

Reid WDK, Close AJ, Humphrey S, Chaloner G, Lacharme-Lora L, Rothwell L,
Kaiser P, Williams NJ, Humphrey TJ, Wigley P, Rushton SP.

[Cytokine responses in birds challenged with the human food-borne pathogen](#)

[*Campylobacter jejuni* implies a Th17 response.](#)

Royal Society Open Science 2016, 3, 150541.

Copyright:

© 2016 The Authors. Published by the Royal Society under the terms of the Creative Commons Attribution License <http://creativecommons.org/licenses/by/4.0/>, which permits unrestricted use, provided the original author and source are credited.

DOI link to article:

<http://dx.doi.org/10.1098/rsos.150541>

Date deposited:

18/03/2016



This work is licensed under a [Creative Commons Attribution 4.0 International License](http://creativecommons.org/licenses/by/4.0/)



Cite this article: Reid WDK *et al.* 2016

Cytokine responses in birds challenged with the human food-borne pathogen

Campylobacter jejuni implies a Th17 response.

R. Soc. open sci. **3**: 150541.

<http://dx.doi.org/10.1098/rsos.150541>

Received: 8 October 2015

Accepted: 16 February 2016

Subject Category:

Cellular and molecular biology

Subject Areas:

immunology/microbiology

Keywords:

gastrointestinal tract, Th1 and Th17 response, structural equation model, bacterial pathogen, gamma-delta T lymphocytes, inflammation

Author for correspondence:

William D. K. Reid

e-mail: william.reid@newcastle.ac.uk

Electronic supplementary material is available

at <http://dx.doi.org/10.1098/rsos.150541> or via

<http://rsos.royalsocietypublishing.org>.

Cytokine responses in birds challenged with the human food-borne pathogen *Campylobacter jejuni* implies a Th17 response

William D. K. Reid¹, Andrew J. Close¹, Suzanne

Humphrey², Gemma Chaloner², Lizeth

Lacharme-Lora², Lisa Rothwell⁴, Pete Kaiser⁴, Nicola J.

Williams³, Tom J. Humphrey⁵, Paul Wigley² and

Stephen P. Rushton¹

¹School of Biology, Newcastle University, Ridley Building, Newcastle upon Tyne NE1 7RU, UK

²Department of Infection Biology, and ³Department of Epidemiology and Population Health, Institute of Global Health, University of Liverpool, Leahurst Campus, Neston, Chester CH64 7TE, UK

⁴The Roslin Institute and Royal (Dick) School of Veterinary Science, University of Edinburgh, Midlothian EH25 9RG, UK

⁵College of Medicine, Swansea University, Singleton Park, Swansea SA2 8PP, UK

Development of process orientated understanding of cytokine interactions within the gastrointestinal tract during an immune response to pathogens requires experimentation and statistical modelling. The immune response against pathogen challenge depends on the specific threat to the host. Here, we show that broiler chickens mount a breed-dependent immune response to *Campylobacter jejuni* infection in the caeca by analysing experimental data using frequentist and Bayesian structural equation models (SEM). SEM provides a framework by which cytokine interdependencies, based on prior knowledge, can be tested. In both breeds important cytokines including pro-inflammatory interleukin (IL)-1 β , IL-6, IL-4, IL-17A, interferon (IFN)- γ and anti-inflammatory IL-10 and transforming growth factor (TGF)- β 4 were expressed post-challenge. The SEM revealed a putative regulatory pathway illustrating a T helper (Th)17 response and regulation of IL-10, which is breed-dependent. The prominence of the Th17 pathway indicates the cytokine

response aims to limit the invasion or colonization of an extracellular bacterial pathogen but the time-dependent nature of the response differs between breeds.

1. Introduction

The epithelial lining of the gastrointestinal tract is a high-risk area where pathogens, especially bacteria, can infect a host. On recognition of a pathogen the host immune response involves a number of cells (e.g. natural killer, helper T (Th) cells) and signalling molecules resulting in a complex network of interactions involving the innate and adaptive immune systems [1,2]. A key component of this immune network is cytokines [3], which help control and regulate the host's immune system. Cytokine activation initiates a cascade of responses dependent on the initial stimuli that can regulate innate and adaptive immune systems [2,4–6]. The cytokine micro-environment during Th-cell activation determines effector T-cell differentiation through selective signal transducer and activator of transcription (STAT) proteins, guided by key transcription factors. Immunopathogenesis, disease progression and development of auto-immune diseases require an understanding of how cytokines interact *in vivo*. The problem that many immunological studies have is capturing the complexity of multiple immunological parameters in an analytical framework.

Innate and adaptive immune responses are initiated sequentially in order to protect against different pathogens [4], although there is feedback and cooperation between the two systems. Innate $\gamma\delta$ T cells are considered the first line of defence and development of adaptive naive CD4T cells often marks a secondary step if the innate immune system fails to cope with the infection [7]. Pro- (e.g. 1 family, interleukin (IL)-6) and anti-inflammatory (transforming growth factor (TGF)- β) cytokines stimulate naive CD4T-cell differentiation into specific effector T-cell subsets or populations. The various Th1, Th2 and Th17 cytokines have the potential to induce, amplify or control innate immune cells, which aid in the clearance or containment of the pathogen. The three main groups of effector T cells tailor their function to the specific nature of the pathogen threat [8]: Th1 cells (e.g. interferon (IFN)- γ) are important for intracellular immunity against microorganisms; Th2 cells (e.g. IL-4, IL-13) respond to extracellular pathogens including helminths and protozoans; and Th17 cells (e.g. IL-17A, IL-17F, IL-22) tackle extracellular bacteria and fungi. Further cytokines (e.g. IL-10) are produced by regulatory T (Treg) cells which aim to restrain the pro-inflammatory nature of the Th response and minimize tissue damage. However, cytokines from one Th population have the potential to inhibit the production of a different Th subset. For example, IL-4 drives Th2 cell production and IL-10 but inhibits the production of IFN- γ [9], while the Th17-Treg interactions have been described as antagonistic and cooperative [10]. While there has been extensive study of the mechanisms and pathways involved for individual cytokines in host immune responses, there have been fewer studies of how these system components interact.

Understanding the response of the immune system is complicated because the individual cytokine responses are not independent but form a network of interacting responses. A great deal of past research on cytokine response to immunological challenge has used cross-sectional data and univariate or multivariate statistical techniques to investigate processes [11]. These do not take account of the interacting nature of the immune system nor do they consider the time-dependent nature of the responses. Structural equation modelling provides a framework by which the time-dependent nature of cytokine interactions can be analysed. A structural equation model (SEM) is an extension of pathway analysis and is subtly different to those cross-sectional and descriptive statistical frameworks in that it seeks to challenge a conceptual model of the pathways in the system based on prior knowledge. The relationships among cytokines are characterized by a series of equations that link multiple response variables to one or more predictors that are defined *a priori*. The goodness of fit of the conceptual pathway model to the data is assessed through analysis of the variance and covariance structure of the putative relationships in the network of pathways [12]. In this case, the equations are defined as hypothetical interactions between cytokines where individual cytokines may up- or downregulate each other. In complex pathways, the response from one equation can act as a predictor for other cytokine responses as defined further down the reaction pathway. A key feature of SEM is that pathways/networks are developed *a priori*, and it is therefore possible to challenge competing models of the system.

Structural equation modelling has been largely overlooked in analysing immunological processes [11]; perhaps because of the relatively small samples sizes that are frequently found in immunological studies. Small sample sizes (less than 100 or less than 10 times the number of connections between variables) are often considered inadequate for developing SEM using maximum-likelihood (ML)

approaches [12]. In order to avoid such limitations, the SEM can be translated into a Bayesian framework [13,14], which has a number of advantages over the frequentist approach. The Bayesian approach incorporates prior knowledge about the parameters and does not rely on asymptotic theory and, therefore, has the potential to produce reliable results with small sample sizes [14,15].

Here, we show that the expression of a number of cytokines in response to *Campylobacter jejuni* challenge in commercial broiler chickens is breed-dependent and that the Th17 response emerges as predominant cytokine response using a combination of experiments and statistical modelling. The ever-increasing production requirement for uniform size and faster growth to harvestable size is believed to have had a negative impact on breed immunocompetence, especially to emerging zoonotic diseases [16]. We developed a conceptual model of cytokine interactions derived from published literature, which we challenged with experimental data from two commercial breeds using structural equation modelling to elucidate cytokine interactions in caecal tissue post-challenge. The primary site of *C. jejuni* colonization is the mucosal layer close to the epithelial cells in the deep crypts of the caecum at the terminal end of the gastrointestinal tract [17]. The relationship between *C. jejuni* and the chicken was originally thought to be commensal [18] with the inflammatory response being localized and self-limiting and not leading to severe pathology [19]. Recent investigations, however, indicate that the chicken–bacterium relationship cannot be considered as a commensal and severe mucosal damage can occur as a result of a prolonged pro-inflammatory response [20]. The expression of IL-1 β and IL-6 had a positive impact on IL-17A emerging as the dominant interacting cytokines in what appears to be a protective response against *C. jejuni*. Th17 cells may act as important sentinels protecting mucosal surfaces in the gastrointestinal tract and as such will be upregulated as a result of the nature of immunological challenge presented by *C. jejuni* [17].

2. Material and methods

2.1. Bacterial strains and culture conditions

Campylobacter jejuni M1 was kindly provided by Dr Lisa Williams (University of Bristol). Bacteria were grown from stocks maintained at -80°C on Columbia blood agar (Lab M, Heywood, Lancashire, UK) supplemented with 5% defibrinated horse blood (Oxoid, Basingstoke, Hampshire, UK) for 48 h in microaerobic conditions (80% N₂, 12% CO₂, 5% O₂ and 3% H₂) at 41.5°C. Liquid cultures were grown for 24 h in 10 ml of Mueller-Hinton broth (MHB) in microaerobic conditions at 41.5°C and adjusted by dilution in fresh MHB to a final concentration of 10⁶ CFU ml⁻¹.

2.2. Animal husbandry and experimental design

The two broiler breeds used for the modelling study were housed in the University of Liverpool high-biosecurity poultry unit. Breed A reaches live slaughter weight (2.2 kg) at 36 days of age while breed B reaches a similar weight by 48 days. All animals were checked a minimum of twice daily to ensure their health and welfare; different breeds and control birds were housed separately; and all individuals were screened for *Campylobacter* using cloacal swabs prior to the experimental infection. Further details of animal husbandry can be found in Humphrey *et al.* [20].

Twenty-one days old broiler chicks, A ($n = 40$) and B ($n = 40$), were orally infected with 10⁵ cells of *C. jejuni* M1 in 0.2 ml of MHB. Control birds, A ($n = 16$) and B ($n = 16$) received 0.2 ml of sterile MHB. At 2, 5 and 12 days post-challenge, 10 infected and five control birds of each breed were chosen at random and killed by cervical dislocation. At post mortem examination, samples of tissue and gut contents were taken and processed for host gene expression analysis and *Campylobacter* enumeration.

2.3. Laboratory analysis

Caecal tissue samples from infected and control birds were collected and stored in 500 μl of RNAlater at -20°C (Sigma, Poole, Dorset, UK). Total RNA was isolated from 20 to 30 mg of tissue using an RNeasy minikit (Qiagen, West Sussex, UK) according to the manufacturer's instructions. Isolated RNA was eluted into 50 μl RNase-free water and stored at -80°C . The yield of total RNA was determined using a Nano-Drop (ND-1000) spectrophotometer. Expression of mRNA for the cytokines in caecal tissues was measured by real-time quantitative reverse transcription PCR (qRT-PCR) as previously described by Humphrey *et al.* [20]. Expression of the following additional cytokines were performed using our previously described methods; IL-13 [21], IL-19 [22], IL-17A

(forward primer: CATGGGATTACAGGATCGATGA, reverse primer: GCGGCACTGGGCATCA, probe ACAACCGCTTCCCCCGCTTGG) and IL-17F (forward primer: TGACCCTGCCTCTAGGATGATC, reverse primer: GGGTCCTCATCGAGCCTGTA, probe: CAGGAATCGGTCTCTCGCTCCTTGG).

Expression of the target gene was determined using the cycle threshold (C_T) value relative to that for the 28S rRNA reference gene (ΔC_T). Results are expressed as fold changes in corrected target gene expression (ΔC_T) in infected animals relative to the control animals ($2^{-\Delta\Delta C_T}$).

2.4. Data analysis

The time-dependent expression of cytokines and subsequent modelling was undertaken in the R statistical programme (v. 3.0.3 [23]). Cytokine fold change was $\log(x + 1)$ transformed prior to analysis. Generalized linear models using quadratic, linear and $\log(\text{time})$ relationships were used to examine the relationship between cytokine fold change and time. The Akaike information criteria with the lowest score was used to select the best-fitting model. Parameter values were then estimated using the R package FME [24] by fitting the best model to the data with upper and lower bounds and, secondly, by performing a separate Markov chain Monte Carlo (MCMC) simulation. In total, 100 000 iterations were used in the MCMC estimation process, with a burn-in of 10 000 and a thinning of 10. The MCMC simulation allowed estimates of the parameter confidence intervals.

A conceptual model was developed from published experiments that brought together knowledge on potential cytokine interactions. The studies included both whole organism and cultured cells from a variety of vertebrate hosts, including humans, mice and chickens. The conceptual model and whether bird genotype has any influence on the immune response was then challenged using an SEM. The current study generated a small sample size ($n = 58$). In order to avoid sample size limitations, we initially generated an SEM using a frequentist approach in order to describe the underlying processes within these data before translating this into a Bayesian framework [13,14]. The frequentist SEM (SEMf) was fit using ML estimation in the lavaan R-package [25]. The two genotypes used in this study have different growth performances, with genotype A gaining a larger maximum size than B. A Gompertz curve is the most appropriate nonlinear regression model for describing chicken growth curves [26]. It was fitted to the commercial breeds' growth performance data and the asymptotic value was used as an indication of maximum bird mass (kg) for the model. The growth curves were fitted using the FME R-package and parameter estimates are presented in the electronic supplementary material, table S1 and figure S1). The fold change values calculated for the cytokine data showed great variability and were $\log(x + 1)$ transformed prior to SEMf analysis. In addition, IL-17F was rescaled prior to $\log(x + 1)$ transformation by dividing all values by 10 because fold changes of more than 600 were recorded. Model adjustment and selection was initiated by removing non-significant parameters and by assessing 'goodness of fit', which was assessed using: (i) a χ^2 -test, where $p > 0.05$ indicates that the observed and expected covariance matrices for the model are not different; (ii) the root mean square approximation (RMSEA), with the lower confidence interval (CI) < 0.05 and upper 90% CI < 0.1 ; (iii) standardized root mean square residual (SRMR) < 0.08 ; and (iv) comparative fit index (CFI) > 0.95 [12,27,28].

The Bayesian SEM (SEMb) analysis was performed in JAGS interfaced with R using the rjags package [29] in order to test the significance and reliability of the SEMf parameter estimates and also to investigate whether small sample sizes could be analysed effectively in this framework JAGS use MCMC simulation based on the Gibbs sampling algorithm to generate a posterior distribution of the model parameters. Non-informative priors were used as there was little prior information [13]. In total, 200 000 iterations were used in the MCMC estimation process, with a burn-in of 100 000 and a thinning of 20. Model convergence was tested using the Gelman–Rubin convergence diagnostics which measures the difference within several chains and the variance between chains of the MCMC simulation by the potential scale reduction factor (psrf) [30,31]. If convergence is achieved then the MCMC chains should be indistinguishable and the Gelman–Rubin convergence diagnostic should return a multivariate psrf of 1. The SEMb parameter estimates were deemed significant when their 95% credible intervals excluded zero.

3. Results

3.1. Response of individual cytokines to *Campylobacter jejuni* challenge

Campylobacter jejuni challenge elicited an immune response in the two breeds. In breed A, all the cytokines examined were upregulated but their responses varied over the duration of the experiment (figure 1). CXCL12, IL-1 β , TGF- β 4, IL-10 and IFN- γ were initially upregulated but began to decrease over time

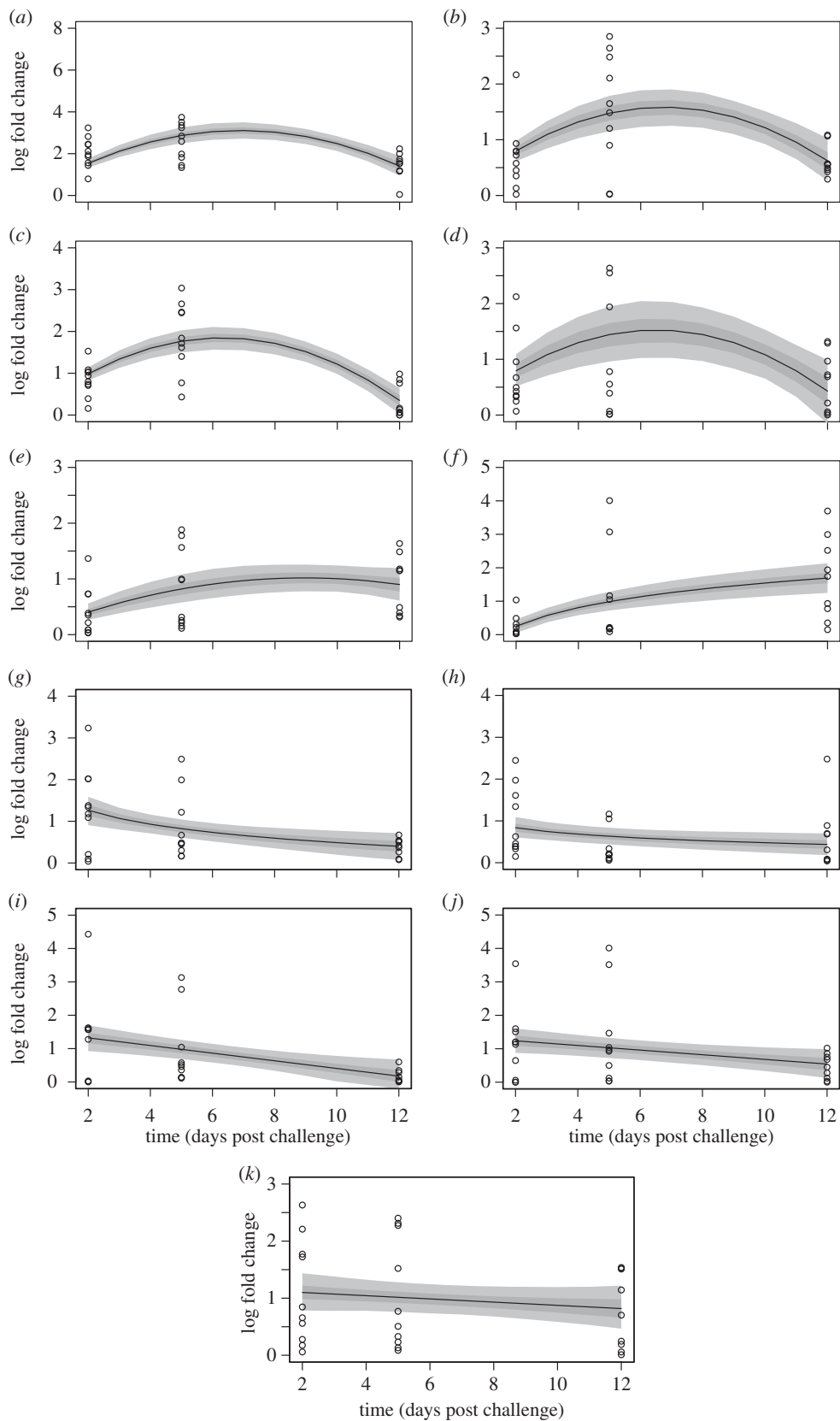


Figure 1. Time-dependent cytokine response post *Campylobacter jejuni* challenge in breed A (a) CXCLi2, (b) IL-1 β , (c) TGF- β 4, (d) IL10, (e) IFN- γ , (f) IL-17F, (g) IL-4 (h) IL-19, (i) IL-17A, (j) IL-6, (k) IL-13. The predictive envelope indicates the 25–75% (dark grey) and 5–95% (light grey) quantiles generated from the MCMC analysis of the parameter estimates.

Table 1. Parameter estimates and upper and lower 95% quantiles using Markov chain Monte Carlo (MCMC) simulation for the individual cytokine responses to *Campylobacter jejuni* challenge in breed A.

parameter	estimate (s.d.)	p-value	MCMC estimate (s.d.)	lower 95% quantile	upper 95% quantile
CXCLi2 = $a \times \text{time}^2 + b \times \text{time}$					
<i>a</i>	−0.065 (0.038)	<0.001	−0.065 (0.0069)	−0.077	−0.054
<i>b</i>	0.90 (0.42)	<0.001	0.90 (0.076)	0.78	1.02
IFN-γ = $a \times \text{time}^2 + b \times \text{time}$					
<i>a</i>	−0.012 (0.025)	<0.05	−0.012 (0.0048)	−0.020	−0.0044
<i>b</i>	0.22 (0.28)	<0.001	0.22 (0.053)	0.13	0.31
IL1-β = $a \times \text{time}^2 + b \times \text{time}$					
<i>a</i>	−0.034 (0.032)	<0.001	−0.034 (0.0061)	−0.044	−0.024
<i>b</i>	0.46 (0.36)	<0.001	0.47 (0.067)	0.36	0.58
IL-10 = $a \times \text{time}^2 + b \times \text{time}$					
<i>a</i>	−0.028 (0.040)	<0.001	−0.028 (0.0071)	−0.039	−0.016
<i>b</i>	0.39 (0.082)	<0.001	0.38 (0.079)	0.25	0.52
TGF-β4 = $a \times \text{time}^2 + b \times \text{time}$					
<i>a</i>	−0.046 (0.027)	<0.001	−0.046 (0.0051)	−0.054	−0.037
<i>b</i>	0.58 (0.29)	<0.001	0.58 (0.056)	0.48	0.67
IL-4 = $a + b \times \log(\text{time})$					
<i>a</i>	1.60 (1.78)	<0.001	1.61 (0.29)	1.11	2.13
<i>b</i>	−0.48 (1.03)	<0.05	−0.49 (0.16)	−0.76	−0.20
IL-17F = $a + b \times \log(\text{time})$					
<i>a</i>	−0.30 (2.52)	0.51	−0.30 (0.16)	−0.55	−0.038
<i>b</i>	0.80 (1.46)	<0.05	0.80 (0.13)	0.59	1.04
IL-19 = $a + b \times \log(\text{time})$					
<i>a</i>	0.99 (1.67)	<0.01	0.99 (0.26)	0.65	1.32
<i>b</i>	−0.22 (0.96)	0.22	−0.22 (0.10)	−0.39	−0.047
IL-6 = $a + b \times \text{time}$					
<i>a</i>	1.38 (1.88)	<0.001	1.38 (0.26)	0.94	1.82
<i>b</i>	−0.07 (0.25)	0.14	−0.071 (0.031)	−0.12	−0.019
IL-17A = $a + b \times \text{time}$					
<i>a</i>	1.56 (1.77)	<0.001	1.55 (0.28)	1.10	2.01
<i>b</i>	−0.11 (0.24)	<0.05	−0.11 (0.038)	−0.17	−0.05
IL-13 = $a + b \times \text{time}$					
<i>a</i>	1.18 (1.52)	<0.001	1.15 (0.24)	0.76	1.54
<i>b</i>	−0.033 (0.20)	0.38	−0.027 (0.030)	−0.076	0.024

while IL-17F continued to increase over the duration of the experiment (figure 1*a–f* and table 1). IL-4, IL-17A, IL-6, IL-19 and IL-13 all decreased from 2 to 12 days post-challenge but the relationship was only significant in IL-4 and IL-17A (figure 1*g–k* and table 1). Similar to breed A, upregulation of all the cytokines was observed in breed B (figure 2 and table 2). However, cytokine response over time varied between the two breeds. IL-10, IFN- γ and IL-17F all showed sustained increases over the experimental duration (figure 2*d–f* and table 2), while IL-19 response was greatest mid-way through (figure 2*h*). Parameter estimation indicated that CXCLi2, TGF- β 4, IL-4, IL-17A, IL-6 and IL-13 all decreased from 2 to 12 days post-challenge (figure 2*b,c,g,i,j* and table 2) but the relationship was only significant for IL-4 and IL-6 (table 2).

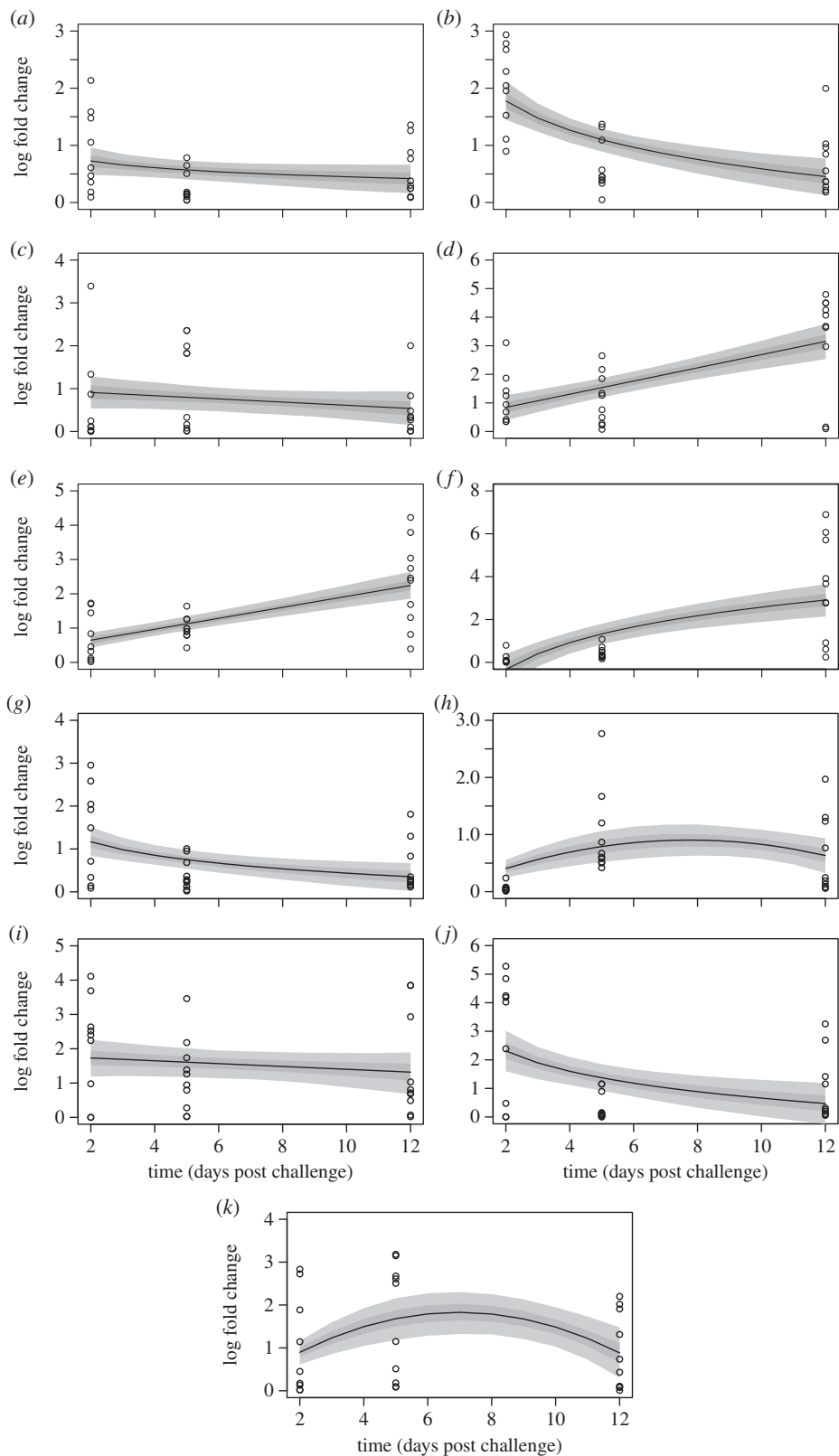


Figure 2. Time-dependent cytokine response post *Campylobacter jejuni* challenge in breed B (a) CXCLi2, (b) IL-1 β , (c) TGF- β 4, (d) IL-10, (e) IFN- γ , (f) IL-17F, (g) IL-4 (h) IL-19, (i) IL-17A, (j) IL-6, (k) IL-13. The predictive envelope indicates the 25–75% (dark grey) and 5–95% (light grey) quantiles generated from the MCMC analysis of the parameter estimates.

Table 2. Parameter estimates and upper and lower 95% quantiles using Markov chain Monte Carlo (MCMC) simulation for the individual cytokine responses to *Campylobacter jejuni* challenge in breed B.

parameter	estimate (s.d.)	p-value	MCMC estimate (s.d.)	lower 95% quantile	upper 95% quantile
CXCLi2 = a + b × log(time)					
a	0.84 (1.33)	<0.01	0.83 (0.21)	0.46	1.16
b	−0.17 (0.75)	0.22	−0.16 (0.12)	−0.36	0.04
IFN-γ = a + b × time					
a	0.32 (1.55)	0.27	0.32 (0.16)	0.058	0.59
b	0.16 (0.20)	<0.001	0.15 (0.026)	0.11	0.20
IL-1β = a + b × log(time)					
a	2.29 (1.64)	<0.001	2.28 (0.30)	1.81	2.80
b	−0.74 (0.92)	<0.001	−0.74 (0.17)	−1.02	−0.46
IL-10 = a + b × time					
a	0.37 (2.34)	0.44	0.38 (0.33)	−0.15	0.94
b	0.23 (0.30)	<0.001	0.22 (0.045)	0.15	0.30
TGF-β4 = a + b × time					
a	0.98 (1.76)	<0.01	0.97 (0.26)	0.53	1.40
b	−0.038 (0.22)	0.37	−0.037 (0.032)	−0.090	0.017
IL-4 = a + b × log(time)					
a	1.47 (1.86)	<0.001	1.47 (0.31)	0.94	2.00
b	−0.44 (1.04)	<0.05	−0.44 (0.17)	−0.73	−0.15
IL-17F = a + b × log(time)					
a	−1.56 (3.75)	<0.05	−1.55 (0.67)	−2.66	−0.44
b	1.80 (2.10)	<0.001	1.79 (0.37)	1.17	2.41
IL-19 = a × time² + b × time					
a	−0.015 (0.029)	<0.01	−0.015 (0.0050)	−0.024	−0.0073
b	0.23 (0.32)	<0.001	0.23 (0.056)	0.14	0.33
IL-6 = a + b × log(time)					
a	3.00 (3.84)	<0.001	2.96 (0.65)	1.93	4.06
b	−1.02 (2.16)	<0.05	−1.00 (0.36)	−1.61	−0.38
IL-17A = a + b × time					
a	1.83 (2.52)	<0.001	1.82 (0.39)	1.17	2.48
b	−0.042 (0.32)	0.49	−0.041 (0.049)	−0.12	0.038
IL-13 = a × time² + b × time					
a	−0.037 (0.052)	<0.001	−0.037 (0.0089)	−0.051	−0.022
b	0.52 (0.58)	<0.001	0.52 (0.10)	0.35	0.68

3.2. Development of a conceptual model to challenge with a structural equation model

Examining the interaction of cytokines in response to *C. jejuni* challenge in chickens using SEM required the development of a conceptual *a priori* model from the published literature (figure 3). When a host recognizes specific bacterial ligands and microbial products an immune response is activated. An initial response produces the interleukins, IL-1β and IL-6 [19,32–34]. IL-1β plays a major role in the innate immune response and induces the production IL-6 and CXCLi2 [17,19,35,36]. The pattern of signals

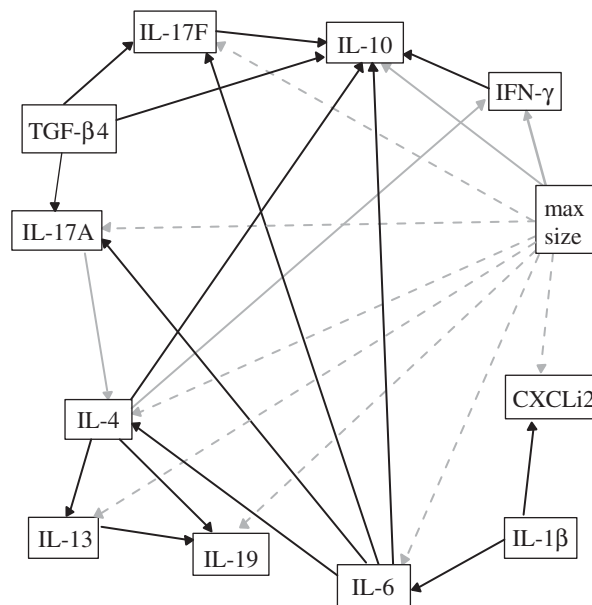


Figure 3. Full model depicting the potential response of cytokines and their interactions developed from peer-reviewed literature after challenge by a bacterial pathogen. This network of interactions is challenged by structural equation modelling. The arrows indicate the order of cytokine regulations: solid black arrows indicate a positive response or upregulation; solid grey arrows indicate a negative response or downregulation; and grey dashed arrows indicate that there is a negative relationship between maximum body size, used as a surrogate for breed type and cytokine response.

received by the host during the interaction with the pathogen will then determine whether a Th1, Th2 or Th17 response will be initiated [32]. The Th2 response is mediated through the upregulation of IL-4 [3,32,33], which can be stimulated by the production of IL-6 [37,38]. Production of IL-4 results in the induction of IL-13 and IL-19 [3,33,39]. IL-4 counteracts the IFN- γ function by suppressing the inflammatory response [40]. Th17 response is initiated by IL-1 β , IL-6 and TGF- β can upregulate IL-17A and IL17F [41] and is negatively related to IL-4 and IFN- γ [42,43]. IL-10 production can be stimulated by cytokines of the Th1, Th2 and Th17 pathways [32,41,44] but its primary role is part of the Treg pathway of which TGF- β is the most important stimulating cytokine. This corresponds to a series of interactions as shown in figure 3.

3.3. Structural equation model of the cytokines response to *Campylobacter jejuni* challenge

The experiment provided data to investigate the cytokine interactions and assess which were involved in the hypothesized regulatory pathways detailed in the *a priori* model (figure 3). Model validation was initiated using the SEMf. The full *a priori* model was not an adequate representation of these data because of the non-significant parameter coefficients within the pathways and based on the RMSEA score (0.07) and the upper bounds of the 90% CI (0.12). IL-13 and IL-19, as well as a series of interaction pathways between these and other cytokines, including IFN- γ regulating IL-10 and a number of the cytokine-maximum size relationships, were removed from the model because they were non-significant. IL-4 ($p = 0.056$) and TGF- β 4 ($p = 0.057$) influence on IL-10 and IL-17A, respectively, were marginally non-significant and therefore remained within the model.

The model adjustments greatly improved the 'goodness of fit'. The final SEMf converged normally after 53 iterations: $\chi^2_{22} = 19.30$, $p = 0.62$; RMSEA = 0.00, 90% CI = 0.00–0.09; SRMR = 0.06; CFI = 1.00. The final SEMf including all the direct and indirect modelled cytokine interaction pathways is shown in figure 4 with the standardized coefficients while the unstandardized coefficients can be found in table 3. IL-17A response to TGF- β 4 and IL-10 response to IL-4 did not improve in the final model (table 3).

The final SEMf was translated into a Bayesian framework. The SEMb was tested for convergence and whether it explained the observed variance in these data. The Gelman–Rubin convergence diagnostic test resulted in a multivariate psrf = 1. This can be visualized with the development of the psrf over the chain iterations (electronic supplementary material, figure S2). The MCMC chains all converged for the parameters and the model explained the variance observed in these data. The parameter estimates

Table 3. Coefficients, credible intervals and significance for the parameter estimates of the final frequentist structural equation model (SEMf) and Bayesian structural equation model (SEMb) for cytokine interactions in response to *Campylobacter jejuni* challenge in chickens. (The SEMf parameter estimates are unstandardized. The SEMb significant parameter estimates are highlighted in bold.)

response	predictor	SEMf parameter estimate (standard error)	p-value	SEMb parameter estimate	95% credible intervals
IL-10	IL17F	0.66 (0.14)	<0.001	0.68	0.26–1.10
	IL-4	0.67 (0.36)	0.06	0.64	–0.45 to 1.73
	IL-6	–0.51 (0.21)	<0.05	–0.50	–1.12 to 0.11
	max. size	–2.34 (0.60)	<0.001	–2.16	–3.95 to –0.36
IFN- γ	max. size	–1.37 (0.46)	<0.01	–1.37	–2.62 to –0.14
IL-4	IL-6	0.41 (0.04)	<0.001	0.41	0.22–0.60
	IL-17A	0.16 (0.04)	<0.001	0.16	–0.05 to 0.38
	max. size	0.77 (0.18)	<0.001	0.77	–0.08 to 1.64
IL-6	IL-1 β	1.28 (0.15)	<0.001	1.28	0.87–1.68
CXCLi2	IL-1 β	0.40 (0.10)	<0.001	0.41	0–0.81
	max. size	3.20 (0.34)	<0.001	3.21	1.85–4.57
IL-17A	IL-6	0.57 (0.08)	<0.001	0.57	0.33–0.81
	TGF- β 4	0.25 (0.13)	0.05	0.25	–0.13 to 0.63
	max. size	–1.02 (0.49)	<0.05	–1.16	–2.61 to 0.26

for the SEMf and SEMb were very similar (table 3) suggesting that the two modelling approaches were providing comparable results. The SEMb, however, indicated that 4 out of 14 parameter estimates were non-significant. The SEMb non-significant pathways were the relationships between: IL-10 and IL-4, IL-17A and TGF- β 4 and the negative relationships between IL-10 and IL-6 and maximum body size and IL-17A (figure 4). Yet these pathways can still be considered marginally non-significant (table 3).

The exogenous variable maximum body size, a surrogate for bird breed, had a major impact on the cytokine response throughout the network. While there was no effect of breed on the cytokines associated with the initial cytokine response (IL-1 β and IL-6), there were other effects deeper in the network. Breed type had an effect on CXCLi2 response to IL-1 β , which can be interpreted as having an impact on a secondary step within a pathway (figure 4); i.e. upregulation of IL-1 β by a pattern recognition receptor is step one while the response of CXCLi2 to IL-1 β is step two. Breed type also had an effect on the third step of a pathway which impacted on IL-4, IL-17A and IL-10 responding to IL-6 (figure 4). There was a positive pathway that linked IL-1 β to IL-4 and IL17A via IL-6 (figure 4). IL-1 β was a positive driver of IL-6 and CXCLi2. IL-6 had a direct effect on IL-4 but also indirectly via IL-17A. IL-10 was influenced by IL-4, IL-6 and IL-17F, with the strongest relationship being between IL-10 and IL-6 which was negative.

4. Discussion

The primary aim of many immunological studies is to investigate changes in some form of immune status at a cellular, tissue or organ level over time in response to variations in the internal environment whether that is elicited by a pathogen, another agent or from an auto-immune perspective. The time-dependent analysis presented here, while based on a relatively small dataset, provided valuable information on which cytokines were up- or downregulated post *C. jejuni* challenge. However, it does not demonstrate cytokine interdependencies which leave knowledge of the system rather fragmented. Undertaking a structural equation modelling approach allowed us to investigate interactions among cytokines which allowed us to identify the most important pathways. We believe that the combined modelling approach provided a deeper understanding of the system because it identified which cytokines were upregulated and then allowed us to challenge a conceptual model of the theoretical relationships among cytokines with real data.

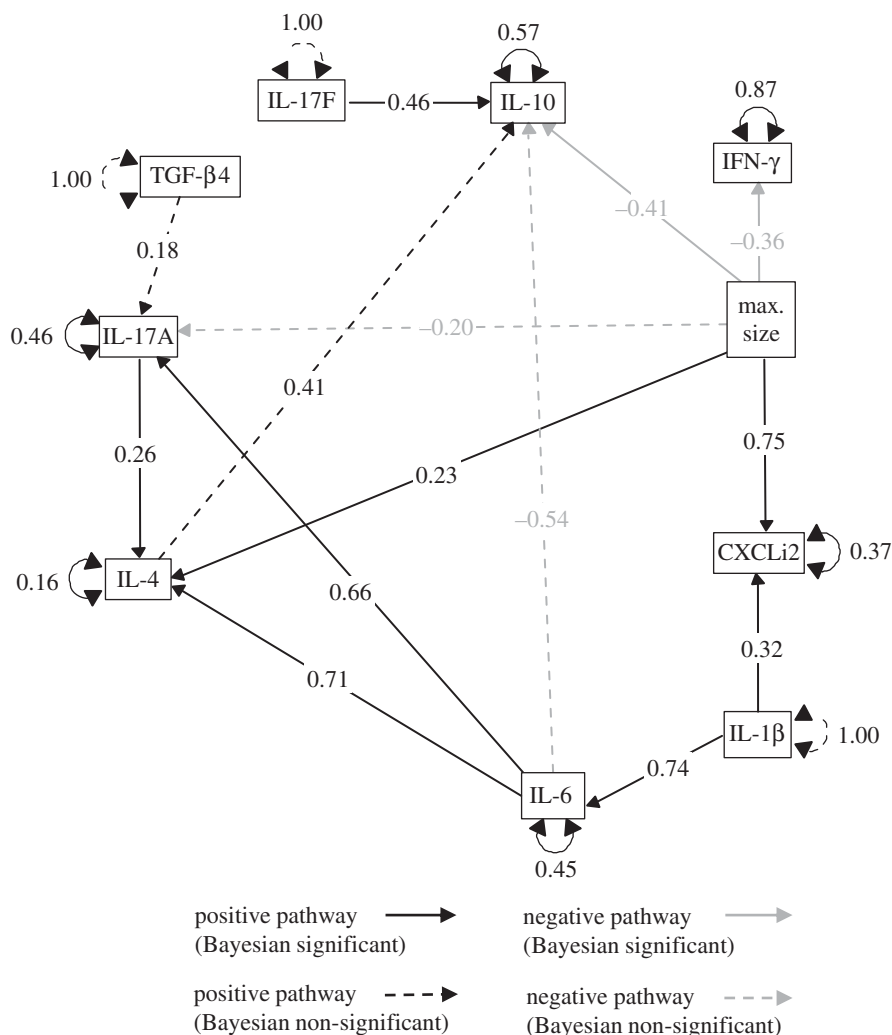


Figure 4. The final path model describing the cytokine interactions post-challenge in two breeds of broiler chicken. Positive relationships are indicated by black arrows, whereas grey arrows indicate a negative response. The standardized parameter estimates for each response are shown on each arrow. The unexplained variation for each of the variables, internally predicted by the model, is shown adjacent to their respective boxes.

The time-dependent response and SEM indicated that there were differences between the faster, larger growing breed A and the slower growing, smaller breed B in their cytokine responses to *C. jejuni* challenge. Both breeds initiated an innate response through the upregulation of CXCLi2, IL-1 β and IL-6. The SEM indicated that the level of CXCLi2 expression was breed-dependent, as there was a positive relationship between the exogenous variable maximum size and expression of CXCLi2, although expression of IL-1 β and IL-6 were not related to maximum size. CXCLi2 is homologous with human IL-8 and is important for an early immune response in the gastrointestinal tract, especially for its chemotactic role in monocyte and heterophil recruitment [3,45]. The upregulation of CXCLi2 suggested that breed A and B initiated a functional innate response to *C. jejuni* [20]. However, the subsequent upregulation of IFN- γ , IL-17A and IL-10 and their levels expressed through time, along with CXCLi2, indicated the course of the immunological response to *C. jejuni* varied between the two breeds. The SEM identified this through a negative relationship between size and IFN- γ , IL-17A and IL-10. Both IFN- γ and IL-17A are important effector T cells and play a crucial role in the clearance of bacterial pathogens [8,19,32]; they are also potent pro-inflammatory mediators. The anti-inflammatory IL-10 is often expressed by T regulatory cells and is important in restraining the pro-inflammatory response. The differential response of IFN- γ , IL-17A and IL-10 has implications for how the two breeds fight infection but also how each breed controls inflammation. We have previously shown that the inflammatory response between these breeds differs, with prolonged inflammation and inflammatory damage occurring in the faster growing breed A, even though invasion and colonization are similar [20].

The Th17 response was clearly identified in the final SEMf and was in agreement with the SEMb along with the upregulation of innate immune system chemokine CXCLi2. In mammals, the Th17 pathway responds to extracellular bacterial infection and while frequently associated with inflammatory conditions, also plays a key role as a sentinel response in the intestinal tract, preventing invasion by bacterial pathogens and maintaining gut integrity [4]. While the role of the Th17 pathway has yet to be fully elucidated in the chicken, it is likely to play a similar role particularly during *C. jejuni* infection which colonizes the host in the mucosal layer of the deep crypts within the caecum. The chemotactic properties of IL-17 make it an important inducer of CXCLi1 and CXCLi2 chemokines. Yet, the SEM did not identify any IL-17-CXCLi2 association within the model's covariance structure. The key cytokines resulting in the induction of IL-17A expressed within the SEM is largely in agreement with that expected from experimental studies [46] but the SEM did not explain all of the variation observed in IL-17A nor did it implicate any cytokines in the upregulation of IL-17F. The SEM indicated that IL-17A production potentially started with IL-1 β which had a positive effect on IL-6 and in turn stimulated IL-17A. IL-1 β combined with IL-6, a STAT3 activator, are important for Th17 cell differentiation. IL-1 β has a crucial role at the initial stages of the Th17 cell differentiation because it upregulates the expression of IL-1 receptors and retinoic acid receptor-related orphan receptors- γ t on the surface of Th17 lymphocytes resulting in the subsequent production of IL-6 and IL-17A [47].

The SEM also indicated that approximately 50% of IL-17A variance was unexplained. The cytokine cascade involving IL-1 β , IL-6 and IL-17A potentially relates to the differentiation of naive CD4T cells which are part of the adaptive immune response. The unexplained variation will undoubtedly be related to interactions with other cytokines or induction pathways that were not captured within the model. IL-23 was not part of the suit of cytokines examined post-experimentation but it has a role in activating and sustaining Th17 cell development [7], leading to the production of IL-17 cytokines that induce localized tissue inflammation [48]. Toll-like receptors (TLR) on innate immune $\gamma\delta$ T cells can produce IL-17A [49]. TLR are a group of receptors that are involved in orchestrating the innate immune response and recognize bacteria and microbial products. *Campylobacter jejuni* activates TLR2 in chickens [50], which in mouse models is believed to interact directly with pathogens resulting in the upregulation of IL-17 [49]. Intestinal epithelium harbour important populations of $\gamma\delta$ T lymphocytes where they act as a first line of defence against pathogens [51]. $\gamma\delta$ T cells are more numerous in the chicken accounting for up to 60% of the peripheral lymphocytes and are numerous in mucosal sites such as the gastrointestinal and reproductive tracts [51–53]. We hypothesize that $\gamma\delta$ T cells may play a key role in avian mucosal immunity producing a substantial amount of IL-17.

There was only a weak relationship between IL-17A and TGF- β 4 while TGF- β 4 appeared to have no effect on IL-17F, IL-10 or IFN- γ . TGF- β is often implicated in conjunction with IL-6 as important for the upregulation of IL-17 cytokines even though Th17 cytokines can differentiate in a TGF- β independent manner [46]. The role of the anti-inflammatory TGF- β in regulating Th17 cell differentiation is complex because there appears to be differences in the relative importance of TGF- β in the Th17 response depending on the system being studied [46,48]. In humans, there is evidence for TGF- β -dependent and -independent [46,54] production of IL-17 in conjunction with other cytokines but in mice TGF- β is a requirement for successful development of Th17 cells [46]. Human and mouse model systems focus on TGF- β 1 but TGF- β 4 in chickens is believed to have the same functional role [3]. The role of TGF- β *in vivo* is likely to be dependent on a number of environmental stimuli not to mention interactions with other cytokines, which may not be evident when particular genes or cytokines are experimentally suppressed. The weak relationship observed in the SEM may indicate the TGF- β 4 is not involved in the dominant IL-17 producing pathway or that TGF- β 4 concentrations are below a particular threshold, which in conjunction with IL-6 and IL-1 β , allows the production of IL-17A instead of Treg [55,56].

Both IL-10 and TGF- β are anti-inflammatory cytokines which have the potential to suppress Th1, Th2 and Th17 responses. IL-10 is crucial for the control of inflammation during the host's response to a pathogen and can be regulated by the co-induction of Th1 [44], TGF- β [57] and IL-4. Neither IFN- γ nor TGF- β appeared to have an effect on IL-10 in the SEM, either directly or indirectly via examination of the model's covariance structure. The SEM indicated that IL-6 had a negative effect on IL-10. IL-6 production has the potential to block Treg activity. In mammals, IL-6 induction through STAT3 phosphorylation downregulates FOXP3 and blocks Treg cell differentiation while maintaining the Th17 response in order to continue dealing with the infection [58,59]. Although our understanding of chicken Tregs is rudimentary a similar process may be happening here. The fact that IL-6 has a negative effect on IL-10 and there is no relationship between IL-10 and IFN- γ or TGF- β 4 provided more evidence that the chicken's response is aimed at fighting an extracellular bacterial infection. However, both IL-4 and IL-17F had a positive effect on IL-10, suggesting these cytokines were interacting in a positive manner. It

may be that the IL-6 positive effect on IL-4 in turn allows the upregulation of IL-10 through a different pathway. Such an interaction would potentially allow the anti-inflammatory properties of IL-10 to control tissue inflammation and protect the host. However, without further experimental studies this assertion is speculative.

The results are of biological as well as analytical interest because the approach provided an opportunity to validate the potential for using an SEM to investigate cytokine interactions [11]. From an infection immunology perspective, these modelling approaches can add new insight into the complex interactions of cytokine responses in a system such as the gut. While we realize that we have analysed only part of the immune system response to challenge with a pathogen, there is no reason why this analytical approach could not be expanded to include other key components of the immune system beyond cytokines, e.g. $\gamma\delta$ T cell or naive CD4T cell proliferation, or measures of inflammation as a latent variable. Structural equation modelling may even assist in identifying which immunological variables result in recurrence of disease [60]. It is clear that these statistical techniques have considerable potential in defining the essential protective responses or crucial differences in responses that underlie host immune responses to any form of infection or challenge to the immune system.

In this study, the overarching hypothesis was that bird genotype impacts on the immunocompetence of commercial broiler chickens and that this would be evident in the cytokine response. We used a combined modelling approach that first assessed the time-dependent cytokine response of two breeds of broiler chicken to *C. jejuni* challenge before examining cytokine interactions in a generalized multi-equation framework facilitated by structural equation modelling [61]. Both SEM approaches demonstrated that bird genotype had an effect on cytokine interactions post *C. jejuni* challenge in caecal tissue as well as identifying interdependences among cytokines, which have the potential to be cytokine cascades. The fact that IL-13 and IL-19 were removed from the final model and IFN- γ appeared to be only related to bird genotype potentially indicated that the Th17 response was the prominent cytokine pathway observed. The prominence of the Th17 pathway suggests that broiler chickens' immune response was directed at controlling an extracellular bacterial pathogen through restricting it to the gut lumen.

Ethics. All work was undertaken in accordance with UK legislation governing experimental animals under project license PPL 40/3063 and was approved by the University of Liverpool ethical review process prior to the award of the license.

Data accessibility. The data used in this study are available from the Dryad Digital Repository and can be found under the following: <http://dx.doi.org/10.5061/dryad.7h3v1>.

Competing interests. We have no competing interests.

Authors' contributions. W.D.K.R., A.J.C., S.H., P.W. and S.P.R. wrote the manuscript and analysed these data; P.W., S.P.R., N.J.W., T.H. and S.H. designed the experiments; P.W., L.R., P.K., G.C., S.H. and L.L. conducted the experiments; N.J.W. and T.J.H. edited the manuscript.

Funding. We would like to acknowledge Biotechnology and Biological Sciences Research Council (BB/I024674/1 and BB/J017159/1) for funding the project and UK food retailers, poultry breeding, production and feed companies.

Acknowledgements. We thank Sue Jopson and Anne Forrester for their excellent work in caring for the chickens and the technicians at the Histology Laboratories, Veterinary Laboratory Services, School of Veterinary Science at the University of Liverpool for excellent technical support. We also thanks two anonymous reviewers for their positive comments and valuable suggestions.

References

- Kamada N, Seo SU, Chen GY, Nunez G. 2013 Role of the gut microbiota in immunity and inflammatory disease. *Nat. Rev. Immunol.* **13**, 321–335. (doi:10.1038/nri3430)
- Sansonetti PJ. 2004 War and peace at mucosal surfaces. *Nat. Rev. Immunol.* **4**, 953–964. (doi:10.1038/nri1499)
- Kaiser P, Staheli P. 2014 Avian cytokines and chemokines. In *Avian immunology* (eds KA Schat, B Kaspers, P Kaiser), pp. 189–204. London, UK: Elsevier.
- Littman DR, Rudensky AY. 2010 Th17 and regulatory T cells in mediating and restraining inflammation. *Cell* **140**, 845–858. (doi:10.1016/j.cell.2010.02.021)
- Peterson LW, Artis D. 2014 Intestinal epithelial cells: regulators of barrier function and immune homeostasis. *Nat. Rev. Immunol.* **14**, 141–153. (doi:10.1038/nri3608)
- Ivanov II, Frutos RD, Manel N, Yoshinaga K, Rifkin DB, Sartor RB, Finlay BB, Littman DR. 2008 Specific microbiota direct the differentiation of IL-17-producing T-helper cells in the mucosa of the small intestine. *Cell Host Microbe* **4**, 337–349. (doi:10.1016/j.chom.2008.09.009)
- Stockinger B, Veldhoen M, Martin B. 2007 Th17 cells: linking innate and adaptive immunity. *Semin. Immunol.* **19**, 353–361. (doi:10.1016/j.smim.2007.10.008)
- Curtis MM, Way SS. 2009 Interleukin-17 in host defence against bacterial, mycobacterial and fungal pathogens. *Immunology* **126**, 177–185. (doi:10.1111/j.1365-2567.2008.03017.x)
- Min W, Kim WH, Lillehoj EP, Lillehoj HS. 2013 Recent progress in host immunity to avian coccidiosis: IL-17 family cytokines as sentinels of the intestinal mucosa. *Dev. Comp. Immunol.* **41**, 418–428. (doi:10.1016/j.dci.2013.04.003)
- Chen X, Oppenheim JJ. 2014 Th17 cells and T-regs: unlikely allies. *J. Leukoc. Biol.* **95**, 723–731. (doi:10.1189/jlb.1213633)
- Genser B, Cooper PJ, Yazdanbakhsh M, Barreto ML, Rodrigues LC. 2007 A guide to modern statistical

- analysis of immunological data. *BMC Immunol.* **8**, 15. (doi:10.1186/1471-2172-8-27)
12. Grace JB. 2006 *Structural equation modeling and natural systems*. Cambridge, UK: Cambridge University Press.
 13. Song XY, Lee SY. 2012 A tutorial on the Bayesian approach for analyzing structural equation models. *J. Math. Psychol.* **56**, 135–148. (doi:10.1016/j.jmp.2012.02.001)
 14. Lee SK. 2007 *Structural equation modeling: a Bayesian approach*. Chichester, UK: Wiley.
 15. Tetard-Jones C, Gatehouse AMR, Cooper J, Leifert C, Rushton S. 2014 Modelling pathways to Rubisco degradation: a structural equation network modelling approach. *PLoS ONE* **9**, 6. (doi:10.1371/journal.pone.0087597)
 16. Swaggerty CL, Pevzner IY, Kaiser P, Kogut MH. 2008 Profiling pro-inflammatory cytokine and chemokine mRNA expression levels as a novel method for selection of increased innate immune responsiveness. *Vet. Immunol. Immunopathol.* **126**, 35–42. (doi:10.1016/j.vetimm.2008.06.005)
 17. Smith CK, AbuOun M, Cawthraw SA, Humphrey TJ, Rothwell L, Kaiser P, Barrow PA, Jones MA. 2008 *Campylobacter* colonization of the chicken induces a proinflammatory response in mucosal tissues. *FEMS Immunol. Med. Microbiol.* **54**, 114–121. (doi:10.1111/j.1574-695X.2008.00458.x)
 18. Beery JT, Hugdahl MB, Doyle MP. 1998 Colonization of gastrointestinal tracts of chickens by *Campylobacter jejuni*. *Appl. Environ. Microbiol.* **54**, 2365–2370.
 19. Smith CK, Kaiser P, Rothwell L, Humphrey T, Barrow PA, Jones MA. 2005 *Campylobacter jejuni*—induced cytokine responses in avian cells. *Infect. Immun.* **73**, 2094–2100. (doi:10.1128/iai.73.4.2094-2100.2005)
 20. Humphrey S, Chaloner G, Kemmett K, Davidson N, Williams N, Kipar A, Humphrey T, Wigley P. 2014 *Campylobacter jejuni* is not merely a commensal in commercial broiler chickens and affects bird welfare. *mBio* **5**, e013364. (doi:10.1128/mBio.01364-14)
 21. Shini S, Kaiser P. 2009 Effects of stress, mimicked by administration of corticosterone in drinking water, on the expression of chicken cytokine and chemokine genes in lymphocytes. *Stress* **12**, 388–399. (doi:10.1080/10253890802526894)
 22. Powell FL, Rothwell L, Clarkson MJ, Kaiser P. 2009 The turkey, compared to the chicken, fails to mount an effective early immune response to *Histomonas meleagridis* in the gut. *Parasite Immunol.* **31**, 312–327. (doi:10.1111/j.1365-3024.2009.01113.x)
 23. R Core Team. 2011 *R: a language and environment for statistical computing*. Vienna, Austria: R Foundation for Statistical Computing.
 24. Soetaert K, Petzoldt T. 2010 Inverse modelling, sensitivity and Monte Carlo analysis in R using package FME. *J. Stat. Softw.* **33**, 28. (doi:10.18637/jss.v033.i03)
 25. Rosseel Y. 2012 lavaan: an R package for structural equation modeling. *J. Stat. Softw.* **48**, 1–36. (doi:10.18637/jss.v048.i02)
 26. Aggrey SE. 2002 Comparison of three nonlinear and spline regression models for describing chicken growth curves. *Poult. Sci.* **81**, 1782–1788. (doi:10.1093/ps/81.12.1782)
 27. Allen VM, Ridley AM, Harris JA, Newell DG, Powell L. 2011 Influence of production system on the rate of onset of *Campylobacter* colonization in chicken flocks reared extensively in the United Kingdom. *Br. Poult. Sci.* **52**, 30–39. (doi:10.1080/00071668.2010.537306)
 28. Kline RB. 2011 *Principles and practices of structural equation modeling*, 3rd edn. New York, NY: Guilford.
 29. Plummer M. 2014 rjags: Bayesian graphical models using MCMC. R package version 3.13 edn. See <https://cran.r-project.org/web/packages/rjags/index.html>.
 30. Brooks SP, Gelman A. 1998 General methods for monitoring convergence of iterative simulations. *J. Comput. Graph. Stat.* **7**, 434–455. (doi:10.2307/1390675)
 31. Gelman A, Rubin DB. 1992 Inference from iterative simulation using multiple sequences. *Stat. Sci.* **7**, 457–472. (doi:10.1214/ss/117701136)
 32. Zhu JF, Paul WE. 2008 CD4T cells: fates, functions, and faults. *Blood* **112**, 1557–1569. (doi:10.1182/blood-2008-05-078154)
 33. Kaiser P. 2010 Advances in avian immunology—prospects for disease control: a review. *Avian Pathol.* **39**, 309–324. (doi:10.1080/03079457.2010.508777)
 34. Hermans D, Pasmans F, Heyndrickx M, Van Immerseel F, Martel A, Deun K, Haesebrouck F. 2012 A tolerogenic mucosal immune response leads to persistent *Campylobacter jejuni* colonization in the chicken gut. *Crit. Rev. Microbiol.* **38**, 17–29. (doi:10.3109/1040841x.2011.615298)
 35. Weining KC, Sick C, Kaspers B, Staeheli P. 1998 A chicken homolog of mammalian interleukin-1 beta: cDNA cloning and purification of active recombinant protein. *Eur. J. Biochem.* **258**, 994–1000. (doi:10.1046/j.1432-1327.1998.2580994.x)
 36. Staeheli P, Puehler F, Schneider K, Gobel TW, Kaspers B. 2001 Cytokines of birds: conserved functions—a largely different look. *J. Interferon Cytokine Res.* **21**, 993–1010. (doi:10.1089/107999001317205123)
 37. He HQ, Genovese KJ, Swaggerty CL, MacKinnon KM, Kogut MH. 2012 Co-stimulation with TLR3 and TLR21 ligands synergistically up-regulates Th1-cytokine IFN-gamma and regulatory cytokine IL-10 expression in chicken monocytes. *Dev. Comp. Immunol.* **36**, 756–760. (doi:10.1016/j.dci.2011.11.006)
 38. Avery S, Rothwell L, Degen WDJ, Schijns V, Young J, Kaufman J, Kaiser P. 2004 Characterization of the first nonmammalian T2 cytokine gene cluster: the cluster contains functional single-copy genes for IL-3, IL-4, IL-13, and GM-CSF, a gene for IL-5 that appears to be a pseudogene, and a gene encoding another cytokine-like transcript, KK34. *J. Interferon Cytokine Res.* **24**, 600–610. (doi:10.1089/1079990042280865)
 39. Hofmann SR, Rosen-Wolff A, Tsokos GC, Hedrich CM. 2012 Biological properties and regulation of IL-10 related cytokines and their contribution to autoimmune disease and tissue injury. *Clin. Immunol.* **143**, 116–127. (doi:10.1016/j.clim.2012.02.005)
 40. Rincon M, Anguita J, Nakamura T, Fikrig E, Flavell RA. 1997 Interleukin (IL)-6 directs the differentiation of IL-4-producing CD4⁺ T cells. *J. Exp. Med.* **185**, 461–469. (doi:10.1084/jem.185.3.461)
 41. McGeachy MJ, Bak-Jensen KS, Chen Y, Tato CM, Blumenschein W, McClanahan T, Cua DJ. 2007 TGF- β and IL-6 drive the production of IL-17 and IL-10 by T cells and restrain Th17 cell-mediated pathology. *Nat. Immunol.* **8**, 1390–1397. (doi:10.1038/nri1539)
 42. Harrington LE, Hatton RD, Mangan PR, Turner H, Murphy TL, Murphy KM, Weaver CT. 2005 Interleukin 17-producing CD4⁺ effector T cells develop via a lineage distinct from the T helper type 1 and 2 lineages. *Nat. Immunol.* **6**, 1123–1132. (doi:10.1038/nri1254)
 43. Park H *et al.* 2005 A distinct lineage of CD4 T cells regulates tissue inflammation by producing interleukin 17. *Nat. Immunol.* **6**, 1133–1141. (doi:10.1038/nri1261)
 44. Cope A, Le Friec G, Cardone J, Kemper C. 2011 The Th1 life cycle: molecular control of IFN- γ to IL-10 switching. *Trends Immunol.* **32**, 278–286. (doi:10.1016/j.it.2011.03.010)
 45. Kaiser P *et al.* 2005 A genomic analysis of chicken cytokines and chemokines. *J. Interferon Cytokine Res.* **25**, 467–484. (doi:10.1089/jir.2005.25.467)
 46. Acosta-Rodríguez EV, Napolitani G, Lanzavecchia A, Sallusto F. 2007 Interleukins 1 beta and 6 but not transforming growth factor-beta are essential for the differentiation of interleukin 17-producing human T helper cells. *Nat. Immunol.* **8**, 942–949. (doi:10.1038/nri1496)
 47. Chung Y *et al.* 2009 Critical regulation of early Th17 cell differentiation by interleukin-1 signaling. *Immunity* **30**, 576–587. (doi:10.1016/j.immuni.2009.02.007)
 48. Gaffen SL, Jain R, Garg AV, Cua DJ. 2014 The IL-23-IL-17 immune axis: from mechanisms to therapeutic testing. *Nat. Rev. Immunol.* **14**, 585–600. (doi:10.1038/nri3707)
 49. Martin B, Hirota K, Cua DJ, Stockinger B, Veldhoen M. 2009 Interleukin-17-producing gamma delta T cells selectively expand in response to pathogen products and environmental signals. *Immunity* **31**, 321–330. (doi:10.1016/j.immuni.2009.06.020)
 50. de Zoete MR, Keestra AM, Roszczenko P, van Putten JPM. 2010 Activation of human and chicken toll-like receptors by *Campylobacter* spp. *Infect. Immun.* **78**, 1229–1238. (doi:10.1128/iai.00897-09)
 51. Fellah JS, Jaffredo T, Nagy N, Dunon D. 2014 Development of the avian immune system. In *Avian immunology* (eds KA Schat, B Kasper, P Kaiser), pp. 45–63. London, UK: Elsevier.
 52. Pieper J, Methner U, Berndt A. 2008 Heterogeneity of avian gamma delta T cells. *Vet. Immunol. Immunopathol.* **124**, 241–252. (doi:10.1016/j.vetimm.2008.03.008)
 53. Wigley P, Barrow P, Schat KA. 2014 The avian reproduction immune system. In *Avian immunology* (eds KA Schat, B Kasper, P Kaiser), pp. 265–274. London, UK: Elsevier.
 54. Manel N, Unutmaz D, Littman DR. 2008 The differentiation of human T(H)-17 cells requires transforming growth factor-beta and induction of the nuclear receptor ROR gamma t. *Nat. Immunol.* **9**, 641–649. (doi:10.1038/nri1610)
 55. Zhou LA, Ivanov II, Spolski R, Min R, Shenderov K, Egawa T, Levy DE, Leonard WJ, Littman DR. 2007 IL-6 programs TH-17 cell differentiation by promoting sequential engagement of the IL-21 and IL-23 pathways. *Nat. Immunol.* **8**, 967–974. (doi:10.1038/nri1488)
 56. Veldhoen M, Hocking RJ, Atkins CJ, Locksley RM, Stockinger B. 2006 TGF beta in the context of an

- inflammatory cytokine milieu supports de novo differentiation of IL-17-producing T cells. *Immunity* **24**, 179–189. (doi:10.1016/j.immuni.2006.01.001)
57. Travis MA, Sheppard D. 2014 TGF-beta activation and function in immunity. *Annu. Rev. Immunol.* **32**, 51–82. (doi:10.1146/annurev-immunol-032713-120257)
58. Bettelli E, Carrier YJ, Gao WD, Korn T, Strom TB, Oukka M, Weiner HL, Kuchroo VK. 2006 Reciprocal developmental pathways for the generation of pathogenic effector T(H)17 and regulatory T cells. *Nature* **441**, 235–238. (doi:10.1038/nature04753)
59. Yang XO *et al.* 2008 Molecular antagonism and plasticity of regulatory and inflammatory T cell programs. *Immunity* **29**, 44–56. (doi:10.1016/j.immuni.2008.05.007)
60. Pripp AH, Stanisic M. 2014 The correlation between pro- and anti-inflammatory cytokines in chronic subdural hematoma patients assessed with factor analysis. *PLoS ONE* **9**, e90149. (doi:10.1371/journal.pone.0090149)
61. Grace JB, Bollen KA. 2005 Interpreting the results from multiple regression and structural equation models. *Bull. Ecol. Soc. Am.* **86**, 283–295. (doi:10.1890/0012-9623(2005)86[283:ITRFMR]2.0.CO;2)