# *Eimeria falciformis* (BayerHaberkorn) and wild derived isolates from house mice: differences in parasite reproduction dynamics, host pathology and immune reactions

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Infections with different species of the genus *Eimeria* (Apicomplexa:Coccidia) can have different dynamics, vary in the immune reactions they induce and in pathology they imposed on their hosts. Here, we investigate dynamics of parasite reproduction (oocyst shedding), cytokine profiles, tissue inflammation and cellular immune response in infections of mice with wild derived isolates of *E. flaciformis* (falW) and *E. ferrisi* Levine and Evens, 1965 (ferW) and *E. flaciformis* BayerHaberkorn (falL)*,* a isolate propagated in the laboratory for roughly 60 years through experimental passaging.

The two different species *E. flaciformis* and *E. ferrisi* have different infection daynamics in NMRI mice: after a pre-patency period of only 3 days oocyct shedding of ferW peaks in intensity at 6 days post infection (dpi). In contrast, after pre-patency of 5 day, oocyst shedding of falL and falW has a later peak intensity at 8 dpi and 9 dpi, respectively. Concomitant impact on the host is parasite species specific and maximal weigh loss occurs before maximal oocyst shedding in infections with *E. ferrisi* but after or simultaneously with maximal shedding in *E. falciformis* (both isolates). mRNA expression of the cytokines CxCL9, IL6, IL10 and IL 12, TGFβ and the transcription activator STAT6 in the spleen is significantly increased in infections with falL. All those genes are detected at significantly lower levels in infections with the two wild derived isolates falW and ferW. Inflammation manifested in a mixture of granulocytes and mononuclear cells in the ceca and the area occupied by leukocyte infiltration is stronger in infections with falL than in those with both wild derived isolates (falW and ferW).

We can conclude that NMRI mice mount a stronger immune and inflammatory response against the laboratory isolate of *E. flaciformis* compared to both wild derived lines of *E. falciformis* an *E. ferrisi*. It seems possible that evolution during laboratory passages produced a parasite isolate, which still resembles a closely related wild derived isolate genetically and in infection dynamics, but differs in the immune response it induces. Immune response against laboratory and wild derived isolates of *E. falciformis* differs to the extend, that the latter is more similar to the response against a different species of the parasite. In most general terms these result show that caution is needed when using laboratory strains of pathogens to draw conclusions about infections in natural systems.

**Introduction**

Maintenance of parasite life cycles via serial passaging is a cornerstone of experimental parasitology. Parasites are propagated under defined and controlled conditions with the aim to provide infective stages for experiments (Lucius and Loos-Frank, 2008). The procedure allows the parasite to evolve due to mutation and genetic drift or adaptation to the passaging host and environment (Burke, 2012). Genetic drift is promoted by the use of small inocula during passaging and drift can act while diversity of a parasite isolate is reduced intentionally to obtain a clonal strain. In clonal strains mutation and drift can continue to act (Theron et al 2008; Chelo et al 2012). Considering adaptive evolution, an important part of the passaging environment is given by living hosts, or host cells (in case of propagation in culture), which usually have low genetic diversity (e.g, clonal or inbred lines, cultures) ( Elena and Lenski, 2003), are immunologically naive due to the absence of previous infections (Mackinnon ‎2004; Abolins et al 2017) and lack co-infections with other parasites (Abolins et al 2017). Procedures for serial passaging of parasites typically collect infective stages at a particular time after infection and use the obtained inoculum to infect new animals or use haphazard infections in dense environments. Iteration of such a static routine may further differ from natural parasite environments with different timing of infections and variable transmission. In most cases both the biotic (host) and abiotic passaging environment thus differs profoundly from the that experienced by the parasite during its life cycle under natural conditions. To summarize, parasite laboratory strains might experience both neutral and adaptive processes and as a consequence be unrepresentative for analogues in the field (Irschick and Reznick 2009).

Serial passage leads in most cases to higher virulence (enhanced growth and reproduction of theThey

parasite and larger impact on the host), especially in the host type used for passaging (reviewed by (Ebert, 1998) Garland and Rose 2009). This can be due to low genetic diversity in host populations (for example inbred line used for passage) that may reduce fitness trade-offs associated with specialization and promote the expansion of highly virulent pathogens. This phenomenon has been long demonstrated in several parasitic system such as viruses (Kubinak et al., 2015) and plant pathogenic parasites (Altermatt and Ebert, 2008) and equally in vertebrate host-eukaryotic parasite systems like the apicomplexa parasite of *Plasmodium* spp in rodents (Mackinnon and Read, 1999 and‎2004; Barclay et al, 2014) and in its mosquito vector (Pigeault et al., 2015) or common laboratory strains of *Toxoplasma gondii* (Khan et al. 2009). These studies collectively suggests adaptation to the passage host in a way that increases parasite virulence.

Consistent with this, serial passage of highly virulent isolates of the apicomplexa parasite *Eimeria* spp can lead to attenuation when only the first oocyst committing to sexual reproductions are selected for the next passage ([Shirley](https://www.ncbi.nlm.nih.gov/pubmed/?term=Shirley MW%5BAuthor%5D&cauthor=true&cauthor_uid=3375582) and [Bellatti](https://www.ncbi.nlm.nih.gov/pubmed/?term=Bellatti MA%5BAuthor%5D&cauthor=true&cauthor_uid=3375582) 1988, [McDonald](https://www.ncbi.nlm.nih.gov/pubmed/?term=McDonald V%5BAuthor%5D&cauthor=true&cauthor_uid=6877863) and [Ballingall](https://www.ncbi.nlm.nih.gov/pubmed/?term=Ballingall S%5BAuthor%5D&cauthor=true&cauthor_uid=6877863) 1983, Matsubayashi et al, 2016). These attenuated strains are called “precocious lines” and are the basis for successful live vaccines used in the poultry industry (Shirley and Long, 1990)(Shirley and Millard, 1986). Given the practical implications of this phenomenon, numerous serial passage experiments focused on changes in parasite life history that arise as a consequence of parasite adaptation to the passage host. It is surprising that only a few empirical studies attempt to correlate these evolutionary changes of attenuation or enhanced virulence in the parasite with physiological (e.g. immune-) responses in the passaging host.

Species of the genus *Eimeria* have usually a small host range, often infecting a single host species (Hashimoto et al., 2014; Hnida and Duszynski, 1999; Kvičerová and Hypša, 2013; Vrba and Pakandl, 2015) and reside at specific sites within the intestines of their hosts (Chapman et al., 2013; Haberkorn, 1970; Owen, 1975). All species have a direct life cycle with asexual expansion and sexual reproduction within epithelial cells of the gastrointestinal tract before diploid transmission stages (oocysts) are released. Oocysts become infective after reductive divisions (sporuation) in the environment (Cacho et al., 2012; Canning and Anwar, 1968)Kheysin 1972.

Different species of *Eimeria* are widespread in diverse host species including all vertebrates. Speciescausing coccidiosis in livestock have long been a focus of veterinary research due to their economical impact (Brake et al., 1997; Cacho et al., 2012; Gadde et al., 2009; Laurent et al., 2001; Swaggerty et al., 2011). *Eimeria* species capable of natural infection of the mouse (*Mus musculus*), the main model system of biomedical research, have been proposed as a model for e.g. host immune response against *Eimeria* (Heitlinger et al., 2014; Schmid et al., 2014). Serial passaging of laboratory isolates of *Eimeria* is conducted by a collection of oocysts at the day of peak shedding, in case of the isolate *E. flaciformis* BayerHaberkorn (Haberkorn, 1970) 7 days post infection (dpi). Oocysts are sporulated an aqueous solution of potassium dichromate and inocula are used for new infections two to six month later, before interactivity decreases (Ref). The isolate *E. flaciformis* Bayer Haberkorn has been isolated in 1960 (Haberkorn 1970) and has been propagated in the laboratory for nearly 60 years through experimental passaging. It has been routinely used to asses basic immunology against *Eimeria* infections (Ehret et al., 2017; Pogonka et al., 2010; Schmid et al., 2014, 2012; Stange et al., 2012; Steinfelder et al., 2005).

Although normally controlled by the host immune system, and self-limiting even in immune-compromised hosts (Ehret et al., 2017), *Eimeria* infection can lead to a variety of outcomes in infected host for different species (Schito et al., 1996). After penetrating the intestine, Eimeria stimulate production of the cytokines, interleukines (IL) and interferon (IFN)-γ by immune cells. Several cytokines promote a strong Th1 response which limits parasite production (Byrnes et al., 1993; Laurent et al., 2001; Lillehoj, 1998; Lillehoj and Choi, 1998; Ovington et al., 1995; Rose et al., 1992). Previous studies has shown that an inflammatory reaction at the site of infection driven by IFNγ is dominant during *E. flaciformis* infection of mouse caecum (Schmid et al. 2014). And cytokines such as TGF-beta, EGF, IL-1 IL6 and IL10 in addition to the chemokines belonging to CXC and CCL family are up-regulated (Ehret et al., 2017; Schmid et al., 2012; Stange et al., 2012). To advance the understanding of this complex infection, both local and systemic studies are required, which complement and direct each other.

Curiously, systemic immune response during *E. flaciformis* infection of the mouse has not been studied in such detail. The spleen is in mammals a secondary lymphoid organ in which innate and adaptive immune responses are controlled (Bronte and Pittet, 2013). Therefore, gene expression in the spleen is commonly used as an indicator of systemic immune response during parasite infections (Mueller et al, 2007, Li et al, 2009; Burk et al 2010; Shen et al; 2015). Few studies have assessed it in *Eimeria* infections e.g. via expression of cytokines in the spleen. Steinfelder et al. (2005) showed that proliferated cells from spleen of *E. flaciformis* infected mice released IFNγ and IL4 and likely contribute to the development of a systemic humoral response in infected mice. *E. tenella* antigen has been shown to induce IFNγ release in spleen cells of from immunized chickens (Prowse & Pallister, 1989). Similarly, Byrnes et al., 1993 illustrated the abilities of splenic macrophages to produce IL1 and TNFα during the primary infection of *E. tenella* and *E. maxima*. The expression of chicken Toll-Like receptors (TLR3,TLR15), signal adaptor (MyD88)*(Zhou et al., 2014)* and IFNy (Rothwell et al., 2000) has been detected in the spleen of chickens as a response to infection with *E. tenella*.

In the 60 plus years since its isolation, the *E. flaciformis* Bayer Haberkorn isolate has become the most commonly used laboratory isolate of rodent *Eimeria*. In the present study, we compare infection of mice (NMRI) with the laboratory isolate *E. flaciformis* BayerHaberkorn (falL), wild derived islotates of *E. flaciformis* (falW) and *E. ferrisi* Levine and Evens, 1965 (ferW). We asses similarities and differences in proliferation of tissue stages, oocyst shedding, induced pathology and immune response between two different *Eimeria* species and use these differences as a background to compare the laboratory isolate of *E. falciformis* (BayerHaberkonrn) with a novel field isolate of the same species.

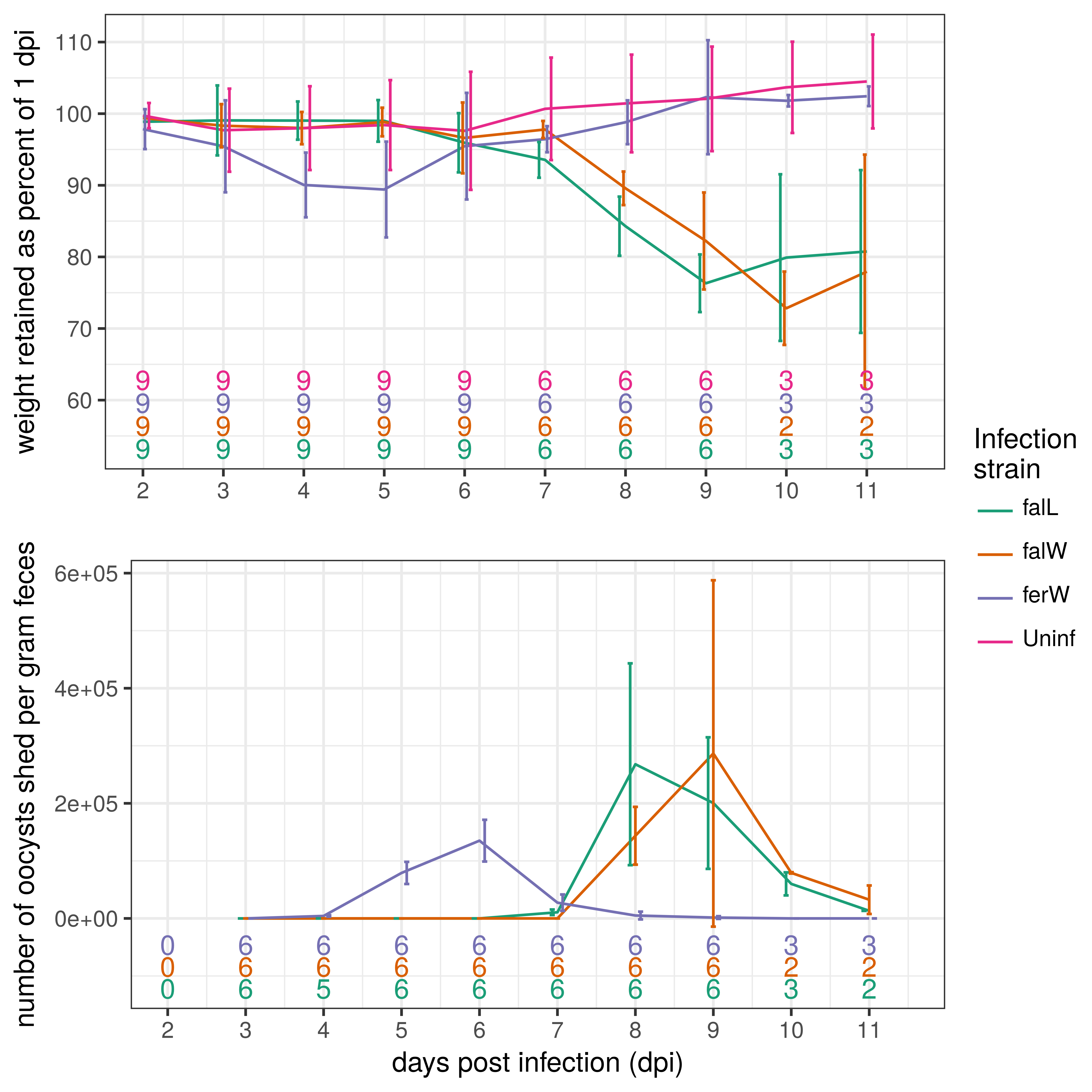
## RESULTS

### **Dynamics of infection and body weight loss differ between Eimeria species**

We infected mice with three different *Eimeria* isolates and followed the infection. A time course of the parasite reproduction assessed via oocyst shedding is shown in Figure 1a. We used the two different species *E. flaciformis* and *E. ferrisi* and show that these have different infection dynamics in NMRI mice: oocyst shedding of *E. ferrisi* has a peak intensity at 6 dpi, was drastically reduced on 7 dpi (n = 12, U = 2.91, p = 0.002) and below detection levels on 10dpi. Oocyst shedding of *E.falciformis* has a peak intensity at 8 dpi for the laboratory isolate BayerHaberkorn (falL) and 9 dpi for a novel, wild derived isolate (falW). The oocyst numbers declined after this peak in both isolates, but shedding was still detectable at 11 dpi when we ended the experiment and sacrificed all mice. For the two *E. falciformis* isolates we observed no difference in shedding intensity of oocyst at the peak day (n = 12, U = 0.24, p = 0.846) and also peak oocyst abundance did not differ significantly between *E. ferrisi* and both *E. flaciformis* isolates (ferW vs. falW, n = 12, U = 0.32, p= 0.777; ferW vs. falL, n = 12, U = 0.96, p= 0.37).

The time of patency (oocyst shedding) was characterized by body weight loss in infected mice in all three investigated infection groups (Figure 1b). Infections with *E. ferrisi* coincided with significant weight loss at 4 dpi (n = 18, U = -2.43, p = 0.013) and 5 dpi (n = 18, U = - 2.52, p = 0.010) in comparison to the control group. Infection with *E. falciformis* was accompanied by significant weight loss at 8 and 9 dpi in both falL (both dpi, n = 12, U = -2.89, p = 0.002) and falW (8dpi, n = 12, U = -2.41, p = 0.013; 9dpi, n = 12, U = -2.89, p = 0.002) isolates as compared to the control group. At later time points sample sizes for infected groups were likely too low for significant results on weight loss. Weight losses in infections with *E. ferrisi* at their maximum (at 5 dpi) were, however, significantly lower compared to weight loss in infections with *E. falciformis* at their maximum (9 dpi; ferW vs. falW, n = 15, U = -2.0, p = 0.049; ferW vs. falL, n = 15, U = -2.59, p = 0.007).

Oocyst shedding and weight loss show different relative timing in *E. falciformis* compared to *E. ferrisi*. In infections with both isolates of *E. falciformis* weight loss coincides with or follows one to two days after oocyst shedding, in infection with *E. ferrisi* weight loss precedes peak oocyst shedding by one day or more.



**Figure 1** – Dynamics of parasite reproductions and weight loss of hosts differ between the species *E. falciformis* and *E. ferrisi*. a) Oocyst shedding of *Eimeria* spp. from experimentally infected mice (NMRI) is displayed from 1 to 11 days post infection (dpi). Mice were infected with 200 sporulated oocyst of *E. ferrisi* (a wild derived isolate; ferW) and *E. flaciformis* (the laboratory isolate BayerHaberkorn; falL) or (a wild derived isolate; falW). b) Body weight loss of the same three groups of mice is depicted as percentage of body weight retained compared to 1 dpi. The number of mice (n) is given at the bottom of the plot, it is reduced at the end of the experiment, because mice were sacrificed for collection of tissue samples. Lines indicate the mean for each group, error bars give the standard deviation.

**Intensity of tissue stages of *Eimeria spp.***

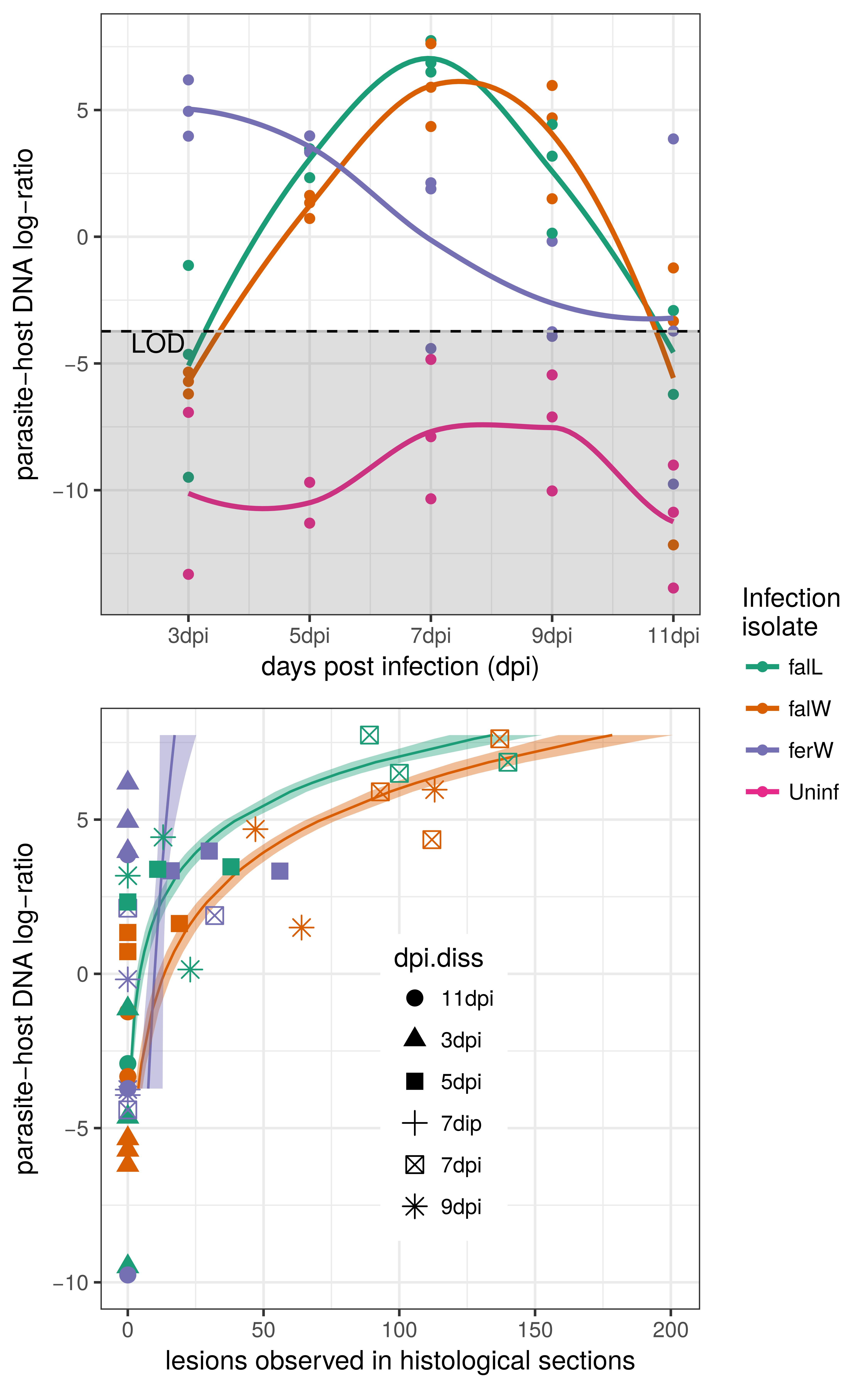
*Eimeria* infections result in a transient presence of parasite stages in epithelial cells of the caecum. The intensity of infection was quantified by quantitative PCR (qPCR) assay. We analysed the ratio of parasite DNA (determined by amplification of a fragment of the Cytochrome C-oxidase subunit I; COI) to host DNA (determined by amplification of a fragment of the nuclear *cdc42* gene). We report this ratio on a native (log2) scale of measurement and further call it the parasite-host DNA log-ratio (Figure 2a). The analysis of infected and control samples (*E. ferrisi* n =15; *E.falciformis*, falW n = 14, falL n = 14; uninfected controls n = 13) allowed us to estimate a limit of detection (LOD; mean + 2 standard deviations of the negative controls) for the assay at a parasite-host DNA log-ratio of -3.73. This corresponds to roughly eight *Eimeria* COI molecules for 100 copies of the mouse nuclear genome. The highest value measured for an individual negative control sample was a parasite-host DNA log-ratio of -4.84. Maximum values for parasite-host DNA log-ratio (observed in the falL isolate) were 7.74 indicating a ratio of 214 parasite COI mDNA copies for each copy of the mouse genome in crude tissue at this point.

Given the low sample size we abstain from a statistical analysis of individual sampling days but describe the observed pattern (Figure 2a). Concerning *E. falciformis,* for all falW and two out three falL samples the parasite-host DNA log-ratio was at 3 dpi was still below the limit of detection, while for *E. ferrisi* the highest value (at 6.19) was measured at this sampling day. For *E. falciformis* (both isolates) parasite-host DNA log-ratio increased to values well above zero (equal number of parasite mitochondrial and host nuclear DNA copies) on 5 dpi. For both isolates of *E. falciformis* the highest values were reached at 7 dpi. Again the amount of DNA measured was similar to that of *E. ferrisi* (at 3 dpi). Except for one *E. ferrisi* outlier-sample, for which a value of 3.86 was measured, at 11 dpi the parasite-mouse DNA log-ratio was reduced to values below zero for all samples and for most samples below the limit of detection.

To test how this parasite-host DNA log-ratio predicts visible tissue stages we counted lesions (thought to be directly caused by parasite tissue stages) in histological sections. In a generalized linear model (Table 1) the parasite-host DNA log-ratio is a significant predictor for lesions (Figure 2b). The effect of the parasite-host DNA log-ratio is similar for both isolates of *E. falciformis*, meaning that a similar number of lesions are found for similar parasite-host DNA log-ratio in this species. For *E. ferrisi,* however, a significant interaction effect indicates that a lower number of lesions is found for similar DNA concentrations. Inspection of Figure 2b makes clear that for *E. ferrisi* early (3dpi) parasite presence did not coincide with lesions.

**Table 1** – A generalized linear model predicts tissue lesions with the amount of parasite DNA relative to host DNA (parasite-host DNA log-ratio).

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  |  | Lesion score | | |
|  |  | *Prediction* | *CI* | *p* |
| (Intercept) |  | 4.43 | 3.18 – 6.03 | <.001 |
| PH.delta |  | 1.55 | 1.48 – 1.64 | <.001 |
| Infection isolate | | | | |
| *Inf. falW* |  | 3.03 | 2.09 – 4.46 | <.001 |
| *Inf. ferW* |  | 2.23 | 1.45 – 3.42 | <.001 |
| PH.delta : Inf. falW |  | 0.90 | 0.84 – 0.95 | <.001 |
| PH.delta : Inf. ferW |  | 0.69 | 0.63 – 0.76 | <.001 |
| Observations |  | 33 | | |



**Figure 2 –** The relative amount of parasite vs. host DNA (parasite-host DNA log-ratio) estimates the intensity of parasite tissue stages. a) The parasite-host DNA log-ratio was calculated from the difference in threshold (cq) values of qPCRs performed on a single copy nuclear gene (CDC42) of the host and on a mitochondrial (COI) gene of the parasite. Lines are drawn using local polynomial regression fitting (a “loess smoother”). b) Predictions of tissue lesions from these qPCR data are shown for all parasite isolates at different times after infections. Lines here represent predictions from a generalized linear model, shaded areas 95% confidence intervals of these predictions. For *E. falciformis* (falW and falL) this model provides better fit than for *E. ferrisi* (ferW).

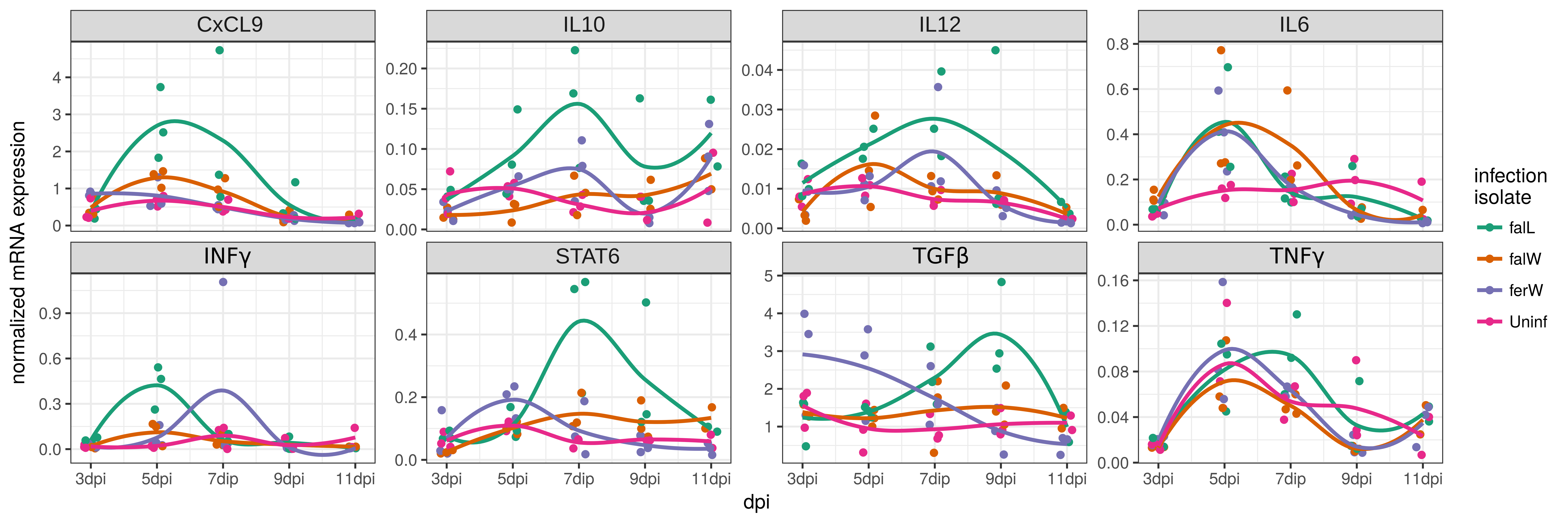
**Differences in immune gene expression between the laboratory and wild derived isolates**

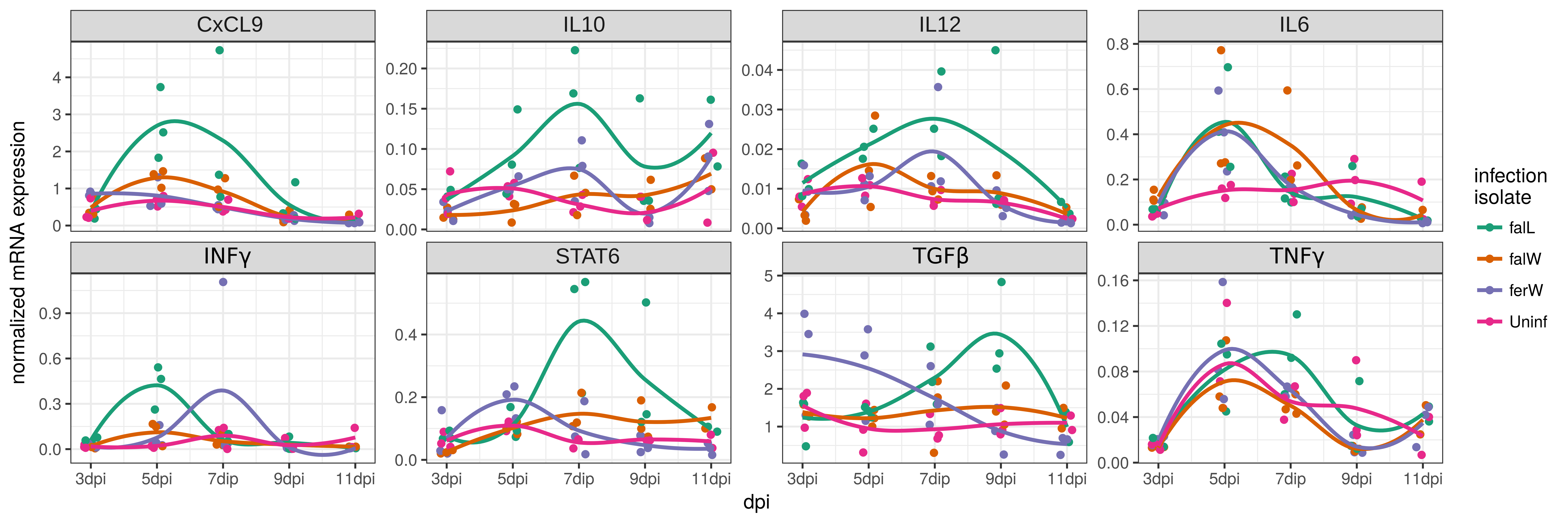
To characterized the immune response of NMRI mice against the *Eimeria* isolates we performed reverse transcription (RT) qPCR assays on relevant cytokines. Expression levels for most genes differed significantly between uninfected controls and mice infected with the *E. falciformis* laboratory (falL**)** isolate (Figure 3). We used linear mixed effect models with the dpi as random effect to “pool” information over multiple dpi, increasing sample sizes for comparisons. Mice infected with falL had significantly higher expression levels of chemokine 9 (CxCL9), interleukines 10 and 12 (IL10 and IL12), tumour growth factor β (TGFβ), and signal transducer and activator of transcription 6 (STAT6). No significant difference in the expression levels compared to control mice was observed for interleukine 6 (IL6), interferon γ (IFNγ) and tumour necrosis factor α (TNFα). In contrast, expression levels did not differ significantly between uninfected controls and mice infected with wild derived stains of both *E. falciformis* (falW) and *E. ferrisi* (ferW) for any of the examined genes. Immune gene expression in infections with the laboratory (falL) isolate showed to be significantly higher than in infections with wild parasite isolates (ferW and falW) for all genes significantly different between falL and uninfected controls (Table 2).

Some genes showed differences in gene expression profiles over the course of infection, in some cases including differences between infections with different parasite isolates (Figure 3). We did not analyse differences on individual days statistically due to the low sample sizes, but give a description of our observations. IL6 showed elevated levels of expression for all infection groups compared to controls as early at 5 dpi. Expression levels for IFNγ seemed elevated only at 5 dpi and only in infections with the *E. falciformis* laboratory isolate (falL). Both cases of potential elevations in expression failed to be detected as significant over controls in our mixed effect models because it was very transient and diminished already at 7 dpi. The interleukins IL10 and IL12 as well as STAT6 and CxCL9 showed elevated expression levels at multiple days of infection for the falL compared to all other infection groups. *TGFβ* showed somewhat elevated expression levels early in infection with EferW (3 and 5 dpi) and late in infections with EfalL (7 and 9 dpi). For TNFα we observed elevated expression levels in all infected but also in uninfected control groups. Taken together these observations add detail on the individual cytokines and underline our general finding of differences between wild derived and laboratory isolates of *E. falciformis*.

In summary expression of genes relevant for immune responses did not differ significantly from uninfected controls during infection with wild derived isolates of both *E. falciformis* (falW) and *E. ferrisi* (falL). In contrast, most genes were expressed significantly higher in infections with the laboratory isolate of *E. falciformis* compared to uninfected controls but also to all other infections including those with the wild derived *E. falciformis* (falW) isolate*.*

**Table 2** – Generalized linear mixed effect models show gene expression differences between wild (falW) and laboratory (falL) isolates of *Eimeria falciformis.* This are submitted as separate files due to its size.

Figure 3 – Hosts infected with wild and laboratory isolate of *E. falciformis* show different patterns of gene expression in the spleen. Mice were sacrificed at different times post infection and mRNA expression was assessed using quantitative PCRs. Points indicate normalized expression values for individual mice. Lines are drawn using local polynomial regression fitting (a “loess smoother”).

**Inflammatory cell infiltration is stronger *E falciformis* laboratory (falL) compared to wild derived (falW) isolate.**

To link our observation of gene expression with independent measures of immune response and tissue damage, we performed a histological scoring of inflammatory cell infiltration (Table 3, Figure 4). At most no infiltration of leukocytes was observed in uninfected control mice except in few samples minimal infiltration was observed.

We again describe observation of individual sampling days without performing statistical tests: In the laboratory isolate of *E. falciformis* a high score of inflammation was already observed during pre-patency (at day 5 dpi), the area occupied by immune infiltration remained high until 9dpi and declined only towards 11 dpi. In contrast, in caeca of mice infected with thewild derived *E. falciformis* isolate only low inflammation was found in the pre-patent period. Infiltration consisted mainly of agranulocyte at this stage of infection. Inflammation then increased at day 7 onward, and during this period, granular leukocytes were seen. Infiltration was marked at 9dpi before decreasing slightly towards 11 dpi.

The quality of the observed infiltration was the same in both infections with wild derived and laboratory isolate of *E. falciformis*. Infiltration are consistently characterised by both granulated and granular leukocytes and the presence of neutrophils.

In infections with *E. ferrisi* milder inflammation was detected on 3dpi and 5dpi. The strongest infiltration was detected at 5 dpi followed by a subsequent decline towards 7 dpi. Making use of information of mice sacrificed at different different days of the experiment in mixed effect models overall inflammation was significantly lower in *E. ferresi* infected mice than in those infected with *E.falciformis* (glmm; falL *vs.* ferW p = 0.001; falW *vs*. ferW p = 0.014).

**Table 3 - Score for the severity of leukocyte infiltration in sections from the mid-part of the caecum from NMRI mice infected with Eimeria spp.**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| Treatment¹ | Severity of leukocyte infiltration in cecum1 | | | | |
| 3dpi | 5dpi | 7pdi | 9dpi | 11dpi |
| ***E. ferrisi* (ferW)** | 2, 1, 1 | 1, 2, 2 | 1, 2, 1 | 1, 1, 1 | 0, 0, 0 |
| ***E.falciformis* (falW)** | 0, 1, 1 | 1, 0, 1 | 2, 2, 2 | 3, 2, 3 | 2, 2 |
| ***E.falciformis* (falL)** | 1, 2, 1 | 3, 3, 3 | 3, 3, 2 | 3, 3, 2 | 1, 1 |

1Leukocyte infiltration was scored on a 0 to 3 scale, where 0 represent no infiltration and 1, 2, 3 represented low, moderate, or high infiltration, respectively. One section from each of the cecum was used for scoring. 3 fields were observed for each section, and infiltration was characterized as a mixture mononuclear and granulocytic cells. A numerical score was assigned to each section. For each infection group and, values are reported for three mice.

**Discussion**

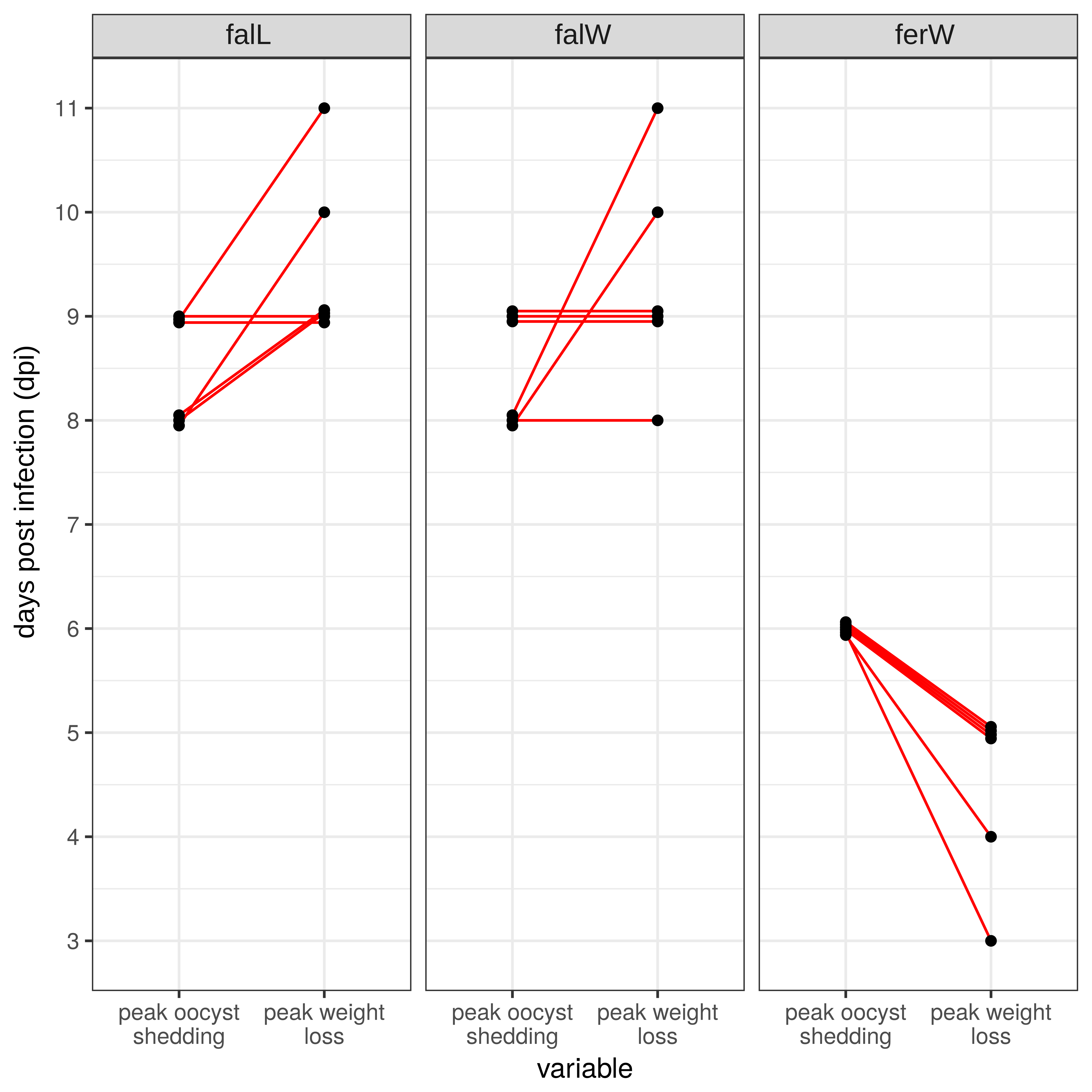
We evaluated in the present study how wild derived isolates of *Eimeria* spp. differ in infection dynamics, in the immune reactions and in the pathology they induce from those of *Eimeria falciformis* BayerHaberkorn (Haberkorn, 1970),a classical laboratory isolate. We found differences in parasite reproduction dynamics between *Eimeria* species (*E. falciformis* vs. *E. ferrisi*). While such differences between species are expected, we also found differences between wild derived isolates of *E. falciformis* and the laboratory isolate of *E. falciformis.* The laboratory isolate of *E. falciformis* induces be stronger immune reactions and pathology in NMRI mice by than wild derived isolates of both *E. ferrisi* and *E. falciformis*.

Between the two *E. falciformis* isolates we observed only slight differences in the length of the pre-patent period (time until oocyst shedding, starting at 6 *versus* 7 dpi). This is agreement with previous reports from the same host (NMRI mice) and parasite isolate (Ehret et al., 2017; Schmid et al., 2014, 2012; Stange et al., 2012). The pre-patent period for the wild derived *E. falciformis* isolate (7 dpi) corresponds to that reported for the parasite isolate *E. falciformis* var praghensis (Kasai et al., 1991; Mesfin et al., 1978). Finally, Mahrt and Shi (1988) and Schito et al. (1996) reported slightly longer pre-patent periods for *E. falciformis* infections (7 or 8 dpi) and while the first study seems to have used the BayerHaberkorn isolate it is unclear for the latter. We note that isolate identity needs to be more carefully reported in future research and that a slight difference in pre-patent time between two isolates of *E. falciformis* in our experiment might recapitulate differences observed between *E. falciformis* isolates by other authors before. Similarly, the output of oocysts in our study (for all isolates) was comparable or slightly lower to that in previous reports (Ehret et al., 2017; Schmid et al., 2014).

Our observation regarding the infection dynamics of *E. ferrisi*, agree with the initial description of the life cycle in *Mus musculus* (Ankrom et al., 1975). Our present work confirms that *E. ferrisi* is characterized by a – especially compared to *E. falciformis* – short life cycle with patency at 3 dpi. It is also noteworthy that the oocyst output of this species differed not significantly when compared to that of both *E. falciformis* isolates (Figure 2), although this might be a result of the low sample size especially at late times of infection (when *E. falciformis* was shedding most oocysts but most mice had already been sacrificed).

Pathogenicity assessed by maximal weight loss during infections with *E. falciformis* was very strong in the present study compared to previous experiments (Ehret et al., 2017; Schmid et al., 2012; Stange et al., 2012). A potential reason for this is an elevated pathology induced by a variation in the inoculum dose. As observed by Haberkorn (1970) higher dosed inocula lead to enhanced pathology, while infection dynamics (the extent and timing of oocyst shedding) are not impacted. In *E. falciformis* infections maximal weight loss was observed at 9 dpi, while infections with *E. ferrisi* induced a significantly lower maximal weight loss at 5 dpi. This observations are likely due to fewer cycles of asexual merogony of the parasite leading to a lower burden of intestinal stages. Interestingly, lines of poultry *Eimeria* with an abbreviated and early life cycle (so called “precocious lines”) show low oocyst output and are less pathogenic to their host (Shirley and Harvey, 2000; Shirley and Long, 1990). While *E. ferrisi* attains substantial oocyst output, it shares this low pathology with precocious lines. This suggests that short phases of asexual expansion might be correlated with low pathogenicity in *Eimeria* infections independent of total oocyst shedding. In other words, parasite fitness for “naturally precocious” species of *Eimeria* might be high, while simultaneously host fitness is relatively little reduced by infections.

Infection with *Eimeria* causes damage in the intestinal mucosa resulting in malabsorption of nutrients and weight loss (Chapman et al., 2013; Haberkorn, 1970). Interestingly, comparisons of weight loss between the two isolates of *E. falciformis* and that of *E. ferrisi* demonstrate that *E. ferrisi* induces most weight loss before the peak of its oocyst shedding, while both *E. falciformis* isolates impact the host after the peak of their oocyst shedding (Figure 5).

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**Figure 5** – Peak weight loss precedes peak oocyst shedding in infections with *E. ferrisi*, while in infections with *E. falciformis* host lost most weight either on the day or after the days parasites shed most oocysts. Points depict the peak day of both oocyst shedding and weight loss respectively and red lines connect both measurement from the same mouse. For raw data and underlying experimental procedures see Figure 1.

These differences suggest that mechanisms underlying pathogenesis might be different between the two parasite species. Weight loss in infections with two isolates of *E. falciformis* occurs simultaneously with and reaches its maximum at or after sexual reproduction of the parasite. Sexual reproduction of *E. falciformis* might cause an exhaust of epithelial cell which burst when oocysts are released into the lumen (Kasai et al., 1991). Additionally, histology indicates that weight loss coincides with immune cell influx in *E. falciformi*s infections. This influx differed slightly in timing starting at 5 dpi in the *E. falciformis* laboratory isolate and 7dpi in the field isolate. Influx of immune cells into the tissue might be associated with immuno-pathology (Baskin et al., 2009; Brant et al., 2014; Stange et al., 2012). Maximal intensity of tissue stages (assessed using a qPCR assay, we established in the present study and counts of histological lesions) precedes for the two *E.falciformis* the release of oocysts from tissues into the intestine and maximal weight loss. For infection with *E. ferrisi*, in contrast, weight loss coincides with the peak abundance of endogenous stages at 3 dpi (Figure 2) it can thus be speculated whether it is simply the extraction of energy of *E. ferrisi* for its own growth causing pathology in this system. Intensity of tissue stages also coincides with immune cell activation at the site of infection and weight loss, suggesting that parasite proliferation cause pathology in host infected with this species.

Our histological analysis link weight loss pathology to tissue damage and can also validate our qPCR approach and measurements of parasite-host DNA log-ratio derived from it. We observed more lesions and stronger immune cell infiltration in infections with the laboratory isolate of *E. falciformis* than in the wild derived isolate. The number of lesions over different dpi correlated well with the parasite-host log-ratio for these isolates. Infections with *E. ferrisi* were characterised by yet lower immune cell infiltration and tissue damage. In infections with this species the parasite-host DNA log-ratio seems to be more sensitive than the histological counting of lesions, which can hardly be observed.

Cellular infiltrations observed within the mucosa during experimental infections of *Eimeria* in many host species including mice have been described by several authors (Gadde et al., 2009; Laurent et al., 2001; Mesfin et al., 1978; Muñoz-Caro et al., 2016; Rose et al., 1992; Schmid et al., 2014). Tissue lesions are considered to be mostly caused by parasites directly, but inflammatory reactions contribute to the process (Muñoz-Caro et al., 2016). In this context, the relatively modest pathology (weight loss) observed during *E. ferrisi* infection might be a cause or consequence of milder immune cell infiltration observed in the infected tissues of this species.

Schmid et al., (2014) demonstrated by immunohistochemical analyses that *E. falciformis* infection in the cecum of NMRI mice leads to tissue infiltration with lymphocytes and macrophages. These changes are accompanied by elevation expression of INFγ and the production of the major chemokines CxC subfamily at the site of infection. Several authors before had described the role of INFγ and these chemokines to leukocyte attraction and its likely involvement in controlling the growth of *Eimeria* in mouse models (Lillehoj, 1998; Pogonka et al., 2010; Rose, 1974) and in *Eimeria* infected chickens (Laurent et al., 2001; Lowenthal et al., 1997; Yun et al., 2000). Inflammatory infiltrates were also slightly more prominent in our experiment in the laboratory isolate of *E. falciformis* than in closely related wild derived *E. falciformis* isolate and we thus asked whether systemic immune response differs between infections.

To investigate a more systemic immune response during infection we measured the expression of pro- and anti-inflammatory cytokines and regulatory chemokines in the spleen. Eight markers for different immune response pathways were investigated: CxCL9 has broad immune-regulatory role, INFγ and TNFα represent the innate pro-inflammatory response (Mosmann et al., 1986) IL12is a marker for Th1-type response against intracellular parasites (Ref), STAT6 for a Th2-type response (Ref). IL6 is a marker for the Th17pathway in the mucosal barrier tissues (Ref), and finally, IL10 and TGFβ have anti-inflammatory roles (reviewed in Terner *et al*, 2014, Stenger and Röllingho 2001).

Only the laboratory isolate of *E. falciformis* significantly influences the mRNA expression levels of most of these markers in the spleen. The wild derived isolates of *E. falciformis* and *E. ferrisi* do not induce expression changes over control levels.

IL6 is the only counter-example for this, we observed it to be elevated (although not significantly) in all infections over control levels early in infection. IL6 synthesized in the initial stage of inflammation at a local lesion, then it moves through the bloodstream to the liver and spleen (Hienrich et al, 1990). There, it exerts promotes specific differentiation of naïve CD4+T cells, linking innate to acquired immune response (Tanaka et al, 2014). Beyond its immune-regulatory function IL6 has a role in stimulating the intestinal epithelial proliferation and repair after injury and an elevated level might thus be a consequence of parasite tissue damage (Kuhn et al., 2014). In experimental infections of mice with *E. falciformis* marked induction of IL6 transcription between 5 and 7 dpi has been reported at the site of infection (Ehret et al., 2017). In the present study IL6 is the only cytokine showing potentially enhanced expression also in the spleen. This might suggest that the immune modulatory role of IL6 in the spleen during *Eimeria* infections could be augmented by elevated mRNA expression within this organ.

We observed elevated mRNA levels of pro-inflammatory Th1 cytokines IFNγ, TNFα and IL12 in the spleens of mice infected with the laboratory isolate of *E.falciformis*. Our data from the spleen mirrors previously published data indicating and induction of these genes at the site of infection between 5 and 7 dpi (Ehret et al., 2017; Schmid et al., 2014). The IL12/IFNγ axis is crucial for the activation cellular immune responses against intracellular parasites including *Eimeria* (Cacho et al., 2012; Chow et al., 2011; Ehigiator et al., 2007; Heinzel et al., 1991; Kulkarni et al., 2011; Lillehoj, 1998; Michailowsky et al., 2001; Ovington and Smith, 1992; Rose et al., 1992).

An increase in the expression of TNFα of was detected in all experimental groups, including the non-infected control group, around 5 dpi. This might be due to a stress response during the course of experiment. TNFα cellular responses involved in systemic inflammation that stimulate the acute phase reaction (Turner et al, 2014) can eradicate infectious agents, but can also lead to harmful systemic effects (Waters et al, 2013).

We observed significantly increased expression of the anti-inflammatory Th1 cytokines IL10 and TGFβ in infections with the laboratory isolate of E*. falciformis*. IL10 can counteract IFNγ and is expressed also in the spleen of chickens infected with *Eimeria* (Rothwell et al., 2000). IL10 expression in the spleen could be indicative for an attempt to balance inflammation during *E.flaciformis* laboratory isolate infection. A failure to established this inflamatory balance can lead to pronounced inflammation (Inagaki-Ohara et al., 2006). TGFβ showed elevated expression levels early in infection with *E. ferrisi* (3 and 5 dpi) and late in infections with the *E.falciformis* laboratory isolate (7 and 9 dpi). Considering an elevated level of IL6 expression, these finding could indicate the involvement of a Th17 pathway to control the infection events. TGFβ and IL6 play non redundant roles in the generation of IL17 from naïve CD4+T cells of mouse (Sehrawat and Rouse, 2017)(Korn et al, 2009. IL17 in turn contributes to both immunopathology and parasite restriction during infection with E.falciformis (Stange, 2013).

Induction of STAT6 in the spleen has been reported in several infections with intestinal parasites (Lee et al., 2013) Lopez et al, 2013). In addition to elevated expression of STAT6 we noticed significantly elevated expression of the major regulatory chemokines CxCL9 in infections with the laboratory isolate of *E. falciformis*. CxCL9 can be induced downstream of INFγ (Djamiatun et al., 2017; Hirako et al., 2016; Schmid et al., 2014) and is involved in recruitment and activation of effector T lymphocytes in the spleen as well as non lymphoid organs such as intestine in disease models including *E.falciformis* (Hardison et al., 2006; Khan et al., 2001; Schmid et al., 2014).

The apparent differences in immune response against the wild derived and the laboratory isolate of *E. falciformis* invite speculations about their origin. Such speculation should be qualified with the notion that we can not be sure what infection phenotype (pathology) the original *E. falciformis* BayerHaberkorn isolate had when it was derived. It is plausible, however, that the pathology before serial passaging resembled that observed in our the isolate we derived newly. In the laboratory the consequences of serial passage can be understood in terms of a selection experiment (Ebert, 1998). In *Eimeria* artificial selection has been used to e.g. create attenuated “precoccious” stains, which undergo a faster development, are less pathogenic but still induce protective immunity against reinfections (McDonald and Ballingall, 1983; Shirley and Bellatti, 1988).

Independent of the ultimate reasons for the difference in immunogenicity the conclusion can be drawn that the infections with the laboratory isolate *E. falciformis* might not be representative for parasite-host interaction in their ecological and evolutionary context. In addition to the description of infection dynamics, induced immune reactions and histopathology for a wild derived isolate of *E. falciformis* in comparison to the BayerHaberkorn we show that *E. ferrisi* possesses a short developmental cycle and low pathology. This update on the life cycle (Ankrom et al., 1975)with details on histopathology and immune reactions will increase the attractivity of this species as rodent infection model for *Eimeria*.

**3. Material and Methods**

**3.1. Wild isolate of *E. flaciformis* (falW) and *E. ferrisi* (ferW)**

The pure inocula of *E. flaciformis* (falW) and *E. ferrisi* (ferW) wild derived isolates were produced in our lab through NMRI infection experiment. Briefly, sporulated oocysts of *Eimeria* were recovered from samples obtained after field collection from house mouse hybrid zone to the north of Berlin in 2016, from individual faeces sample in which each genotype predominated 300 oocysts for E64 and 600 oocysts for Efwild were inoculated into 16 weeks old -NMRI female. All mice were reared individually in wire cages in isolation rooms and provided with food and water *ad libidum*. The faeces from those mice were collected daily during the period of oocyst release from 1 to 12 days post-inoculation [pi]. Oocysts in faeces were harvested by screening, sedimentation- flotation in saturated NaCl salt solution, and washings. They were then placed in 2% potassium dichromate and incubated at 25C for 4 days to permit oocyst sporulation. Sporulated oocyst were examined repeatedly under light microscopy to ensure its purity, and were then stored at 4 °C for about 1 months prior to use.

**3.2. Infection protocol, oocyst counting and sample collection:**

The cleaned inocula of the wild derived *E. flaciformis* (falW) and *E. ferrisi* (ferW) isolates produced from the previously described experiment. The inoculum of *E. flaciformis* was originally isolated in Wupperthal by Haberkorn working at Bayer Animal Health company in 1960 and described in 1970, it was since then propagated through experimental passaging in NMRI mice every 3 months.

15 female NMRI mice (10 to 12 weeks old) were randomly assigned to four groups, including a group maintained as study controls that were not inoculated. The remaining 45 mice were inoculated via oral gavage with 0.1 ml of inoculum containing a single dose of 200 sporulated oocyst. The inoculum had been prepared counting the total number of oocysts in 10µl directly on a standard microscope slide.

Complete faeces were collected every day of the experiment. After weighting of the faeces, flotation was performed as follows: Saturated salt (NaCl) solution was added, the mixture was stirred and

centrifuged at 3175g. It was washed twice with physiological salt solution and after the last washing 2ml of 2% potassium dichromate solution were added to the pellet and 10µl of the solution were loaded into a “Neubauer-improved chamber”. Oocyst were counted in eight grid squares. Then the number of oocysts per gram faeces was obtained according to the (0.1µl) volume of a grid square: *Concentration (oocyst/g) = total #of oocyst / #squares counted \** *10.000 ml-1 \* 2ml / g (faeces)*

The weight of mice was recorded every day for each mouse during the 11 days of the experiment. Three mice were sacrificed from each group on 3, 5, 7, 9, and 11 dpi. Immediately after death the viscera were exposed, the spleen and caeca removed. Caecal contents were gently washed away with physiological salt solution and the tissue was cut longitudinally into two pieces. One piece was transferred into a 30 ml polypropylene tube containing 20 µl of RNAlater® (Life Technologies; Carlsbad, CA, USA). Samples were stored for 4h at 4°C before transfer to and storage at−20 °C until use. The second piece of caecum tissue was fixed in 4% formalin and stored at room temperature until used for histological examination.

**3.3. Quantification of *Eimeria* load in infected mouse cacum tissue**

For DNA extraction frozen tissue was manually homogenized by addition of liquid N2 and grinding. Genomic DNA was immediately extracted using innu PREP DNA Mini Kit® (Analytika jena) according to the standard manufacturer protocol, incorporating proteinase K digestion. Purified DNA stored at -20 until subjected to qPCR for host and parasite DNA quantification.

Primers used in this study to amplify the mitochondrial COI region of *Eimeria* spp. are Eim-COI-forward 5’TGTCTATTCACTTGGGCTATTGT3’ and Eim-COI-reverse5’GGATCACCGTTAAATGAGGCA 3’. For host genomic DNA amplification we used a primer targeting the *Mus*-*cdc42* gene with the sequence: *Mus*-*cdc42-*forward 5’CTCTCCTCCCCTCTGTCTTG3’ and *Mus*-*cdc42-*reverse 5’TCCTTTTGGGTTGAGTTTCC3’.

60 DNA samples were added for qPCR to Multiplate™ 96-Well PCR plates (BioRad), with reactions performed in duplicate for each sample. Each plate also contained a non template control and a plate control sample (ddH2O). All pipetting steps took place in a clean and sterile flow box, designated for working with DNA. The qPCR mixture of 10 μL was prepared using the iQ™ SYBR® Green PCR Kit (Bio-Rad): 5 μL of 2X iQ™ SYBR® Green Master Mix, 0.3 μL of 20 μM forward and reverse primers, and 4 μL of 10ng/μL template DNA. The thermal cycling protocol was set as follows: initial denaturation for 15 minutes at 95°C, followed by 40 cycles of 15 seconds at 95°C, 30 seconds annealing at 60°C for Eim-COI primer or 58°C for *Mus*-*cdc42,* and 30 seconds at 68°C and measuring the fluorescence signal at the end of every step.

qPCR amplifications and analysis were performed using Bio-Rad CFX96, Thermalcycler1000 system, which determined the cycle of quantification (Cq). To assess the validity of our protocol we examined assay specificity, efficiency and repeatability on three separate RT-PCR amplifications. To confirm the specificity of the assay a melting curve was generated during RT-PCR by adding a stepwise temperature increase from of 65.0°C to 95.0°C, with 0.5°C increment after amplification. After calculating mean Cp between technical replicates, the abundance of *Eimeria* relative to host DNA was estimated as the ∆Cq between mouse and parasite DNA. As a log of a ratio is equivalent to subtractions between log values, this represents a log(2)-ratio between mouse host (*Mus*-*cdc42*) and *Eimeria* parasite (Eim-COI) DNA copies.

**3.4. RNA extractions and reverse transcription**

Before RNA isolation, frozen spleen tissue was homogenized by grinding after addition of liquid nitrogen. Total RNA was isolated using the PureLink™ RNA Mini Kit ([Thermo Fisher Scientific](https://www.google.de/url?sa=t&rct=j&q=&esrc=s&source=web&cd=1&cad=rja&uact=8&ved=0ahUKEwi__puD3IfYAhXGKlAKHYjaDEAQFgg4MAA&url=https%3A%2F%2Fwww.thermofisher.com%2Forder%2Fcatalog%2Fproduct%2F12183018A&usg=AOvVaw0FVD2VXdCZI8KVSaguQkxK)). Briefly, frozen homogenized sample was transferred with a sterile scalpel blade into tubes with 1ml lysis solution with 1% 2-Mercaptoethanol and 1.4 mm zirconium oxide beads (Peqlab GmbH, Erlangen, Germany). Subsequently samples were homogenized at room temperature (RT) using a Precellys® 24 tissue homogenizer twice at 6,000 rpm for 20 sec interrupted by a 30 sec cooling break. All further steps took place in a clean and sterile flow box, designated for RNA extraction only to eliminate most of the aerosol which developed during the shaking. All samples were centrifuged for 1 min at maximum speed (13,400 rpm) (Eppendorf) to eliminate the foam. The pure supernatant was collected and mixed with 1:1 ratio of 70% EthO. Afterwards, 600 µl of the previous mixture was added onto the Spin Filter in a 2.0 ml tube and centrifuged at 13,400 rpm for 30 sec. The filter binds all double stranded genomic DNA (gDNA), whereas single stranded RNA remains in solution. To get red of gDNA, an on-column DNA digestion was accomplished by PureLink DNase Set ([Thermo Fisher Scientific](https://www.google.de/url?sa=t&rct=j&q=&esrc=s&source=web&cd=1&cad=rja&uact=8&ved=0ahUKEwi__puD3IfYAhXGKlAKHYjaDEAQFgg4MAA&url=https%3A%2F%2Fwww.thermofisher.com%2Forder%2Fcatalog%2Fproduct%2F12183018A&usg=AOvVaw0FVD2VXdCZI8KVSaguQkxK)) according to the manufacturer protocol. To purify the RNA a washing solutions were added to the samples on the column and were centrifuged at 13,400 rpm for 30 sec.

After RNA isolation, the synthesis of complementary DNA (cDNA) was performed using the RevertAid H Minus First Strand cDNA Synthesis Kit (Thermo Fisher Scientific, Braunschweig, Germany). The Kit included the genetically engineered RevertAid™ H Minus M-MuLV Reverse Transcriptase (200 U/µl) which lacks the ribonuclease H activity to prevent RNA digestion. Therefore, degradation of RNA in RNA-DNA hybrids during synthesis of the first strand cDNA did not occur.

Nuclease-free H2O was added to 1µg template RNA to a total volume of 22 µl and 10 µl 0.1 pg/µl. 2 µl Oligo (dT)18 primer (100 µM, 0.5 µg/µl) were added to synthesize only RNAs with 3’-poly(A) tails (as those are mostly mRNAs). To denature potential secondary structures, the mixture was heated to 65°C for 5 min using the 2720 Thermal Cycler (Applied Biosystems) and rapidly cooled on ice afterwards to prevent renaturation. Subsequently, the reverse transcriptase mix was added.

The reaction was carried out by incubation for 60 min at 42°C followed by heating (termination) at 70°C for 10 min. two separated cDNA synthesis reaction was carried out for each individual sample. Thereby resulting first strand cDNA was then pooled from which aliquots was then drawn for subsequent gene expression study.

**3.5. Gene expression quantification**

To make an overall measure of the function of the immune response during the infection, we measured the expression levels of eight genes of interest, along with three reference genes. Genes of interest were CxCL9, IL10, IL12, TGF-β, STAT6, IL6, INFγ and TNFα. Primers for these regions were

2- Supplementary material for gene expression:

|  |  |  |  |
| --- | --- | --- | --- |
| Gene | Primer Sequence (5´- 3´) | Amplicon size(nt) | Source |
| *Mm*CDC42-F  *Mm*CDC42-R | CTCTCCTCCCCTCTGTCTTG  TCCTTTTGGGTTGAGTTTCC | 96 | This study |
| *Mm*Ppia-F  *Mm*Ppia-R | ACCGTGTTCTTCGACATCAC  ATGGCGTGTAAAGTCACCAC | 198 | This study |
| *Mm*Ppib-F  *Mm*Ppib-R | CAAAGACACCAATGGCTCAC  TGACATCCTTCAGTGGCTTG | 161 | Ehret et al. 2017 |
| *Mm*IFNg-F  *Mm*IFNg-R | ACAGCAAGGCGAAAAAGGATG  TGGTGGACCACTCGGATGA | 106 | Primer Bank  ID 145966741c2 |
| *Mm*IL6-F  *Mm*IL6-R | TAGTCCTTCCTACCCCAATTTCC  TTGGTCCTTAGCCACTCCTTC | 88 | Primer Bank  ID 13624311a1 |
| *Mm*IL10-F  *Mm*IL10-R | CCCATTCCTCGTCACGATCTC  TCAGACTGGTTTGGGATAGGTTT | 110 | Primer Bank  ID 6680389a1 |
| *Mm*IL12-F  *Mm*IL12-R | ATGGCTGCTGCGTTGAGAA  AGCACTCATAGTCTGTCTTGGA | 108 | Primer Bank  ID 6680399a1 |
| *Mm*TGFb-F  *Mm*TGFb-R | TACGTCAGACATTCGGGAAGCAGT  AAAGACAGCCACTCAGGCGTATCA | 186 | This study |
| *Mm*CXCL9  *Mm*CXCL9 | GGAGTTCGAGGAACCCTAGTG  GGGATTTGTAGTGGATCGTGC | 82 | Primer Bank  ID 162287427c1 |
| *Mm*STAT6  *Mm*STAT*6* | CTCTGTGGGGCCTAATTTCCA  CATCTGAACCGACCAGGAACT | 135 | Primer Bank  ID 6678155a1 |
| *Mm*TNFa  *Mm*TNFa | CATCTTCTCAAAATTCGAGTGACAA  CCTCCACTTGGTGGTTTGCT | 63 | This study |

A total of 60 cDNA samples were split between PCR plates (Multiplate™ 96-Well, BioRad) with reactions performed in duplicate for each sample. If the standard deviation of Cq values between duplicates was > 0.4, corresponding samples were repeated. Each plate contained a non template control sample and negative controls.

The qPCR mixture of 10 μL was prepared using the iQ™ SYBR® Green PCR Kit (Bio-Rad): 5 μL of 2X iQ™ SYBR® Green Master Mix, 3 μL of 10 μM forward and reverse primers, and 4 μL of 10ng/μL template cDNA. qPCR amplifications were performed using Bio-Rad CFX96, Thermalcycler1000 system as follows: initial denaturation for 15 minutes at 95°C, followed by 40 cycles of 15 seconds at 95°C, 30 seconds at 60°C and 30 seconds at 68°C with a measuring of the fluorescence signal at the end of every step. The cycle of quantification (Cq) was determined by the amplification plot in CFX96-Bio-Rad software. Finally, a melting curve was generated to confirm the specificity of the reaction by adding a cycle of 65.0°C to 95.0°C in 0.5°C increments.

Accurate normalization of gene expression is essential for the production of reliable data in RT-PCR experiments, with the optimal reference genes being specific to a particular set of experimental conditions. To select the most appropriate reference genes for normalization, we performed a geNorm analysis and tested four candidate genes among frequently described genes in the literature (*gabdh*6, *cdc42*, *ppia*, *ppip*). A total of four primer pairs for these four genes were evaluated using cDNAs synthesized from 16 tissue samples randomly selected. Analysis of the gene stability measure was performed using qbase+ (Biogazelle) in Bio-Rad CFX96 Thermalcycler1000 system, which identified *cdc42*, *ppia* and *ppip* as the most stable reference genes for this study. For all three reference genes, normalization factors (NF) were calculated using the geometric mean of the corresponding expression values (See Vandesompele et al. 2002).

Relative expression values for each tested sample of each gene of interest were then calculated using the ∆Cq method, adjusted for the amplification efficiencies of each primer pair and standardized against the normalization factors (NF) of each sample.

### **3.6. Histological examination and scoring**

Formalin fixed samples from the mid-part of the caeca were washed by water and dehydrated by Ethanol gradient concentration series. Thenembedded in paraffin and sections (one section from each part), with 3 μm thickness, and stained with hematoxylin and eosin. Stained tissues were examined at 20×, 40× and 100× magnification with a bright field microscope. An indication of the cellular infiltration in response to infection was obtained by subjectively assessing the extent and nature of leukocyte infiltration into the mucosa and sub mucosa of the entire section. A numerical score was assigned where 0, 1, 2, and 3 represented no leukocyte infiltration and mild, moderate, or severe infiltration, respectively. The nature of the leukocyte infiltrate was characterized as mononuclear, granulocytic, or a mixture of granulocytes and mononuclear cells based on the distinct morphological characteristics of these cell types.

A lesion score for the investigated *Eimeria* species was based on detection of characteristic developmental stages of this parasite (Goodwin, 1996). We used Cell® image analysis application in order to collect high resolution images of caecum sections and count the parasite at any stages in 6 fields of view.

**3.7. Statistical analyses and visualisation**

All statistical analyses and visualisations were performed in R (R Development Core Team, 2008). An “exact” version of the Mann-Whitney U-tests available in the package “coin” was used to account for ties in all comparisons of weight loss or oocyst shedding. A generalized linear model of the poisson family (log-link) was used to predict histological lesions with the parasite-host DNA log-ratio and parasite isolate used for infection allowing for different intercept and slopes for each isolate.

Linear mixed effect models (function “lmer” of the package lme4) were used to test for differences in gene expression. For each gene (as response variable) these models used the infecting *Eimeria* isolate as only fixed effect and the time of infection (dpi) as a random intercept. Similarly, linear mixed effect models for leukocyte infiltration scores (as response variable) were used with infection isolate as fixed effect and dpi as a random intercept. For visualisations the package ggplot2 was used, including the default “loess” smoother as indicated in figure legends.

**Ethics statement**

All Animal procedures in this investigation were performed according to the German Animal Protection Laws as directed and approved by the overseeing authority Landesamt für Gesundheit und Soziales (Berlin, Germany) under permit number H0098/04.

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