Physiological response of mammary glands to *Escherichia coli* infection: A conflict between glucose need for milk production and immune response

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The mammary immune and physiological responses to distinct mammary-pathogenic *E. coli* (MPEC) strains were studied. One gland in each of ten cows were challenged intra-mammary and milk composition (lactose, fat, total protein, casein), biochemical (glucose, glucose-6-phosphate (Glu6P), oxalate, malate, lactate, pyruvate and citrate, malate and lactate dehydrogenases, lactate dehydrogenase (LDH), nitrite, lactic peroxidase, catalase, albumin, lactoferrin, immunoglobulin) and clotting parameters were followed for 35 days post-challenge. Challenge lead to clinical acute mastitis, with peak bacterial counts in milk at 16–24 h post-challenge. Biochemical and clotting parameters in milk reported were partially in accord with lipopolysaccharide-induced mastitis, but increased Glu6P and LDH activity and prolonged lactate dehydrogenase and Glu6P/Glu alterations were found. Some alterations measured in milk resolved within days after challenge, while others endured for above one month, regardless of bacterial clearance, and some reflected physiological responses to mastitis such as the balance between aerobic and anaerobic metabolism (citrate to lactate ratios). The results suggest that *E. coli* mastitis can be divided into two stages: an acute, clinical phase, as an immediate response to bacterial infection in the mammary gland, and a chronic phase, independent of bacteria clearance, in response to tissue damage caused during the acute phase.

*Escherichia coli* is one of the major pathogen causing acute mastitis in cattle, a disease of significant economic and animal welfare impact in dairy production worldwide. Upon entry into the mammary gland, mammary pathogenic strains of *E. coli* (MPEC) rapidly grow using milk as a nutrient source, in a manner highly dependent on the expression of the ferric-dicitrate system by these bacteria1–3. The resulting infection leads to an acute inflammation that causes a sharp decrease in milk yield and deep changes of milk properties and constituents4. Even after clinical healing and infection clearance in the mammary gland, indicated by cessation of bacterial secretion in the milk, milk yield and quality might remain impaired for a long period in a significant number of affected cows. The degree and length of the post-acute inflammation effect on udder health and milk quality vary greatly and may depend on the health status of the individual animal, time of diagnosis and treatment, as well as the infecting *E. coli* strain5.

Much has been studied about the immune and inflammatory reaction to *E. coli* in the mammary gland, yet, less is known about the physiological and biochemical changes that occur in the gland and in the milk during and following intra-mammary infections by *E. coli* bacteria. In a previous study by our group6, it was shown that under acute mammary inflammation induced by intra-mammary challenge of cows with *E. coli* lipopolysaccharide...

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(LPS), the passage of glucose-derived carbons shifts to the pentose phosphate pathway, leading to an epithelial cell metabolism shift to glycolysis at the expense of mitochondrial respiratory activity. A negative correlation between milk secretion and lactose, glucose and citrate concentrations and inflammation parameters was described. In parallel, the concentrations of glucose-6-phosphate (Glu6P), malate, oxaloacetae, lactate and pyruvate, and the activities of the respective enzymes glucose-6-phosphate dehydrogenase (Glu6PD), malate dehydrogenase (MDH) and lactate dehydrogenase (LDH) increased and, in general, positively correlated with inflammation parameters. A main difference between LPS and live bacteria intra-mammary challenge is the fact that bacteria represent a continuous stimulant of the mammary immune system due to bacteria replication in the gland, which may remain for days after the infection commences, whereas a single LPS injection would be a one-time stimulus. There is limited knowledge on the physiological and biochemical changes induced by live distinct mammary pathogenic *E. coli* and how these changes correlate to LPS-induced mastitis. Recently, an intra-mammary challenge experiment with MPEC strains (VL2874, VL2732 and P4) and with the non-mammary pathogenic *E. coli* strain K1 was conducted in order to study whether the dynamics and intensity of the immune response in bovine mammary glands is different and depends on the infecting *E. coli* strain. These strains were chosen for representing distinct presentations of *E. coli* mastitis: per-acute (VL2874), persistent (VL2732) and clinical (P4), respectively.

Cellular and chemokine responses and bacterial culture follow-up were performed for 35 days post-challenge (DPC). Cows challenged by any of the MPEC strains developed clinical acute mastitis, with peak bacterial counts in milk at 16–24 h post-challenge. During the 35 days of the study, differences related to the *E. coli* strains were found in the intensity and duration of the response in various of the parameters studied, namely the somatic cells count (SCC), leukocytes distribution, secreted cytokines (TNF-α, IL-6 and IL-17) and levels of membrane TLR4 on leukocytes in the milk.

The objective of the present work was to study the physiological and biochemical responses measured in the milk and mammary gland tissue of cows challenged with the above mentioned three distinct mammary-pathogenic strains of *E. coli*.

### Materials and Methods

#### Animals.

Ten Israeli Holstein cows at the Agricultural Research Organization (A.R.O.) dairy farm at lactations 1–6, 133–442 days in lactation, and with milk yield of 22–38.4 L/day were used for intra-mammary challenge with mammary pathogenic strains of *E. coli*. The cows were milked thrice daily (05:00, 13:00 and 20:00) in a dairy parlor equipped with an on-line computerized AfiFarm Herd Management data acquisition system, including the AfiLab milk analyzer, which provides on-line data on gross milk composition - fat, protein and lactose (Afmilk, Afikim, Israel; [http://www.afiMLK.com](http://www.afiMLK.com)). The average milk yield in the herd was >11,000 L during 305 days of lactation. Animals were fed a typical Israeli total mixed ration containing 65% concentrate and 35% forage (17% protein). Food was offered *ad lib* in mangers located in sheds. At the beginning of the study, all glands of the 10 cows were free of infection as tested by three consecutive bacteriological tests and had SCC <1 x 10⁵ cell/mL milk.

#### Bacteria challenge.

Three different *E. coli* strains were used for the intra-mammary challenge: VL2874 (isolated from per-acute mastitis), VL2732 (isolated from persistent mastitis) and P4 (widely used as model strain for mammary pathogenic *E. coli* isolated from clinical mastitis in the UK). Strains typing, phenotypic and genomic characteristics were extensively studied and described before. Bacteria were recovered from frozen stocks (kept at −80°C in a mixture of brain-heart infusion and 25% glycerol) on blood agar (Tryptose Blood Agar Base; Becton-Dickinson, Sparks, MD, USA, with 5% washed sheep erythrocytes) and incubated aerobically at 37°C overnight. For inoculum preparation, bacteria were harvested and washed in pyrogenic-free saline (PFS), suspended in PFS and stored at 4°C for 10 h. Aliquots were taken for colony forming units (CFU) count on blood agar. CFU number in the inoculum ranged from 10–30 CFU/mL. Final CFU number in the inoculum ranged from 10–30 CFU/mL, as determined from aliquots separated just prior to challenge. The small CFU counts in the inoculum aimed to better represent a natural infection in which it is expected that only a small number of bacteria enter the mammary gland and initiate the infection.

#### Study layout.

The 10 cows were challenged each in one gland, while all the other three glands served as control. Before challenge, milk yield, milk composition, SCC and leukocytes distribution were determined at ~2 days and on day 0, separately for the intended challenged gland and composite milk of the other three glands. At the day of challenge, after morning milking, a single gland (rear left or right in nine cows and front right in one cow) was infused via the teat canal with one of the *E. coli* strains (3–4 cows each). Clinical symptoms were assessed for the first 24 h, including every 4 h measurements of rectal temperature and presence of general signs of illness such as changes in respiratory rate, loss of appetite, diarrhea and dehydration. After challenge, milk yield was recorded separately for the challenged gland and the three control glands. Milk was sampled at: 4, 8, 12, and 16 h post-challenge, and then at 1, 2, 4, 7, 10, 14, 17, 21, 29, and 35 DPC. A week after the termination of the experiment (at day 45 PC) the cows were slaughtered in an abattoir. Post slaughter, teats were externally cleaned and disinfected with 70% alcohol, then the nipples and ~10 cm of parenchymal tissue above it were removed as sterile as possible, stored individually in sterile bags and placed on ice. Samples were transferred to the laboratory for histological and PCR analyses for the presence of the bacteria.

#### Histology.

Samples for histological analysis were fixed in neutral buffered 4% formaldehyde and embedded in paraffin. Sections were cut (4 µm) and stained with hematoxylin and eosin (H&E). Slides were viewed using a light microscope (Nikon Eclipse E600, Melville, NY, USA), and images were captured using the MagnaFire Digital...
Milk sample collection and analyses. For bacteriological tests, teats were cleaned, disinfected, and milk was dispensed, and 3 mL samples of milk were aseptically collected in sterile assay tubes. For the other tests, the infected glands and then composite milk of the other three glands were milked separately into containers, and milk volume was recorded and gently mixed and 0.5–1.0 L was taken for analysis as follows: SCC with the Fossomatic 360 (Foss Electric, Hillerod, Denmark) and gross milk composition - fat, protein, casein and lactose contents with the MilkoScan FT6000 (Foss Electric). These analyses were performed at the Israel Cattle Breeders Association Laboratory (Caesarea, Israel). Rennet clotting time (RCT) and curd firmness (CF) were tested using the Optigraph (Ysebaert, Frepillon, France) as previously described. From 15 TL (Gist-Brocades nv, Delft, The Netherlands) was diluted 1:100 from the stock solution and 0.5 mL/well were added to achieve clotting at about 900 s in non-infected milk. Tests were performed within 24 h after sample collection with milk stored at 4 °C.

The activity of malate dehydrogenase and lactate dehydrogenase in milk samples were performed. The final activity of diaphorase and the concentration of resazurin used for these enzyme-activity determinations were the same as used for the metabolite determinations. The basic reaction conditions for MDH activity in milk were as described by Silanikove et al. Differences in NADH concentration (between 10 and 1000 μM) at the linear stage of the reaction were divided by time and were converted to activity where 1 U of MDH will convert 1.0 μmol/L of oxaloacetate and NADH to L-malate and NAD+ per minute at pH 7.5 at 25°C. The reaction conditions for LDH activity were as described by Larsen. One international unit of LDH activity was defined as the amount of enzyme that catalyzes the conversion of pyruvate into lactate to generate 1.0 μmol/L of NAD+ per minute at 37°C.

Nitrite concentration was determined by the fluorometric assay with DAN reagent as described by Sonoda et al., after precipitating the casein by ultracentrifugation. Lactic peroxidase (LPO) assay was assayed by the oxidation of 2,2 V-azino-bis (3-ethylbenzothiazoline-6-sulfuric acid) diammonium salt, as described by Silanikove et al. Catalase activity was determined with the aid of fluorescent substrate as previously described. Additional sub-sets of milk samples were defatted under cold conditions and analyzed according to previously described procedures for albumin, lactoferrin and immunoglobulin type G (IgG) by ELISA assays.

Statistical analysis. Due to the low number of cows, and the lack of statistical differences among cows challenged with different bacteria strains in a preliminary statistical analysis (ANOVA, repeated measures), all cows were analyzed as a single group. The 10 cows were tested for 38 different variables during the time course of the experiment. Each cow was tested in 21 different time points, from day 0 to 35 days post-challenge during the study. The ProcMixed procedure of SAS (version 9.2, SAS Institute Inc., Cary, NC) for repeated measures in time, with cow controlled as random effect, was used for analysis with the general form: Dependent variable = time during the experiment + error. Where: time = 21 different time points from bacterial inoculation time to 35 days later. The dependent variables were: log SCC, milk (infected), milk (cow), fat, protein, % casein, lactose, RCT, CF, IgG, BSA, bovine lactoferrin, lactate, malate, lactate + malate, nitrite, LDH, glucose, Glu6P, Glu6PD, oxalate, lactoperoxidase, catalase, pyruvate and citrate. To compare levels within a variable, we ran the Bonferroni adjustment for multiple comparisons. Data are presented as means and SEM.

Ethics approval. Animal experimentation was approved by the Committee of Animal Experimentation of the A.R.O., the Volcani Center, Bet Dagan and followed the committee’s ethical guidelines (Permit no. 59315).
the challenged glands fully recovered during the study period, showing approximately 40–60% loss of milk yield compared to ~20% loss in some control glands, while in others a compensation (gain in milk yield) was recorded. Overall, cow’s milk yield decreased by 10–30% at the end of the study (Fig. 2). The loss of milk yield per average cow during the 35 days of the study was 336 L (32%). Milk yield of infected glands significantly correlated with casein, lactose, coagulation properties, citrate, and significantly negatively correlated with SCC on the cow level.

**Biochemical response to *E. coli* challenge.** No significant results were found between the distinct MPEC strains; therefore, the following results are presented and discussed for the three MPEC strains combined.

The statistical significance ($P$ values) of the ANOVA results for the Day effect, $R^2$ and % variance between cows from the overall variance in the trial are summarized in Table 1. Fat in milk decreased during the first 2 h and then significantly increased ($P<0.001$) up to 7–10 DPC and gradually returned to pre-challenged level only at 35 DPC (Fig. 3). A significant decrease ($P<0.001$) in total protein was already noticed at 6–8 h PC, after which total protein levels in milk increased up to a peak at 7–10 DPC, returning to pre-challenged levels at 35 DPC. In parallel,
% casein decreased significantly from ~78% to ~68% between 8–24 h PC, then gradually increased afterwards but remained significantly lower than pre-challenged levels until the end of the study (Fig. 3).

Milk coagulation properties assessed before challenge showed a mean RCT ~28 min and CF ~9 V. After challenge, milk coagulations properties were deeply affected and for the 14 DPC, none of the milk samples coagulated. From day 21, milk from 4–5 challenged glands coagulated but formed a weak coagulum, while the other cows’ infected glands milk did not coagulate until day 35 (Fig. 4).

Milk lactose of the infected glands decreased significantly (P < 0.05) at 12–16 h from ~50 g/L before E. coli inoculation to ~33 g/L. Only in 60% of the infected glands milk lactose increased gradually from 7–10 DPC up to pre-challenge level on days 28–35, while remaining low in the other glands (Fig. 5). The kinetics of changes in lactose concentration closely followed the kinetics of milk yield and the inflammatory response, as reflected by significant negative correlations with log SCC and the above-described inflammatory indices and oxidative-stress markers. A significant positive correlation was found between protein level and inflammation parameters, while a significant negative correlation was found between % casein and lactose and inflammation. Regarding clotting markers. A significant positive correlation was found between protein level and inflammation parameters, while significant negative correlations with log SCC and the above-described inflammatory indices and oxidative-stress markers were measured earlier, at ~12 h, whereas increase of malate was detected at 1 DPC. Lactate and malate levels started to elevate at ~12 h, peaked on 2–4 DPC and declined thereafter (Fig. 8). A sharp increase of lactate was measured earlier, at ~12 h, whereas increase of malate was detected at 1 DPC. Lactate and malate levels remained about 6 and 3 folds higher than pre-challenge levels until the end of the experiment, respectively.

| Variable          | R-Square (day) | P[F] | Variance between cows |
|-------------------|----------------|------|-----------------------|
| Log SCC           | 0.749          | <0.001 | NS                    |
| Milk (infected gland) L/d | 0.403          | <0.001 | NS                    |
| Milk (cow) L/d    | 0.121          | NS    | 15.2%                 |
| Fat (g/L)         | 0.170          | NS    | 18.8%                 |
| Protein (g/L)     | 0.509          | <0.001 | 29.3%                 |
| % Casein          | 0.399          | 0.008 | 76.1%                 |
| Lactose (g/L)     | 0.521          | <0.001 | 25.8%                 |
| RCT (sec)         | 0.755          | <0.001 | NS                    |
| CF (V)            | 0.630          | <0.001 | NS                    |
| IgG (mg/mL)       | 0.335          | <0.001 | NS                    |
| BSA (µM/mL)       | 0.499          | <0.001 | NS                    |
| Lactoferrin (µM/mL)| 0.551          | <0.001 | NS                    |
| Lactate (µM)      | 0.588          | <0.001 | NS                    |
| Malate (µM)       | 0.589          | <0.001 | NS                    |
| Nitrite (nM)      | 0.528          | <0.001 | NS                    |
| LDH (U/mL)        | 0.648          | <0.001 | NS                    |
| Glucose (µM)      | 0.228          | 0.016  | NS                    |
| Glu6P (µM)        | 0.392          | <0.001 | NS                    |
| Glu6PD (µM/mL)    | 0.362          | <0.001 | NS                    |
| Oxalate (mM)      | 0.215          | 0.06   | NS                    |
| Lactoperoxidase (U/mL)| 0.227    | 0.04   | NS                    |
| Catalase (U/mL)   | 0.267          | 0.007  | NS                    |
| Pyrovate (µM)     | 0.337          | 0.001  | NS                    |
| Citrate (mM)      | 0.318          | 0.004  | NS                    |

Table 1. The significance (P values) of the ANOVA results for the day effect and R² of the variables tested.

Abbreviations used: SCC – somatic cell count; RCT – rennet clotting time; CF – curd firmness; IgG – immunoglobulin G; BSA – bovine serum albumin; LDH – lactate dehydrogenase; Glu6P – glucose 6 phosphate; Glu6PD - glucose 6 phosphate dehydrogenase; NS = Variance between cows not significant (P > 0.05).
Pyruvate behaved differently, compared to the previous parameters. Pyruvate levels fluctuated during the first day post-challenge; gradually increasing until a peak >1400 µM at 7 DPC, after which pyruvate levels sharply declined to ~750 µM at 10 DPC. By the end of the study, pyruvate levels were lower than pre-challenge levels (~600 µM and ~900 µM, respectively) (Fig. 9). Citrate concentration dynamics mirrored that of lactate and malate above. Citrate levels sharply decreased at 16 h from 8.9 mM to lowest levels (3.8 mM) at 2 DPC, then gradually increased but remained lower on 35 DPC compared to pre-challenge levels (Fig. 10). Consequently, the ratio citrate/lactate + malate dropped sharply, starting at 12 h from ~17 to <1 at 2 DPC and then started to rise to the pre-challenged level ratio at 28 DPC (Fig. 11).

Catalase and lactoperoxidase activity, LDH, BSA and nitrite concentrations displayed similar patterns. Their levels started to elevate at ~12 h PC, peaked at 2–4 DPC and declined to about pre-challenged levels at the end of the study (Fig. 12A–E). These proteins, enzymes and ion which originate from blood, significantly positively correlated with SCC and milk protein, and significantly negatively correlated with infected glands and cow milk yield,
lactose, % casein and CF. Lactoferrin concentration increased at 16–24 h from ~300 to 1,600 µg/mL, peaked at 7 DPC and declined to the end of the study, but remained ~2 fold higher from the pre-challenge level (Fig. 12F). IgG concentration increased within 12 h from 0.2 to >2 mg/mL, peaked at 3 DPC >7, remained in this level up to 21 DPC and only then declined, but also remained ~2 fold higher from the pre-challenge level at 35 DPC (Fig. 12G).

Histology at 42 days post-challenge. Histological changes in infected and control glands were compared within each and among cows. Overall, the major differences in the control glands was the level of alveolar cuboidal epithelial structure in full lactating glands in the high yielding cows (Fig. 13A), and increased interlobular collagen rich areas with fibrous stroma and fat in low yielding cows (Fig. 13B). In challenged glands, regardless if bacteria were isolated on 35 DPC or not, increased interlobular collagen rich areas with fibrous stroma and fat were observed, and in the high yielding cows, normal alveolar structure. Moreover, in most challenged glands, leucocytes infiltration was observed, mainly inter-lobular mononuclear infiltration, and in two of the 10 glands PMN were observed within normal alveoli (Fig. 13C).

Discussion

_**E. coli** mastitis is characterized by acute inflammation, including local clinical signs of inflammation, such as udder swelling and pain, and sharp decrease in milk production along with significant changes in its composition, such as increased SCC and passage of blood elements into milk. Systemic signs, including increased body_
temperature, diarrhea and dehydration are sometimes also observed. *E. coli* growth in milk in the mammary gland is fast, and increased bacterial counts are detected as early as within 4 h from challenge even when a minimal inoculum is used to best represent natural infections (10–30 CFU in the present study). The present study focused on the physiological and biochemical changes in milk following challenge with three different MPEC strains isolated from distinct mastitis presentations (VL2874, VL2732 and P4). Although some significant differences were found in the inflammatory and immune responses in the mammary gland between these strains, the physiological and biochemical markers revealed only minor differences without statistical significance. Therefore, cows challenged with any of the three strains were combined in a single group for analysis.

Cows did not receive medication, thus bacteria clearing in milk was natural, starting on 7 DPC. In 50% of the glands, bacteria shedding was intermittent until the end of the study. Regardless of the clearing rate and bacteria numbers in milk, no significant correlation was found with all the parameters tested, suggesting that the recovery process depends mainly on the severity of the infection and the inflammatory response elicited, and possibly tissue damage, during the first hours or days after *E. coli* entry to the mammary gland.

Two clear infection stages were observed: 1: Initiation and establishment of the infection and consecutive inflammation, which started at 4–10 h PC and ended by 7–15 DPC; 2: Resolution of infection, as expressed by

![Figure 7. Oxalate in milk of glands infected with pathogenic *Escherichia coli*. Mean and SE of oxalate (mM) measured in the infected glands of 10 cows challenged with different mammary pathogenic strains of *E. coli*. * and ** significantly different from time 0 to 0.6 d and on 7 DPC (*P < 0.001).](image1)

![Figure 8. Lactate and malate in milk of glands infected with pathogenic *Escherichia coli*. Mean and SE of lactate (○) and malate (□) measured in the infected glands of 10 cows challenged with different mammary pathogenic strains of *E. coli*.](image2)
most of the parameters tested recovering to pre-challenge levels, although with different dynamics. A schematic presentation of the two-phase proposal for the development of E. coli mastitis is presented in Fig. 14. However, of note, SCC and milk yield at the infected glands and at the cow level was not restored until the end of the study at 35 DPC, similar to our observations in natural E. coli mastitis5.

Milk production in the mammary epithelial cells exists under osmotic balance between blood and the epithelial cells20. Acute signaling of E. coli LPS and immune cytokines like IL8, TNF and others occur within hours, starting a cascade of events, including infiltration of leucocytes, mainly PMN, and selectively opening of the tight junctions21 as has been showed in our earlier study4.

The shift of mammary epithelial cells metabolism to anaerobic glycolysis as a tradeoff between use of glucose to support lactose synthesis and liberation of glucose to support the immune system was thoroughly discussed by Silanikove et al.6, using E. coli LPS as a model for mammary gland infection. LPS is a major immunogenic determinant of E. coli and Gram-negative bacteria in general, but there are important differences between LPS inoculation vs. live E. coli bacteria challenge. First, a single inoculation of LPS leads to a single triggering event of the inflammatory response, whereas live bacteria replicate in the mammary gland and therefore represent a longer trigger for inflammation. Second, LPS by itself does not cause direct damage to the gland tissues whereas...

Figure 9. Pyruvate in milk of glands infected with pathogenic Escherichia coli. Mean and SE of pyruvate (μM) measured in the infected glands of 10 cows challenged with different mammary pathogenic strains of E. coli. *significantly different from time 0 to 5 d and on 20 DPC (P < 0.05).

Figure 10. Citrate in milk of glands infected with pathogenic Escherichia coli. Mean and SE of citrate (mM) measured in the infected glands of 10 cows challenged with different mammary pathogenic strains of E. coli. **significantly different from time 0 to 0.8–5.0 DPC (P < 0.001).
live bacteria have different virulence factors that can interact with and damage the mammary epithelial cells, leading to a longer recovery process. Third, the amount of LPS often used to induce mastitis is often higher than the amount of LPS found in a small bacterial inoculum, such as the one used here. Here we observed that the inflammation caused by \textit{E. coli} could be divided into two stages: an acute phase, with clinical symptoms and a chronic phase, independent of bacteria clearance and in response to the tissue damage caused in the first phase (Fig. 14). The present results suggest that the tight junctions in the mammary alveoli were “open” at 12–24 h PC. This is expressed by infiltration of glucose, IgG, BSA, nitrite and LDH from blood into the milk for 2–3 d after LPS inoculation\(^2\) and 6–10 d after live bacteria inoculation. However, the free transfer of the immune cells (SCC) remained for weeks, suggesting active cell infiltration.

Overall, the dynamics of biochemical parameters evaluated was similar to that of LPS induced mastitis,\(^6\) but levels achieved were either higher (e.g. glucose-6-phosphate and lactate dehydrogenase activity) or changes remained longer (e.g. lactate dehydrogenase, Glu6P/glucose). The present results are in agreement with those published by Moyes \textit{et al}.\(^2\) although most of the parameters tested in that study returned to the pre-challenge levels (\(\sim 20–40\) CFU of live \textit{E. coli} - Danish field isolate k2bh2) within 2–4 d. For instance, peak glucose and GLU6P, drop in lactose and increase in fat measured following intra-mammary infusion of \textit{E. coli} were quite similar to other findings\(^2\), but in the present study, fat increased after about 5–10 DPC, compared to 1–2 DPC\(^2\). The ratio citrate/lactate + malate, which represents the milk-reflected mitochondrial/cytosol metabolic index\(^1\), significantly decreased during the acute phase of inflammation as described in LPS induced mastitis. Here, however, this change remained for a longer period, and the ratio did not return to pre-challenge levels by the end of the study at 35 DPC. Cows tested 30 days after an episode of natural \textit{E. coli} mastitis (“post \textit{E. coli}”) showed similar results\(^2\).

Peak bacterial counts in milk were observed at 24 h\(^4\), following a steep decrease in citrate between 16 to 24 h after challenge and concomitant to a significant increase in lactoferrin in milk. The ability to assimilate iron from citrate is a key feature of MPEC\(^3\) and therefore it is possible that the exchange of citrate to lactoferrin as main iron chelator in milk contributes to reduce bacterial load in the mammary gland\(^2\).

Interestingly, regardless of bacteria presence in the challenged mammary glands, milk yield was lower than pre-challenge, the number of the immune cells remained high (\(>1 \times 10^6\) cell/mL) and the quality of the milk was low as expressed by impaired clotting parameters, even though fat and protein levels were actually higher than pre-challenge ones. This could be related to other milk constituents, but also to the quality of fat and protein fractions present in milk following inflammation, which deserve further investigation. Histology at 42 DPC allowed the observation of disrupted mammary gland tissues, affected alveolar cuboidal epithelial structure and increased interlobular collagen rich areas with fibrous stroma and fat. In most gland, leukocytes infiltration, mainly mononuclears among lobules and PMN within the normal alveoli was present. This correlates with the chronic, long-term effects reminiscent of mastitis caused by \textit{E. coli}, indicating tissue damage and the process of cleaning of damaged tissue, and which result in affected milk quality and yield\(^7\). Overall, the present study adds information about the physiological and biochemical changes observed in milk during and following acute intra-mammary inflammation induced by infection by \textit{E. coli} bacteria. Moreover, the present results highlight the shift between the acute phase and a chronic phase of \textit{E. coli} mastitis, the latter being independent of bacterial shedding in milk; and is therefore independent from an ongoing infection. This chronic phase of inflammation is characterized by continuous changes in milk properties for a long period after clinical recovery, and the present results may explain
Figure 12. (A–G) Physiological components in milk of glands infected with pathogenic *Escherichia coli*. Mean and SE of catalase (A), lactoperoxidase (B), lactate dehydrogenase (C), bovine serum albumin (D), nitrite (E), lactoferrin (F) and IgG (G) in the infected glands of 10 cows challenged with different mammary pathogenic strains of *E. coli*. Catalase and lactoperoxidase activity, LDH, BSA and nitrite concentrations displayed similar patterns. Their levels started to elevate at ~12 h PC, peaked at 2–4 DPC and declined to about pre-challenged levels at the end of the study (A–E). Lactoferrin concentration increased at 16–24 h, peaked at 7 DPC and declined to the end of the study, but remained ~2 fold higher from the pre-challenge level (F). IgG concentration increased within 12 h, peaked at 3 DPC >7, remained in this level up to 21 DPC and only then declined, but also remained ~2 fold higher from the pre-challenged level at 35 DPC (G). **significantly different (P < 0.001).
Figure 13. (A–C) Histological sections of udder tissues of glands infected with pathogenic *Escherichia coli*. Representative hematoxylin-eosin histological sections of control and infected glands of 10 cows challenged with different mammary pathogenic strains of *E. coli*. Note the level of alveolar cuboidal epithelial structure in full lactating glands in high yielding cows (A), and increased interlobular collagen rich fibrous stroma and fat in low yielding cows (B). In challenged glands, increased interlobular collagen rich areas with fibrous stroma and fat was observed, with mononuclear leucocytes among lobules and PMN within normal alveoli in two out of 10 glands (C). Images were captured using the MagnaFire Digital Camera controlled by NIS-Elements F3.0 software (https://www.microscope.healthcare.nikon.com/products/software).
the long-term effects on milk quality that have been previously described5. This study also emphasizes some differences between LPS and live bacteria-induced mastitis, which are important to be accounted when choosing the right model for mastitis research.

Conclusions
Escherichia coli challenge induces acute conversion of epithelial cells metabolism from principally mitochondrial-oxidative to principally cytosolic (glycolytic), which allows diversion of metabolic resources normally used to synthesize milk and to support the immune system. In turn, reduction in lactose concentration is consistent with the concept that it is part of the defense mechanism of the mammary gland. We suggest a two-phase mechanism for the development of E. coli mastitis. In phase 1, the acute phase, bacterial infection in the mammary gland or in LPS induced mastitis, triggers an immediate immune and inflammatory responses in which the endothelial-mammary barriers are open, allowing for cellular and humoral infiltration from blood to milk. In phase 2, the chronic phase, tight junctions close and infiltration is limited to areas of deep damage in the gland, which may perpetuate alterations that are measured in milk regardless of bacterial clearance off the mammary gland.

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**Author contributions**

S.B., D.H. and G.L. were responsible for performing the experiments, collecting data, experimental design, data analysis and preparing the manuscript draft; S.J. was responsible for animal husbandry and care; O.K. performed the laboratory work; Y.L. performed the statistical analyses; N.E. performed histological exams, provided histological pictures and descriptions; U.M., N.S.B. and G.L. analyzed the data and helped in writing and editing the manuscript. All authors read and approved the final manuscript.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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