Dietary Epidermal Growth Factor Supplementation Alleviates Intestinal Injury in Piglets with Intrauterine Growth Retardation via Reducing Oxidative Stress and Enhancing Intestinal Glucose Transport and Barrier Function

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Abstract: EGF plays an important role in the intestinal repair and nutrients transport of animals. However, the effect of EGF on the intestinal health of piglets with IUGR has not been reported. Thus, the present study was performed to investigate the effects of EGF on the intestinal morphology, glucose absorption, antioxidant capacity, and barrier function of piglets with IUGR. A total of 6 NBW piglets and 12 IUGR piglets were randomly divided into three treatments: NC group (NBW piglets fed with basal diet, n = 6), IC group (IUGR piglets fed with basal diet, n = 6), and IE group (IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF, n = 6). Growth performance, serum biochemical profile, jejunum histomorphology, jejunum glucose absorption and antioxidant capacity, and jejunal barrier function were measured. The results showed that EGF supplementation significantly increased the final body weight (FBW), average daily gain (ADG), and average daily feed intake (ADFI) of piglets with IUGR; EGF supplementation significantly increased the total protein (TP), glucose (GLU), and immunoglobulin G (IgG) levels compared with the IUGR piglets in the IC group; EGF administration effectively exhibited an increased jejunal villus height (VH) and the villus-height-to-crypt-depth ratio (V/C) of IUGR piglets compared with the IC group; EGF supplementation significantly increased sodium/potassium-transporting adenosine triphosphatase (Na+/K+-ATPase) activity, intestinal alkaline phosphatase (AKP) activity, glucose transporter 1 (SGLT1), glucose transporter 2 (GLUT2), and AMP-activated protein kinase α1 (AMPK-α1) mRNA expressions in the jejunum of IUGR piglets compared with the IC group; EGF supplementation exhibited increased superoxide dismutase (SOD) activity and total antioxidant capacity (T-AOC) levels, tended to increase glutathione peroxidase (GSH-Px) and catalase (CAT) activities, and tended to decrease the malondialdehyde (MDA) level in the jejunum of IUGR piglets compared with the IC group; EGF supplementation significantly increased ZO-1, Claudin-1, Occludin, and MUC2 mRNA expressions and improved secreted immunoglobulin A (sIgA) secretion in the jejunum of IUGR piglets compared with the IC group and tended to decrease the interleukin 1β (IL-1β), IL-6, and tumor necrosis factor α (TNF-α) levels in the jejunum of IUGR piglets compared with the IC group. Pearson’s correlation analysis further showed that EGF can promote intestinal...
development and nutrient absorption by promoting intestinal barrier function, thus improving the growth performance of IUGR piglets.

**Keywords:** epidermal growth factor; glucose absorption; intestinal barrier function; intrauterine growth retardation; oxidative stress; piglets

### 1. Introduction

IUGR usually refers to impaired growth and development of the embryo and/or its organs during pregnancy [1,2]. Pigs, a kind of multifetal mammal, exhibit a high incidence of IUGR, which seriously affects the postnatal growth and intestinal health of piglets and ultimately affects the economic benefit of pig production [3,4]. Studies have confirmed that piglets with IUGR are associated with feeding intolerance and intestinal dysfunction [4,5], which leads to a retard growth and poor health status of pigs. Therefore, strategies to alleviate intestinal injury and promote intestinal development may help to reverse the growth retardation in IUGR pigs. Nowadays, many strategies have been tested to reduce the incidence of IUGR at the sow level [6,7] or through nutritional interventions at the piglet level to increase IUGR piglets’ neonatal survival and growth [8,9].

The intestinal tract is an important site for the digestion and absorption of nutrients, and also functions as an important barrier to prevent the entry of an exogenous harmful substance into the circulation system [10,11]. Thus, maintaining a healthy gut is crucial for an animal’s overall health. IUGR is associated with intestinal injury of piglets, which can be demonstrated by the impaired intestinal morphology, raised apoptosis of intestinal epithelial cells, and increased oxidative injury and inflammatory response [4,8,12–14]. Emerging evidence suggests that EGF is a promising therapeutic treatment for intestinal injury due to its multiple biological functions [15–19]. EGF exhibits potential antioxidant and anti-inflammatory activities, and it can protect enterocytes against alcohol-induced inflammation [20], hydrogen peroxide [19] and acetaldehyde [21] caused barrier dysfunction, and lipopolysaccharide (LPS) [16] and ischemia/reperfusion [22]-induced oxidative injury. However, there are little data on EGF as a nutritional additive to alleviate intestinal injury in IUGR offspring. Our previous study showed that EGF treatment can promote glucose absorption of damaged intestinal or intestinal epithelial cells of piglets [18,23], which means that glucose plays an important role in intestinal repair. However, whether EGF can promote intestinal repair of IUGR piglets by increasing intestinal glucose absorption has not been reported. Therefore, it has a certain application value to study the effect of EGF on the intestinal health of IUGR piglets.

Accordingly, we hypothesized that dietary EGF supplementation can alleviate intestinal injury of piglets with IUGR by reducing intestinal oxidative stress and enhancing intestinal glucose transport and barrier function. To test the above hypothesis, we studied the effects of EGF on intestinal oxidative damage, barrier function, and the glucose absorption capacity of piglets with IUGR. This research can provide a reference for the application of EGF in IUGR piglets and provide some reliable reference for intestinal disorders treating in IUGR human neonates.

### 2. Materials and Methods

#### 2.1. Animals and Experimental Design

In this study, 6 NBW piglets (1.55 ± 0.05 kg; within one SD of the mean birth weight) and 12 IUGR piglets (0.94 ± 0.06; two SD below the mean birth weight) were selected from 6 sows with similar parity. All piglets suckled by their mother sows. At weaning (21 days of age), 6 NBW piglets were assigned to the NC group (NBW piglets fed with basal diet, n = 6), and 12 IUGR piglets were randomly divided into the IC group (IUGR piglets fed with basal diet, n = 6) and the IE group (IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF, n = 6). The EGF product (pig source) was obtained from Zyme Fast
(Changsha, China) Biotechnology (Changsha, China) with egg yolk powder as the carrier, and its purity was 4000 mg/kg. The composition and nutrient levels of basal diet (Table 1) met the nutrient specifications for 5 to 10 kg BW pig according to the NRC (2012) [24]. The experimental piglets were housed individually and had free access to feed and water. The room temperature was maintained at an ambient temperature range of 25–28 °C, and the room lighting was natural. The experiment lasted for 14 days.

**Table 1.** Composition and nutrient levels of the basal diet.

| Ingredients                  | Content (%) | Item          | Nutrient Levels |
|------------------------------|-------------|---------------|-----------------|
| Corn                         | 57.30       | DE/(MJ/Kg)    | 14.08           |
| Squeezed soybean meal        | 14.20       | Crude protein (%) | 20.14           |
| Expanded soybean meal        | 13.53       | Total Lys (%) | 1.50            |
| Whey powder                  | 5.03        | Total Thr (%) | 0.75            |
| Fish meal                    | 4.60        | Total Met (%) | 0.82            |
| Glucose                      | 2.00        | Met + Cys (%) | 0.73            |
| Limestone                    | 0.84        | Calcium (%)   | 0.86            |
| CaHPO4                       | 0.81        | Available Pi (%) | 0.45            |
| L-lysine-HCl (98.0%)         | 0.34        |               |                 |
| L-methionine (98.0%)         | 0.11        |               |                 |
| NaCl                         | 0.24        |               |                 |
| Premix ^2                    | 1.00        |               |                 |
| Total                        | 100.00      |               |                 |

1 Corn (8.5% CP), squeezed soybean meal (47.9% CP), expanded soybean (35.5% CP), fish meal (67.0% CP), whey powder (11.55% CP). ^2 The premix provided the following per kilogram of diet: VA, 10,000 IU; VB1, 1.8 mg; VB2, 5 mg; VB6, 7 mg; VB12, 0.02 mg; VD3, 1500 IU; VE, 60 IU; VK3, 3 mg; choline chloride, 1000 mg; niacin, 30 mg; an anti-microbial agent, 1 mg. ^3 Crude protein and calcium levels were measured values, and other nutrient levels were calculated values.

The body weights of piglets were weighed at the beginning and the end of the animal experiment to calculate the ADG of piglets. The feed intake was recorded weekly to calculate the ADFI of piglets. The feed-to-gain ratio (F/G) of piglets was calculated by ADFI/ADG [25].

2.2. Sample Collection

All animals were fasted for 12 h before sampling. All piglets were anesthetized by injection of 50 mg kg⁻¹ BW sodium pentobarbital, and 10 mL of blood was collected from the jugular vein and then centrifuged at 3000×g for 15 min at 4 °C to obtain serum samples. After blood collection, all pigs were killed by exsanguinations to death, after which the abdomens of the piglets were quickly opened for the removal of the entire intestine to isolate the jejunum. The middle jejunum segment (approximately 2 cm) was fixed in 10% neutral buffered formalin for intestinal morphology measurement. The intestinal mucosa of the middle jejunum tissue was gently scraped down and then rapidly frozen in liquid nitrogen and stored at −80 °C for subsequent analysis.

2.3. Serum Biochemical Profile Detection

Serum alanine aminotransferase (ALT), aspartate aminotransferase (AST), urea nitrogen (BUN), TP, GLU, T, immunoglobulin A (IgA), IgG, and IgM were measured using an automatic blood analyzer, Mindray BS-420 (Shenzhen Mindray Bio-Medical Electronics Co., Ltd., Shenzhen, China), in accordance with the manufacturer’s instructions.

2.4. Intestinal Histomorphology

Jejunum samples were fixed in 10% neutral buffered formalin for 24 h, then dehydrated and embedded in paraffin. Specimens were sliced into a thickness of 4 μm and stained with hematoxylin and eosin for morphological evaluation. A light microscope with a computer-assisted morphometric system (BioScan Optometric, BioScan Inc., Edmonds, WA, USA) was used to measure VH and crypt depth (CD) [26]. V/C was calculated by VH/CD.
2.5. Jejunal Mucosal Antioxidant and Immune Indices Detection

Jejunal mucosa samples (about 250 mg) were homogenized with a high-speed bench-top homogenizer (Tekmar, Cincinnati, OH, USA) in saline solution (1:4, weight/volume), then centrifuged at 3500 r/min for 15 min at 4 °C. The supernatants were collected for subsequent analysis. The MDA content was detected by thiobarbituric acid (TBA) method (kit no. A003-1-2), GSH-Px activity was detected by measuring the consumption of reduced glutathione in enzymatic reaction (kit no. A005-1-2), SOD activity was detected by hydroxylamine method (kit no. A001-1-2), CAT activity was detected by ammonium molybdate method (kit no. A007-1-1), and trolox equivalents were used for T-AOC measured (kit no. A015-2-1). All kits were purchased from Nanjing Jiancheng Bioengineering Institute (Nanjing, China). sIgA (kit no. CSB-E12063p), IL-1β (kit no. CSB-E09725p), IL-6 (kit no. CSB-E06786p), and TNF-α (kit no. CSB-E16980p) were measured using porcine-specific enzyme-linked immunosorbent assay (ELISA) kits according to the manufacturer’s instructions (Cusabio Biotech Co., Wuhan, China).

2.6. AKP and Na+/K+-ATPase Activity in Jejunum Mucosa

ATPase can decompose ATP to produce ADP and inorganic phosphorus, and the Na+/K+-ATPase activity can be judged by measuring the amount of inorganic phosphate. The intestinal Na+/K+-ATPase activity (kit no. A070-2-2) was measured using a commercial kit purchased from Nanjing Jiancheng Bioengineering Institute (Nanjing, China). AKP can decompose benzene disodium phosphate to produce free phenol and phosphoric acid. The phenol reacts with 4-aminobiphenyl in alkaline solution and oxidizes to red quinone derivatives by potassium ferricyanide. The AKP activity can be measured according to the red color. The intestinal AKP activity (kit no. A059-2-2) was measured using a commercial kit purchased from Nanjing Jiancheng Bioengineering Institute (Nanjing, China).

2.7. Real-Time PCR

Total RNA extraction and reverse transcription were performed as described previously [27]. The primers used in this study (Table 2) were synthesized by Sangon Biotech (Shanghai, China). The formula was $2^{-\Delta \Delta Ct}$, where $\Delta \Delta Ct = (Ct_{Target} - Ct_{\beta\text{-actin}})$ treatment $- (Ct_{Target} - Ct_{\beta\text{-actin}})$ control was used to calculated the relative gene expression [4].

| Gene       | Primers Sequence                | Product Length | Reference |
|------------|---------------------------------|----------------|-----------|
| AMPK-α1    | F:5′-GGTGAAAACCTCGGCCACTACA-3′  | 72 bp          | [23]      |
|            | R:5′-TGGGCAACCTCTACATCCG-3′    |                |           |
| SGLT1      | F:5′-ATATGGCCTTATAATCTCGCTC-3′  | 138 bp         | [23]      |
|            | R:5′-AAATAGTGGTCTTGGATGACG-3′  |                |           |
| GLUT2      | F:5′-CAGGCTATTTCTAGTGACG-3′    | 151 bp         | [23]      |
|            | R:5′-AAATAGTGGTCTTGGATGACG-3′  |                |           |
| MUC2       | F:5′-AGCCATCTGCTGGTGTAAGC-3′   | 121 bp         | [28]      |
|            | R:5′-AGCGTGCCGTCGGACTG-3′      |                |           |
| ZO-1       | F:5′-TGCTGCGCACTGACGATG-3′     | 129 bp         | [4]       |
|            | R:5′-CAGCCTGCAATCTGACGCA-3′   |                |           |
| Claudin-1  | F:5′-TCCTGCTGGGACGACGAT-3′     | 102 bp         | [4]       |
|            | R:5′-CAATCGACGACGACGATG-3′    |                |           |
| Occludin   | F:5′-CATCCTGGCTGGTGACGATG-3′   | 140 bp         | [4]       |
|            | R:5′-GCTGGCACTGAAATGACGATG-3′ |                |           |
| β-actin    | F:5′-CAAATGTCACGGACGATTCC-3′   | 116 bp         | [16]      |
|            | R:5′-TTATGTCACGGACGATTCC-3′   |                |           |

2.8. Statistical Analysis

Data were presented as the mean ± SEM. Data analysis was performed by one-way ANOVA procedure of SPSS Statistics 21.0 (SPSS, Inc., Chicago, IL, USA). Differences among treatment mean were determined using a Tukey multiple comparison test procedure of
SPSS Statistics 21.0 (SPSS, Inc., Chicago, IL, USA). The results of all data analyses were input into the GraphPad Prism 9.0 (GraphPad Software, Inc., San Diego, CA, USA) software for graphical display [4]. The correlations analysis was performed by the genescloud tools, a free online platform for data analysis (https://www.genescloud.cn (accessed on 25 June 2022)). p < 0.05 was considered to be statistically significant.

3. Results

3.1. Growth Performance

As shown in Table 3, the initial body weight (IBW), FBW, ADG, and ADFI were significantly different among three treatments (p < 0.01). Compared with the NC group, the IC group had a lower IBW, FBW, ADG, and ADFI (p < 0.05) during the first 2 weeks after weaning. Compared with the IC group, the IE group had a higher (p < 0.05) FBW, ADG, and ADFI, although significantly lower (p < 0.05) than the CN group.

Table 3. Effects of dietary EGF supplementation on the growth performance of piglets with IUGR.

| Items      | NC   | IC    | IE    | SEM | p Value |
|------------|------|-------|-------|-----|---------|
| IBW (kg)   | 6.77 | 5.02  | 5.04  | 0.20| <0.001  |
| FBW (kg)   | 9.74 | 6.95  | 7.63  | 0.30| <0.001  |
| ADG (g/d)  | 212.14| 137.14| 184.88| 8.25| <0.001  |
| ADFI (g/d) | 301.71| 216.13| 268.47| 10.30| <0.001  |

Values are expressed as means ± SEM, n = 6 (one pig in the IC group was dead, n = 5 in the IC group); NC: NBW piglets fed with basal diet; IC: IUGR piglets fed with basal diet; IE: IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF. a,b,c means that values within a row with different superscript letters were significantly different. p < 0.05 was taken to indicate statistical significance.

3.2. Serum Biochemical Profiles

As shown in Table 4, the concentrations of serum AST, TP, GLU, IgG, and IgM were markedly changed in this study (p < 0.05). The IC group had a higher (p < 0.05) serum AST level and lower (p < 0.05) TP, GLU, IgG, and IgM levels compared with the NC group. EGF supplementation significantly increased (p < 0.05) the TP, GLU, and IgG levels of IUGR piglets compared with the IC group and showed no difference compared with the NC group.

Table 4. Effects of EGF on serum biochemical profiles of piglets with IUGR.

| Items     | NC     | IC     | IE     | SEM | p-Value |
|-----------|--------|--------|--------|-----|---------|
| ALT (U/L) | 50.28  | 64.16  | 62.30  | 3.48| 0.214   |
| AST (U/L) | 110.88 | 163.31 | 137.78 | 7.86| 0.015   |
| TG (mmol/L)| 0.57  | 0.43  | 0.48  | 0.03| 0.168   |
| TP (g/L)  | 44.19  | 33.10  | 41.31  | 1.83| 0.032   |
| BUN (mmol/L)| 3.87 | 3.21  | 4.28  | 0.27| 0.294   |
| GLU (mmol/L)| 4.29 | 2.30  | 3.91  | 0.29| 0.005   |
| IgA (g/L) | 1.23   | 0.96  | 1.14  | 0.08| 0.368   |
| IgG (g/L) | 2.83   | 1.76  | 2.54  | 0.16| 0.011   |
| IgM (g/L) | 0.49   | 0.34  | 0.40  | 0.02| 0.025   |

Values are expressed as means ± SEM, n = 6 (one pig in the IC group was dead, n = 5 in the IC group); NC: NBW piglets fed with basal diet; IC: IUGR piglets fed with basal diet; IE: IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF. a,b,c means that values within a row with different superscript letters were significantly different. p < 0.05 was taken to indicate statistical significance.

3.3. Intestinal Morphology and Its Correlations Analysis with Growth Performance

The effects of EGF on the intestinal morphology of piglets with IUGR are presented in Figure 1. It shows that VH (Figure 1A) and V/C (Figure 1C) were markedly changed in this study (p < 0.05); however, there was no difference in CD (Figure 1B) among the three groups. IUGR reduces VH (p < 0.01) and V/C (p < 0.01) significantly. Meanwhile, EGF administration exhibited increased (p < 0.05) VH and V/C in the jejunum of IUGR piglets.
IgA (g/L) 1.23 0.96 1.14 0.08 0.368
IgG (g/L) 2.83 a 1.76 b 2.54 a 0.16 0.011

* p < 0.05, ** p < 0.01, *** p < 0.001.

IgA (g/L) 1.23 0.96 1.14 0.08 0.368
IgG (g/L) 2.83 a 1.76 b 2.54 a 0.16 0.011

* p < 0.05, ** p < 0.01, *** p < 0.001.

3.4. Glucose Absorption Capacity and Its Correlation Analysis with Growth Performance and Intestinal Morphology

The effects of EGF on the intestinal glucose absorption capacity of piglets with IUGR are presented in Figure 2. The activities of Na+/K+-ATPase activity (Figure 2A) and AKP activity (Figure 2B) and the gene expression of SGLT1 (Figure 2C), GLUT2 (Figure 2D), and AMPK-α1 (Figure 2E) were significantly different among the three treatments (p < 0.05). The IUGR piglets in the IC group had a lower intestinal Na+/K+-ATPase activity (p < 0.01), AKP activity (p < 0.01), and SGLT1 mRNA expression (p < 0.01;) compared with the NBW piglets in the NC group; IUGR did not affect GLUT2 and AMPK-α1 mRNA expression. EGF supplementation significantly increased Na+/K+-ATPase activity (p < 0.05), AKP activity (p < 0.01), SGLT1 (p < 0.01), GLUT2 (p < 0.01), and AMPK-α1 (p < 0.01) mRNA expression in the jejunum of IUGR piglets compared with IUGR piglets without EGF supplementation. Compared with the NC group, IUGR piglets in the IE group had a higher (p < 0.01) GLUT2 and AMPK-α1 mRNA expression, and there was no difference in Na+/K+-ATPase activity, AKP activity, and SGLT1 mRNA expression between the IE group and the NC group. Pearson’s correlation analysis showed that the Na+/K+-ATPase activity (p < 0.01; p < 0.001) and AKP activity (p < 0.001, p < 0.001; p < 0.05) were positively correlated with the ADFI, ADG, and FBW of piglets; and SGLT1 gene expression (p < 0.01, p < 0.01) was positively correlated with the ADFI and ADG of piglets (Figure 2F). Na+/K+-ATPase
activity \((p < 0.01, p < 0.01)\), AKP activity \((p < 0.05, p < 0.01)\), and SGLT1 mRNA expression \((p < 0.01, p < 0.01)\) were positively correlated with VH and V/C (Figure 2G).

Figure 2. Effects of EGF on the intestinal glucose absorption capacity of piglets with IUGR. (A) Na\(^+\)/K\(^+\)-ATPase activity, (B) AKP activity, (C) SGLT1 mRNA, (D) GLU2 mRNA, (E) AMPK-\(\alpha\)1 mRNA, (F) correlations analysis between intestinal glucose absorption capacity and growth performance, (G) correlations analysis between intestinal glucose absorption capacity and intestinal morphology. Values are expressed as means ± SEM, \(n = 6\) (1 pig in the IC group was dead, \(n = 5\) in the IC group); NC: NBW piglets fed with basal diet; IC: IUGR piglets fed with basal diet; IE: IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF. * \(p < 0.05\), ** \(p < 0.01\), *** \(p < 0.001\).

3.5. Antioxidant Activities and Its Correlation Analysis with Growth Performance and Intestinal Morphology

As shown in Figure 3, there was a significant difference \((p < 0.05)\) of SOD activity (Figure 3A), GSH-Px activity (Figure 3B), CAT activity (Figure 3C), T-AOC level (Figure 3D), and MDA level (Figure 3E) among the NC, IC, and IE groups. It showed that the IC group had a lower SOD activity \((p < 0.01)\), GSH-Px activity \((p < 0.01)\), CAT activity \((p < 0.01)\), and T-AOC level \((p < 0.05)\) and a higher MDA level \((p < 0.01)\) compared with the NC group. The IE group exhibited an increased SOD activity \((p < 0.05)\) and T-AOC level \((p < 0.05)\) compared with the IC group; EGF supplementation tended to increase the GSH-Px activity \((p = 0.053)\) and CAT activity \((p = 0.072)\) and tended to decrease the MDA level \((p = 0.090)\) in the jejunum of IUGR piglets compared with IUGR piglets without EGF supplementation. No differences in GSH-Px, SOD, CAT, T-AOC, and MDA were observed between the IE group and the NC group. Pearson’s correlation analysis showed that SOD activity \((p < 0.01, p < 0.01, p < 0.001)\), GSH-Px activity \((p < 0.01, p < 0.01, p < 0.01)\), CAT activity \((p < 0.01, p < 0.001, p < 0.001)\), and T-AOC level \((p < 0.01, p < 0.01, p < 0.05)\) were positively
correlated with the ADFI, ADG, and FBW of piglets, and the MDA level \( (p < 0.01, p < 0.05, p < 0.01) \) was negatively correlated with the ADFI, ADG, and FBW of piglets (Figure 3F). SOD activity was positively correlated with VH and V/C \( (p < 0.05) \), CAT activity was positively correlated with V/C \( (p < 0.05) \) and negatively correlated with CD \( (p < 0.05) \), T-AOC level was positively correlated with VH \( (p < 0.05) \), and MDA level \( (p < 0.05) \) was negatively correlated with VH and V/C (Figure 3G).

Figure 3. Effects of EGF on the intestinal antioxidant capacity of piglets with IUGR. (A) SOD, (B) GSH-Px, (C) CAT, (D) T-AOC, (E) MDA, (F) correlation analysis between intestinal antioxidant capacity and growth performance, (G) correlation analysis between intestinal antioxidant capacity and intestinal morphology. Values are expressed as means ± SEM, \( n = 6 \) (1 pig in the IC group was dead, \( n = 5 \) in the IC group); NC: NBW piglets fed with basal diet; IC: IUGR piglets fed with basal diet; IE: IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF. * \( p < 0.05 \), ** \( p < 0.01 \), *** \( p < 0.001 \).

3.6. Intestinal Barrier Function and Its Correlation Analysis with Growth Performance and Intestinal Morphology

As shown in Figure 4, MUC2 (Figure 4A), ZO-1 (Figure 4B), Claudin-1 (Figure 4C), and Occludin (Figure 4D) mRNA expression and the secretion of sIgA (Figure 4E), IL-1β (Figure 4F), IL-6 (Figure 4G), and TNF-α (Figure 4H) were significantly different among the three treatments \( (p < 0.05) \). Results showed that IUGR did not affect the MUC2 mRNA expression in jejunal mucosa. The IC group had a lower \( (p < 0.05) \) ZO-1, Claudin-1, and Occludin gene expression and a lower sIgA \( (p < 0.01) \) secretion in jejunal mucosa compared with the IC group; the IC group had a higher IL-1β \( (p < 0.01) \), IL-6 \( (p < 0.01) \), and TNF-α level \( (p < 0.05) \) in jejunal mucosa compared with the IC group. EGF supplementation
(IE group) significantly \((p < 0.01)\) increased the ZO-1, Claudin-1, Occludin, and MUC2 mRNA expression and improved the sIgA secretion in jejunal mucosa compared with the IC group, and significantly increased ZO-1 \((p < 0.01)\) and MUC2 \((p < 0.05)\) mRNA expression compared with the NC group, and tended to decrease the IL-1\(\beta\) \((p = 0.072)\), IL-6 \((p = 0.062)\), and TNF-\(\alpha\) level \((p = 0.073)\) in jejunal mucosa compared with the IC group. Pearson’s correlation analysis showed that the ZO-1 \((p < 0.05)\), MUC2 \((p < 0.05)\), and Occludin \((p < 0.01)\) gene expressions were positively correlated with the ADFI of piglets; the Claudin-1 gene expression \((p < 0.05, p < 0.05, p < 0.05)\) and sIgA \((p < 0.01, p < 0.01, p < 0.001)\) were positively correlated with the ADFI, ADG, and the FBW of piglets; and TNF-\(\alpha\) \((p < 0.01, p < 0.05, p < 0.01)\), IL-6 \((p < 0.001, p < 0.001, p < 0.001)\), and IL-1 level \((p < 0.01, p < 0.001, p < 0.001)\) were negatively correlated with the ADFI, ADG, and the FBW of piglets (Figure 4). Claudin-1 gene expression \((p < 0.05, p < 0.05)\) and sIgA \((p < 0.001, p < 0.001)\) were positively correlated with VH and V/C, and the TNF-\(\alpha\) level \((p < 0.001)\) was negatively correlated with V/C (Figure 4).

Figure 4. Effects of EGF on the intestinal barrier function of piglets with IUGR. (A) MUC2 mRNA, (B) ZO-1 mRNA, (C) Claudin-1 mRNA, (D) Occludin mRNA, (E) sIgA level, (F) IL-1\(\beta\) level, (G) IL-6 level, (H) TNF-\(\alpha\) level, (I) correlation analysis between intestinal barrier function and growth performance, and (J) correlation analysis between barrier function and intestinal morphology. Values are expressed as means ± SEM, \(n = 6\) (1 pig in the IC group was dead, \(n = 5\) in the IC group); NC: NBW piglets fed with basal diet; IC: IUGR piglets fed with basal diet; IE: IUGR piglets fed with basal diet supplemented with 2 mg/kg EGF. * \(p < 0.05\), ** \(p < 0.01\), *** \(p < 0.001\).
3.7. The Correlation Analysis of Intestinal Barrier Function with Glucose Absorption Capacity

Pearson’s correlation analysis showed that MUC2 ($p < 0.05, p < 0.001, p < 0.001$), ZO-1 ($p < 0.01, p < 0.001, p < 0.001$), Claudin-1 ($p < 0.05, p < 0.01, p < 0.001, p < 0.01$), and Occludin ($p < 0.01, p < 0.001, p < 0.001, p < 0.001$) gene expression were positively correlated with AKP activity, GLUT2, AMPK $\alpha$1, and SGLT1 mRNA expression; sIgA ($p < 0.05, p < 0.001, p < 0.01$) was positively correlated with AKP activity, Na$^+$/K$^+$-ATPase activity, and the gene expression of SGLT1; and TNF-$\alpha$ ($p < 0.05, p < 0.05$) was negatively correlated with Na$^+$/K$^+$-ATPase activity and the gene expression of SGLT1; the IL-6 level ($p < 0.01$) was negatively correlated with Na$^+$/K$^+$-ATPase activity; and the IL-1$\beta$ level ($p < 0.05, p < 0.05, p < 0.05$) was negatively correlated with AKP activity, Na$^+$/K$^+$-ATPase activity, and the gene expression of SGLT1 (Figure 5).

![Figure 5](image-url) Correlation analysis between intestinal barrier function and intestinal glucose absorption capacity. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$.

4. Discussion

4.1. Effects of EGF on the Growth Performance of IUGR Piglets

Animals with IUGR are characterized by low birth weight and postnatal growth retardation [4,29]. Our previous study showed that IUGR significantly reduced the average body weight gain of piglets in the whole sucking period [4]. In this study, IUGR piglets showed a lower ADG and ADFI compared with the NBW pigs, which was consistent with previous studies [9,30]. EGF has a positive effect on the growth performance of weaned piglets to some extent [17,31], but whether EGF could promote the growth performance of IUGR piglets has not been reported yet. In this study, EGF supplementation significantly increased the FBW, ADG, and ADFI of IUGR piglets compared with IUGR piglets without EGF supplementation. These results indicate that EGF has a certain effect on the alleviating growth inhibition of weaned piglets caused by IUGR.

4.2. Effects of EGF on the Serum Biochemical Indices of Piglets with IUGR

Serum biochemical indexes are important indexes reflecting the metabolic status of animals [25,26]. For example, the measurement of the serum activities of the hepatic enzymes ALT and AST was considered a reliable indicator to evaluate the liver function of animals [32]. The serum TP level can reflect the status of protein synthesis and metabolism [33]. Glucose is essential for maintaining the energy and health status of animals and regulating the absorption and transport capacity of various nutrients [18,23]. Immunoglobulins (IgA, IgG, and IgM) play important roles in immunity, which are considered important indicators reflecting the health status of animals [34,35]. In the present
study, the AST, TP, Glu, IgA, and IgM indexes were changed remarkably in response to piglets fed with EGF, which means that EGF is beneficial to the growth, nutrient absorption, immunity, protein synthesis, and body health of piglets with IUGR.

4.3. Effects of EGF on the Intestinal Morphology of IUGR Piglets

The integrity of the intestinal tract provides a guarantee for the utilization of nutrients [4]. Intestinal VH, CD, and V/C can reflect the healthy state of intestinal function, the decreased CD, and increased VH and V/C of intestines indicate a better intestinal function of animals [36]. Studies have shown that IUGR can cause intestinal villus shedding, crypt hyperplasia, and intestinal mucosal atrophy and damage intestinal mucosal barrier function, thereby resulting in the decline of intestinal absorption function [4,9]. EGF, as an intestinal trophic factor, plays important roles in intestinal development and intestinal repair [15,16,23]. The results of this study showed that dietary EGF supplementation significantly increased VH and V/C, which indicated that EGF could repair the intestinal morphology of piglets with IUGR. Similarly, our previous study showed that EGF treatment could repair intestinal morphology in LPS-stimulated piglets [23]. Pearson’s correlation analysis showed that VH and V/C were positively correlated with the ADFI, ADG, and FBW of piglets, which means that a good intestinal morphology is essential for animal growth.

4.4. Effects of EGF on the Glucose Absorption Capacity of Piglets with IUGR

Glucose is an important energy source in the body and is absorbed mainly in the small intestine [36]. It has been proved that the glucose absorption in the intestinal tract is achieved through either the paracellular or transcellular route, in which the Na\(^+\)/glucose cotransporter (SGLT1)-mediated transcellular route is the major route for the glucose transport [4,23,37,38]. GLUT2, a facilitated Na\(^+\)-independent monosaccharide transporter, also contributes to glucose absorption; it can be recruited transiently to the apical membrane and assist in glucose absorption in response to a sugar-rich meal [39] or intestinal repair process [23]. Therefore, the factors that influence SGLT1 and GLUT2 expression would also affect glucose absorption and metabolism. The transmembrane Na\(^+\) gradient and the membrane potential generated by Na\(^+\)/K\(^+\)-ATPase are the driving forces of SGLT1-mediated glucose transport [40]. AKP speeds up the absorption and transfer of nutrients, indirectly providing energy to the body [23]. It has been shown that sustained impairment of intestinal development in piglets results in decreased intestinal glucose absorption capacity [4,23]. The present study showed that IUGR significantly decreased Na\(^+\)/K\(^+\)-ATPase and AKP activity and SGLT1, GLUT2, and AMPK-\(\alpha\)1 mRNA expression compared with the NC group, which indicated that IUGR significantly impaired the intestinal glucose absorption capacity of piglets during the weaning period. Dietary EGF supplementation significantly increased Na\(^+\)/K\(^+\)-ATPase and AKP activity and SGLT1, GLUT2, and AMPK-\(\alpha\)1 mRNA expression compared with the IC group, which demonstrated that EGF can improve the glucose absorption capacity of the damaged intestine caused by IUGR. Our previous studies demonstrated that EGF can promote the glucose absorption capacity of injured IPEC-J2 and the jejunum of piglets induced by LPS [18,23]. Pearson’s correlation analysis showed that glucose absorption capacity was closely related to the intestinal morphology and growth performance of piglets. Thus, this study suggested that one way of EGF to alleviate intestinal injury and promote the growth performance of IUGR piglets is by improving intestinal glucose absorption capacity.

4.5. Effects of EGF on the Antioxidant Activities of Piglets with IUGR

An increased MDA level and decreased SOD, CAT, and GSH-Px levels are generally considered markers of intestinal oxidative injury [4,16,41]. Substantial evidence indicates that IUGR impairs the antioxidant defense system by increasing MDA production and decreasing antioxidant enzyme activity [4,8,13,42]. Similarly, in the present study, the IC group had a higher MDA content and lower GSH-Px, SOD, CAT, and T-AOC levels than
the CN group, which indicated that IUGR results in jejunum oxidative damage in piglets during the first 2 weeks after weaning. EGF is an important cell protective peptide, which has certain antioxidant function and can alleviate oxidative injury induced by LPS [16] and intestinal ischemia/reperfusion [22]. Our previous study showed that EGF exhibited potent protective effects on IPEC-J2 cells against LPS-induced cell oxidative damage by enhancing GSH-Px, SOD, and CAT expressions [16]. In this study, the addition of EGF significantly increased SOD activity and T-AOC level, tended to increase GSH-Px and CAT activity, and tended to decrease MDA level compared with the IC group, which indicated that EGF could alleviate intestinal oxidative damage of IUGR piglets. Pearson’s correlation analysis showed that intestinal redox status was closely related to the growth performance and intestinal morphology of piglets. Thus, the results of this study indicated that EGF can improve growth performance and alleviate the intestinal injury of IUGR piglets by enhancing intestinal antioxidant capacity.

4.6. Effects of EGF on the Intestinal Barrier Function of Piglets with IUGR

Intestinal mucus layer, intestinal epithelial cells, immune cells, normal microorganisms, and their metabolites in the intestinal lumen constitute the intestinal barrier of animals, and the dynamic balance between the barriers is the key to maintaining the homeostasis of the intestinal environment [10,11]. As a key composition of the intestinal mucosal epithelial barrier, ZO-1, claudin-1, and occludin are closely related to intestinal permeability [10]. In the present study, the IC group had lower ZO-1, Claudin-1, and Occludin gene expressions, which is consistent with previous studies [4,42]. EGF is an important regulatory factor of intestinal mucosal homeostasis and has a strong repair effect on intestinal morphological damage caused by weaning [43], intestinal ischemia/reperfusion [22], and necrotizing enterocolitis [44]. Similarly, the results of this study also indicated that EGF could promote the intestinal barrier function of piglets suffered from IUGR by increasing the expressions of ZO-1, Claudin-1, and Occludin genes.

The mucus layer provides the first defense line of the gastrointestinal tract, which can effectively prevent the colonization of pathogenic microorganisms [45]. Mucins (MUCs), mainly MUC2, forms the bulk of the intestinal mucus, which can not only lubricate the intestine and antagonize pathogenic bacteria, but also participate in intercellular signal transduction and regulation of intestinal mucosal related immune factors [10,46]. EGF could repair intestinal mucosal injury by promoting intestinal MUC2 secretion [28,44,47]. Similarly, in the present study, the addition of EGF in diet significantly increased the gene expression of MUC2 in the jejunum mucosa of IUGR piglets, which suggested that EGF can repair the intestinal barrier function damage caused by IUGR by promoting mucin secretion.

The gut immune system provides an important barrier to prevent pathogens from penetrating the intestinal epithelium [11]. It was reported that IUGR could result in intestinal inflammatory injury, which was manifested by the increased secretion of proinflammatory cytokines [4,41] and the decreased secretion of slgA [4]. Our present study showed that the IC group had a lower slgA level and higher IL-1β, IL-6, and TNF-α levels compared with the NC group, which also suggested that IUGR caused intestinal inflammatory injury. EGF can promote intestinal immune function and alleviate intestinal inflammation by promoting the secretion of immunoglobulins and reducing the expression of proinflammatory factors [20,33,46]. For example, Chen et al. [20] showed that EGF treatment effectively reduced alcohol-induced inflammatory factors (IL-1β, IL-10, and TNF-α) in rats. Similarly, our present study showed that dietary EGF supplementation significantly increased the slgA level and tended to decrease the IL-1β, IL-6, and TNF-α levels of IUGR piglets, which means that EGF alleviated the inflammatory injury caused by IUGR.

IUGR disrupts intestinal barrier function, while EGF provides protection for the intestinal barrier function of animals. It is generally believed that the integrity of intestinal barrier function is the basis of intestinal nutrients absorption, which determines the intestinal health and animal growth and development [11,48]. Pearson’s correlation analysis of the
present study showed that indicators related to mechanical barrier (ZO-1, Occludin and Claudin-1), an indicator related to chemical barrier (MUC2), and an indicator related to immune barrier (sIgA) were positively correlated with the growth performance, glucose absorption, and intestinal morphology of piglets; and proinflammatory cytokines (IL-1β, IL-6, TNF-α) were negatively correlated with the growth performance, glucose absorption capacity, and intestinal morphology of piglets. These results suggest that EGF can promote intestinal development and nutrient absorption by promoting intestinal barrier function, thus improving the growth performance of IUGR piglets.

5. Conclusions

In summary, EGF has a positive effect on the growth performance and serum biochemical indexes of IUGR piglets; meanwhile, EGF could alleviate the intestinal injury of piglets caused by IUGR, which is indicated by the better intestinal morphology, enhanced intestinal glucose absorption capacity and antioxidant capacity, the improved intestinal barrier function, and the decreased intestinal inflammatory injury.

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Institutional Review Board Statement: The study was conducted in accordance with the Chinese Guidelines for Animal Welfare, and approved by the Ethics Committee of Guizhou Normal University (protocol code GZNU-2021-0054; June 2021).

Informed Consent Statement: Not applicable.

Data Availability Statement: The data presented in this study are available on request from the corresponding author.

Conflicts of Interest: The authors declare no conflict of interest.

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