids using Mag Max extraction kit with the automatic KingFisher Duo Prime Purification
157 System extractor (ThermoFisher Scientific, Illkirch-Graffenstaden, France). A final volume of 80µl
158 of eluted RNA/DNA was stored at -80◦C.

159 ***Bat species identification***

160 All positive *Astrovirus* bat species were identified by sequencing *cytochrome b* [28]. *S*equences were
161 then compared to available bat sequences in the GenBank database using *Basic Local Alignment*
162 *Search Tool* (BLAST) program.
163

164 ***Astrovirus detection***

165 Reverse Transcription (RT) was done using random hexamers were done on 5µl of RNA sample
166 template using 1µl random hexamers, 0.5µl Oligo dT primer, 0.4µl of dNTPs (10mM) (ThermoFisher
167 Scientific, Illkirch-Graffenstaden, France) and 5.5µl molecular grade water incubated at 65°C for 5
168 min. This was followed by addition of 4µl of Buffer 5X, 2µl of 0.1M DTT (M-MLV Reverse
169 Transcriptase, Invitrogen, ThermoFisher Scientific) and 1µl of RNAse OUT, incubated at 37°C for 2
170 min. A volume of 1µl of M-MLV (M-MLV Reverse Transcriptase, Invitrogen, ThermoFisher
171 Scientific, Illkirch-Graffenstaden, France) reverse transcriptase was added to the mixture followed
172 by incubation at 25ºC for 10 min, 37ºC for 50 min and 70ºC for 15 min. The cDNA obtained was then
173 used to partially amplify the *Astrovirus RNA-dependent-RNA polymerase* gene (*RdRp*) by using a
174 semi-nested Pan-Astrovirus PCR system developed by Chu et *al* [13]. Positive PCR products (422
175 bp) were gel-agarose purified (Geneclean Turbo Kit, MP Biomedicals, Illkirch-Graffenstaden, France)
176 and then sequenced in both 5' and 3' directions (LGC, Berlin, Germany) by using Sanger method.

For the community approach, purified PCR products were cloned by using Topo PCR Cloning kit according to the manufacturer's protocol (ThermoFisher Scientific, Illkirch-Graffenstaden, France). Ten clones per PCR product were sequenced in both 5' and 3' direction using the Sanger method (Eurofins, Germany).

**Phylogenetic analyses**

Overlapping sequences were assembled into contiguous sequences using Geneious software package V. 2021.2.2 (Biomatters Ltd, Auckland, New Zealand). Partial non-concatenated nucleic acid sequences of the new *Astrovirus* were aligned using MEGA 7 [29], with minor manual adjustments. Sites that could not be unambiguously aligned and divergent regions were excluded from subsequent analyses. Phylogenies were inferred using Maximum Likelihood (ML) method implemented in PhyML [30]. The suited evolution model was selected by Akaike's Information criterion (AIC) using Topali software [31]. The reliability of branching orders was tested using the bootstrap approach (1000 replicates) [32] and the GTR + F+ I substitution model was determined as the best suited evolution model.

**Temporal variations of Astrovirus prevalence and bat reproductive phenology**

In the same way as in Chidoti et *al*, the prevalence of astrovirus infection was calculated at the community level for small insectivorous bat species from the two longitudinal study sites, Magweto and Chirundu farm [26]. The proportion of RNA AstV-positive samples were estimated per month and site with 95% confidence intervals (CI) using Wilson score test [33]. A descriptive analysis of AstVs prevalence per month and corresponding reproductive season at each site was done.

The influence of the different phases of the bat reproductive cycle (periods of pregnancy, parturition/lactation, weaning, weaned juveniles of 4 to 6 months old) and of *Coronavirus* infection/shedding on the prevalence of astroviruses was tested by running a generalized linear mixed model (GLMM) for each site (Magweto and Chirundu) as described in Chidoti et *al* [26]. Parturition and lactation were analysed as one season because the observations did not allow the separation of the two periods as they were interlinked. The response variable with a binomial distribution was the *Astrovirus* PCR result of the samples, and the explanatory variables with fixed effects were the different phases of the reproduction cycle. Given *Coronavirus* PCR results on the same samples were available through our previous study (see below), *Coronavirus* status was also integrated into the model as fixed variable to investigate any effect of co-infection on *Astrovirus* infection/shedding. The reproductive phases were coded as 1 if it was during the corresponding reproduction phase and 0 if it was not. A session identification code was included as a random effect to control for repeated measures from the same sampling session to account for clustered samples collected.

212

213 ***Astrovirus and Coronavirus co-infection***

214 *Coronavirus* data, produced by Chidoti et *al* [26], on the same sample sets were used in this study to
215 assess the *Astrovirus* and *Coronavirus* co-infection in bat communities from Magweto and Chirundu
216 sites and a descriptive analysis of prevalences was carried out.

217

218 ***GenBank Accession Numbers***

219 The *Astrovirus* sequences have been deposited to the GenBank under the following numbers:
220 OQ271049 - OQ271203

221 The Cytochrome B sequences have been deposited to the GenBank under the following numbers:
222 OM487705-OM488020

223

224 **Results**

225 ***Astrovirus detection in pooled sampling sites***

226 The sampling at these sites varied from June 2016 to December 2020 (Table 1). At Magweto 24
227 sampled were collected. In Bashongwe collection was carried out in March, July and December 2018,
228 and 251 samples were collected. In Mabura cave site, sampling was done once in 2016 and 2017 and
229 a total of 58 samples were collected. In Mugunza cave site, sampling was done once in 2017 and 50
230 samples were collected. In the Honde Valley site, 336 samples were collected under 9 *Eidolon helvum*
231 roost trees (Table 1). Five (62.5%) pooled samples from Magweto site were positive for AstVs out of
232 8 pooled samples analysed and all 72 pooled samples from Bashongwe cave were negative for AstV
233 (Table 1). Six (31.6%) out of 19 pooled samples from Mabura cave tested for AstVs were positive
234 and all of the 13 samples from Mugunza tested for AstVs were negative (Table 1). In Honde Valley,
235 173 pooled samples were tested for AstVs and only four (2.3%) were positive (Table 1). Only one
236 (4.8%) out of the 21 samples collected in Chirundu Baobab tree site was positive for AstV.

237

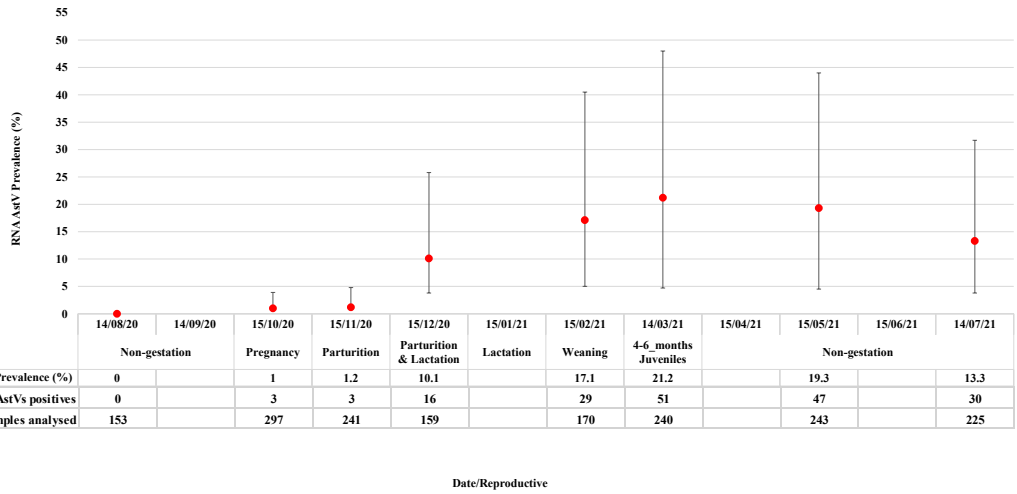
238 ***Prevalence and seasonality of Astroviruses at Chirundu and Magweto site***

239 We analysed 1559 samples from insectivorous bats from the Magweto cave site and a total of 1728
240 faecal samples from bats at the Chirundu farm site (Table 2).The overall prevalence of *Astrovirus* at
241 bat community level of Chirundu was 10.4% [95% CI: 9.0-11.8] (179 positives out of the 1728
242 samples). The highest prevalence of 21.3% [95% CI: 16.5-26.8] was observed in March 2021 and
243 corresponded to the 4-6 months old juvenile period (Table 2). The prevalence of *AstV* increased from
244 December 2020 to March 2021, which corresponded to the lactation, weaning and 4-6 months
245 juvenile periods. The lowest prevalence was observed in August (0%); October (1% [95% CI: 0.3-

2.9]) and November (1.2% [95% CI: 0.4-3.6]), which correspond to the transition between non-gestation and gestation period, as well as the beginning of the parturition period (Table 1, Figure 2a).

**Figure 2:** Bat community *Astrovirus* prevalence per month at Chirundu site. **Fig 2a & 2b** show estimation of the astrovirus prevalence (with CI 95%) at Chirundu and Magweto sites respectively for the months sampled. The graphs were plotted with y-axes as the number of AstVs positive samples, and the prevalence of AstVs and x-axis as the sampling date and corresponding reproductive season.

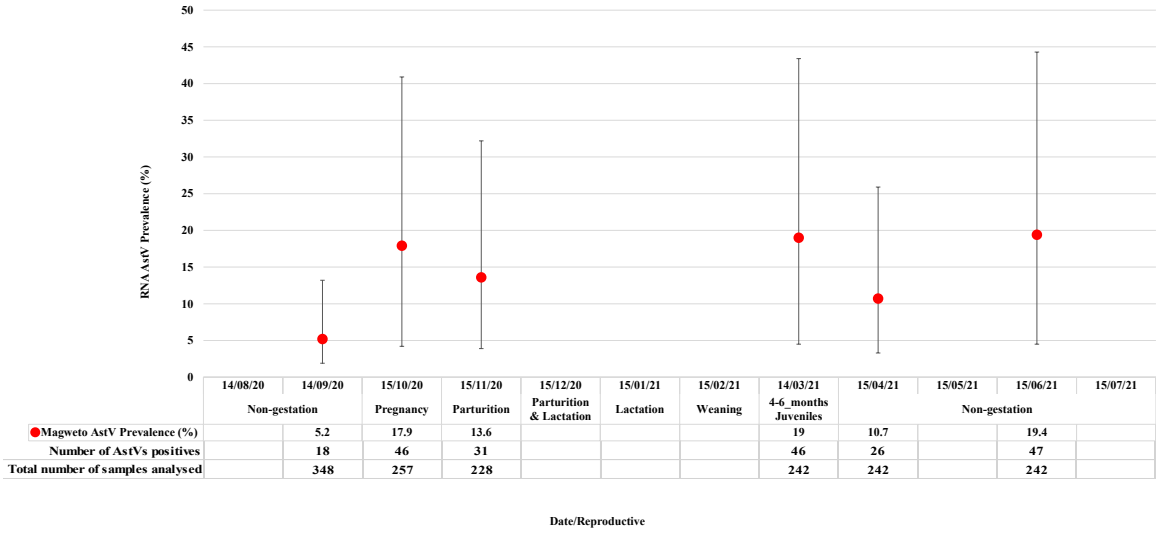
Figure 2a




| | 14/08/20 | 14/09/20 | 15/10/20 | 15/11/20 | 15/12/20 | 15/01/21 | 15/02/21 | 14/03/21 | 15/04/21 | 15/05/21 | 15/06/21 | 14/07/21 |
|---|---|---|---|---|---|---|---|---|---|---|---|---|
| | Non-gestation | | Pregnancy | Parturition | Parturition & Lactation | Lactation | Weaning | 4-6_months Juveniles | Non-gestation | | | |
| Chirundu AstV Prevalence (%) | 0 | | 1 | 1.2 | 10.1 | | 17.1 | 21.2 | | 19.3 | | 13.3 |
| Number of AstVs positives | 0 | | 3 | 3 | 16 | | 29 | 51 | | 47 | | 30 |
| Total number of samples analysed | 153 | | 297 | 241 | 159 | | 170 | 240 | | 243 | | 225 |

In Magweto site, the overall prevalence of Astroviruses for the insectivorous bat community was 13.7% [95% CI:12.1-15.5] (214 positives out of 1559 samples). There was no clear pattern for the prevalence of AstVs according to reproductive seasons in this site (Table 2, Figure 2b).The highest prevalence of 19.4% [95% CI: 14.9-24.9] was observed in June 2021 during the non-gestation period , and very close prevalences observed during the pregnancy (17.9% [95% CI: 13.7-23]) and 4-6 months old juvenile (19% [95% CI: 14.5-24.4]) periods (Table 2). Lower prevalence was observed in September at the end of the non-gestation period (5,2% [95% CI: 3.3-8]).

Figure 2b




Results from the GLMM didn't show an effect bats seasonal reproduction periods on detection of RNA-AstVs positive samples at Magweto site (Table 3a). For Chirundu site, the GLMM showed significantly higher probability of being positive to astrovirus during the period of weaned 4-6 months old juveniles (odds ratio (OR) = 5.088, 95% CI = 1.33- 36.32, p= 0.016), and significantly lower probability of astrovirus positivity during the pregnancy period (odds ratio (OR) = 0.136, 95% CI = 0.03- 0.81, p=0.006) (Table 3b).

**Table 3*:*** Results of the GLMM with the following explanatory variables (a/ Magweto, b/Chirundu): PCR_Pan-*Coronavirus*, Pregnancy, Weaning, Weaned juveniles 4 to 6 months. With references OR odds ratio, CI95 95% confidence interval.

***a/***

| **Variables** | **Odds ratios** | **p-value** | **OR CI95** |
|---|---|---|---|
| Intercept | 0.235 | 0.0005 | 0.088-0.627 |
| PCR_Pan-*Coronavirus* | 1.238 | 0.290 | 0.825-1.825 |
| Pregnancy | 0.473 | 0.145 | 0.138-1.548 |
| Weaned_juveniles 4 to 6 months | 0.678 | 0.448 | 0.202-2.256 |

277 ***b/***

| **Variables** | **Odds ratios** | **p-value** | **OR CI95** |
|---|---|---|---|
| Intercept | 0.05 | 6.68e-09 | 0.010- 0.121 |
| PCR_ Pan-*Coronavirus* | 1.116 | 0.536 | 0.785 -1.574 |
| Pregnancy | 0.4 | 0.006 | 0.027-0.810 |
| Weaning | 3.91 | 0.094 | 0.721-39.593 |
| Weaned_juveniles 4 to 6 months | 5.09 | 0.016 | 1.333-36.322 |

278

279 ***Prevalence and seasonality of coinfection with Astroviruses and Coronavirus at Chirundu and***
280 ***Magweto site.***

281 The prevalence of co-infection of bat communities from Chirundu and Magweto sites was fairly low
282 however it showed peaks that correlated with peaks in coronavirus and astrovirus infections as
283 observed in the graphs. The overall prevalence of co-infection of insectivorous bats from Chirundu
284 site by Astroviruses and Coronaviruses was 3.6% [95% CI: 3.8-5.8] (Table 2). The highest co-
285 infection prevalence observed in this site was 10.6% [95% CI: 6.8-16.1] in February 2021 during the
286 weaning period while the lowest prevalence of 0% was observed in August [95% CI: 0.0-0.2] and
287 October [95% CI: 0.0-1.3] during the end of the non-gestation and the pregnancy period (Table 2,
288 Figure 3a).

289 The overall prevalence for co-infection of insectivorous bats in Magweto site by coronaviruses and
290 astroviruses was 2.6% [95% CI: 1.8-3.5] (Table 2, Figure 3b). Co-infection in this site was recorded
291 in all the months, from September 2020 to June 2021, except from December to February during
292 which no sampling took place (Table 2, Figure 3b). The highest prevalence of co-infection was 8.3%
293 [95% CI: 5.4-12.4] in March during the 4-6 months old juvenile period; while the lowest 0.3% [95%
294 CI: 0.05-1.6] was observed in September 2020 during the non-gestation period (Table 2, Figure 3b).

295

Figure 3a

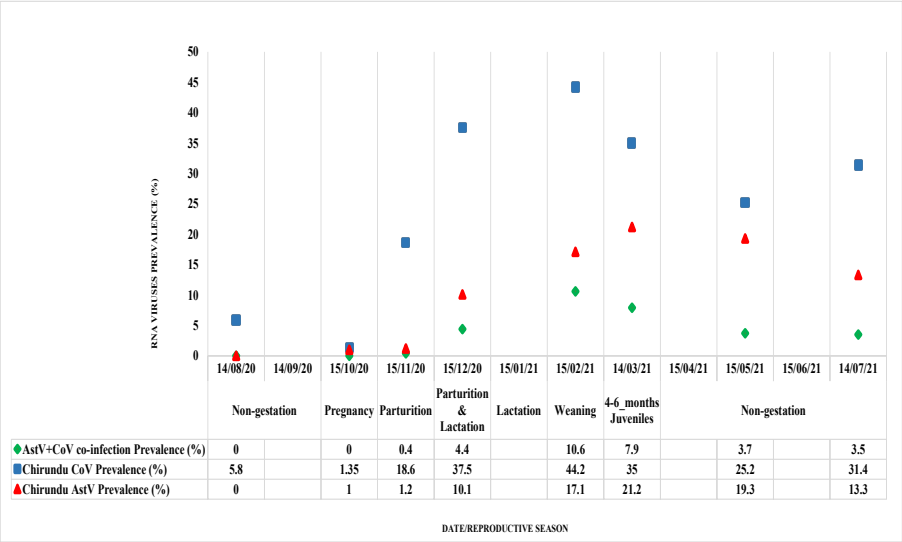



Figure 3b

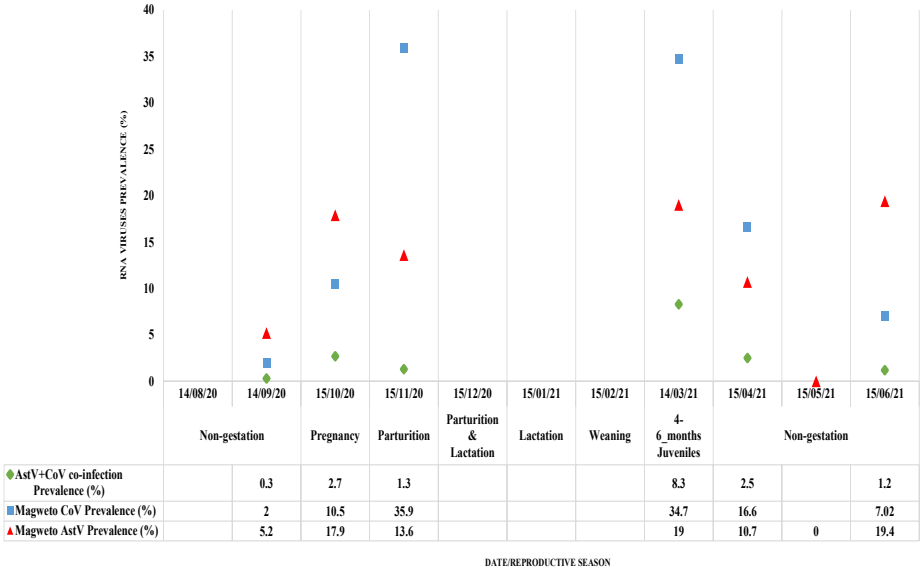



**Figure 3***:* Prevalences of astroviruses and coronavirus co-infection (in green), coronaviruses (in blue) and astroviruses (in red) per month in bat communities for a/ Chirundu site and b/ Magweto site.

301 ***Genetic diversity of RNA Astroviruses***

302 In this study we sequenced all *Astrovirus* positive samples from 5 sites: (Chirundu farm N=179;
303 Magweto cave N=229; Mabura cave N=7; Honde Valley N=7 and Baobab tree N=1). Of the 423
304 samples, 214 (50.6%) were amplified from *Hipposideros* spp, 63 (14.9%) from *Rhinolophus* spp, 53
305 (12.5%) from *Miniopterus*, 10 (2.4%) from *Nycteris* spp, 7 (1.7%) from *Eidolon* spp and for the
306 remaining 76 samples, the bat genus / species could not be determined mainly owing to bad sequence
307 qualities.

308 Phylogenetic analysis of the 155 *RdRp AstV* nucleotide sequences generated for this study showed
309 that they all clustered with the genus *Mamastrovirus* of the *Astroviridae* family (Figure 4).

310 Two sequences isolated from Magweto cave, 1 from *Rhinolophus spp* (MAG-573) and 1 from
311 *Hipposideros spp* (MAG-292) clustered with a group of Human astroviruses (*HuAstVs*) (Figure 4).
312 This cluster was well sustained (Bootstrap >90%) and the nucleotidic acid identities between Bat
313 Astroviruses MAG-573, MAG-292 and HuAstV-2 were 96 and 93% respectively (Data not shown).

314 The other specific cluster of bat astrovirus showing a close relation to human astroviruses, is a
315 sequence isolated from *Hipposideros spp* (MAG-1236) in Magweto, it clustered with *HuAstV-MLB-*
316 *1; -2* and *3* (Figure 4). Six sequences isolated from the frugivorous bats *E.helvum* found in Honde
317 Valley and one from *Rhinolophus spp* collected in a Baobab tree formed a sister clade with another
318 group of Human astroviruses *HuAstV-VA-1;-2;-3;-4* and *HuAstV-HMO-A* (Figure 4). The majority of
319 the sequences, comprises sequences closely related to insectivorous Bat Astroviruses (Figure 4). The
320 majority of these insectivorous bat related astroviruses sequences were amplified from *Hipposideros*
321 (HC-1 to -9) and *Rhinolophus* (Rhi-1 to -4) bat species. One cluster comprised of sequences isolated
322 from *Rhinolophus spp*. Most of these sequences were isolated from Chirundu and a few of them from
323 Magweto, and clustered together with *AstV* strains from *Nyctalus* and *Verpitillio* bat species isolated
324 in Czech Republic (references Y & Z in the phylogenetic tree) (Figure 4). Another cluster comprising
325 of majority of sequences from Chirundu and some from Magweto site, isolated from *Miniopterus* and
326 *Rhinolopus* spp showed phylogenetic clustering with strains of bat astroviruses isolated from
327 *Miniopterus* spp from Madagascar and China (references N, O, R, U, T), *Rhinolophus* spp from Korea
328 (references P, Q) as well as *Rousettus* and *Paratrianops BAstV* from Madagascar (references V, W)
329 (Figure 4)

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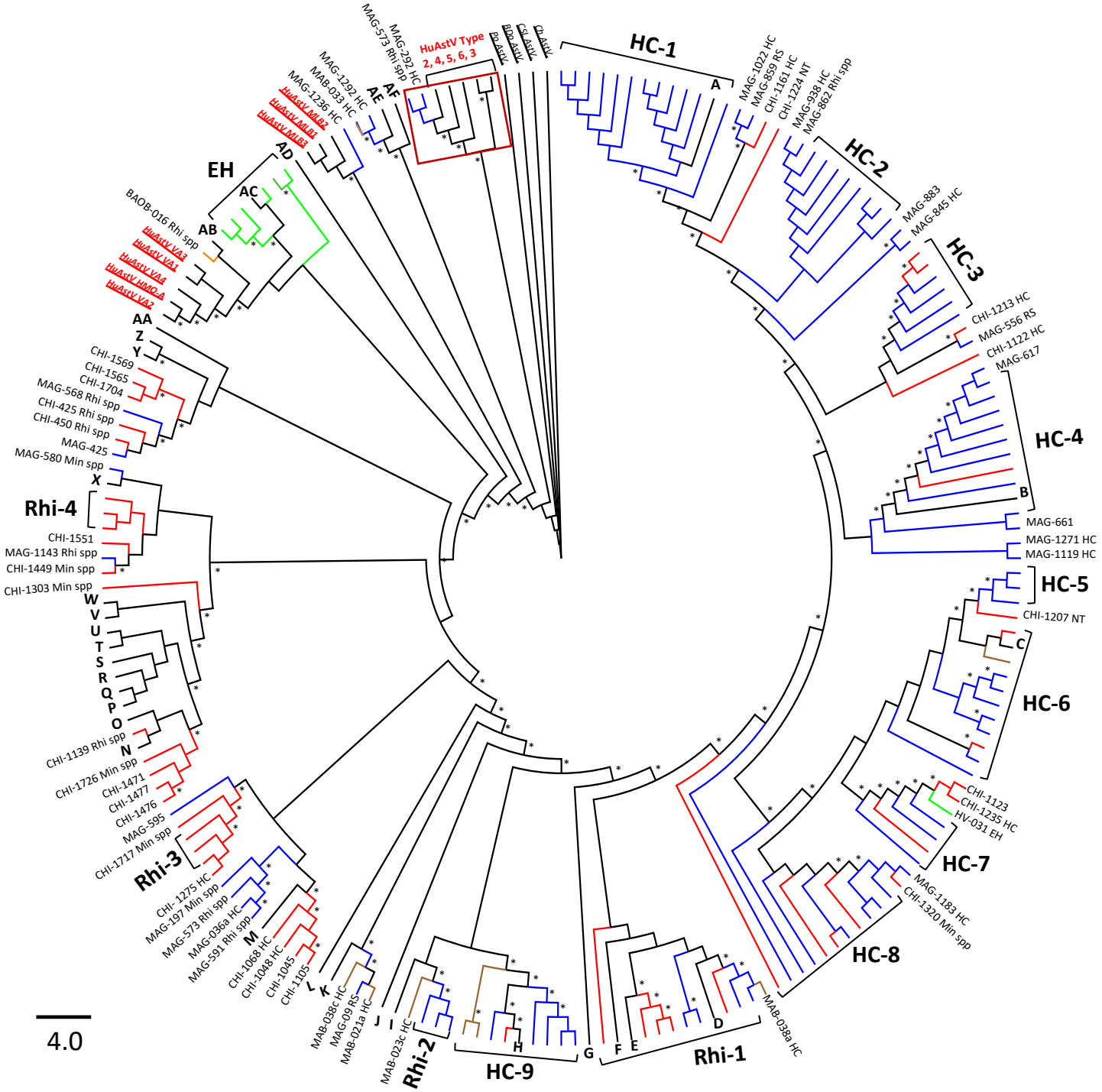
HC-1
A
MAG-1022 HC
MAG-859 RS
CHI-1161 HC
CHI-1224 NT
MAG-938 HC
MAG-862 Rhi spp
HC-2
MAG-883
MAG-845 HC
HC-3
CHI-1213 HC
MAG-556 RS
CHI-1122 HC
MAG-617
HC-4
B
MAG-661
MAG-1271 HC
MAG-1119 HC
HC-5
CHI-1207 NT
C
HC-6
CHI-1123
CHI-1235 HC
HV-031 EH
HC-7
MAG-1183 HC
CHI-1320 Min spp
HC-8
MAB-038a HC
D
Rhi-1
E
F
G
HC-9
H
Rhi-2
MAB-023c HC
I
J
MAB-038c HC
MAG-09 RS
MAB-021a HC
K
L
Rhi-3
CHI-1275 HC
MAG-197 Min spp
MAG-573 Rhi spp
MAG-036a HC
MAG-591 Rhi spp
M
CHI-1068 HC
CHI-1048 HC
CHI-1045
CHI-1105
N
CHI-1139 Rhi spp
CHI-1726 Min spp
CHI-1471
CHI-1477
CHI-1476
MAG-595
CHI-1717 Min spp
O
P
Q
R
S
T
U
V
W
CHI-1303 Min spp
Rhi-4
CHI-1551
MAG-1143 Rhi spp
CHI-1449 Min spp
X
MAG-580 Min spp
MAG-425
CHI-450 Rhi spp
CHI-425 Rhi spp
MAG-568 Rhi spp
CHI-1704
CHI-1565
CHI-1569
Y
Z
AA
HuAstV VA2
HuAstV HMO-A
HuAstV VA4
HuAstV VA1
HuAstV VA3
BAOB-016 Rhi spp
AB
AC
EH
AD
HuAstV MLB3
HuAstV MLB1
HuAstV MLB2
MAG-1236 HC
MAB-033 HC
MAG-1292 HC
AE
AF
MAG-573 Rhi spp
MAG-292 HC
HuAstV Type
2, 4, 5, 6, 3
Po AstV
BDo AstV
CSL AstV
Ch AstV
4.0

**Figure 4*: Phylogenetic tree of Astroviruses* partial *RdRp* gene**. The sequences detected at Chirundu site are represented by red branches, at Magweto site by blue branches, at Honde Valley in by green branches, at Mabura cave by brown branches and at Baobab site by an orange branch. The red rectangle highlighted the phylogenetic relationship between MAG-292, MAG-573 and *HuAstV* strains. The tree was built using the maximum likelihood method based on the GTR + G4 + I model. The robustness of nodes was assessed with 1000 bootstrap replicates. Bootstrap values >70 are in asterisk and those <70 are not shown. HC=*Hipposideros caffer*; RS=*Rhinolophus simulator*; NT=*Nycteris thebaica*; EH= *Eidolon helvum*; Min=*Miniopterus spp*; Rhi=*Rhinolophus species*. MAG=Magweto site; CHI=Chirundu site; BAOB= Baobab; MAB= Mabura site; HV=Honde valley site.

Human *Astrovirus* are highlighted in red and bold letters represented the different bat AstV references used to build this tree:

**Bat Astrovirus references:**

**A** - MH013972.1: Hipposideros caffer, Mozambique; **B** - MH013971.1: Hipposideros caffer, Mozambique; **C** - MH013970.1: Hipposideros caffer, Mozambique; **D** - KX858349.1: Rhinolophus spp., Laos; **E** – MC841039.1: Rhinolophus Ferrumquinum, Korea; **F** - MH013989.1: Trianop afer, Mozambique ; **G** - MH013986.1: Trianop afer, Mozambique ; **H** - MH013971.1: Hipposideros caffer, Mozambique ; **I** - KX858371.1: Hipposidoros larvatus, Laos **J** - KY575656.1: Myotis goudoti, Madagascar ;

**K** - KY575655.1: Myotis goudoti, Madagascar; **L** - MH013988.1: Trianops afer, Mozambique

**M** - MH013076.1: Nycteris thebaica, Mozambique ; **N**- KY575649.1: Miniopterus griveaudi, Madagascar ; **O** - KY575647.1: Miniopterus griveaudi, Madagascar ; **P** - MC840967.1: Rhinolophus ferrumquinum, Korea ; **Q** - MC840969.1: Rhinolophus ferrumquinum, Korea ; **R** - KY575648.1: Miniopterus griveaudi, Madagascar ; **S** - EU847151.1 : Miniopterus magnater, Chine ; **T** - KY575674.1: Miniopterus griveaudi, Madagascar ; **U**- KY575666.1: Rousettus madagascariensis, Madagascar ; **V** - KY575661.1 : Paratrianops furculus, Madagascar ; **W** - KY575644.1 : Miniopterus griveaudi, Madagascar ; **X** - EU847155.1: Miniopterus pussilus, Chine ; **Y**- KP843561.1: Vespertillio Murinus, Czech ; **Z**- KP843558.1: Nyctalus noctule, Czech ; **AA** - KX858513.1: Megaderma lyra, Cambodge ; **AB** - HM368175.1: Myotis myotis, Germany ; **AC** - KX858367.1: Roussetus spp., Laos ; **AD** - FJ571067.1: Taphozous melanopogon, Chine ; **AE** - KX858377.1 : Roussetus spp, Cambodge ; **AF** - KX858378.1 : Roussetus spp, Laos

**Human Astrovirus references:**

GQ502195.1: Human Astrovirus VA2; JQ673585.1: Human Astrovirus HMO-A; JX857869.1: Human Astrovirus VA4; KJ920196.1: Human Astrovirus VA1; GQ502196.1: Human Astrovirus VA3; JX857870.1: Human Astrovirus MLB3; JQ673581.1: Human Astrovirus MLB1; GQ502188.1: Human Astrovirus MLB2; L23509.1: Human Astrovirus Type 2; AY20891.1: Human Astrovirus Type 4; DQ028633.1: Human Astrovirus Type 5; GQ495608.1: Human Astrovirus Type 6; JF491430.1: Human Astrovirus Type 3

**Other Astrovirus references:**

GQ914773.1: Porcine Astrovirus (Po AstV); FJ890355.1: Bottlenose Dolphin Astrovirus (BDo AstV); JN 420351.1: California Sea Lion Astrovirus (CSL AstV); æEU669001.1: Chicken Astrovirus (Ch AstV)

Two major clades from Chirundu and one from Magweto, isolated from *Miniopterus, Rhinolophus* and *Hipposideros spp* of bats also phylogenetically clustered with a strain from Mozambique isolated from *Nycteris thebaica* (reference M). One sequence from Magweto isolated from *Rhinolophus spp* and two from Mabura cave isolated from *Hipposideros* spp*,* clustered with a strain of bat astrovirus isolated from *Myotis* spp in Madagascar (references K, L) (Figure 4). A cluster, comprising two clades from Magweto and one sequence from Chirundu and three sequences from Mabura cave isolated from *Rhinolophus* and *Hipposideros,* were closely related to bat astrovirus strains from *Hipposideros* spp of Mozambique and Laos (references H, I) (Figure 4). The last small group comprising one clade from Chirundu, four sequences from Magweto, all isolated from *Rhinolophus* spp*,* and one sequence from Mabura isolated from *Hipposideros* spp clustered together with bat astrovirus strains from *Rhinolophus* spp of LAOS and Korea (references D, E, F) (Figure 4).

One of the large clusters shows the phylogenetic relationship amongst sequences derived from, *Hipposideros* spp (Clades HC-5 to -8), these making up the majority of the sequences in this clade

389 and one isolated from *Miniopterus* spp, one from *E. helvum* and one from a *Nycteris* spp. The bat
390 astroviruses in this clade, all showed a close relation to a bat astrovirus strain isolated from
391 *Hipposideros spp* of Mozambique (reference C). The largest cluster constituted mainly by Magweto
392 site sequences and some Chirundu sequences, from *Hipposideros* spp (clades HC-1 to 4) and a few
393 *Rhinolophus* and *Nycteris* spp, clustered with a strain from *Hipposideros* spp of Mozambique
394 (references A, B) (**Figure 4**).
395
396 **Discussion**
397 The overall prevalence of *AstVs* in bats was 10% and 13.7% in Chirundu and Magweto sites,
398 indicating a fairly high circulation of this family of viruses in bats in Zimbabwe. However, compared
399 to studies in other African countries, the observed prevalence was low: Hoarau et *al* and
400 Lebarbenchon et *al* detected *AstV* in 20-22% of individual bats in Mozambique and Madagascar,
401 respectively [17,19]. This difference in prevalence of *AstV* compared to other countries can be
402 attributed to the lack of longitudinal survey in the Mozambican and Madagascar studies as sampling
403 effort was done for periods of two separate months in both studies [17,19].
404 In Chirundu site increasing RNA-*AstV* prevalences were observed from the lactation to the 4-6
405 months old juvenile seasons with 21.2% of RNA -*AstV* detection during the latter. Astroviruses have
406 been described to display seasonal variations in prevalence [17]. Drexler et al observed different
407 peaks of astrovirus detection associated with different stages of the reproductive season with peaks
408 correlating to maternal aggregations during breeding season and after parturition season due to
409 establishment of a susceptible subpopulation of weaned new-born bats who did not yet have their
410 own adaptive immunity [34]. In our study the prevalence was high during some of the months
411 compared to overall prevalence, which might be related to the effect of reproductive seasons on the
412 shedding of viruses, as demonstrated for other viruses. During the reproductive season with juveniles
413 and immature individuals, the prevalence of CoV was very high as compared to prevalence in the
414 absence of juveniles and presence of sexually mature bats [35,36]. In this study we observed a similar
415 trend for AstVs for one of the study sites. Furthermore, Mendenhall et *al*, also identified the bat
416 juvenile stage as exhibiting a greater *Astrovirus* viral burden than any other stage of the reproductive
417 season [37]. The range of prevalence observed during the highest peaks coincides with the ≥20%
418 overall AstV detection reported by Hoarau et *al* [19] during the time of their sampling in February
419 and May. Here we observe a peak during the 4-6 months old juvenile season in Chirundu site which
420 corresponds to a high influx of immunologically immature individuals in the bat population, similar
421 to the what is observed in the Chidoti et *al*, (2022) for CoV infection [26].

422 During this period, pups are known to contribute to shedding pulses of viruses as they develop
423 productive infections during the acute phase [24], which coincides with the waning protection of
424 maternal antibodies [26], thus increasing the susceptibility and rate of infection in the young.
425 We could not observe a clear trend in Magweto site, which might be related to the absence of samples
426 from December 2020 to March 2021, which correspond to the lactation and weaning periods.
427 In our study we also investigated co-infection of bats by coronaviruses and astroviruses. We compared
428 the co-infection prevalence with the prevalence of astroviruses and coronaviruses described at each
429 site during the same reproductive cycles. The co-infection prevalence showed a similar trend to that
430 observed in the individual viruses, whereby during observed peaks of high AstVs and CoVs
431 detections, the co-infection also peaked at similar phases of the reproductive cycle. Thus the co-
432 infection of bats by both CoVs and AstVs is high when there is high infection of bats by either virus.
433 The overall prevalence of co-infection was 3.5% and 2.6 % in Chirundu and Magweto respectively
434 while the bats from Madagascar and Mozambique respectively showed a *AstV-CoV* co-infection of
435 5% (±2.7%) and of 2.7% (±1,1%) [17,19]. Co-infection and recombination of viruses in bats have
436 been reported on several occasions including co-infection of bats by coronaviruses and astroviruses
437 [17,23]. Bats infected with either *CoV* or *AstV* were shown to be more likely co-infected with the
438 respective virus [23]. In our study, no effect of coronavirus infection/shedding on *Astrovirus*
439 infection/shedding was observed.
440 In the current study we detected bat Astroviruses from *Hipposideridae, Rhinolophidae*, *Pteropodidae,*
441 *Nycterida*e and *Miniopterida*e families. Astroviruses are known to show no host restriction and are
442 widespread within the Chiroptera order [13,20]. The majority of the astroviruses amplified were from
443 *Hipposideros* and *Rhinolophus* species and these clustered with *AstV* sequences derived from
444 Mozambique, Madagascar and China of the same families [17,19,20]. The observed trend is due to
445 the two genera being the most dominant species of bats at both sites where sampling was done.
446 Therefore, the probability of higher detection of *AstV* in these species is expected and evidently
447 observed to be higher than other rare or less dominant species. Active transmission events of
448 astroviruses amongst *Rhinolophidae* and *Hipposideridae* bat species are known to occur [20]. In this
449 study, a high degree of species-specific tropism, especially in bat astrovirus related clusters was
450 observed with astrovirus strains isolated from *Hipposideros spp* clustering with each other and similar
451 trends for *Rhinolophus* and *Miniopterus spp*. However, the clusters were not site-specific as
452 sequences from Magweto, Mabura and Chirundu sites formed clusters with each other. There is a
453 significant phylogenetic clustering of isolated bat astrovirus according to specific species. Previous
454 studies on astroviruses in bats have also described this family of viruses to be involved solely in
455 tropism transmission which is species specific [38–40]. *Astrovirus* sequences isolated from different
456 bat species of the same site clustered together. This indicates potential cross species transmission of

457 astroviruses within each site, as reported by Xiao et *al* in bats sharing the same habitat [40]. We also
458 observed phylogenetic clustering of astroviruses from Chirundu and Magweto into same clades
459 showing a great diversity of astroviruses in circulation within the studied bat communities. Novel
460 astroviruses species have been reported in bats hosts [21], and in this study it is evident due to the
461 diverse astroviruses described. The *AstV* isolated from *E. helvum*, frugivorous bats, clustered together
462 and showed close phylogenetic relation to *BAstV* isolated from *Rousettus spp.* bats in China and
463 *Taphozous spp.* bats in Laos. One strain isolated from *E. helvum* clustered with *Hipposideros* bat
464 astroviruses sequences isolated in Chirundu site. This could mean that frugivorous bats and
465 insectivorous bats can be infected by the same type of AstVs strain thus indicative of the multi-host
466 spectrum for the Astroviridae family. Detection of *Astrovirus* in the frugivorous bats was low
467 compared to insectivorous bats, and this can be attributed to the insectivorous bats ability to adapt
468 and harbour more astroviruses than frugivorous bats [40].
469 Two strains, Mag-292 *H. Caffer*, Mag-573 *Rhinolophus spp* clustered within the *HuAstv* type 1 to 6,
470 that are known to cause infection and diarrhoea in children and infants [40]. Six of our strains isolated
471 from *E. helvum* and one from *Rhinolophus spp* formed sister clades to Human *AstV* HMO, associated
472 with severe extra-intestinal illnesses in humans [12] and VA-1-4 strains. Another *AstV* strain detected
473 in *Hipposideros* bat species was close and formed a sister clade with *HuAstv* MLB-1, -2 and -3 all
474 known to cause gastro-intestinal and neurological diseases with mild-severe symptoms in humans
475 [41]. This phylogenetic clustering could suggest that *HuAstV* and *BAstV* may have a shared common
476 ancestor. However, with limited sequence data the evolutionary history is merely a speculation that
477 needs to be further investigated through ORFs sequencing and MRCA reading. Another hypothesis
478 for the observed relation is that, since bats are known to be important reservoir hosts of astroviruses
479 with great diversity, they could be ancestors of the *HuAstV* due to the high prevalence of *AstV* detected
480 in bats. *HuAstV* transmission occurs via the faecal-oral route and contaminated food or water [2],
481 therefore it is unlikely that anthroponosis may have occurred. Third, this observed relation could be
482 due to a rare event or mutation, or a new strain of bat astrovirus that has not been described yet.
483 Further full genomic data as well as epidemiological data on these bat colonies are needed to reach a
484 full and true conclusion.
485 As novel strains and host species continue to be discovered, understanding *AstV* transmission and
486 their zoonotic potential is essential. Disseminating *HuAstV* and *AstV* from zoonotic reservoir hosts
487 remain significant threats to public health [4]. Astroviruses have a high genetic diversity, multiple
488 mechanism of generating additional diversity, and infect a wide range of host species therefore
489 understanding their prevalence in their wildlife hosts can help to predict or prevent the emergence of
490 novel astrovirus strains into domestic animals and human populations.

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**Author's contribution: Conceptualization**: VC, MA, HdN, MB, LG and FL.; **Methodology**: VC, MA, LG, HdN, DP, EG, GM, EM and FL.; **Software:** VC, MA, HdN and FL; **Validation**: VC, MA HdN, JC, and FL; **Formal Analysis**: VC, HdN, JC, and FL.; **Investigation**: VC, VP, NC, GM, and FL.; **Resources**: DP, GM, MP, MB, HdN, LG, and FL.; **Data Curation**: VC, HdN, and FL.; **Writing – Original Draft Preparation**: VC and FL; **Writing – Review & Editing**: HdN, MA, JC, MB, LG, VP, EG, DM, DP, EM, GM and FL; **Supervision**: HdN, DP, GM and FL.; **Project Administration**: LG, MB, HdN and FL; **Funding Acquisition**: MB, HdN, LG and FL.

**Author Approvals:**

All authors have seen and approved the manuscript, and that it hasn't been accepted or published elsewhere

512 **Reference**


513 [1] Bosch A, Pintó RM, Guix S. Human Astroviruses. Clin Microbiol Rev . 2014;27:1048–1074.
514 doi/10.1128/CMR.00013-14.

515 [2] Donato C, Vijaykrishna D. The Broad Host Range and Genetic Diversity of Mammalian and
516 Avian Astroviruses. Viruses . 2017;9:102. doi/10.3390/v9050102.

517 [3] Wu H, Pang R, Cheng T, et al. Abundant and Diverse RNA Viruses in Insects Revealed by
518 RNA-Seq Analysis: Ecological and Evolutionary Implications. Cristea IM, editor. mSystems.
519 2020;5. doi/10.1128/mSystems.00039-20.

520 [4] Roach SN, Langlois RA. Intra- and Cross-Species Transmission of Astroviruses. Viruses.
521 2021;13:1127. doi/10.3390/v13061127.

522 [5] El Taweel A, Kandeil A, Barakat A, et al. Diversity of Astroviruses Circulating in Humans,
523 Bats, and Wild Birds in Egypt. Viruses . 2020;12:485. doi/10.3390/v12050485.

524 [6] Vu D-L, Cordey S, Brito F, et al. Novel human astroviruses: Novel human diseases? J Clin
525 Virol . 2016;82:56–63. doi/10.1016/j.jcv.2016.07.004.

526 [7] Johnson C, Hargest V, Cortez V, et al. Astrovirus Pathogenesis. Viruses . 2017;9:22.
527 doi/10.3390/v9010022.

528 [8] Frémond M-L, Pérot P, Muth E, et al. Next-Generation Sequencing for Diagnosis and Tailored
529 Therapy: A Case Report of Astrovirus-Associated Progressive Encephalitis. J Pediatric Infect
530 Dis Soc . 2015;4:e53–e57. doi/10.1093/jpids/piv040.

531 [9] Cortez V, Meliopoulos VA, Karlsson EA, et al. Astrovirus Biology and Pathogenesis. Annu
532 Rev Virol . 2017;4:327–348. doi/10.1146/annurev-virology-101416-041742.

533 [10] Janowski AB, Klein RS, Wang D. Differential In Vitro Infection of Neural Cells by Astrovi-
534 ruses. Griffin DE, editor. MBio . 2019;10. doi/10.1128/mBio.01455-19.

535 [11] Boujon CL, Koch MC, Seuberlich T. The Expanding Field of Mammalian Astroviruses: Op-
536 portunities and Challenges in Clinical Virology. Adv Virus Res. 2017. p. 109–
537 137.10.1016/bs.aivir.2017.07.002

538 [12] Wohlgemuth N, Honce R, Schultz-Cherry S. Astrovirus evolution and emergence. Infect Genet
539 Evol . 2019;69:30–37. doi.org/10.1016/j.meegid.2019.01.009.

540 [13] Chu DKW, Poon LLM, Guan Y, et al. Novel Astroviruses in Insectivorous Bats. J Virol .
541 2008;82:9107–9114. doi/10.1128/JVI.00857-08.

542 [14] Dufkova L, Straková P, Širmarová J, et al. Detection of Diverse Novel Bat Astrovirus Se-
543 quences in the Czech Republic. Vector-Borne Zoonotic Dis . 2015;15:518–
544 521.doi/10.1089/vbz.2015.1813.

545 [15] Rougeron V, SuquetT E, Maganga GD, et al. Characterization and phylogenetic analysis of
546 new bat astroviruses detected in Gabon, Central Africa. Acta Virol . 2016;60:386–392.
547 doi/10.4149/av_2016_04_386.

548 [16] Lacroix A, Duong V, Hul V, et al. Diversity of bat astroviruses in Lao PDR and Cambodia.
549 Infect Genet Evol . 2017;47:41–50. doi/10.1016/j.meegid.2016.11.013.

550 [17] Lebarbenchon C, Ramasindrazana B, Joffrin L, et al. Astroviruses in bats, Madagascar. Emerg
551 Microbes Infect . 2017;6:1–3. doi/10.1038/emi.2017.47.

552 [18] Waruhiu C, Ommeh S, Obanda V, et al. Molecular detection of viruses in Kenyan bats and
553 discovery of novel astroviruses, caliciviruses and rotaviruses. Virol Sin . 2017;32:101–114.
554 doi/10.1007/s12250-016-3930-2.

555 [19] Hoarau F, Le Minter G, Joffrin L, et al. Bat Astrovirus in Mozambique. Virol J . 2018;15:104.
556 doi/10.1186/s12985-018-1011-x.

557 [20] Lee S-Y, Son K-D, Yong-Sik K, et al. Genetic diversity and phylogenetic analysis of newly
558 discovered bat astroviruses in Korea. Arch Virol . 2018;163:3065–3072. doi/10.1007/s00705-
559 018-3992-6.

560 [21] Bergner LM, Mollentze N, Orton RJ, et al. Characterizing and Evaluating the Zoonotic Poten-
561 tial of Novel Viruses Discovered in Vampire Bats. Viruses. 2021;13:252.
562 doi/10.3390/v13020252.

563 [22] Fischer K, Zeus V, Kwasnitschka L, et al. Insectivorous bats carry host specific astroviruses
564 and coronaviruses across different regions in Germany. Infect Genet Evol . 2016;37:108–116.
565 doi/10.1016/j.meegid.2015.11.010.

566 [23] Seltmann A, Corman VM, Rasche A, et al. Seasonal Fluctuations of Astrovirus, But Not Coro-
567 navirus Shedding in Bats Inhabiting Human-Modified Tropical Forests. Ecohealth.
568 2017;14:272–284. doi/10.1007/s10393-017-1245-x.

569 [24] Plowright RK, Peel AJ, Streicker DG, et al. Transmission or Within-Host Dynamics Driving
570 Pulses of Zoonotic Viruses in Reservoir–Host Populations. Remais J V., editor. PLoS Negl
571 Trop Dis . 2016;10:e0004796. doi/10.1371/journal.pntd.0004796.

572 [25] Bourgarel M, Pfukenyi DM, Boué V, et al. Circulation of Alphacoronavirus, Betacoronavirus
573 and Paramyxovirus in Hipposideros bat species in Zimbabwe. Infect Genet Evol.
574 2018;58:253–257. doi/10.1016/j.meegid.2018.01.007.

575 [26] Chidoti V, De Nys H, Pinarello V, et al. Longitudinal Survey of Coronavirus Circulation and
576 Diversity in Insectivorous Bat Colonies in Zimbabwe. Viruses. 2022;14:781.
577 doi/10.3390/v14040781.

578 [27] Monadjem A, Taylor PJ, Schoeman FPD, et al. Bats of Southern and Central Africa: A Bioge-
579 ographic and Taxonomic Synthesis. African Zool. 2011;46:437–437.
580 doi/10.1080/15627020.2011.11407657.

581 [28] Kocher TD, Thomas WK, Meyer A, et al. Dynamics of mitochondrial DNA evolution in ani-
582 mals: amplification and sequencing with conserved primers. Proc Natl Acad Sci.
583 1989;86:6196–6200. doi/10.1073/pnas.86.16.6196.

584 [29] Kumar S, Stecher G, Tamura K. MEGA7: Molecular Evolutionary Genetics Analysis Version
585 7.0 for Bigger Datasets. Mol Biol Evol . 2016;33:1870–1874. doi/10.1093/molbev/msw054.

[30] Guindon S, Dufayard J-F, Lefort V, et al. New Algorithms and Methods to Estimate Maximum-Likelihood Phylogenies: Assessing the Performance of PhyML 3.0. Syst Biol. 2010;59:307–321. doi/10.1093/sysbio/syq010.

[31] Milne I, Lindner D, Bayer M, et al. TOPALi v2: a rich graphical interface for evolutionary analyses of multiple alignments on HPC clusters and multi-core desktops. Bioinformatics. 2009;25:126–127. doi/25/1/126/302670.

[32] Lemoine F, Wilkinson E, Correia D, et al. Renewing Felsenstein's Phylogenetic Bootstrap in the Era of Big Data. Nature. 2018. doi/10.1038/s41586-018-0043-0

[33] Wilson, E. B. Probable inference, the law of succession, and statistical inference. Journal of the American Statistical Association. 1927; **22** (158): 209–212. doi/10.2307/2276774.

[34] Drexler JF, Corman VM, Wegner T, Tateno AF, Zerbinati RM, Gloza-Rausch F, et al. Amplification of emerging viruses in a bat colony. Emerg Infect Dis. 2011;17:449–56. doi/10.3201/eid1703.100526.

[35] Wacharapluesadee, S.; Duengkae, P.; Chaiyes, A.; Kaewpom, T.; Rodpan, A.; Yingsakmongkon, S.; Petcharat, S.; Phengsakul, P.; Maneeorn, P.; Hemachudha, T. Longitudinal study of age-specific pattern of coronavirus infection in Lyle's flying fox (Pteropus lylei) in Thailand. Virol. J. 2018, 15, 1–10. doi/10.1186/s12985-018-0950-6.

[36] Cappelle J, Furey N, Hoem T, Ou TP, Lim T, Hul V, Heng O, Chevalier V, Dussart P, Duong V. Longitudinal monitoring in Cambodia suggests higher circulation of alpha and betacoronaviruses in juvenile and immature bats of three species. Sci Rep. 2021;11(1):24145. doi/10.1038/s41598-021-03169-z.

[37] Mendenhall IH, Skiles MM, Neves ES, Borthwick SA, Low DHW, Liang B, Lee BPYH, Su YCF, Smith GJD. Influence of age and body condition on astrovirus infection of bats in Singapore: An evolutionary and epidemiological analysis. One Health .2017; 4: 27-33. doi.org/10.1016/j.onehlt.2017.10.001.

[38] George DB, Webb CT, Farnsworth ML, et al. Host and viral ecology determine bat rabies seasonality and maintenance. Proc Natl Acad Sci. 2011;108:10208–10213. doi/10.1073/pnas.1010875108.

[39] Baxendale W, Mebatsion T. The isolation and characterisation of astroviruses from chickens. Avian Pathol . 2004;33:364–370. doi/10.1080/0307945042000220426.

[40] Xiao J, Li J, Hu G, et al. Isolation and phylogenetic characterization of bat astroviruses in southern China. Arch Virol . 2011;156:1415–1423. doi/10.1007/s00705-011-1011-2.

[41] Kapoor A, Li L, Victoria J, et al. Multiple novel astrovirus species in human stool. J Gen Virol. 2009;90:2965–2972. doi/10.1099/vir.0.014449-0.