

Blood glucose, insulin and glycogen profiles in Sprague-Dawley rats co-infected with *Plasmodium berghei* ANKA and *Trichinella zimbabwensis* - A laboratory animal model for malaria and tissue-dwelling nematodes co-infection (#69406)

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Blood glucose, insulin and glycogen profiles in Sprague-Dawley rats co-infected with *Plasmodium berghei* ANKA and *Trichinella zimbabwensis* - A laboratory animal model for malaria and tissue-dwelling nematodes co-infection

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Background. *Plasmodium falciparum* and tissue dwelling helminth parasites are endemic in sub-Saharan Africa (SSA). The geographical overlap in co-infection is a common phenomenon. However, there is continued paucity of information on how the co-infection influence the blood glucose and insulin profiles in the infected host. Animal models are useful to elucidate the effects of co-infection on the outcome of the disease caused by the parasite and hence we assessed blood glucose, insulin and glycogen profiles in Sprague-Dawley rats co-infected with *Plasmodium berghei* ANKA (Pb) and *Trichinella zimbabwensis* (Tz), a tissue-dwelling nematode. **Methods.** One-hundred-and-sixty-eight male Sprague-Dawley rats (weight range 90-150g) were randomly divided into four separate experimental groups; Control (n=42), Pb-infected (n=42), Tz-infected (n=42) and Pb- + Tz-infected group (n=42). ~~Trichinella zimbabwensis~~ (Tz) muscle larvae were administered orally at 3 muscle larvae/g body weight at day 0 while Pb (10^5 parasitised RBCs) was administered intra-peritoneally at day 28 of the 42-day experimental study protocol. Control group was given an equal volume of phosphate buffered saline vehicle. Measurement of Pb percentage parasitaemia was done daily throughout the experimental study period for the Pb and the Pb + Tz group. Measurement of blood glucose was done every 3rd day in all experimental groups throughout the experimental study period. Animals were humanely sacrificed on day 0, 7, 14, 28, 35 and 42 (n=6 in each group) post-infection for collection of serum for the determination of insulin, and muscle and liver for glycogen concentration. Number of adults in the intestines and muscle larvae of ~~T. zimbabwensis~~ were also determined. Liver and skeletal muscle were harvested, snap frozen and stored at -70 °C for determination of glycogen concentration. **Results.** Blood glucose concentration in the malaria mono-infected group was significantly reduced ($P <$

0.05) in comparison to the control group, at day 7 post Pb infection and in comparison, to all other experimental groups, at day 21 post Pb infection. Liver and muscle glycogen concentration of all experimental groups, at day 0 post Pb infection, were significantly reduced ($P < 0.05$). Reduced insulin concentration in both Tz mono-infected and co-infected experimental groups coincided with reduced muscle glycogen concentration at day 0 post Pb infection. Gradual elevation of insulin concentration in both trichinella mono-infected and co-infected experimental groups, coincided with a concomitant gradual elevation of muscle glycogen concentration at day 7 and 14 post Pb infection. Serum insulin concentration in Pb mono-infected group persisted until day 7 post infection.

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ABSTRACT

Background. *Plasmodium falciparum* and tissue dwelling helminth parasites are endemic in sub-Saharan Africa (SSA). The geographical overlap in co-infection is a common phenomenon. However, there is continued paucity of information on how the co-infection influence the blood glucose and insulin profiles in the infected host. Animal models are useful to elucidate the effects of co-infection on the outcome of the disease caused by the parasite and hence we assessed blood glucose, insulin and glycogen profiles in Sprague-Dawley rats co-infected with *Plasmodium berghei* ANKA (Pb) and *Trichinella zimbabwensis* (Tz), a tissue-dwelling nematode.

Methods. One-hundred-and-sixty-eight male Sprague-Dawley rats (weight range 90-150g) were randomly divided into four separate experimental groups; Control (n=42), Pb-infected (n=42), Tz-infected (n=42) and Pb- + Tz-infected group (n=42). *Trichinella zimbabwensis* (Tz) muscle larvae were administered orally at 3 muscle larvae/g body weight at day 0 while Pb (10^5 parasitised RBCs) was administered intra-peritoneally at day 28 of the 42-day experimental study protocol. Control group was given an equal volume of phosphate buffered saline vehicle. Measurement of Pb percentage parasitaemia was done daily throughout the experimental study period for the Pb and the Pb + Tz group. Measurement of blood glucose was done every 3rd day in all experimental groups throughout the experimental study period. Animals were humanely sacrificed on day 0, 7, 14, 28, 35 and 42 (n=6 in each group) post-infection for collection of serum for the determination of insulin, and muscle and liver for glycogen concentration. Number of adults in the intestines and muscle larvae of *T. zimbabwensis* were also determined. Liver and skeletal muscle were harvested, snap frozen and stored at -70 °C for determination of glycogen concentration.

Results. Blood glucose concentration in the malaria mono-infected group was significantly reduced ($P < 0.05$) in comparison to the control group, at day 7 post Pb infection and in comparison, to all other experimental groups, at day 21 post Pb infection. Liver and muscle glycogen concentration of all experimental groups, at day 0 post Pb infection, were significantly reduced ($P < 0.05$). Reduced insulin concentration in both Tz mono-infected and co-infected experimental groups coincided with reduced muscle glycogen concentration at day 0 post Pb infection. Gradual elevation of insulin concentration in both trichinella mono-infected and co-infected experimental groups, coincided with a concomitant gradual elevation of muscle glycogen concentration at day 7 and 14 post Pb infection. Serum insulin concentration in Pb mono-infected group persisted until day 7 post infection.

Keywords: Co-infection, Blood glucose, Tissue-dwelling helminths, Insulin, Liver glycogen, Muscle glycogen, *Plasmodium berghei*, *Trichinella zimbabwensis*

INTRODUCTION

Co-infection and poly-parasitism remain a major health challenge in sub-Saharan Africa (SSA) (Onkoba et al., 2015). There is a geographical overlap in the endemicity of malaria and helminth infections in SSA (Onkoba et al., 2015) which include tissue-dwelling helminth (TDH) parasites (Onkoba et al., 2015), and 90% of all global malaria related deaths also occur on SSA (Hotez and Kamath, 2009). Undoubtedly, malaria-helminth co-infections are a common phenomenon in SSA (Onkoba et al., 2015). *Ascaris lumbricoides*, *Taenia solium* cysts, *Echinococcus* spp cysts, filarial worms, *Schistosoma* spp, *Fasciola* spp and *Trichinella* spp are some of the common TDHs reported worldwide and have either their larvae or adults stages passing through host tissues other than the gastrointestinal tract and are common in SSA (Onkoba et al., 2015). Glucose homeostasis during *Plasmodium* spp mono-infection is well understood and documented under laboratory conditions. However, there is paucity of information on glucose homeostasis during *Plasmodium* spp-TDHs co-infection. *Trichinella zimbabwensis* (Tz) has been successfully used in laboratory animal model experiments for TDHs (Onkoba et al., 2015a) and in determining chemokine, cytokine and haematological profiles in Sprague-Dawley rats co-infected with *Plasmodium berghei* ANKA (Murambiwa et al., 2020) and we propose extending the use of this parasite to determine the blood glucose, insulin and glycogen profiles during co-infection with *P. berghei* ANKA (Pb).

During *Trichinella* spp mono-infection, it has been reported that trichinella-induced hypoglycaemia could be mediated via three possible mechanisms, viz; high glucose consumption by trichinella parasites, reduction in the absorptive capacity of the intestine or impairment of glucose production by the liver (Wu et al., 2009). Increase of glucose uptake in infected muscle cells of the host has also been reported as a trichinella-induced hypoglycemia mechanism (Wu et al, 2009). *Trichinella*-induced hypoglycaemia in humans, mice and dogs has also been reported in other studies (Nishina et al., 2004, Reina et al., 1989, Steward, 1983, Montgomery et al., 2003). During trichinella infection, there is increased demand for glucose by both trichinella muscle larvae and infected muscle cells for their rapid growth and metabolism (Wu et al., 2009). However, in Nile crocodile (*Crocodylus niloticus*) experimentally infected with *T. zimbabwensis* no hypoglycaemia was detected in infected groups contrary to previous findings in mammals (La Grange and Mukaratirwa 2014). In high infection groups, peak blood glucose concentration was recorded 35-49 days post infection (La Grange and Mukaratirwa 2014). Significant increase in blood glucose concentration 30 days post infection was also reported in pigs (Oltean et al., 2012).

An increase in glycogen concentration was observed at day 8 post-infection and day 18 post-infection in mice infected with *T. spiralis* and *T. pseudospiralis* infections (Wu et al., 2009). However, at day 28 and 48 post-infection with *T. spiralis* and *T. pseudospiralis* infections, the study showed depletion of glycogen stores (Wu et al., 2009). Increased glucose uptake through insulin signaling pathways has been speculated to be the possible mechanism of increased glycogen accumulation post-infection (Wu et al.,

2009). Infection of mice with *T. spiralis* and *T. pseudospiralis* has been reported to cause an initial decrease in insulin concentration, which is restored back to normal (Wu et al., 2009). Increased insulin concentration in mice infected with *T. zimbabwensis* compared to the control has also been reported (Onkoba et al., 2016). However, the effect of Pb + Tz co-infection on insulin, liver and muscle glycogen concentration remains obscure. We recently reported that Tz infection predisposed the co-infected animals towards rapid development of Pb parasitaemia during co-infection (Murambiwa et al., 2020). However, there is still paucity of information on how this phenomenon influences glucose homeostasis during co-infection.

Therefore, glucose homeostasis during mono-infection of the two parasites is fairly understood and documented, however, glucose homeostasis during co-infection, remain obscure to date. It is against this background that we attempt to use an animal model aimed to determine the blood glucose, insulin profiles and glycogen concentration in Sprague-Dawley rats co-infected with Pb ANKA and Tz and have an insight of the impact of tissue-dwelling helminths in the epidemiology of malaria in geographical areas where the two overlap.

MATERIALS & METHODS

Study animals and study design

Male Sprague Dawley rats (90-150g) used in this study were bred and maintained at the Biomedical Research of the University of KwaZulu-Natal, South Africa (Murambiwa et al., 2020). The animals were fed rat chow (Meadow feeds, Pietermaritzburg, South Africa) with free access to water and maintained following standard laboratory conditions (Murambiwa et al., 2020). The animals were divided into four separate experimental groups were randomly divided into control (n = 42), *T. zimbabwensis*-infected (Tz) (n = 42), *Plasmodium berghei*-infected (Pb) (n = 42) and Pb + Tz-infected (n = 42) group) using a simple random method (Murambiwa et al., 2020) with the only variation in the parameters which were measured in this study. Severe weight loss, anaemia and body temperature <32 °C were considered as an endpoint for Pb- and Pb + Tz infected groups.

At the end of the experimental period, animals were humanely sacrificed using CO₂ on day 0, 7, 14, 28, 35 and 42 (n=6 in each group) post Tz-infection for collection of sera and rat carcasses for determination of serum insulin and trichinella adult worm and muscle larvae load respectively (Murambiwa et al., 2020). Liver and skeletal muscle were harvested, snap frozen and stored at -70 °C for determination of glycogen concentration. ARRIVE Guidelines for reporting animal research were followed in this experiment as reported by Kilkenney et al. (2010) (Kilkenny et al., 2010). The completed ARRIVE Guidelines checklist is attached.

Ethical statement

Experimental procedures and protocols for the study were reviewed and approved by the University of KwaZulu-Natal Animal Ethics Committee (AREC/018/016).

Induction and determination of Pb and Tz infection

The procedures followed for the induction and determination of Pb and Tz are as described by Murambiwa *et al.* (2020).

Determination of serum insulin

An ultrasensitive rat insulin ELISA kit (DRG Instruments GmbH, Marburg, Germany) was used for the determination of serum insulin. This immunoassay is a quantitative method that utilizes two monoclonal antibodies specific for insulin and procedures were followed as described by the manufacturer. The lower limit of detection was 1.74 pmol/L.

Determination of liver and gastrocnemius muscle glycogen concentration

Glycogen was determined as described by Ngubane *et al.* (2011). Homogenization of pre-weighed (0.25-0.5g) liver and gastrocnemius muscle tissue samples was done in 2 mL of 30% potassium hydroxide (300 g/L) and boiled at 100°C for 30 minutes and then cooled in ice-saturated sodium sulfate. Ethanol was used to precipitate glycogen, which was then pelleted and resolubilised in deionized water. Treatment with anthrone reagent was done for glycogen content determination, which was done at 540nm using a Spectrostar Nano microplate reader (BMG Labtech Germany).

Data Analysis

GraphPad InStat Software (version 4.00, Statistical comparison of the differences between the means of the experimental groups means with the 95% upper and lower confidence intervals, and/or box plots with the median and the 25% and 75% quartiles was done using GraphPad Software, San Diego, CA, USA. Effects of co-infection on blood glucose, serum insulin as well as liver and gastrocnemius muscle glycogen concentration were determined using two-way analysis of variance (ANOVA), followed by Bonferroni post hoc test. A value of $p < 0.05$ was considered statistically significant.

RESULTS

There were no adverse events in this experimental protocol.

Effect of co-infection on Pb parasitaemia and Tz adult worms and muscle larvae load

The effect of coinfection on Pb parasitaemia and Tz adult worms and muscle larvae load has been reported elsewhere by Murambiwa et al (2020) (Figure 1 and 2) where it is shown that percentage parasitaemia of the Pb mono-infected group was generally lower than in the co-infected group (Figure 1).

Adult worm counts were recuperated in both experimental groups at day 7 and 14 post-infection and no adult worms were recovered as from day 21 post infection) while muscle larvae (ML) counts were detected from day 28 post-infection and relatively higher in the co-infected group (Figure 2).

Effects of Tz and Pb co-infection on blood glucose concentration

Effects of Tz and Pb co-infection on blood glucose concentration is shown in Figure 3. *Plasmodium berghei* (Pb) mono-infected group had a significant reduction in blood glucose concentration at day 7 post Pb infection ($P<0.05$) in comparison to control group. Also, at day 14 post Pb infection, there was a further reduction in blood glucose concentration in Pb mono-infected group ($P<0.001$) in comparison to control group. Blood glucose concentration in Pb mono-infected group was significantly reduced, in comparison to Tz mono-infected ($P<0.001$) and co-infected ($P<0.001$) experimental groups at day 14 post Pb infection. There were no significant differences in blood glucose concentration among experimental groups at day 21 post Pb infection.

Effects of Tz and Pb co-infection on serum insulin concentration

There was a significant reduction in serum insulin concentration in Tz mono-infected ($P<0.05$) and co-infected ($P<0.05$) experimental groups at day 0 post Pb infection in comparison to both the control and Pb infected groups (Table 1). The co-infected group serum insulin concentration was significantly reduced ($P<0.05$) at day 14 Pb infection in comparison to the control group. There were no significant differences in serum insulin concentration of all experimental groups 21 days post Pb infection.

Effects of Tz and Pb co-infection on liver and gastrocnemius muscle glycogen concentration

There was a significant reduction in liver glycogen concentration in Tz mono-infected ($P<0.001$) and co-infected ($P<0.001$) experimental groups at day 7 post Pb infection in comparison to the control group (Table 1). A significant reduction in liver glycogen concentration in Tz mono-infected ($P<0.001$) and co-infected ($P<0.01$) experimental groups was also observed at day 7 post Pb infection in comparison to Pb mono-infected group (Table 1). Additionally, liver glycogen concentration in co-infected group was significantly

reduced ($P < 0.05$) at day 21 post Pb infection in comparison to the control group. There were no significant differences in liver glycogen concentration among all experimental groups at day 0 post Pb infection.

There was a gradual increase in muscle glycogen concentration at day 0, 7 and 14 post Pb infection although the differences among groups were not statistically significant (Table 1). Co-infected group had elevated muscle glycogen concentration at day 7 and 14 post Pb infection in comparison to Tz mono-infected group, although the differences were not statistically significant.

DISCUSSION

Elevated Pb percentage parasitaemia in Pb mono-infected group coincided with reduced blood glucose concentration in comparison to control group at day 7 post Pb infection. Blood glucose concentration in the Pb mono-infected group was significantly reduced in comparison to control group at day 7 post Pb infection and in comparison, to all other experimental groups at day 21 post Pb infection. Indeed, blood glucose lowering effects of *Plasmodium* parasites have been previously reported (Mehta et al., 2005). Malaria parasites have been reported to precipitate hypoglycaemia through unclear mechanisms (English et al., 1998, Thien et al., 2006). There are studies which have reported that hypoglycemia is induced through utilization of glucose and gluconeogenic substrates such as thiamine in an effort to meet the parasite's increased energy demands (Krishna et al., 1999).

During *P. falciparum* infection, studies have reported depressed aerobic glycolysis and increased lactic acid acidosis in the host following parasite-induced depletion of vital gluconeogenic substrates such as thiamine (Krishna et al., 1999). Furthermore, other studies have reported that malaria parasites are completely dependent on the host for all energy requirements (Phillips, 1989). On the other hand, the liver plays a major role in glucose homeostasis and induced hypoglycaemia may also be due to *P. falciparum* induced hepatocellular damage (Dekker et al., 1997) which causes loss of intracellular fluid components from the hepatocytes such as liver enzymes to the extracellular fluid compartment (Dekker et al., 1996; Dekker et al., 1997; Kauser et al., 2010). Hepatocellular damage may also lead to slow insulin receptor recycling (Onyesom and Agbo, 2011), thereby aggravating malaria induced hypoglycaemia.

Increased glucose demands during trichinella infection also coincide with NBL larvae migration, penetration, establishment and encystment in the striated muscle nurse cells (Wu et al., 2009). Studies have also reported a temporary decrease in blood glucose concentration between day 8 and 28 post *T. spiralis* and *T. pseudospiralis* infection, coinciding with peak larval growth and development (Wu et al., 2009). The same study further reported increased glycogen accumulation in infected muscle cells during transient hypoglycemia phase (Wu et al., 2009). Indeed, it is speculated that trichinella parasite may regulate glycogen synthesis in the infected muscle cells in line with its glucose requirements for its growth and metabolism (Wu et al., 2009).

Hyperinsulinemia was observed in the current study and results are in agreement with *Wu et al.* (2009), who have reported that the transient hypoglycemia reported between 8 and 28 days post *T. spiralis* and *T. pseudospiralis* infection was a result of increased glucose uptake by infected muscle cells mediated through up regulation of insulin signaling pathway related genes (*Wu et al.*, 2009), and not necessarily through increased blood insulin concentration (*Wu et al.*, 2009). However, there were no statistical differences in insulin concentration of all the experimental groups in the current study, 14 days post *P. berghei* infection. This could be suggestive of differences in blood glucose uptake mechanisms following muscle cells formation completion. In comparison to control group, reduction in serum insulin concentration in Pb mono-infected group persisted at day 7 post Pb infection in comparison to control group.

Glucose transport is insulin mediated, via specific insulin signaling pathways in both the skeletal muscle and adipose tissue (*Wu et al.*, 2009). Insulin mediated glucose transport has also been reported to enhance conversion of glucose to glycogen storage molecule (*Wu et al.*, 2009). Studies have also shown that infection of mice with *T. spiralis* and *T. pseudospiralis* causes an initial decrease in insulin concentration, which is restored back to normal (*Wu et al.*, 2009). Interestingly, increased insulin concentration in mice infected with *T. zimbabwensis* compared to the control has also been reported (*Onkoba et al.*, 2016). Also, studies have shown a difference in glucose metabolism handling between children and adults in *falciparum* malaria through unclear mechanisms (*Dekker et al.*, 1996). In the current study, weanling male Sprague-Dawley rats (90-150g) were used for all experimental protocols.

There was elevated liver glycogen concentration which coincided with reduction of blood glucose and plasma insulin through unclear mechanisms. Significantly reduced insulin concentration in both Tz mono-infected and co-infected experimental groups in comparison to both the control and Pb infected groups coincided with reduced muscle glycogen concentration at day 0 days post Pb infection. Gradual elevation of insulin concentration in both Tz mono-infected and co-infected experimental groups coincided with a concomitant gradual elevation of muscle glycogen concentration in both Tz mono-infected and co-infected experimental groups at day 7 and 14 post Pb infection. Additionally, elevated Tz infection coincided with elevated muscle glycogen concentration. Although these differences did not reach statistical significance, they could be of biological importance. Interestingly, there was a significant reduction in glycogen concentration in Tz mono-infected and co-infected groups in comparison to both control and Pb mono-infected group. Indeed, Tz infection could cause hypoglycemia through three possible mechanisms; high glucose consumption by developing stages of *Trichinella* parasites, reduction in the absorptive capacity of the intestine or impairment of glucose production by the liver (*Wu et al.*, 2009). *Nishina et al.* (2004) has previously reported hypoglycemia through unknown mechanisms in mice. Studies have also reported hypoglycemia in dogs experimentally infected with *T. spiralis* (*Reina et al.*, 1989). Several *Trichinella* induced hypoglycemia mechanisms have been postulated, such as increased consumption of

glucose by rapidly growing *Trichinella* larvae within muscle cells (Wu et al., 2009). Steward (1983) reported that trichinella induced hypoglycemia was mediated via increased total glycogen in infected muscle tissue as well as increased glycogen content in trichinella larvae. Other researchers have ascribed the trichinella induced hypoglycemia to increased metabolic activity associated with sugar metabolism of nurse cells (Montgomery et al., 2003).

Interestingly, there were no significant differences in the gastrocnemius muscle glycogen concentration of the experimental groups of animals throughout the experimental period. However, the observed increase in gastrocnemius muscle glycogen concentration of the co-infected group at day 7 and 14 post Pb infection could be of biological importance since glucose is transported in the muscle via insulin mediated GLUT 4 transporters (Azpiazu et al., 2000, Ferrer et al., 2003).

CONCLUSIONS

To the best of our knowledge, there is paucity of studies that have investigated the effects of Pb and Tz (representing a tissue-dwelling nematode) co-infection on blood glucose, glycogen concentration and insulin profiles in male Sprague-Dawley rats as an animal model for human infection. We conclude from our study that Tz mono-infection and Tz + Pb co-infection did not have blood glucose lowering effect in the host. There was hypoinsulinemia and increase in liver glycogen content in Tz mono-infection and Tz + Pb co-infection groups but the triggering mechanism remains unclear. We postulate that there are other possible mechanisms through which tissue-dwelling parasite up-regulates the glucose store without decreasing the blood glucose concentration as exhibited by the absence of hypoglycaemia in Tz + Pb co-infection group. Limitations of the study is in the use of *T. zimbawensis* which has striated muscles as site of predilection and future studies should include other tissue-dwelling helminths such as *Taenia taeniformis* which has strobilocercus as the metacestode in the liver to mimic infections such as hydatid disease in humans.

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ADDITIONAL INFORMATION AND DECLARATIONS

Authors' contributions

- Pretty Murambiwa conceived and designed the experiments, performed the experiments, analyzed and interpreted the data and wrote the paper.
- Achasih Quinta Nkemzi performed the experiments, analyzed and interpreted the data, and wrote the paper.
- Samson Mukaratirwa conceived and designed the experiments, coordinated the research, analyzed and interpreted the data, contributed reagents, materials, analysis tools, and wrote the paper

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Competing interests

The authors declare that they have no competing interests.

Additional information

No additional information is available for this paper

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Figure 1

Percentage parasitaemia in male Sprague-Dawley rats infected with *Plasmodium berghei* (Pb) only and co-infected with Pb and *Trichinella zimbabwensis* (Pb + Tz) (Murambiwa et al., 2020).

Day 0 represents the day of Pb infection when Tz muscle larvae were in the rat muscle at day 28 post Tz infection. Values are presented as means and vertical bars indicate standard error of mean (SEM). N = 6 for each group). ** = $P < 0.01$, *** = $P < 0.001$

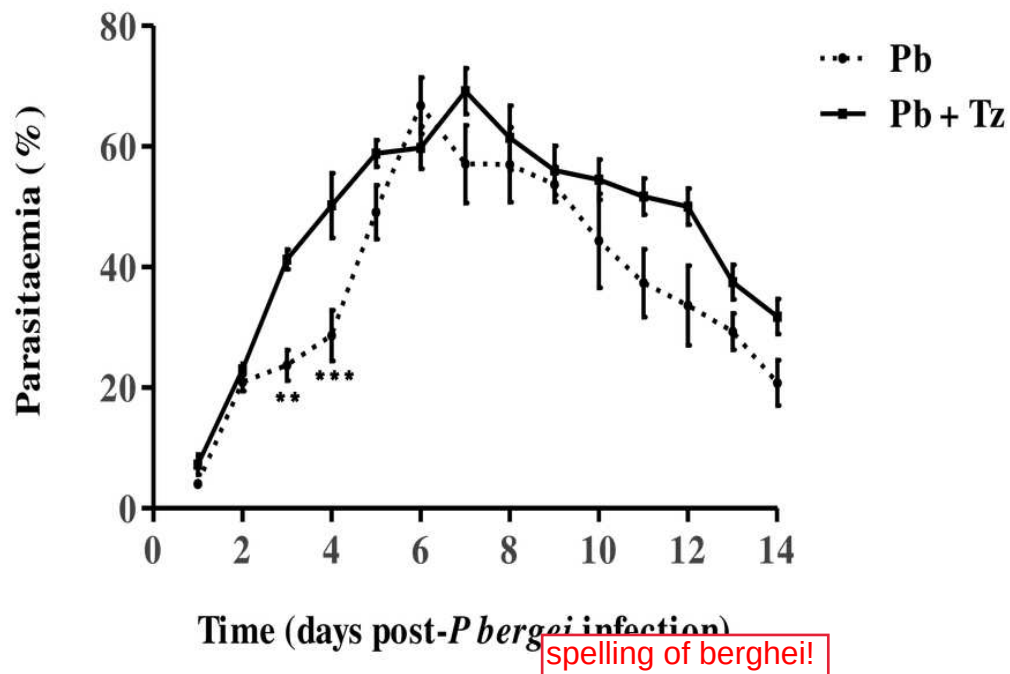


Figure 2

Mean number of intestinal adult worms (AW) and muscle larvae counts (ML) per gram of muscle (lpg) recovered from rats infected with *Trichinella zimbabwensis* (Tz) and the group co-infected at day 28 post-infection (Murambiwa et al, 2020).

Values are presented as means and vertical bars indicate standard error of mean (SEM). N = 6 for each group.

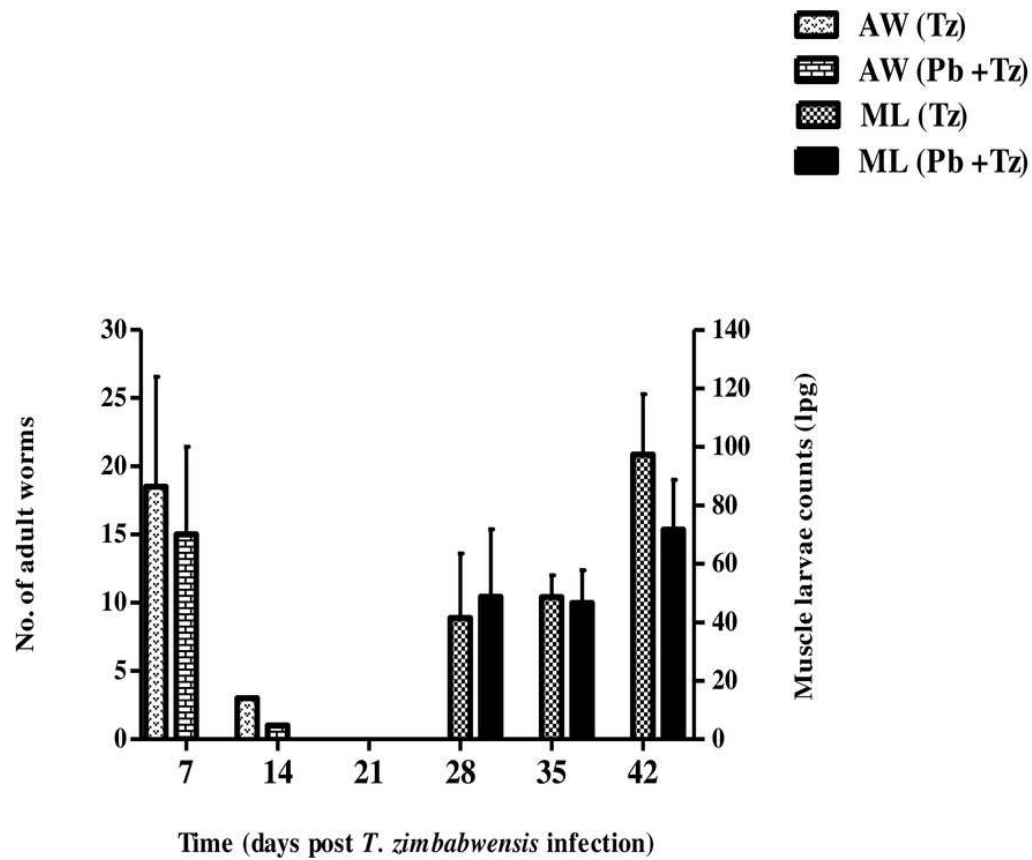


Figure 3

Comparison of the effects of *Plasmodium berghei* (Pb) and *Trichinella zimbabwensis* (Tz) mono-infection and co-infection (Pb + Tz) on blood glucose concentration.

Day 0 represents the day of Pb infection when Tz muscle larvae were in the rat muscle at day 28 post Tz infection. Values are presented as means and vertical bars indicate standard error of mean (SEM). N = 6 for each group. ** = $P < 0.01$, *** = $P < 0.001$.

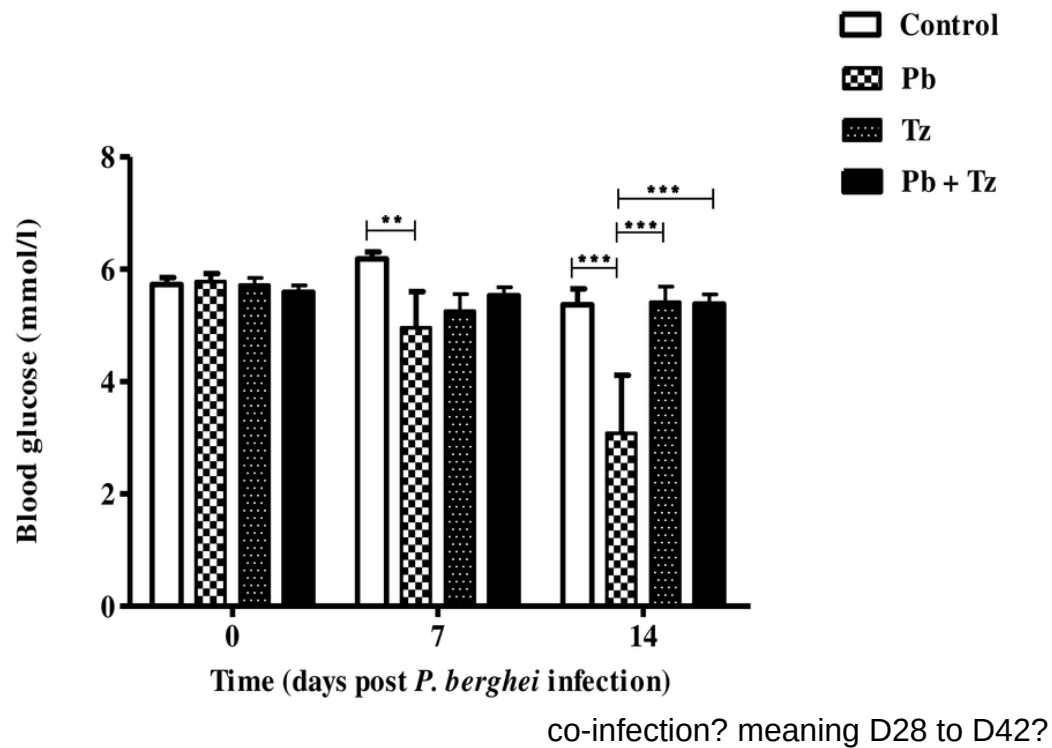


Table 1 (on next page)

Median values (25% -75% quartiles) of serum insulin concentration, liver and muscle glycogen (mg/g tissue) in Sprague-Dawley rats co-infected with *Plasmodium berghei* ANKA (Pb) and *Trichinella zimbabwensis* (Tz) (Pb + Tz), Pb mono-infection and

1

Parameter		Days post <i>P. berghei</i> infection	Experimental Groups			
			Control	Pb	Tz	Pb + Tz
Insulin (pmol/l)		Day 0	21.93 (19.70-46.89)	21.93 (19.70-46.89)	2.69 (2.48-4.17)	2.69 (2.48-4.17)
		Day 7	21.93 (19.70-46.89)	25.66 (8.75-39.96)	4.35 (2.26-25.84)	5.39 (3.04-12.92)
		Day 14	21.93 (19.64-46.89)	15.93 (9.37-25.71)	27.06 (15.97-40.54)	18.62 (14.14-26.97)
Liver glycogen (mg/g tissue)		Day 0	0.73 (0.52-0.77)	0.57 (0.49-0.50)	0.64 (0.45-0.75)	0.68 (0.41-0.79)
		Day 7	0.62 (0.53-0.75)	0.65 (0.54-0.72)	0.28 (0.20-0.33)***	0.37 (0.28-0.39)***
		Day 14	0.63 (0.43-0.76)	0.60 (0.49-0.75)	0.58 (0.47-0.62)	0.35 (0.25-0.53)*
Muscle glycogen (mg/g tissue)		Day 0	-	-	0.11 (0.10-0.13)	0.14 (0.09-0.16)
		Day 7	-	-	0.18 (0.12-0.30)	0.25 (0.17-0.30)
		Day 14	-	-	0.18 (0.18-0.22)	0.27 (0.21-0.36)

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