Integrated signaling pathway and gene expression regulatory model to dissect dynamics of *Escherichia coli* challenged mammary epithelial cells

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**Abstract**

Cells transform external stimuli, through the activation of signaling pathways, which in turn activate gene regulatory networks, in gene expression. As more omics data are generated from experiments, eliciting the integrated relationship between the external stimuli, the signaling process in the cell and the subsequent gene expression is a major challenge in systems biology. The complex system of non-linear dynamic protein interactions in signaling pathways and gene networks regulates gene expression. The complexity and non-linear aspects have resulted in the study of the signaling pathway or the gene network regulation in isolation. However, this limits the analysis of the interaction between the two components and the identification of the source of the mechanism differentiating the gene expression profiles. Here, we present a study of a model of the combined signaling pathway and gene network to highlight the importance of integrated modeling.

Based on the experimental findings we developed a compartmental model and conducted several simulation experiments. The model simulates the mRNA expression of three different cytokines (*RANTES, IL8* and *TNFα*) regulated by the transcription factor NFκB in mammary epithelial cells challenged with *E. coli*. The analysis of the gene network regulation identifies a lack of robustness and therefore sensitivity for the transcription factor regulation. However, analysis of the integrated signaling and gene network regulation model reveals distinctly different underlying mechanisms in the signaling pathway responsible for the variation between the three cytokine’s mRNA expression levels. Our key findings reveal the importance of integrating the signaling pathway and gene expression dynamics in modeling. Modeling infers valid research questions which need to be verified experimentally and can assist in the design of future biological experiments.

**Keywords:**
Integrated network
Signaling pathway
Gene regulatory network
ODE
NFκB
Mastitis

1. Introduction

Gene expression is the result of the perturbation of a hierarchically organized and tightly controlled network of interacting elements in signaling pathways and gene regulation networks in the cell. These large number of interacting biochemical reactions show the emergent properties such as homeostasis and robustness with respect to perturbations (Ling et al., 2013). As more interactions between these signaling elements are identified, it becomes clear that signaling does not necessarily occur through parallel, linearly independent processes (Hornberg et al., 2006). Interactions can occur at many hierarchical levels and signaling proteins can influence gene network regulation, which leads to complex behavior (Bhalla and Iyengar, 1999). To understand these properties we need to study the system rather than the individual components using computational, mathematical techniques and biological knowledge, a systems biology approach (Suresh Babu et al., 2006). Ordinary differential equations (ODEs) are the preferred technique for modeling the dynamics of quantitative and qualitative aspects of signaling pathways and gene network regulation over time (de Jong and Ropers, 2006).

External stimuli to cells activate signal transduction pathways to initiate transcription-factor (TF) driven gene expression in gene regulatory networks. Transcription factors are often pleiotropic and involved in gene expression profiles of multiple genes and
therefore multiple biological processes and phenotypes. Due to complexities involved in intertwined signaling processes, the mathematical studies of signaling pathways (e.g., Chen et al., 2009; Goldstein et al., 2004; Suresh Babu et al., 2006; Vera et al., 2008; Vera et al., 2007; Wolkenhauer et al., 2005) and gene regulatory pathways (e.g., Schlitt et al., 2007; Xie et al., 2007) are conducted separately even though these pathways often have interdependent interactions that significantly affect the inter- and intra-cellular functionalities. In several cases the gene expression is influenced by the dynamics of the translocation of the transcription factor to the nucleus (Hoffmann et al., 2002; Sillitoe et al., 2007). Altering the dynamics of the signaling pathway influences the gene network regulation and therefore the gene expression profile which result in altered protein production and phenotype. Therefore, understanding the interaction between the signaling and gene network regulation will improve our understanding of the gene expression profiles and the underlying dynamics.

Here, we integrate the modeling and analysis of the signaling pathway and gene regulatory network. We show that the origins of underlying dynamics differentiating the cytokine's gene expression following a perturbation can be found in the signaling pathway but not in the gene regulatory network. To this end, we use cytokine mRNA expression profiles in bovine mammary epithelial cells. Mammary epithelial cells invoke the immune response in mastitis. Mastitis is the result of an inflammatory event in the mammary gland usually caused by a variety of bacteria. Bovine mastitis is one of the major diseases in the dairy industry worldwide and causes distress for the animal (De Ketelaere et al., 2006). The economic impact leads to a worldwide cost of US $25 billion per annum (Parke et al., 2005). In humans, mastitis is associated with the increased transmission of bacterial infections (Wang et al., 2007) and human immunodeficiency virus (HIV) passing from mother to child (John et al., 2001).

1.1. Cytokine mRNA expressions in mammary epithelial cells

Gene expression studies of mammary epithelial cells identified pro-inflammatory cytokines such as RANTES, IL8 and TNFa, expressed at a higher level in mammary epithelial cells challenged by Escherichia coli (Griesbeck-Zilch et al., 2008; Lahouassa et al., 2007; Lutzow et al., 2008; Pareek et al., 2005; Tao et al., 2007). Previously, variations in mRNA expressions of the cytokines, such as RANTES, IL8 and TNFa in mastitis have been reported (De Schepper et al., 2008; Griesbeck-Zilch et al., 2009; Rainard et al., 2006). The precise regulation of cytokine expression is essential for the regulation of the response to the infection. It is not completely understood how the cytokine expression is regulated.

1.2. Toll like receptor signaling

In mammary epithelial cells challenged with E. coli the toll like signaling activates the translocation of NFkB to the nucleus, which in turn initiates cytokine expression (Bannerman et al., 2004). Briefly, on the membrane of the epithelial cells, toll like receptors (TLR) recognizes the E. coli bacteria because of the molecular pattern, the endotoxin lipopolysaccharide (LPS) on the bacterial wall (Kawai and Akira, 2006) (Fig. 1). TLRs are the primary line of defense against invading pathogens (Doyle and O’Neill, 2006) initiating the toll like receptor signaling (Petzl et al., 2008). Experimental studies by Werner et al. showed that E. coli bacteria, which engages TLR4, elicit a small increase in IKK activity in the first 30 min, followed by a larger increase between 45 and 90 min, attenuating in the late phase (Werner et al., 2005). This allows for stimulus specific signal processing of NFkB regulated genes such as the cytokines studied in this paper (Hoffmann et al., 2006; Werner et al., 2005). The TLR signaling pathway therefore triggers gene expression initiating inflammatory and immune responses in the fight against E. coli infection (Akira et al., 2006) by activating the translocation of nuclear factor-kappa-B (NFkB) transcription factor to the nucleus (Strandberg et al., 2005) (Fig. 1). NFkB is a principal transcription factor in mammalian signaling (Cheong et al., 2008) and has been recognized as the ‘master switch’ in regulating the expression of various cytokines (Hayden et al., 2006). NFkB translocation does not require protein synthesis for its activation, allowing for fast reaction, within minutes, to inflammation (Hoffmann and Baltimore, 2006). The pattern and the timing of the translocation of NFkB to the nucleus lead to specific transcriptional outputs in NFkB regulated genes (Sillitoe et al., 2007).

1.3. NFkB signaling

In the cytoplasm NFkB is inactive as an IxB–NFkB heterodimer (Fig. 1). NFkB activity is largely controlled by three IxB isoforms, IxBa, IxBB and IxBC, which bind to NFkB in the cytoplasm preventing transport of NFkB to the nucleus (Hoffmann et al., 2002). TLRs send signals to IxB kinase (IKK) and IKK phosphor-ylates IxB, which results in degradation of the IxB-NFkB heterodimer and free NFkB. NFkB can then translocate to the nucleus and bind to DNA to function as a transcription factor for a large number of genes. In addition, NFkB initiates IxB transcription, which therefore acts as a strong negative feedback loop in NFkB activity (Fig. 1). Negative feedback loops can provide stability, linearity, and influence the frequency response or change the response into step response simulating an on/off switch.

Fig. 1. Conceptual model of TLR-IKK-NFkB signaling. The TLR receptor on the cell membrane recognizes the bacterial challenge. The signaling pathway activates the kinase IKK which breaks the IxB–NFkB dimer. As a result, the transcription factor NFkB translocates to the nucleus initiating gene expression. Among the genes expressed are the IxB isoforms (IxBa, IxBB, IxBC) which bind with NFkB in the cytoplasm to prevent translocation of NFkB to the nucleus. This process creates a negative feedback loop for the translocation of NFkB to the nucleus. For clarity, only IxB is shown in the picture representing three isoforms. In the model all three isoforms IxBa, IxBB and IxBC are included as individual reactions. In addition the cytokines initiated by the transcription factor NFkB studied in this work, RANTES, IL8 and TNFa are shown.
(Brandman and Meyer, 2008). IkBα is also induced by NFκB, with a delay relative to IkBα. The two feedback loops are in anti-phase and the role of IkBα is to dampen IkBα mediated oscillations during long lasting NFκB activity (Kearns and Hoffmann, 2008).

Stability of the NFκB response is essential for the regulation of gene expression. Several diseases, including diabetics (Bragt et al., 2009) cancer and chronic inflammation (Fraser, 2008) have been related to the impairment of the NFκB regulation. In addition, immune responses such as inflammation, cell proliferation, apoptosis (Viator et al., 2005) and milk protein levels (Connelly et al., 2010) are regulated by NFκB. NFκB levels as result of chronic mastitis were raised in milk (Boulanger et al., 2003). Targeted inhibition of NFκB signaling reduced milk loss and apoptotic signaling, which are of great concern during mastitis (Connelly et al., 2010). However, lack of regulation of the NFκB response can lead to severe diseases such as sepsis (Liew et al., 2005). Therefore, intricate knowledge of the NFκB regulation and the effect on the cytokine gene expression is of importance for the treatment and understanding of mastitis.

1.4. Mathematical models

Several models using ODEs to simulate the TLR–IKK–NFκB signaling have been published (Covert et al., 2005; Hoffmann et al., 2002; Kearns et al., 2006; Lipniacki et al., 2004; Werner et al., 2005). These models have played a critical role in understanding the innate immune response aspects in TLR–IKK–NFκB signaling (Hoffmann and Baltimore, 2006) because the dynamics of biological networks are often difficult to identify with in vivo or in vitro experiments (Thakar et al., 2007). For example, some signaling pathways encode information not just as protein concentrations or location, but via temporal changes in the dynamics of those concentrations (Kell, 2005; Nelson et al., 2004). In these cases, an in silico model can provide additional insights into the dynamics of the network.

Sensitivity analysis can be used to analyze the role of signaling proteins, identify potential drug targets and plan future experiments. For instance, the in silico simulation of NFκB pathway dynamics as a result of inhibitor drugs indicated the potential for inhibition of upstream events with low drug concentrations (Sung et al., 2004). Total inhibition of proteins can be modeled with knockout simulations. Biological experiments cannot always simulate knockout due to lethality, cost and ethical considerations. In silico knock out simulations are therefore a good alternative to investigate the influence of specific model components.

Robustness analysis can simulate the effect of the change in bacterial load, e.g., as a result of milking, over time and investigate the effect on the cytokine expression levels. While mathematical models are informative, model development can be time consuming and costly. One way to reduce time and experimental cost is to use a modular approach, extending an existing model. Werner et al. combined the knowledge of the TLR–IKK–NFκB signaling models by Hoffmann et al. (2002) and Kearns et al. (2006) and demonstrated that experimental IKK activity profiles of E. coli infection can be used to explain the effect of the feedback regulation of the two out of phase feedback loops established by the three IkB isofoms on the NFκB activity with an in silico model (Werner et al., 2005). The patterns and timing of the translocation of NFκB lead to different transcriptional outputs in NFκB regulated genes (Sillitoe et al., 2007) and different subsets of NFκB target genes are activated by the changes in the time-dependent kinetic profiles of NFκB signaling (Vanden Berghe et al., 2006). However, this model does not investigate the relationships between the mechanisms in the signaling pathway and the NFκB dynamics which result in the gene expression regulated by NFκB.

2. Objectives

The goal of this research is to demonstrate the necessity for integrated modeling of signaling and gene network regulation when studying cellular behaviors. We examine this paradigm with the analysis of an integrated model for the signaling pathway and gene regulatory network of mammary epithelial cells challenged by E. coli. To this end, we add a model of the gene network regulation of cytokine mRNA expression in mammary epithelial cells to the well-studied TLR–IKK–NFκB signaling pathway model developed by Werner et al. (2005) The developed model facilitates the investigation of the relationship between the signaling pathway variation and gene expression.

First, we discuss the model for the TLR–IKK–NFκB signaling pathway and incorporation of new ODEs accounting for the gene regulatory network for cytokine mRNA expression to form the integrated model, followed by the description of parameter estimation. Secondly, we perform sensitivity analysis and in silico knockout simulations to identify the mechanisms influencing the gene expression. Thirdly, we investigate the robustness of the gene network regulation, especially the influence of the variation of the transcription factor NFκB time profiles on the cytokine expression.

3. Model development and analysis

The need for an integrated model presented in this study is shown with the analyses of an integrated signaling and gene network regulation model of cytokine expression in primary bovine mammary epithelial cells challenged with LPS. The results are organized as follows: we develop the integrated model, and with the integrated model we perform sensitivity analysis and analyze in silico knockout models to show some of the mechanisms underlying the gene expression and the effect of possible drug targets. We then use robustness analysis to look at the effect of the perturbation of the model input, simulating a variation in bacterial load, on the cytokine expression.

3.1. Integrating the signaling pathway and gene regulation network

Based on the conceptual model as shown in Fig. 1, we developed the model for the TLR–IKK–NFκB signaling pathway and gene regulation network for cytokine expression. The activation of the TLR–IKK–NFκB signaling and the translocation of NFκB from the cytoplasm to the nucleus was modeled with 24 differential equations by Werner et al. (2005). The translocation of NFκB is the output of the signaling pathway and the input, the transcription factor, for the gene regulation network. In order to develop the combined model we extended the model with the gene regulation network. The gene network regulation is modeled with ODEs representing the mRNA expressions of RANTES, IL8 and TNFα as a result of the translocation of the transcription factor NFκB into the nucleus.

The reactions in the model are formulated as uni-, bi- and tri-molecular processes according to the law of mass action. The model is divided in two compartments, cytoplasm and nucleus. Compartmentalization is achieved by representing a single protein as multiple species, one for each compartment. Protein transport is modeled as the movement of species between the compartments with first order kinetics making the process computationally tractable. The model input is represented with a piecewise linear function representing IKK stimulation (Fig. S3 in the Supplement) (Werner et al., 2005).

The concentration of cytokine mRNA can be described by the difference between the mRNA synthesis (rsr_x) and degradation (d_n_x) (Eqs. (1)–(6)).
factors (with xenity, such as cooperative binding and multiple transcription rate.

mRNA (transcription factor NF

In Fig. 2, the model simulation with the estimated parameters are shown with the three experimental values (cells challenged with LPS from cow 1, cow 2, and cow 3, see Supplement

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
<th>Units</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>d_n_r</td>
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<td>min^{-1}</td>
<td>Degradation RANTES mRNA induced</td>
</tr>
<tr>
<td>rtf_r</td>
<td>5.28555</td>
<td>μM^{-2} min^{-1}</td>
<td>mRNA synthesis</td>
</tr>
<tr>
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<td>Degradation IL8 mRNA synthesis</td>
</tr>
<tr>
<td>rtf_8n</td>
<td>5.28555</td>
<td>μM^{-2} min^{-1}</td>
<td>NFkB induced IL8 mRNA synthesis</td>
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<td>d_n_TNFa</td>
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<td>min^{-1}</td>
<td>Degradation TNFα mRNA synthesis</td>
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<td>1.67</td>
<td>μM^{-2} min^{-1}</td>
<td>NFkB induced TNFα mRNA synthesis</td>
</tr>
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3.2. Gene network regulation parameter estimation

In Fig. 2, the model simulation with the estimated parameters are shown with the three experimental values (cells challenged with LPS from cow 1, cow 2, and cow 3, see Supplement

Section S1 for a detailed description of the experiment). The input function is a piecewise linear function of IKK and NFκB expression level for RANTES and IL8 than predicted by the model. Individual differences between cows can result in individual differences of cytokine expression levels and, as a result, mastitis resistance. Genetically determined differential expression levels of RANTES mRNA to pathogens between mastitis resistant and non resistant cows have been indicated earlier (Griesbeck-Zilch et al., 2009). The RANTES expression level difference between cows has been indicated as a selection option for mastitis resistant animals. However, the trend of the expression levels is the same and therefore the model can supply qualitative information on the underlying kinetics of the mRNA expression levels.

The model allows us then to explore the influence of the dynamics of TLR–IKK–NFκB signaling and NFκB gene network regulation on the mRNA expression levels of the cytokines.
3.3. Sensitivity analysis

3.3.1. Sensitivity for initial species values

Sensitivities are calculated using time independent sensitivity as described in Section 4.3.2 (Eq. (12)). Initial species values and parameters that influenced the mRNA concentration over the 360 min simulation of the integrated model are ranked to identify the highest sensitivity (Supplement Table S1). The cytokines RANTES, IL8, TNFa are predominantly sensitive to the changes in the initial species values of IkBaIKKNFkB. The sensitivity for TNFa increases towards the end of the simulation period (Supplement Fig. S4). Nuclear and cytoplasmic NFkB are also sensitive for initial species values of IkBaIKKNFkB. However, while RANTES and IL8 have similar top 4 rankings to nuclear NFkB (NFkBn), TNFa follows cytoplasmic NFkB indicating a difference between the three cytokines and the need for the inclusion of the signaling pathway to elicit cytokine specific sensitivities.

3.3.2. Sensitivity analysis for gene network regulation parameters

Time independent sensitivity (Section 4.3.2, Eq. (12)) for degradation parameters is higher than sensitivity for changes in NFkB transcription factor induced synthesis for each cytokine (Fig. 3).

Time dependent sensitivity (Section 4.3.1, Eq. (11)), the study of the effect of sensitivity over the simulation time, on the variation in the mRNA concentration indicates the largest variation in TNFa mRNA degradation parameter (Fig. 4).

A plot of the change in concentration over 360 min of simulation time for the changes of the synthesis and degradation

![Fig. 3. Time independent sensitivities of the model parameters. Time independent sensitivities of the model parameters for degradation and synthesis of the cytokine mRNA expression levels with parameter increase or decrease of 40%. Model predictions with the estimated parameters were compared with model predictions of parameters increased or decreased by 40% and time independent sensitivity calculated. (A) TNFa shows the highest sensitivity of the three cytokines. TNFa is more sensitive to changes in parameters for degradation than NFkB induced synthesis. (B) The expression of IL8 shows a higher sensitivity to degradation than to NFkB induced synthesis of mRNA, while in RANTES changes in either parameter have a similar influence on the expression levels.](#)

![Fig. 4. Time dependent sensitivities of model parameters for degradation and synthesis of mRNA cytokine expression levels. Time dependent parameter sensitivities for degradation (d_n) (A, C, and E) and synthesis (r sr) (B, D, and F) of mRNA cytokine expression levels were calculated for parameter changes from −40% to +40% and the gradual change over the range is shown in the shaded areas. Sensitivity as result of changes in degradation parameters increases over model simulation time (A, C, and E), while sensitivity as result of change in synthesis parameters stays constant (B, D, and F). TNFa shows the largest variation in model sensitivity for the degradation parameter changes and the highest sensitivity values for the model (E).](#)
parameters clearly shows a changing influence over time on the cytokine mRNA concentration levels as a result of the parameter changes (Fig. 5).

**RANTES** degradation parameters have a small influence over this range on the concentration. The influence increases toward 360 min simulation predictions, while synthesis rates have larger but stable influence from 100 min onward (Fig. 5A and B). For the IL8 mRNA cytokine, the changes in degradation rate increase their influence toward 360 min, while the changes in synthesis parameter show a stable influence from 100 min onward (Fig. 5C and D). **TNFα** synthesis and degradation parameters have a large influence at 90 min while the parameter changes have less influence on the changes in concentration levels at 360 min (Fig. 5E and F). This indicates a difference in sensitivity for synthesis and degradation parameters for each cytokine over time, however, parameter changes do not change the trend.

### 3.3.3. Time independent sensitivity in signaling pathway

Time independent sensitivity analysis (Section 4.3.2) is used to rank the cytokine sensitivity for the large number of parameters in the signaling pathway (Fig. 6, Supplement Table S2).

The three most sensitive parameters identified in the time independent sensitivity analysis (Section 4.3.2, Eq. (12)) that influence the total mRNA concentration are analyzed in depth with time dependent sensitivity analysis.

Protein synthesis of IkBα (rd_a), protein degradation of IkBα (pd_c_3ain) influencing NFκB and IKK concentration and IkBα mRNA degradation and synthesis are the major sensitivities in the model for IL8 and TNFα mRNA expression (Supplement Table S2). **TNFα** shows the highest sensitivity to the parameter for protein degradation of IkBαIKKNFκB (pd_c_3ain). **TNFα** also showed a high sensitivity for the initial value of this complex. Earlier sensitivity studies of NFκB signaling as a result of TNFα challenge have identified parameters influencing these proteins most sensitive in the NFκB signaling process (Ihekwaba et al., 2007; Yue et al., 2008). However, **RANTES** mRNA expression shows the largest sensitivity to NFκB import (in_n) into the nucleus followed by protein degradation of IkBα in the cytoplasm and transport of IkBα into the nucleus. RANTES is a late gene that is activated only after prolonged exposure to NFκB in TNFα challenges (Ting et al., 2002). Therefore, finding the sensitivity to the parameter influencing nuclear NFκB import for mRNA expression of RANTES is in line with prior biological knowledge.

### 3.3.4. Time dependent sensitivity in signaling pathway

When the sensitivity (Section 4.3.1, Eq. (11)) of the identified parameters over time is plotted for the three species with the different parameters, a distinct difference for the time of the highest sensitivity between the cytokines **TNFα**, **RANTES** and IL8 mRNA is noticed (Fig. 7).
RANTES is sensitive for the parameter change of parameter representing the transport of NFκB to the nucleus during the entire simulation time (Fig. 7A). This indicates sensitivity to parameter changes influencing exposure of NFκB over the entire simulation. IL8 showed a change in sensitivity from 180 min onward (Fig. 7B) while TNFα showed sensitivity early in the simulation (Fig. 7C). It is clear that each cytokine expression levels is sensitive for changes in different components of the signaling pathway, something that would not have been possible with the study of the gene network in isolation. Neither would the parameter sensitivity in the signaling pathway of NFκB been able to identify the mechanism for the sensitivity in the cytokine expressions. This will become clearer with the study of parameter sensitivity of NFκB and the lack of the relationship between the ratio of nuclear and cytoplasmic NFκB and the cytokine expression level.

3.3.5. Comparison of signaling pathway parameter sensitivity for cytokines and nuclear and cytoplasmic NFκB

We compared the results of the time independent sensitivity ranking for the three cytokines with the time independent sensitivity ranking for nuclear and cytoplasmic NFκB (Supplement Table S2). The top three time independent sensitivity parameter rankings for nuclear NFκB is the same as the ranking for IL8 mRNA. Cytoplasmic NFκB is similar to TNFα mRNA and the ranking for RANTES mRNA differs from both cytoplasmic and nuclear NFκB. While IL8 and nuclear NFκB are sensitive to IkκBα mRNA degradation rate changes, TNFα and cytoplasmic NFκB are sensitive to protein degradation of IkκNfκBαIkκBα, releasing IkκBα. Although the proteins located in the gene network and signaling pathway respectively, the sensitivity indicates a high sensitivity for the role of the negative feedback loop IkκBα provides for these proteins. However, RANTES showed sensitivity for the import of NFκB into the nucleus. Interaction between IkκBα and IkκBβ has been indicated to be responsible for the translocation of NFκB from the cytoplasm to the nucleus. Between the three cytokines RANTES shows the highest ranking of parameters related to IkκBα (Supplement Table S2, RANTES 8th highest, with IL8 17th and TNFα 13th). Indicating that RANTES is more sensitive to the influence of the interaction between IkκBα and IkκBβ and the influence of IkκBβ on the regulation of the negative feedback loop. There is therefore a distinct difference in the underlying mechanistic, originating in the signaling pathway, responsible for the variation in expression levels between the three cytokines. This cannot be explained by the nuclear or cytoplasmic NFκB sensitivity.

While one of the cytokines shows sensitivity similar to the nuclear NFκB and another to the cytoplasmic NFκB would it be possible to use the ratio of nuclear and cytoplasmic NFκB to identify the cytokine expression variation? The change in the ratio of NFκB in the cytoplasm and nucleus (NFκB:NFκBn) was plotted over time for the range of −40% to +40% changes of the sensitive parameters identified above (Fig. 8).

RANTES shows sensitivity to change in translocation of NFκB from cytoplasm to the nucleus (in n,over the 360 min (Fig. 8A). A variation in the NFκB:NFκBn ratio can also be seen with the changes of the parameter values for the translocation of NFκB to the nucleus (Fig. 8A) and the change in the concentration (Fig. 9A). In the first 50 min, the ratio varies with varying parameter values (Fig. 8A) and the sensitivity is high while there is no change in concentration (Fig. 9A).

Sensitivity then drops, however, the variation in sensitivity is the same while the variation in the ratio increases and then becomes stable and the variation in the concentration increases after 100 min and is stable for the remaining simulation period. However, there is no clear trend between the ratio and concentration of RANTES mRNA or the parameter sensitivity.

IL8 shows sensitivity to the change in ratio after 120 min as a result of change in the parameter for RNA degradation of IkκBα (rd_a) (Fig. 8B). During this period, the ratio of NFκB between cytoplasm and nucleus changes significant but is stable while the sensitivity of IL8 increases and therefore unlikely to be related.

TNFα mRNA expression is most sensitive in the early stages of the model simulation where there is a sharp decline in the NFκB concentration (Fig. 8A) and the sensitivity is high while there is no change in concentration (Fig. 9A).

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knockouts but return to wild type level at 360 min of simulation. (F) IκBα knockouts but return to wild type level at 360 min of simulation. (F) IκBα knockouts but return to wild type level at 360 min of simulation.

It is therefore clear that neither the sensitivity of NFκB nor the NFκB:IκBα ratio can be used to explain the mechanism of the sensitivity. It is important to note that the sensitivity of NFκB is higher than the sensitivity of IκBα. However, there is no variation in the NFκB:IκBα ratio as a result of the parameter change and it is therefore unlikely to cause of the sensitivity.

It is therefore clear that neither the sensitivity of NFκB nor the NFκB:IκBα ratio can be used to explain the mechanism of the regulation or predict the cytokine expression levels. This indicates that the signaling pathway and the gene network need to be studied together to elicit the effect of intervention on the variation in cytokine mRNA expression. In addition, the effect of intervention, such as knocking out a signaling protein, need to be studied for each cytokine and the change in expression of one cytokine cannot be used to predict the others.

3.4. In silico knockout experiments

Some pharmaceutical products knock out the component in the signaling pathway rather than changing the rate of production or degradation. Prior in silico simulations of the integrated model can reduce the number of biological targets to investigate because biological networks often show a degree of redundancy. Several examples exist where the manipulation of one enzyme does not lead to the desired effect because of the redundancy in the system (van Someren et al., 2002). Inhibition of IκBβ was considered as a likely anti-inflammatory therapy (Greten et al., 2007). However, further studies revealed that IκBβ inhibition increased LPS susceptibility caused by IκBβ/NFκB dependent signaling of the negative feedback function of the NFκB induced cytokines (Park et al., 2005).

Investigation of the model sensitivity revealed that the highest ranked parameters influence the components NFκB and IκBα. To a lesser extent, the model is sensitive to parameter changes influencing the concentrations of IκBε and IκBβ (Fig. 6). We thus investigated the effect of knockouts on the cytokine expression with in silico simulations in the integrated model.

3.4.1. NFκB knockout simulation

Blocking NFκB is often suggested as a treatment in infections but has been shown to lead to sepsis (Liew et al., 2005). In silico simulation of the knockout of NFκB reduced the concentration of IκBα and IκBε (Fig. 10D and E), while it increased the concentration of IκBβ (Fig. 10F).

Gene activation of IκBα and IκBε is NFκB dependent. RANTES, IL8 and TNFα show an immediate and sustained sharp decrease to 0 over the simulation period with NFκB knockout (Fig. 10C–E), which is to be expected.

Because we study the combined signaling and gene network regulation we can investigate the effects of varying other signaling proteins to vary, rather than block, NFκB level in the cytoplasm and nucleus and investigate redundancy.

3.4.2. IκBα knockout simulation

In silico simulation of the protein IκBα or the IκBα mRNA knockout models show a substantial increase of mRNA levels of

![Fig. 9. mRNA concentration over 360 min of simulation with parameter changes (−40 to +40). The variation of the simulation over time for RANTES, IL8 and TNFα for the parameters with the highest time independent sensitivity for each cytokine are shown. RANTES shows sensitivity for the transport of NFκB into the nucleus (in_n). IL8 is sensitive to the mRNA degradation of IκBα (rd_a), specifically after 180 min and TNFα is sensitive to IκBα mediated protein degradation (pd_c_3ain) in the initial phase of the model. In addition TNFα showed high sensitivity to the initial value of this protein.](image)

![Fig. 10. Simulations of the model with IκBα, IκBε mRNA, NFκB, IκBε, and the double knockout IκBα/IκBε. Knockout models were generated from the wild type model by setting the initial value and the rate of expression to zero. (A and B) RANTES and IL8 expression levels for the knockout models showed attenuation of the IκBα/IκBε double knockout and raised levels for IκBα knockouts, while NFκB knockouts reduced the levels. (C) TNFα expression is raised with IκBα/IκBε knockouts but returns to a stable level, while IκBα knockouts are raised but return to wild type level at 360 min and NFκB knockouts reduces the expression levels. (D) IκBε levels are raised by IκBα and IκBε knockout but return to wild type level at 360 min of simulation. (F) IκBβ levels were raised by the knockouts apart from the IκBε knockout. IκBε does not influence the level of IκBβ. Some knock outs IκBα, IκBε mRNA result in the same effect and do not show separately in the figures.](image)
IkBa −

NFkB −

IkBe −

IkBa/IkBe

model

RANTES, IL8 and TNFα (Fig. 10A–C). The ratio of NFkB in the cytoplasm and nucleus changes. The NFkB in the cytoplasm is reduced as a result of the knockout of IkBα, while the nuclear NFkB increases (Fig. 12D and E).

This is expected since IkBα acts as a negative feedback loop. In the cytoplasm IkBα associates with NFkB preventing the movement of NFkB from the cytoplasm to the nucleus. Reduction of IkBα therefore increases the movement of NFkB to the nucleus where cytokine expression is initiated (Fig. 1).

Both RANTES and IL8 show a prolonged increase in mRNA expression levels while the cytokine TNFα increases with a higher peak but reduces quickly to the wild type (WT) value when the NFkB reduces (Fig. 12A–C). The model therefore indicates a difference in kinetic response to the IkBα knockout between RANTES, IL8 and TNFα mRNA. Due to the difference in magnitude in the degradation parameters between RANTES and IL8 and TNFα this can be expected and is similar to earlier findings in the sensitivity analysis. The above sensitivity analysis (Section 3.3) identified different sensitivities for parameter changes for the individual cytokines. These parameters have different effects on the NFkB levels at different times and a different effect on the individual cytokines is to be expected. However, the mechanism is more complex and not influenced by IkBα alone. In addition to change in IkBα levels in the knockout simulation, the IKK concentration increases (Fig. 12F).

Increased concentration of IKK results in greater degradation of IkBα. The increased degradation then increases concentration of NFkB in the cytoplasm free to trans-locate to the nucleus. Although the reduction of NFkB in the nucleus shows a reduction of TNFα it does not explain the expression levels of RANTES or IL8.

The ratio of NFkB between cytoplasm and nucleus is more stable with the IkBα knockout explaining sustained gene expression of RANTES and IL8 but not TNFα, neither does it explain the difference in the return to the wild type for TNFα. Looking at the NFkB ratio between cytoplasm and nucleus (Fig. 11C), the IkBα knockout reduces the ratio, stabilizing after 70 min with increased concentration of NFkB in the nucleus and reduced concentration of NFkB in the cytoplasm. This confirms that the ratio of NFkB between cytoplasm and nucleus cannot be used to predict cytokine levels and shows a level of redundancy in the model. However it shows that the manipulation of components in the signaling pathway, other than NFkB, can be used to manipulate distinctly different outcomes for the individual cytokines.

3.4.3. Multiple knockout simulations

The parameter sensitivity analysis did not indicate high sensitivity for the parameters influencing IkBe and IkBa, however, knockout simulations show that the cytokine expression can be manipulated through multiple knockouts attenuating the effect of the IkBα knockout. In silico simulation of the IkBe and IkBa

Fig. 11. In silico knockout simulations and the effect on NFkB. (A) NFkB (B) Nuclear NFkB (C) NFkB ratio.

Fig. 12. Simulations of IkBα knockout and the effect on the different species. (A and B) RANTES and IL8 do not return to the model values, while TNFα (C) does return to the model values. While nuclear NFkB (D) is higher than the model value, cellular NFkB (E) is lower than the model value. IKK (F) is increased and stays at increased level.
knockout models show almost no change of mRNA levels of RANTES, IL8 and TNFα (Fig. 10A–C). The knockouts increase the IkBα concentration raising the peak minimally, returning back to the WT model values. NFkB in the cytoplasm and nucleus are marginally affected by the knockouts.

Simulating a double knockout of IkBα and IkBβ is not different from an IkBα knockout. However, a double knockout of IkBα and IkBβ attenuate the increase of NFkB in the nucleus but only marginally in the cytoplasm. The increase of NFkB in the nucleus results in a sustained elevation of RANTES, IL8 and TNFα expression (Fig. 10A–C). The cytokine TNFα stabilizes, while the IL8 and RANTES sustain increase. The knockout of the two negative feedback loops, IkBα and IkBβ, lead to an increase in the concentrations of nuclear NFkB that attenuate during the simulation period (Fig. 11B). There is no difference between IkBα knockout and IkBα IkBβ knockout with respect to NFkB in the cytoplasm, both reduce NFkB. The inhibitory role of IkBα and IkBβ in the form of negative feedback loops for the DNA-binding activity of NFkB of the TLR activated IkBs as result of TNFα simulation has been described earlier (Hoffmann et al., 2006). IkBα provides a negative feedback loop and is responsible for the post-inductional down regulation of NFkB activation. The delayed IkBα function is in an anti-phase to IkBα. It is proposed that the anti-phase regulation of IkBα stabilizes the NFkB activity without reducing the ability to terminate NFkB activation after the removal of the stimulus (Hoffmann et al., 2006). The two kinases, IkBα and IkBβ, work in tandem to rapidly repress NFkB translocation after TNFα stimulation. A similar effect as result of E. coli stimulation is seen in this study. Pharmaceutical targets knocking out IkBα and IkBβ would therefore not achieve a reduction in cytokine levels but an increase.

3.5. Robustness in cytokine expression levels

We conducted several simulation experiments to investigate the effect of the change of the bacterial load, the model input, on the gene expression. We used the simulation output for robustness analysis to identify the source, in both the signaling and gene network, for the change in gene expression. Bacterial loads vary in mastitis due to milking. The model input, expressed in the function ikkm (Fig. 13), represent IKK profiles and simulate external perturbations e.g., bacterial loads. If IKK is knocked out no immune response will be evoked and the bacterial infection will continue to increase.

Table 2
IKK input profiles generating 36 different nuclear NFkB time profiles (a = rising phase, b = first plateau, c = falling phase, x = concentration first plateau, y = concentration second plateau in Fig. 13).

<table>
<thead>
<tr>
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<th>a (min)</th>
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Rather than knocking out IKK it is therefore more informative to vary the IKK profiles (Fig. 13). Previously, simulations with varying IKK time profiles through variation in the ikkm function had identified 36 distinct nuclear NFkB time profiles (Werner et al., 2005). Nuclear NFkB is believed to be responsible for the cytokine expression level (Hayden et al., 2006) however, our sensitivity analysis showed that variation of NFkB does not influence the cytokines uniformly and is unique for each cytokine (Section 3.3). We therefore investigated the effect of the distinct nuclear NFkB time profiles on the cytokine expression levels with robustness analysis. Our simulation ran for 360 min. Increasing the range of values for the times (a–c in Fig. 13) or concentration levels (x and y in Fig. 13) creating more than 2500 different input profiles did not increase the coverage of the input space or the number of different nuclear NFkB time profiles. Thus we chose one time profile from each cluster (Table 2) to simulate distinctly different external perturbations. We then clustered the cytokine profiles, using k-means clustering in MATLAB (Figs. 14 and 15).

The in silico simulations identified different correlations between the nuclear NFkB profiles and the cytokine IL8 and RANTES and TNFα expression levels. In addition, the highest correlation between the cytokine concentration at 360 min and the...
model species was not with nuclear or cytoplasmic NFκB for any of the cytokines but with other species in the signaling pathway.

3.5.1. Robustness in gene network regulation

In the gene network, RANTES and IL8 showed the highest correlation with nuclear IκBαNFκB ($r = 0.96$ and 0.97, $p < 0.05$), while TNFα showed the highest correlation with IκBα mRNA ($r = 0.99$, $p < 0.05$). The correlation with nuclear NFκB was distinctly lower ($r = 0.80$ and 0.82, $p < 0.05$, Table 3).

Clustering the cytokine profiles and plotting the result together with the nuclear IκBαNFκB (Fig. 14) shows a distinct difference between RANTES and IL8 and TNFα. While TNFα will return to a stable state, RANTES and IL8 will either continue to increase, for those clusters with a value higher than the original model, or decrease if the value is lower or equal than the model value and therefore depending on the cluster.

3.5.2. Robustness in signaling pathway

In the signaling pathway IκBα showed the highest correlation with RANTES and IL8 ($r = 0.95$ and 0.97, $p < 0.05$), while IκBα showed the highest correlation with TNFα ($r = 0.99$, $p < 0.05$) (Fig. 15).

The correlation with cytoplasmic NFκB was distinctly lower ($r = 0.88$ and 0.91, $p < 0.05$, Table 4).

The results indicates the need for a combined signaling and gene network regulation model because the underlying correlation with the IκBα and IκBα mRNA showed the highest correlation with (r = 0.99, p < 0.05).

Because we looked at the integrated model of the signaling pathway and gene network regulation we can also speculate on additional mechanisms. With our sensitivity analysis of gene network regulation for the cytokines we compared the relative sensitivity of synthesis and degradation parameter changes (Fig. 4E and F) with the change in concentration values for TNFα mRNA expression (Fig. 5E and F). The sensitivity for parameter values increased, the concentration returned to the model levels, especially for variation in synthesis parameters at 360 min. It can be speculated that the robustness for variation in the synthesis
and degradation rates for TNFα mRNA in the mammary epithelial cells influences the robustness to variation in bacterial load at 360 min. It is thus possible that the robustness for synthesis and/or degradation parameter changes, rather that the robustness for change in NFκB regulation is the cause for robustness at 360 min observed in biological experiments. Changes in cytokine degradation rates as a result of disease have been seen earlier in other cells (Li and Bever, 2001) but need to be verified with biological experiments.

4. Methods

The experimental data used in this manuscript is described in the Supplementary information Section S1. In short, primary bovine mammary epithelial cells were taken from three cows and grown in culture. The cells were challenged with LPS simulating E. coli mammary infection. Gene expression, using Affymetrix bovine microarray time series, was measured and used for the model parameter estimation.

4.1. Mathematical modeling

The cytokine mRNA expression has been modeled in a modular way based on mass action kinetics using ODE. We based the signaling pathway model on a previously published model for TLR–IKK–NFκB signaling (Hoffmann et al., 2002; Werner et al., 2005). We implemented the ODE representing the mass balances of 24 components in the TLR–IKK–NFκB signaling in SBToolbox (Schmidt et al., 2006) in MATLAB (R2007a). The toolbox uses the MATLAB numerical differential equation solver, ode15s, to solve the equations. The RANTES, IL8 and TNFα mRNA expression levels are represented by 3 additional ODEs. The integrated model consists of 27 ODEs and 89 reaction rates. Initial values were calculated by running the model with a basal input level (0.1 μM) until no more changes in concentration could be detected. The MATLAB code is available upon request.

4.2. Parameter estimation

In order to verify the feasibility of the estimation of parameters for the differential equations with the measured data in our experiment, identifiability analysis is performed in SBToolbox (Schmidt et al., 2006) with the method explained by Jacquez and Greif (1985). Parameter estimation and simulations for this model were performed using SBToolbox (Schmidt et al., 2006) in MATLAB (R2007a). Data processing of the experimental data prior to parameter estimation is described in the Supplement (Section S2). In this paper a fast scatter search method is used (Egea et al., 2007; Rodriguez-Fernandez et al., 2006a). Fast scatter search is a combination of local and global optimization techniques, which aims to find the unknown parameters of the model that give the best goodness of fit to the experimental data.

We estimated the parameters and initial values to fit the converted mRNA expression levels of RANTES, IL8 and TNFα. Parameter estimations for models that include delay functions are slow and therefore we took the following steps: (i) the parameters for each cytokine were estimated in individual estimation runs, since identifiability indicates that the parameters between the cytokines were not correlated (see Supplement Section S2); (ii) the model was run with the estimated parameters for the individual cytokine and the model simulation results were compared with the experimental values for all three cytokines; (iii) iteratively the parameters were then estimated with the simulated values of the model that incorporated the estimated parameter values until no further optimization of the parameter values could be achieved with the fast scatter search (Rodriguez-Fernandez et al., 2006a); (iv) a final estimation round combining the parameters of the three ODEs was then performed with the fast scatter search algorithm as described in the methods until the value to satisfy the cost function was reached; and (v) the parameters were fine-tuned using manual tuning (Table 1, Section 3.1).

The parameter space is further explored in the sensitivity analysis as described in Section 3.3.

4.3. Sensitivity analysis

In local sensitivity analysis one parameter is changed at a time while the other parameter values are kept to their nominal values. The derivative vector, \( s_j(t) \), is calculated with Eqs. (10) and (11) to obtain a set of values for the finite parameter or initial value changes \( \delta \), which allows us to compare the sensitive regions of the output of interest \( X \) for each parameter or initial value. The output of interest \( X \) can be any observable such as the concentration at time \( t \) of component \( X_j \) or a combination of the concentrations of several components \( \{X_1, X_2, \ldots, X_d\} \) at time \( t \). \( s_j(t) \) stands for the incremental change in \( X \) due to the incremental change in the parameter \( \theta \) or initial value \( X(0) \).

\[
s_j(t) = \frac{\delta X(t)/X(t)}{\delta \theta_j/\theta_j}
\]

(10)

\[
s_j(t) = \frac{\delta X(t)/X(t)}{\delta X(0)/X(0)}
\]

(11)

4.3.1. Time dependent sensitivity

The local normalized sensitivity of \( s_j(t) \) is calculated for each time step \( t \) of the change in the \( j \)th component \( X_j(t) \) with respect to the change in the \( i \)th parameter \( \theta_i \) or initial value \( X(0) \) (Ihekwa et al., 2004). Because of our interest in the fit of the model, the component \( X \) was chosen as the concentration of the cytokine mRNA at time \( t \) in the 360 min simulation period and evaluated for the synthesis and degradation parameter changes.

An uniform distribution of parameter values was created by changing the value of each parameter with incremental steps of 10% from the model parameter and the corresponding change in mRNA cytokine levels recorded. The value \( s_j(t) \) will give a sensitivity index for each time step of the model simulation. However, time independent sensitivities would allow us to identify parameters with the

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<td>The highest correlation (( p &lt; 0.05 )) between the change cytokine concentration and the signaling (IkBa, IkBc) components at 360 min. The correlation is calculated for the results of the 36 input variations simulating different NFκB profiles.</td>
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<td>The highest correlation (( p &lt; 0.05 )) between the change cytokine concentration and the signaling (IkBa, IkBc) components at 360 min. The correlation is calculated for the results of the 36 input variations simulating different NFκB profiles.</td>
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And finally, the correlation is calculated for the results of the 36 input variations simulating different NFκB profiles.
highest influence on the cytokine mRNA levels for the total simulation period.

4.3.2. Time independent sensitivity
Integration of the sensitivities $s_{ij}(t)$ gives a time-independent value that allows ranking of the individual sensitivities of each cytokine as a result of parameter changes Eq. (12). This is the final time point and absolute value of the integrand prevents positive and negative values cancelling to zero under the integral $s_{ij}(t)$ (Chen et al., 2009). The quantity $S_j$ measures the change in the concentration of the $j$th component with respect to the $j$th parameter normalized by $T$ and therefore, captures variations in concentration level between parameter changes over time.

\[
s_{ij} = \frac{1}{T} \int_{0}^{T} |s_{ij}(t)| \, dt
\]

(12)

4.4. In silico knock out simulations
The in silico IκBα, NFκB, IκBε and IκBβ knockout models were generated from the wild type model by setting the initial value and the rate of expression of IκBα, NFκB, IκBε and IκBβ, respectively to zero.

4.5. Robustness analysis
In order to represent variation in bacterial load the input profile (a in Fig. 13) of the model is varied. Each profile contains a rising phase (a in Fig. 13), a first plateau (b in Fig. 13) and a second plateau (c in Fig. 13) with varying time levels (x and y in Fig. 13). To create a large set of diverse input profiles a computer program was developed. During a total simulation time of 360 min the duration of the rising phase was simulated for 0, 60, 120 and 240 min. The rise of the first plateau (x) was simulated with 0.04, 0.12, 0.34 and 1.01 μM. The duration of the first plateau (b) was 0, 5, 15, 30, 60 or 120 min. The falling phase (c) had duration of 0, 60, 120 or 240 min. The second plateau was equal or lower than the first plateau and varied between 0.01, 0.04, 0.12, 0.34 and 1.01 μM.

The results of the simulations are clustered for nuclear cytokine concentrations at 360 min using the $k$-means clustering algorithm implemented in Statistics Toolbox in MATLAB. Clusters are then compared with the components in the signaling pathway.

5. Conclusion
We set out to highlight the necessity for an integrated modeling approach of signaling pathway and gene networks. Gene expression, as a result of and external stimulus to cells, is investigated with a model combining the signaling pathway and gene network regulation. Signaling pathways and gene networks frequently have interdependent interactions that affect gene expression. Our integrated modular approach allows for investigation into a larger class of models without the need for extensive additional experiments, reducing cost and time.

We illustrated the value of the analysis of an integrated model with an example of the cytokines RANTES, IL8 and TNFα mRNA expression regulation as a result of TLR–IRκK–NFκB signaling and gene network regulation in E. coli challenge of mammary epithelial cells. It is known that TNFα, IL8 and RANTES are induced by the transcription factor NFκB as a result of TLR–IRκK–NFκB signaling. NFκB has been long been recognized as the ‘master switch’ in regulating cytokine expression (Hayden et al., 2006).

Analysis of the model confirmed that the cytokine expression is not robust for variation to the NFκB time profile but the mechanisms could not be identified with the gene network regulation model alone. However, the model identified signaling pathway components with higher sensitivity than NFκB in the regulation of the cytokine expression. In addition, time averaged sensitivity analysis of the integrated signaling and gene network regulation model identified sensitivity for different parameters and different times in the TLR–IRκK–NFκB signaling cascade for each individual cytokine. Intuitively one would have expected variation of nuclear NFκB could explain the variation in each of the cytokines. Our simulations and analysis have proven otherwise.

The challenges of modeling biological systems lie in the decision of the appropriate abstraction level to focus on (Szallasi et al., 2006). In “Therefore all models are wrong . . . some more than others” Wolkenhauer and Ullah explains that it is a means of reducing complexity that motivates modeling (Wolkenhauer and Ullah, 2007). Integrating signaling pathway and gene network regulation increases complexity, however, with our model, we have shown that the increase is not prohibitive and the analysis identifies emerging properties underlying the differentiation of gene expression.

The model facilitates the fine-tuning of the individual cytokine expression levels through the manipulation of the components in the signaling cascade and the identification of the effects on the other cytokines. These effects and timing need to be taken into consideration when developing drugs or planning future experiments. As the model analysis has indicated, the optimum experimental time differs between the cytokines. In addition drugs targeting these parameters will have a different effect over time on the cytokines. As the predictions are based on in silico models, the validity of the results should be experimentally tested, however, this is beyond the scope of the current work.

Acknowledgements
Nicoline Y. den Breems was funded by the Patrick Shannon Scholarship from LIC and the EU FP7 BIOMICS project (Grant No. 318202). Lan K. Nguyen was supported by funding provided by SFI (Grant No. 06/CE/B1129) and PRIMES (FP7-HEALTH-2011-278568) and University College Dublin’s Seed Funding program. We would like to thank Professor Hans-Martin Seyfert for the use of the microarray data.

Appendix A. Supplementary data
Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.biosystems.2014.09.011.

References
Boulanger, D., Bureau, F., Melotte, D., Mainil, J., Lekeux, P., 2003. Increased nuclear factor κB time profile with an example of the cytokines RANTES, IL8 and TNFα mRNA expression regulation as a result of TLR–IRκK–NFκB signaling and gene network regulation in E. coli challenge of mammary epithelial cells. It is known that TNFα, IL8 and RANTES are induced by the transcription factor NFκB as a result of TLR–IRκK–NFκB signaling. NFκB has been long been recognized as the ‘master switch’ in regulating cytokine expression (Hayden et al., 2006).


