Integrated analysis of a lncRNA-mRNA network reveals a potential mechanism underlying necrotizing enterocolitis

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Received August 29, 2019; Accepted March 25, 2020

DOI: 10.3892/mmr.2020.11083

Abstract. Previous studies have shown that long non-coding RNAs (lncRNAs) serve important roles in necrotizing enterocolitis (NEC). However, the underlying mechanisms remain largely unknown. In order to examine the potential role of lncRNAs in NEC, the present study investigated lncRNA and mRNA expression profiles in NEC lesions and adjacent intestinal tissues using Next Generation Sequencing. A total of 4,202 differentially expressed lncRNAs (fold-change >2; P<0.05) and 7,860 differentially expressed mRNAs (fold-change >2; P<0.05) were identified. Moreover, 5 dysregulated lncRNAs and 5 mRNAs were randomly selected, and further assessed by reverse transcription-quantitative PCR in vitro. Gene Ontology and Kyoto Encyclopedia of Genes and Genomes analyses demonstrated that the differentially expressed lncRNAs were closely associated with NEC, and were enriched in ‘inflammatory response’, ‘Toll-like receptor binding’, ‘PPAR signaling pathway’, ‘PI3K-Akt signaling pathway’, ‘transforming growth factor-β signaling pathway’ and ‘hypoxia-inducible factor 1 signaling pathway’. In addition, co-expression analysis demonstrated that these lncRNAs, including lncRNA ENST00000623580, lncRNA NONHSAT087855.2, may mediate the pathogenesis and development of NEC via lncRNA-mRNA network interactions. Therefore, the present study provided a novel insight into the role of lncRNAs in NEC.

Introduction

Necrotizing enterocolitis (NEC) is an acute life-threatening gastrointestinal disease that predominantly affects premature neonates (1,2). The morbidity of NEC is rising rapidly with the continuing increased rate of survival of preterm infants (3). Patients that survive severe NEC may develop serious sequelae, including gastrointestinal complications and abnormal neuro-development (4). Multiple factors contribute to NEC, including premature delivery, intestinal bacterial colonization and infant formula feeding (5). Furthermore, NEC is characterized by severe intestinal inflammation, impaired mucosal healing and intestinal necrosis (4). However, the etiology and pathogenesis of NEC are not fully understood.

Long non-coding RNAs (lncRNAs) are autonomously transcribed non-coding RNAs, which are >200 nucleotides in length (6). lncRNAs can act at the transcriptional, post-transcriptional and epigenetic levels via cis- or trans-acting pathways (7,8). Previous studies have reported the potential importance of lncRNAs in maintaining the intestinal barrier and mediating inflammatory lesions (9-11). For example, lncRNA Uc.173 was revealed to enhance intestinal epithelial barrier function by increasing the expression of claudin 1 (9). However, lncRNA Uc.173 silencing leads to dysfunction of the intestinal epithelial barrier in vitro (9). Another study showed that lncRNA SPRY4-intronic transcript 1 (SPRY4-IT1) regulates the intestinal epithelial barrier by affecting the expression levels of claudin-1, junctional adhesion molecule 1 and occludin (10). Moreover, lncRNA SPRY4-IT1 overexpression in intestinal mucosa in mice protects the intestinal barrier against infection (10). In addition, lncRNA H19 promotes the proliferation of intestinal epithelial cells by directly binding to tumor protein p53 protein, microRNA34a and let-7 (11). Therefore, these previous results indicated that lncRNAs directly affect the expression of mRNAs via co-expression networks. However,
the role of lncRNA-mRNA networks in the pathogenesis of NEC is not fully understood. Thus, the construction of such networks may increase the understanding of the mechanisms underlying NEC.

In the present study, the expression profiles of lncRNAs and mRNAs in NEC lesion and adjacent intestinal tissues obtained from patients with NEC were analyzed by Next Generation Sequencing (NGS). Additionally, a number of these genes were validated by reverse transcription-quantitative PCR (RT-qPCR) in vitro. Moreover, Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) analyses were performed to assess the potential functions of the differentially expressed lncRNAs and mRNAs. In addition, a lncRNA-mRNA network was constructed in order to identify the potential underlying mechanism of NEC. The present results suggested that lncRNAs in intestinal tissues may serve an important role in the development of NEC.

Materials and methods

Sample collection and ethics statement. Preterm infants (25-36 weeks of age; one female and two males) with stage III acute NEC undergoing bowel resection were enrolled in the present study between October 2017 and October 2018 at The Affiliated Wuxi Children's Hospital of Nanjing Medical University. The diagnosis of NEC was based on Bell's stages (12). Infants with other severe intestinal diseases, such as congenital megacolon, were excluded. Tissues of the small bowel segment with NEC lesions that exhibited perforation or necrosis, and adjacent healthy tissue [referred to as the control (CTL)] were resected and stored at -80°C in liquid nitrogen until analysis. The study protocol was approved by the Institutional Review Board of The Affiliated Wuxi Children's Hospital of Nanjing Medical University (approval no. 2017-EYLL-115). Informed consent was obtained from the parents or guardians of all patients.

RNA isolation, quality control and RNA-Seq. Total RNA was extracted from the tissues using a mirVana microRNA isolation kit (cat. no. AM1561; Ambion; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. RNA integrity was subsequently assessed using an Agilent Bioanalyzer 2100 system (Agilent Technologies, Inc.). RNA was further purified using an RNeasy Clean XP kit (Beckman Coulter, Inc.) and an RNase-Free DNase set (Qiagen GmbH). The samples were then subjected to NGS. The mRNAs and lncRNAs were identified using an Illumina platform (HiSeq 2500; Illumina, Inc.). The gene abundance was calculated using StringTie (version no. 1.3.0) (13). NGS data acquisition and processing were performed by Shanghai Biotechnology Corporation.

NEC cell model construction. The human normal colorectal FHC cell line (American Type Culture Collection) was cultured in RPMI-1640 medium (Gibco; Thermo Fisher Scientific, Inc.) containing 5% FBS (Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin/streptomycin solution at 37°C and 5% CO2. In order to establish the NEC cell model, FHC cells were treated with 200 μg/ml lipopolysaccharide (LPS; Sigma-Aldrich; Merck KGaA) at 37°C for 6 h, referring to the dose and time of LPS treatment described in previous study (14). LPS used in the present study was obtained from Escherichia coli O111:B4.

RNA preparation and RT-qPCR. A total of 5 mRNAs and 5 lncRNAs were randomly selected and further assessed by RT-qPCR. Total RNA from FHC cells was extracted using TRIzol® reagent (Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. RT was performed using a PrimeScript RT reagent kit (TaKaRa Bio, Inc.) according to the manufacturer's instructions (37°C for 15 min; 85°C for 5 sec; 4°C for 10 min). qPCR was subsequently performed using a SYBR Green PCR kit (TaKaRa Bio, Inc.). The following thermocycling conditions were used: 95°C for 10 min; 95°C for 10 sec, 60°C for 30 sec, 40 cycles. The primer pairs used for qPCR are presented in Table S1. Gene expression was calculated using the 2-ΔΔcq method (15), and normalized to the expression of the internal reference gene GAPDH. Data are presented as the average of 3 independent experiments.

Protein extraction and western blotting. Total protein from FHC cells was extracted using RIPA lysis buffer (Beyotime Institute of Biotechnology) containing 1% phenylmethylsulfonyl fluoride. The protein concentration was measured by a BCA assay kit (Thermo Fisher Scientific, Inc.). A total of 20 µg protein/lane was separated via SDS-PAGE on a 10% gel and transferred onto a PVDF membrane. The membrane was subsequently blocked with 5% skimmed milk for 1 h at room temperature and incubated with primary antibodies against Toll-like receptor-4 (TLR4; 1:1.000; cat. no. bs-20594R; BIOSS) and β-actin (1:1.000; cat. no. CST-4970; Cell Signaling Technology, Inc.) for 12 h at 4°C. Following primary antibody incubation, the membrane was incubated with a horseradish peroxidase conjugated-secondary antibody (1:5,000; cat. no. CST-7074; Cell Signaling Technology, Inc.) for 30 min at room temperature. Proteins were visualized using an enhanced chemiluminescence detection system (Pierce ECL Plus Western Blotting Substrate; Thermo Fisher Scientific, Inc.). Protein bands were analyzed by ImageJ software (version 1.47; National Institutes of Health).

Hematoxylin and eosin staining. Intestines were fixed in 4% paraformaldehyde overnight at room temperature, dehydrated with various concentrations of ethanol (50, 75, 85, 95 and 100% ethanol) and xylen, and embedded in paraffin. Paraffin-embedded small intestinal specimens were cut into 5-μm sections. Sections were then deparaffinized and rehydrated. Following which, the sections were stained with Mayer's hematoxylin for 30 sec and 1% eosin Y solution for 10-30 sec at room temperature. After dehydration and mounting, the sections were examined under a light microscope, at magnification, x200, and images were taken.

GO and KEGG analysis. GO (version 14; www.geneontology.org) was used to analyze the biological processes, cellular components and molecular functions of the differentially expressed mRNAs and host genes of the lncRNAs. In addition, KEGG (version 4.0; www.genome.jp/kegg) pathway analysis was performed to investigate the associated pathways. GO terms and KEGG pathways were ranked by the enrichment factor.
lncRNA-mRNA co-expression network. A lncRNA-mRNA network was constructed using Cytoscape software (version 3.4.0; https://cytoscape.org/). A total of 38 differentially expressed mRNAs and 44 lncRNAs, whose target genes are same to those mRNAs, were selected. In total, 80 co-expression relationships were identified.

Statistical analysis. Statistical differences were evaluated using the two independent samples t-test. Data are presented as the mean ± standard deviation. The edgeR software package (version 3.28.1) (16) was used to identify differentially expressed lncRNAs and mRNAs, between NEC and CTL tissues. P<0.05 was considered to indicate a statistically significant difference.

Results

Detection of lncRNA and mRNA profiles in NEC tissues. NGS was used to detect the expression of mRNAs and lncRNAs in NEC and CTL tissues. Moreover, differentially expressed mRNAs and lncRNAs were analyzed using a lncRNA-mRNA co-expression network (Fig. 1A). Furthermore, hematoxylin and eosin staining identified a disrupted mucosal architecture in NEC tissues (Fig. 1B).

Differentially expressed gene analysis. A total of 4,202 lncRNAs were significantly differentially expressed between NEC and CTL tissues (fold-change >2; P<0.05), which included 1,657 upregulated lncRNAs and 2,545 downregulated lncRNAs (Fig. 2A; Table S1I). In addition, it was identified that 7,860 mRNAs were significantly differentially expressed between NEC and CTL tissues (fold-change ≥2; P<0.05), which included 4,241 upregulated mRNAs and 3,619 downregulated mRNAs (Fig. 2B; Table S1II). The top 50 significantly downregulated and upregulated lncRNAs and mRNAs are presented in Tables I and II, respectively.

Construction of the NEC cell model and validation of genes by RT-qPCR in vitro. The pathology underlying NEC development requires the activation of the innate immune TLR4 on enterocytes (17). Moreover, TLR4 activation by bacterial LPS inhibits enterocyte proliferation and leads to disease progression (18,19). It was demonstrated that LPS exposure increased the protein expression of TLR4 in FHCs, thus suggesting that it is a reliable cell model for studying NEC in vitro (Fig. 3A). In total, 5 mRNAs and 5 lncRNAs were randomly selected to assess the NGS results by RT-qPCR. The present results indicated that the mRNA expression levels of interleukin (IL)-6 and IL-1β were upregulated, whereas the expression levels of transmembrane 4 L six family member 20, solute carrier family 5 member 12 and leucine rich repeat containing 19 were downregulated (Fig. 3B-F). Furthermore, NONHSAT022134.2 was downregulated, while NONHSAT076868.2, NONHSAT159417.1, NONHSAT094312.2 and NONHSAT142596.2 were upregulated (Fig. 3G-K). Therefore, the RT-qPCR results were consistent with the NGS results.

Differentially expressed lncRNAs are associated with the inflammatory response and the transforming growth factor-β (TGF-β) signaling pathway. GO and KEGG analyses of the differentially expressed lncRNAs were performed. It was identified that the terms ‘intestinal absorption’, ‘IL-18 production’, ‘apoptotic process involved in morphogenesis’ and ‘inflammatory response’ were highly enriched in the biological process category (Fig. 4A). In addition, ‘immunological synapse’, ‘actin filament’, ‘cell-cell contact zone’ and ‘receptor’ were highly enriched in the cellular component category (Fig. 4B). Furthermore, it was demonstrated that ‘extracellular ATP-gated cation channel activity’, ‘chemorepellent activity’, ‘PI3K activity’ and ‘cAMP binding’ were highly enriched in the molecular function category (Fig. 4C). The KEGG pathway analysis results identified that the
Table I. Top 50 significantly upregulated and downregulated long non-coding RNAs.

| IncRNA_ID     | log2FC  | P-value    | Q-value    |
|---------------|---------|------------|------------|
| NONHSAT155609.1 | 11.61383132 | 1.43x10^{-21} | 1.22x10^{-20} |
| ENST00000602813 | 11.55252438 | 1.30x10^{-9} | 7.94x10^{-8} |
| NONHSAT059749.2 | 10.91351071 | 3.06x10^{-12} | 1.92x10^{-11} |
| NONHSAT118562.2 | 10.66845241 | 2.81x10^{-10} | 1.06x10^{-9} |
| ENST00000620266 | 10.60925657 | 2.84x10^{-10} | 5.92x10^{-9} |
| NONHSAT209475.1 | 10.52975989 | 1.54x10^{-17} | 4.47x10^{-15} |
| NONHSAT159678.1 | 10.44687578 | 2.78x10^{-17} | 7.58x10^{-15} |
| NONHSAT169788.1 | 10.30207377 | 8.74x10^{-11} | 6.80x10^{-9} |
| NONHSAT123206.2 | 10.20925898 | 2.57x10^{-10} | 1.81x10^{-9} |
| NONHSAT094312.2 | 10.08888094 | 2.95x10^{-27} | 4.47x10^{-24} |
| ENST00000499624 | 9.681004925 | 6.54x10^{-28} | 7.20x10^{-22} |
| NONHSAT177823.1 | 9.583016227 | 1.06x10^{-36} | 2.59x10^{-14} |
| NONHSAT159417.1 | 9.438955909 | 1.51x10^{-24} | 1.26x10^{-20} |
| NONHSAT173659.1 | 9.328552802 | 1.54x10^{-17} | 4.47x10^{-15} |
| NONHSAT137151.2 | 9.243165734 | 2.22x10^{-08} | 1.04x10^{-06} |
| NONHSAT206983.1 | 9.22839375 | 2.26x10^{-22} | 1.47x10^{-19} |
| NONHSAT178474.1 | 9.08635215 | 1.08x10^{-29} | 8.23x10^{-19} |
| ENST00000628476 | 9.043520963 | 4.70x10^{-11} | 3.93x10^{-09} |
| NONHSAT081637.2 | 9.023472571 | 1.02x10^{-08} | 0.000234023 |
| NONHSAT076868.2 | 8.9920343 | 1.31x10^{-32} | 3.36x10^{-29} |
| NONHSAT155904.1 | 8.93691199 | 7.85x10^{-14} | 1.12x10^{-11} |
| NONHSAT180405.1 | 8.90507266 | 3.21x10^{-07} | 1.13x10^{-05} |
| NONHSAT071404.2 | 8.875787946 | 2.04x10^{-10} | 9.59x10^{-08} |
| NONHSAT183239.1 | 8.869833121 | 3.85x10^{-15} | 6.90x10^{-13} |
| NONHSAT083185.2 | 8.855681215 | 2.60x10^{-20} | 1.20x10^{-17} |
| NONHSAT055522.2 | 8.835207264 | 2.66x10^{-13} | 3.40x10^{-11} |
| ENST00000573951 | 8.78394133 | 3.14x10^{-16} | 6.98x10^{-14} |
| NONHSAT142596.2 | 8.728768517 | 1.32x10^{-19} | 5.25x10^{-17} |
| NONHSAT055581.2 | 8.726034495 | 2.06x10^{-10} | 1.48x10^{-08} |
| NONHSAT050230.2 | 8.724625325 | 8.74x10^{-11} | 6.80x10^{-09} |
| NONHSAT056662.2 | 8.670344103 | 2.25x10^{-16} | 5.17x10^{-14} |
| ENST00000505532 | 8.58704266 | 1.64x10^{-14} | 2.64x10^{-12} |
| NONHSAT009555.2 | 8.577747998 | 3.39x10^{-9} | 9.05x10^{-05} |
| ENST00000635687 | 8.488271168 | 1.13x10^{-18} | 3.91x10^{-16} |
| ENST00000523099 | 8.4516672 | 2.38x10^{-06} | 6.72x10^{-05} |
| NONHSAT050198.2 | 8.410023146 | 1.50x10^{-15} | 0.000327563 |
| NONHSAT064287.2 | 8.405723447 | 7.03x10^{-06} | 0.000168779 |
| ENST00000558776 | 8.383710584 | 0.004874773 | 0.039194634 |
| ENST000010340.2 | 8.380591242 | 3.29x10^{-25} | 0.000064125 |
| ENST00000441790 | 8.379233397 | 1.26x10^{-07} | 4.90x10^{-06} |
| ENST00000426699 | 8.371776115 | 7.03x10^{-06} | 0.000168779 |
| ENST00000614899 | 8.359796918 | 1.26x10^{-07} | 4.90x10^{-06} |
| NONHSAT182506.1 | 8.346803842 | 8.05x10^{-10} | 5.21x10^{-08} |
| ENST00000615722 | 8.342773399 | 1.72x10^{-07} | 6.44x10^{-06} |
| NONHSAT215498.1 | 8.331225239 | 7.54x10^{-09} | 3.93x10^{-07} |
| NONHSAT095885.2 | 8.32856903 | 8.52x10^{-07} | 2.65x10^{-05} |
| NONHSAT175395.1 | 8.31998397 | 1.37x10^{-12} | 1.50x10^{-10} |
| NONHSAT108374.2 | 8.314575441 | 2.86x10^{-04} | 4.35x10^{-12} |
| NONHSAT207205.1 | 8.305343553 | 9.36x10^{-08} | 3.77x10^{-06} |
Table I. Continued.

| IncRNA_ID     | log2FC       | P-value     | Q-value     |
|---------------|--------------|-------------|-------------|
| NONHSAT183597.1 | -16.48090468 | 8.79x10^{-9} | 2.93x10^{-5} |
| NONHSAT168464.1 | -14.21176751 | 2.63x10^{-8} | 2.92x10^{-4} |
| NONHSAT213758.1 | -12.54912544 | 8.00x10^{-8} | 1.90x10^{-4} |
| NONHSAT152066.1 | -12.17786195 | 3.23x10^{-9} | 1.73x10^{-5} |
| NONHSAT024359.2 | -11.60824268 | 8.00x10^{-9} | 3.42x10^{-5} |
| MSTRG.44916.1  | -11.87327964 | 3.83x10^{-9} | 3.68x10^{-5} |
| NONHSAT181735.1 | -11.43001449 | 6.93x10^{-9} | 1.00x10^{-5} |
| NONHSAT160167.1 | -11.41835274 | 3.88x10^{-9} | 3.49x10^{-5} |
| NONHSAT165927.1 | -11.3077161  | 1.29x10^{-9} | 1.23x10^{-5} |
| NONHSAT159286.1 | -11.10183428 | 9.99x10^{-9} | 1.33x10^{-5} |
| NONHSAT091106.2 | -11.07224496 | 4.78x10^{-9} | 7.97x10^{-5} |
| NONHSAT175707.1 | -11.04329678 | 1.25x10^{-9} | 1.61x10^{-5} |
| ENST0000504773  | -10.84721244 | 4.46x10^{-9} | 3.30x10^{-5} |
| NONHSAT091102.2 | -10.66881414 | 1.16x10^{-9} | 3.45x10^{-5} |
| NONHSAT159894.1 | -10.62188109 | 1.57x10^{-9} | 9.02x10^{-5} |
| NONHSAT031256.2 | -10.60253283 | 1.47x10^{-9} | 1.22x10^{-5} |
| NONHSAT123764.2 | -10.50424036 | 8.82x10^{-9} | 2.45x10^{-5} |
| ENST0000507311  | -10.4756391  | 1.96x10^{-9} | 4.36x10^{-5} |
| NONHSAT171172.1 | -10.4117159  | 9.23x10^{-9} | 6.28x10^{-5} |
| NONHSAT162645.1 | -10.33478373 | 3.39x10^{-9} | 9.04x10^{-5} |
| NONHSAT091107.2 | -10.32292548 | 2.57x10^{-9} | 2.04x10^{-5} |
| NONHSAT139922.2 | -10.31278929 | 5.33x10^{-9} | 1.97x10^{-5} |
| NONHSAT183589.1 | -10.28237601 | 5.84x10^{-9} | 6.71x10^{-5} |
| NONHSAT196167.1 | -10.27748084 | 6.07x10^{-9} | 4.31x10^{-5} |
| MSTRG.7793.3    | -10.20152795 | 2.73x10^{-9} | 1.75x10^{-5} |
| NONHSAT188793.1 | -10.11670026 | 1.02x10^{-9} | 6.43x10^{-5} |
| NONHSAT152222.1 | -10.03914993 | 5.21x10^{-9} | 2.10x10^{-5} |
| ENST0000606993  | -10.00747342 | 1.58x10^{-9} | 5.28x10^{-5} |
| NONHSAT157362.2 | -9.985583318 | 2.27x10^{-9} | 2.94x10^{-5} |
| NONHSAT157008.1 | -9.951490231 | 7.85x10^{-9} | 1.12x10^{-5} |
| NONHSAT160671.1 | -9.908059683 | 3.39x10^{-9} | 6.15x10^{-5} |
| ENST0000427901  | -9.906883277 | 4.70x10^{-9} | 3.93x10^{-5} |
| NONHSAT192263.1 | -9.864619513 | 3.19x10^{-9} | 1.01x10^{-5} |
| NONHSAT182621.1 | -9.857221425 | 2.57x10^{-9} | 2.28x10^{-5} |
| NONHSAT094167.2 | -9.856028479 | 1.67x10^{-9} | 2.24x10^{-5} |
| ENST0000419296  | -9.82058918 | 1.29x10^{-9} | 1.23x10^{-5} |
| NONHSAT041891.2 | -9.794594044 | 2.11x10^{-9} | 1.24x10^{-5} |
| NONHSAT176544.1 | -9.772352515 | 1.65x10^{-9} | 9.95x10^{-6} |
| ENST0000455071  | -9.756925085 | 1.60x10^{-9} | 1.48x10^{-5} |
| NONHSAT208752.1 | -9.667998866 | 2.92x10^{-9} | 9.36x10^{-6} |
| NONHSAT022134.2 | -9.65276041 | 7.91x10^{-9} | 5.27x10^{-5} |
| NONHSAT017148.2 | -9.551998808 | 3.83x10^{-9} | 3.29x10^{-5} |
| NONHSAT176002.2 | -9.549948279 | 1.26x10^{-9} | 4.90x10^{-6} |
| NONHSAT143306.2 | -9.53507668 | 3.95x10^{-9} | 1.88x10^{-5} |
| NONHSAT152860.1 | -9.531346389 | 1.03x10^{-9} | 3.63x10^{-6} |
| NONHSAT091105.2 | -9.513378928 | 1.40x10^{-9} | 5.47x10^{-7} |
| ENST0000584749  | -9.484529301 | 8.05x10^{-9} | 5.21x10^{-8} |
| NONHSAT060224.2 | -9.35289754 | 4.70x10^{-11} | 3.93x10^{-10} |
| NONHSAT032678.2 | -9.348901378 | 1.43x10^{-13} | 1.95x10^{-11} |

FC, fold change.
Table II. Top 50 significantly upregulated and downregulated mRNAs.

| A. Upregulated |
|----------------|
| Gene ID       | Gene name               | log2FC | P-value       | Q-value       |
| ENSG000000228536 | RP11-392017.1         | 9.13   | 2.33x10^{-17} | 3.87x10^{-16} |
| ENSG000000129988 | LBP                    | 8.87   | 9.55x10^{-30} | 3.12x10^{-28} |
| ENSG00000121053 | EPX                    | 8.64   | 1.62x10^{-33} | 6.40x10^{-32} |
| ENSG00000267653 | RP1-193H18.3          | 8.63   | 1.30x10^{-09} | 1.16x10^{-08} |
| ENSG00000244437 | IGKV3-15              | 8.47   | 4.11x10^{-07} | 2.73x10^{-06} |
| ENSG00000145850 | TIMD4                 | 8.34   | 7.39x10^{-23} | 1.69x10^{-21} |
| ENSG00000259183 | MED28P6               | 8.33   | 0.001268      | 0.004697      |
| ENSG00000229308 | AC010084.1            | 8.32   | 7.04x10^{-08} | 5.18x10^{-07} |
| ENSG00000279686 | ESCR                   | 8.25   | 1.97x10^{-12} | 2.29x10^{-11} |
| ENSG00000231123 | SPATA2BP1             | 8.17   | 8.35x10^{-06} | 4.60x10^{-05} |
| ENSG00000226012 | AP001434.2            | 8.16   | 0.000667      | 0.002621      |
| ENSG00000230024 | RP11-95P13.1          | 8.13   | 8.35x10^{-06} | 4.60x10^{-05} |
| ENSG00000273153 | RP11-406H21.2         | 8.07   | 0.002421      | 0.008418      |
| ENSG00000164821 | DEF4                   | 8.05   | 2.27x10^{-07} | 1.56x10^{-06} |
| ENSG00000267436 | AC005786.7            | 8.04   | 7.45x10^{-07} | 4.79x10^{-06} |
| ENSG00000271656 | RP11-114N19.5         | 7.99   | 0.000667      | 0.002621      |
| ENSG00000211970 | IGHV4-61              | 7.93   | 2.86x10^{-05} | 0.00146       |
| ENSG00000211942 | IGHV3-13              | 7.92   | 5.33x10^{-05} | 0.000258      |
| ENSG00000259626 | MTND3P12              | 7.83   | 0.001268      | 0.004697      |
| ENSG00000279903 | RP11-349F21.5         | 7.82   | 4.54x10^{-06} | 2.60x10^{-05} |
| ENSG00000231613 | RPS-94333.1           | 7.78   | 2.86x10^{-05} | 0.000146      |
| ENSG00000235821 | IFITM4P               | 7.73   | 0.000187      | 0.00082       |
| ENSG00000214402 | LCNL1                 | 7.71   | 6.67x10^{-48} | 4.38x10^{-46} |
| ENSG00000225938 | RP4-575N6.4           | 7.67   | 0.000667      | 0.002621      |
| ENSG00000211946 | IGHV3-20              | 7.62   | 0.000667      | 0.002621      |
| ENSG00000256039 | RP11-291B21.2         | 7.62   | 1.26x10^{-07} | 8.98x10^{-07} |
| ENSG00000158104 | HPD                   | 7.55   | 2.07x10^{-21} | 4.43x10^{-20} |
| ENSG00000235151 | AC114730.2            | 7.51   | 5.33x10^{-05} | 0.000258      |
| ENSG00000239839 | DEFA3                 | 7.51   | 8.35x10^{-06} | 4.60x10^{-05} |
| ENSG00000275558 | RN78KP175             | 7.49   | 0.008941      | 0.02691       |
| ENSG00000239353 | RP11-492E3.51         | 7.45   | 0.002421      | 0.008418      |
| ENSG00000149516 | MS4A3                 | 7.45   | 2.99x10^{-27} | 8.62x10^{-26} |
| ENSG00000243066 | RN7SL842P             | 7.44   | 0.008941      | 0.02691       |
| ENSG00000196415 | PRTN3                 | 7.39   | 8.74x10^{-55} | 7.16x10^{-53} |
| ENSG00000211950 | IGHV1-24              | 7.35   | 0.002421      | 0.008418      |
| ENSG00000275791 | TRBV10-3              | 7.34   | 0.004643      | 0.014979      |
| ENSG00000211664 | IGLV2-18              | 7.31   | 0.004643      | 0.014979      |
| ENSG00000226660 | TRBV2                 | 7.31   | 0.002421      | 0.008418      |
| ENSG00000250579 | CTD-2297D10.2         | 7.31   | 1.60x10^{-06} | 8.41x10^{-06} |
| ENSG00000166819 | PLIN1                 | 7.31   | 4.63x10^{-02} | 1.30x10^{-09} |
| ENSG00000273006 | RP11-314C9.2          | 7.31   | 0.017298      | 0.047391      |
| ENSG00000258082 | RP11-443B7.3          | 7.31   | 4.54x10^{-06} | 2.60x10^{-06} |
| ENSG00000240864 | IGKV1-16              | 7.30   | 0.004643      | 0.014979      |
| ENSG00000279180 | RP11-417018.1         | 7.30   | 1.60x10^{-06} | 8.41x10^{-06} |
| ENSG00000184811 | TUSC5                 | 7.26   | 2.61x10^{-40} | 1.33x10^{-38} |
| ENSG00000234436 | AC008984.7            | 7.20   | 0.004643      | 0.014979      |
| ENSG00000232286 | RP11-80K6.2           | 7.19   | 0.002421      | 0.008418      |
| ENSG00000211747 | TRBV20-1              | 7.17   | 0.004643      | 0.014979      |
| ENSG00000254673 | RP11-598P20.5         | 7.16   | 0.000187      | 0.00082       |
| ENSG00000231863 | RP3-428L16.1          | 7.16   | 0.000667      | 0.002621      |
Table II. Continued.

| Gene ID | Gene name | log2FC | P-value  | Q-value  |
|---------|-----------|--------|----------|----------|
| ENSG00000147676 | MAL2 | -9.98 | 2.32x10^{-71} | 3.10x10^{-69} |
| ENSG00000140832 | MARVELD3 | -9.97 | 4.72x10^{-93} | 1.09x10^{-90} |
| ENSG00000158125 | XDH | -9.97 | 1.84x10^{-153} | 1.60x10^{-150} |
| ENSG00000272141 | RP11-465B22.8 | -9.97 | 7.32x10^{-33} | 2.79x10^{-31} |
| ENSG00000235122 | RP3-417L20.4 | -9.96 | 2.50x10^{-16} | 3.88x10^{-15} |
| ENSG00000234155 | RP11-30P6.6 | -9.93 | 5.74x10^{-57} | 4.96x10^{-55} |
| ENSG00000160868 | CYP3A4 | -9.92 | 9.78x10^{-59} | 9.35x10^{-56} |
| ENSG00000103534 | TMC5 | -9.89 | 1.11x10^{-42} | 7.63x10^{-40} |
| ENSG00000235523 | RP11-63P12.7 | -9.86 | 9.90x10^{-15} | 1.37x10^{-13} |
| ENSG00000165841 | CYP2C19 | -9.84 | 2.86x10^{-62} | 2.90x10^{-60} |
| ENSG00000253313 | Clorf210 | -9.83 | 1.62x10^{-47} | 1.05x10^{-45} |
| ENSG00000084674 | APOB | -9.82 | 1.30x10^{-200} | 1.93x10^{-200} |
| ENSG00000118094 | TREH | -9.81 | 8.09x10^{-90} | 1.75x10^{-87} |
| ENSG00000172016 | REG3A | -9.80 | 4.24x10^{-130} | 2.03x10^{-127} |
| ENSG00000044012 | GUCA2B | -9.78 | 4.31x10^{-20} | 8.54x10^{-19} |
| ENSG00000081051 | AF5 | -9.77 | 4.20x10^{-70} | 5.49x10^{-68} |
| ENSG00000169876 | MUC17 | -9.77 | 1.80x10^{-99} | 4.11x10^{-92} |
| ENSG00000173467 | AGR3 | -9.69 | 3.30x10^{-56} | 1.48x10^{-54} |
| ENSG00000131910 | NR0B2 | -9.68 | 8.08x10^{-32} | 2.92x10^{-30} |
| ENSG00000122711 | SPINK4 | -9.64 | 3.00x10^{-23} | 7.07x10^{-21} |
| ENSG00000165556 | CDX2 | -9.63 | 8.60x10^{-100} | 2.83x10^{-97} |
| ENSG00000224916 | APOC4-APOC2 | -9.63 | 8.05x10^{-52} | 6.09x10^{-50} |
| ENSG00000167183 | PRR15L | -9.53 | 3.86x10^{-70} | 1.19x10^{-68} |
| ENSG00000232400 | RAD17P1 | -9.53 | 1.93x10^{-42} | 1.06x10^{-40} |
| ENSG00000278505 | CI7orf78 | -9.53 | 1.27x10^{-54} | 1.04x10^{-52} |
| ENSG00000225329 | LHFPL3-AS2 | -9.53 | 7.93x10^{-49} | 5.37x10^{-47} |
| ENSG00000143278 | F13B | -9.52 | 4.39x10^{-49} | 3.00x10^{-47} |
| ENSG00000187664 | HAPLN4 | -9.51 | 5.32x10^{-60} | 5.03x10^{-58} |
| ENSG00000127831 | VIL1 | -9.50 | 1.33x10^{-163} | 1.65x10^{-160} |
| ENSG00000108242 | CYP2C18 | -9.49 | 3.21x10^{-52} | 2.46x10^{-50} |
| ENSG00000136872 | ALDOB | -9.44 | 4.47x10^{-704} | 1.66x10^{-700} |
| ENSG00000160182 | TFF1 | -9.38 | 6.20x10^{-16} | 9.32x10^{-15} |
| ENSG00000105398 | SULT2A1 | -9.38 | 5.95x10^{-128} | 2.76x10^{-125} |
| ENSG00000241388 | HNF1A-AS1 | -9.35 | 2.35x10^{-62} | 2.40x10^{-60} |
| ENSG00000224163 | RP11-309L24.6 | -9.35 | 2.45x10^{-15} | 3.57x10^{-14} |
| ENSG00000257084 | U47924.27 | -9.33 | 3.00x10^{-10} | 2.85x10^{-9} |
| ENSG00000166391 | MOGAT2 | -9.28 | 4.57x10^{-140} | 2.95x10^{-137} |
| ENSG00000140297 | GCNT3 | -9.28 | 4.25x10^{-76} | 6.45x10^{-74} |
| ENSG00000275410 | HNF1B | -9.27 | 6.98x10^{-76} | 1.06x10^{-73} |
| ENSG00000132437 | DDC | -9.27 | 6.68x10^{-100} | 1.80x10^{-97} |
| ENSG00000171431 | KRT20 | -9.26 | 1.15x10^{-99} | 3.07x10^{-97} |
| ENSG00000162460 | TMEM82 | -9.26 | 7.32x10^{-33} | 2.79x10^{-31} |
| ENSG00000197408 | CYP2B6 | -9.25 | 1.08x10^{-99} | 1.39x10^{-97} |
| ENSG00000249948 | GBA3 | -9.25 | 2.62x10^{-99} | 6.81x10^{-97} |
| ENSG00000100604 | CHGA | -9.19 | 1.03x10^{-130} | 5.09x10^{-128} |
| ENSG00000198944 | SOWHA | -9.18 | 1.05x10^{-73} | 1.49x10^{-71} |
| ENSG00000203858 | HSD3BP2 | -9.17 | 9.98x10^{-24} | 2.40x10^{-22} |
| ENSG00000166869 | CHP2 | -9.17 | 4.15x10^{-112} | 1.45x10^{-109} |
| ENSG00000060566 | CREB3L3 | -9.13 | 9.98x10^{-136} | 5.58x10^{-133} |
| ENSG00000260934 | CTA-363E6.7 | -9.10 | 1.19x10^{-12} | 1.40x10^{-11} |

FC, fold change.
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Figure 2. Differentially expressed mRNAs and lncRNAs in necrotizing enterocolitis and adjacent tissues. (A) A volcano plot was used to assess the differentially expressed lncRNAs. Of the 37,521 non-redundant lncRNAs, 4,202 lncRNAs were significantly differentially expressed (fold-change >2; P<0.05). A total of 1,657 and 2,545 lncRNAs were upregulated and downregulated, respectively. (B) A volcano plot was used to assess differentially expressed mRNAs. Of the 136,137 non-redundant mRNAs, 7,860 mRNAs were significantly differentially expressed (fold-change >2; P<0.05). A total of 4,241 and 3,619 mRNAs were upregulated and downregulated, respectively. lncRNA, long non-coding RNA.

Figure 3. NEC cell model construction, and assessment of the dysregulated lncRNAs and mRNAs by reverse transcription-quantitative PCR in vitro. (A) TLR4 protein expression in CTL FHC cells, and FHC cells treated with LPS. Relative mRNA expression levels of (B) IL-6, (C) IL-1β, (D) TM4SF20, (E) SLC5A12 and (F) LRRC19 in the CTL and NEC groups. Relative lncRNA expression levels of (G) NONHSAT022134.2, (H) NONHSAT076868.2, (I) NONHSAT159417.1, (J) NONHSAT094312.2 and (K) NONHSAT142596.2 in the CTL and NEC groups. Data are presented as mean ± standard deviation. *P<0.05, **P<0.01, ***P<0.001. CTL, control; lncRNA, long non-coding RNA; IL, interleukin; TM4SF20, Transmembrane 4 L six family member 20; SLC5A12, solute carrier family 5 member 12; LRRC19, leucine rich repeat containing 19; NEC, necrotizing enterocolitis.
differentially expressed lncRNA host genes were involved in ‘TGF-β signaling pathway’, ‘hypoxia-inducible factor 1 (HIF-1) signaling pathway’ and ‘tight junction and regulation of actin cytoskeleton’ (Fig. 4D).

Differentially expressed mRNAs are involved in TLR binding and the PI3K-Akt signaling pathway. GO and KEGG analysis were also performed for the differentially expressed mRNAs, which were significantly enriched in ‘myeloid dendritic cell chemotaxis’ (biological process; Fig. 5A), ‘inflammasome complex’ (cellular component; Fig. 5B) and ‘TLR binding’ (molecular function; Fig. 5C). Furthermore, KEGG pathway analysis results demonstrated that the differentially expressed mRNAs were enriched in ‘peroxisome proliferator-activated receptors (PPAR) signaling pathway’, ‘leukocyte transendothelial migration’, ‘extracellular matrix-receptor interaction’ and ‘PI3K-Akt signaling pathway’ (Fig. 5D).

lncRNA-mRNA network reveals a potential mechanism for NEC. The lncRNA-mRNA co-expression network results suggested that lncRNAs had a complex interaction with mRNAs (Fig. 6). A total of 80 associations were identified...
among the 44 lncRNAs and 38 mRNAs. Moreover, it was identified that lncRNAs ENST00000623580, NONHSAT180418.1, NONHSAT125636.2 and NONHSAT087855.2 interacted with mRNAs that were closely associated with nec. Therefore, the present results suggested that these lncRNAs may serve important roles in the pathogenesis of nec.

Discussion

To the best of our knowledge, no previous studies have examined lncRNAs associated with nec, and there is no functional evidence from studies investigating lncRNAs with nec and intestinal function. The present study analyzed the differentially expressed lncRNAs and mRNAs in ileal samples obtained from premature neonates with nec. Furthermore, an in vitro cell model of nec was investigated. The aims of the present study were to predict the potential functions of the differentially expressed lncRNAs and mRNAs, and to identify a potential mechanism for nec via lncRNA-mRNA network analyses. While the pathophysiological mechanism of NEC remains unknown, previous studies have reported that the expression of TLR4 is upregulated in NEC (20,21). The activation of TLR4 on intestinal epithelial cells induces a significant proinflammatory response (22). Furthermore, TLR4 activation also results in increased intestinal necrosis and apoptosis, which is the main feature of NEC (19). In addition, it has been shown that LPS-induced reactive oxygen species accumulation is involved in NF-κB activation and subsequent release of proinflammation cytokines, including IL-6 and IL-1β, in the pathogenesis of nec (23,24). A previous study revealed that excessive inflammation of the immature intestine predisposes premature infants to NEC (25). Therefore, elevated levels of IL-6 and IL-1β may be used as an index of successful nec cell model construction (26-28). The present results suggested that the expression levels of IL-1β, IL-6 and TLR4 in FHcs were significantly increased following LPS exposure, thus indicating that this was a reliable cell model for studying NEC in vitro.
GO and KEGG analyses were performed to identify potential mechanisms underlying NEC. The GO analysis results demonstrated that both the differentially expressed lncRNA host genes and mRNAs were associated with inflammation, which is a typical feature of NEC (29). Moreover, it was identified that TLR binding was a highly enriched molecular function, thus indicating the importance of the TLR in NEC. The results of KEGG pathway analysis identified pathways that were mainly associated with the progression of NEC. The PPAR pathway has been shown to affect mucosal immunity, commensal homeostasis and intestinal inflammation (30). Moreover, PPARs downregulation in mice contributes to an imbalance of intestinal symbiosis, resulting in enhanced susceptibility to intestinal inflammation and increased production of inflammatory cytokines (30). Previous studies have shown that the PI3K-Akt signaling pathway is altered in an experimental NEC animal model (31,32). Furthermore, berberine inhibits apoptosis and inflammation via the PI3K/AKT signaling pathway to attenuate the progression of NEC (32). Macrophages exhibited increased Smad7 expression in areas with high bacterial load and severe tissue damage (33). In addition, elevated Smad7 expression inhibits TGF-β signaling in intestinal macrophages, and promotes NF-κB activation and cytokine production during NEC (33). The PI3K-Akt signaling pathway has also been reported to regulate HIF-1 to protect against intestinal injury during NEC (34).

The lncRNA-mRNA network constructed in the present study revealed the lncRNAs and mRNAs that may be potentially involved in NEC. IncRNA ENST00000623580 was identified to interact with 6 mRNAs, including ADAM metallopeptidase with thrombospondin type 1 motif 1 (ADAMTS1), complement C1r, apolipoprotein L domain containing 1 (APOLD1), cytochrome B reductase 1 (CYBRD1), acetyl-CoA carboxylase β (ACACB) and matrix Gla protein (MGP). Among them, ADAMTS1 is involved in various inflammatory processes and is required for maintaining organ function, including the intestine (35,36). Moreover, C1r encodes several members of the peptidase S1 protein family, which are proteins that serve as a proteolytic subunit in the complement system C1 complex (37). The complement system acts as a mediator in the innate immune response by triggering phagocytosis, inflammation and rupture of bacterial cell walls, which may affect the progression of NEC (38). APOLD1 is an endothelial cell early response protein that affects vascular function and regulates endothelial cell signalling, which are closely associated with the NEC process (39). As a member of the cytochrome family, CYBRD1 encodes an iron-regulated protein and exhibits ferric reductase activity (40). Furthermore, CYBRD1 has been demonstrated to serve a key role in dietary iron absorption (41). It has been shown that NEC may affect the expression of CYBRD1 in the duodenal brush border membrane and contributes to the severe anemia associated with NEC (42). ACACB is currently considered to be one of the potential targets for regulating human disease in diabetes, obesity and cancer (43). It was demonstrated that continuous oxidation of fatty acid in ACACB knockout mice can lead to high insulin sensitivity (44). Moreover, a change in tissue ACACB levels via transcriptional regulation may be important in the affected intestine in NEC (45). MGP encodes a member of the osteocalcin/matrix Gla family of proteins (46). The encoded vitamin K-dependent protein is secreted by chondrocytes and vascular smooth muscle cells, and functions as a physiological inhibitor of ectopic tissue calcification (46). Furthermore, the present results suggested that IncRNA ENST00000623580 may be involved in the pathogenesis of NEC by regulating the expression of the aforementioned mRNAs.

SAM and SH3 Domain Containing 1 (SASH1) was identified to be associated with IncRNA NONHSAT180418.1. SASH1 is expressed on all microvascular beds and serves as a scaffold molecule to bind TGF-β-activated kinase 1 (TAK1), TNF receptor associated factor 6 (TRAF6), IκB kinase α and IκB kinase β (47). This interaction promotes the ubiquitination of TAK1 and TRAF6, and promotes LPS-induced activation of NF-κB, p38 and JNK, which leads to increased expression of proinflammatory cytokines and increased endothelial migration (47). Moreover, SASH1 has been shown to be involved in downstream signaling of TLR4 and to activate early endothelial responses to receptor activation (48). Therefore, IncRNA NONHSAT180418.1 was hypothesized to regulate the expression of SASH1 to affect TLR4 and the progression of NEC. In addition, IncRNAs NONHSAT125636.2 and NONHSAT087855.2 were both associated with LYZ. LYZ encodes human lysozyme, whose natural substrate is the bacterial cell wall component peptidoglycan (49). It was reported that increasing the expression of LYZ may control the microbial load in the intestine (50). Therefore, IncRNA NONHSAT125636.2 and NONHSAT087855.2 may exert protective effects in NEC. It has been revealed that IncRNAs can function as competing endogenous RNAs to indirectly regulate the expression levels of target genes via sponging microRNAs, and they are implicated in numerous biological processes (51,52). Thus, further studies on the lncRNA-associated competing endogenous RNA regulatory network may be useful for understanding the pathogenesis of NEC.

In conclusion, the present study identified differentially expressed IncRNAs and mRNAs between NEC and adjacent intestinal tissues by NGS. A number of these genes were randomly selected and assessed by RT-qPCR in vitro. GO and KEGG analyses were performed, and indicated that IncRNAs ENST00000623580, NONHSAT180418.1, NONHSAT125636.2 and NONHSAT087855.2 may have bioactive effects in NEC. Therefore, the present study may provide novel insights into the pathological mechanisms underlying NEC. However, further functional investigation is required to confirm these results.

Acknowledgements

Not applicable.

Funding

The present study was supported by grants from the Wuxi Key Medical Disciplines(grant no.ZDXK12),the Medical Innovation Team of Jiangsu Province (grant no. CXTDB2017016), the Wuxi Medical Development Discipline (grant no. FZXK001), the Wuxi Young Medical Talents (grant nos. QNRC094 and QNRC089), the Wuxi Hospital Management Centre Key Project (grant no. YGZXZ1513) and the Wuxi Hospital Management
Centre Joint Research Project (grant no. YGZXL1319). The study was also supported by the Young Project of Wuxi Health and Family Planning Commission (grant no. Q201815), the National Natural Science Foundation of China (grant nos. 81741089 and 81971427), the Jiangsu Maternal and Child Health Research Project (grant no. F2018525), the National Natural Science Foundation for Youth Scholars of China (grant nos. 81501296, 81901512 and 81901517) and the Nanjing Medical Science and Technological Development Program (grant no. YKK17176).

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions

WC, XY, TT, RY, XW, ZY and YL collected and analyzed the data. LZ and SH designed the study and wrote the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The study protocol was approved by the Institutional Review Board of The Affiliated Wuxi Children’s Hospital of Nanjing Medical University (approval no. 2017-EYLL-115). Informed consent was obtained from the parents or guardians of all patients.

Patient consent for publication

Informed consent was obtained from the parents or guardians of all patients.

Competing interests

The authors declare that they have no competing interests.

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