RNA-Seq Profiling of Circular RNAs During Development of Hindgut in Rat Embryos With Ethylenethiourea-Induced Anorectal Malformations

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Anorectal malformations (ARMs) are among the most common congenital terminal digestive tract malformations. Circular RNAs (circRNAs), a novel type of endogenous non-coding RNAs, play roles in the development of the digestive system; however, their contributions to the pathogenesis of ARMs are not well-established. In this study, we explored the mechanism underlying ethylenethiourea (ETU)-induced ARMs by profiling circRNA expression via RNA-seq and constructing a regulatory circRNA-miRNA-mRNA network. Nine pregnant rats were gavage-fed a single dose of 125 mg/kg 1% ETU (ARM group) on gestational day 10 (GD10), and another 9 pregnant rats received a similar dose of saline (normal group) as a control. Embryos were obtained by cesarean section on the key time-points of anorectal development (GD14, GD15, and GD16). Hindgut samples isolated from the fetuses were evaluated by high-throughput sequencing and differentially expressed circRNAs were validated by reverse transcription-quantitative polymerase chain reaction, agarose gel electrophoresis, and Sanger cloning and sequencing. A total of 18295 circRNAs were identified in the normal and ARM groups. Based on the 425 differentially expressed circRNAs (|Fc| > 2, p < 0.05), circRNA-miRNA and miRNA-mRNA pairs were predicted using miREAP, miRanda, and TargetScan. A total of 55 circRNAs (14 up- and 41 downregulated in the ARM group compared to the normal group) were predicted to bind to 195 miRNAs and 947 mRNAs. Competing endogenous RNA networks and a Kyoto Encyclopedia of Genes and Genomes analysis revealed that novel_circ_001042 had the greatest connectivity and was closely related to ARM-associated signaling pathways, such as the Wingless Type MMTV integration site family, mitogen-activated protein kinase, and transforming growth factor-β pathways. These results provide original insight into the roles of circRNAs in ARMs and provide a valuable resource for further analyses of molecular mechanisms and signaling networks.

Keywords: anorectal malformation, high-throughput RNA sequencing, circRNA, circRNA-miRNA-mRNA network, competing endogenous RNA
INTRODUCTION
Anorectal malformations (ARMs) are the most common congenital terminal digestive tract malformations, with an incidence of approximately 1 in 5,000 births and a slightly lower incidence in females than in males (Cuschieri and Eurocat Working Group, 2001; Wood and Levitt, 2018). The wide spectrum of ARM phenotypes includes stenotic anus, ectopic anus, rectourethral fistula, and rectovestibular fistula (Endo et al., 1999; Bai et al., 2004; Wijers et al., 2014). More than half of patients with ARM also suffer from congenital anomalies of the urogenital system, cardiovascular system, skeleton, and gastrointestinal tract (Stoll et al., 2007; Nah et al., 2012). Despite sophisticated surgical reconstruction operations, many patients continue to experience complications, such as fecal incontinence and constipation, which adversely affect their quality of life (Grano et al., 2013; Rintala, 2016). Recent studies have identified genes and genetic signaling pathways involved in the remarkable pathophysiology of ARMs; however, the specific etiology remains unclear (Draaken et al., 2012; Wijers et al., 2014). Therefore, it is essential to investigate the pathogenesis of ARMs.

Circular RNAs (circRNAs) are a novel type of endogenous non-coding RNA that predominate in the mammalian cytoplasm or exosomes (Szabo et al., 2015). CircRNA forms a covalent, closed-loop structure lacking both 5′-3′ polarity and a poly-A tail (Lasda and Parker, 2014). This special structure makes circRNAs resistant to degradation by RNA exonucleases, RNase R, and debranching enzymes, more stable than linear transcripts, and less prone to degradation (Suzuki and Tsukahara, 2014; Vicens and Westhof, 2014). The first circRNAs to be identified, viroid, was found in RNA viruses in 1976 (Sanger et al., 1976). Previous studies have focused on the biological functions of circRNAs, including their roles as "sponges" for microRNAs via competing endogenous RNA (ceRNA) networks, the circRNA-miRNA-mRNA axis, alternative mRNA splicing patterns via RNA-binding proteins, interactions with RNA polymerase II to regulate transcription, and translation by ribosomes and encoding proteins (Abe et al., 2013; Hansen et al., 2013; Zhang et al., 2013; Du et al., 2016). Recent studies have shown that circRNAs play important regulatory roles in embryonic development, apoptosis, differentiation, and proliferation (Lukiw, 2013; Bachmayr-Heyda et al., 2015; Maiese, 2016). Numerous differentially expressed circRNAs are involved in the development of gastrointestinal diseases and are potential diagnostic and prognostic biomarkers (Torre et al., 2015; Li et al., 2020). Our current understanding of the exact molecular mechanisms by which circRNAs affect the development of ARMs remains limited.

Ethynenethioureia (ETU) is commonly used to create animal models of ARM in rat embryos; typical phenotypes include an abnormally large distance between the urorectal septum and the cloacal membrane, postponed tailgut regression, abnormal apoptosis in the cloacal wall, and maldevelopment of the dorsal cloaca and membrane (Qi et al., 2002; Macedo et al., 2007). Our research group has conducted a series of studies on miRNA and mRNA expression profiles in the development of hindgut in rat embryos (Tang et al., 2014; Long et al., 2018, 2020). However, very little is known about circRNAs in ETU-induced ARMs in rats, and sequence data for circRNA in ETU-induced ARMs have never been systematically described. In the present study, circRNA expression patterns in the hindguts of rat embryos with ETU-induced ARMs were profiled from gestational day (GD)14 to GD16, the key time period for anorectal development (Qi et al., 2002; Bai et al., 2004; Macedo et al., 2007).

To determine the molecular mechanism by which circRNAs operate in ARMs, high-throughput sequencing was used to detect differentially expressed circRNAs in embryonic hindgut tissues of rats with ETU-induced ARMs and normal hindgut tissues. Numerous previously unreported circRNAs were identified, which may improve our understanding of the roles of circRNAs in ARM development. We also constructed ceRNA networks based on target predictions. The aberrant expression of circRNAs detected in the present study improves our understanding of the mechanisms underlying ARMs and provides a valuable basis for future studies of this condition. Furthermore, our data will serve as a useful resource for the development of therapeutic targets and novel diagnostics in the future.

MATERIALS AND METHODS

Ethics Statement
The present study was approved by the Ethics Committee of the China Medical University, Shenyang, China (no. 2015PS213K), and all procedures involving animals were performed in accordance with the guidelines for the care and use of laboratory animals.

Materials Used and Tissue Collection
Mature female Wistar rats (number: 18; age: 7–8 weeks old; body weight: 230–280 g) were acquired from the Experimental Animal Center, Shengying Hospital of China Medical University (Shenyang, China). The Key Laboratory of the Health Ministry for Congenital Malformations (Shenyang, China) provided a specific pathogen-free animal laboratory (room temperature: 22 ± 2°C; humidity: 55 ± 5%; 12 h light/12 h dark cycle; free access to water and food) for rodents. The ARM model and hindgut isolation procedure were constructed as per our previous protocol (Tang et al., 2014). In total, 9 pregnant rats were gavage-fed a single dose of 125 mg/kg 1% ETU (Sigma-Aldrich; Merck Millipore, Darmstadt, Germany) on GD10, and the remaining pregnant rats received the same dose of saline without ETU as a control. Cesarean sections were performed on GD14-GD16 to obtain embryos, and the presence of ARMs was determined by light microscopy. The specimens were stored in

Abbreviations: ARMs, anorectal malformations; ETU, ethylenethioureia; GD, gestational day; Rpm, reads per million mapped reads; circRNAs, circular RNAs; ceRNA, competing endogenous RNA; tRNA, ribosomal RNA; GO, Gene Ontology; KEGG, Kyoto Encyclopedia of Genes and Genomes; FDR, false discovery rate; qRT-PCR, quantitative reverse-transcription polymerase chain reaction; WNT, wingless-type MMTV integration site family; TGF-, transforming growth factor-β; MAPK, mitogen-activated protein kinase; GEO, Gene Expression Omnibus; PCA, principal component analysis.

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liquid nitrogen until use. All surgeries were performed after the intraperitoneal injection of sodium pentobarbital for anesthesia. The control group comprised of 252 saline-treated embryos without malformations. All 230 embryos with ETU-induced ARMs had short tails or no tails, and 15 died in utero. ARMs were detected in 86.1% (198/230) of ETU-treated embryos.

**RNA Extraction and RNA-Seq**

Total RNA was extracted from the specimens using TRIzol (Life Technologies, Carlsbad, CA, United States) according to the manufacturer’s instructions. The RNA was qualified and quantified using an Agilent 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA, United States). After rRNAs were removed, the rest of RNAs were fragmented using a fragmentation buffer and reverse-transcribed into cDNA with random primers. Second-strand cDNA was synthesized using DNA polymerase I, RNase H, dNTP (dUTP instead of dTTP), and buffer. Next, the cDNA fragments were purified using a QIAquick PCR Extraction Kit (Qiagen, Hilden, Germany), end-repaired, polyadenylated, and ligated to Illumina sequencing adapters. Uracil-N-glycosylase was used to digest the second-strand cDNA. After PCR amplification, the digested products were purified using AMPure XP Beads and a High Sensitivity DNA Assay Kit (Agilent Technologies, United States) was used for library quality inspection. RNA sequencing was carried out by Gene Denovo Biotechnology Company (Guangzhou, China) using a HiSeq 2500 system (Illumina, San Diego, CA, United States).

**RNA-Seq Analysis and Identification of circRNAs**

To obtain high-quality clean reads, raw reads were filtered as follows: (1) reads containing adapters were removed, (2) reads containing more than 10% unknown nucleotides (N) were removed, and (3) low-quality reads containing more than 50% low-quality bases (Q-value ≤ 20) were removed. The efficiency of experimental ribosome RNA removal is affected by the species and sample quality. Therefore, the Bowtie2 short read alignment tool (Langmead and Salzberg, 2012) was used to map the reads to the ribosomal RNA (rRNA) database. The rRNA-mapped reads were removed, followed by mapping to the reference genome using TopHat2 (version 2.1.1) (Kim et al., 2013). After alignment with the reference genome, the reads that could be mapped to the genome were discarded, and the unmapped reads were collected for circRNA identification. The 20-mers from both ends of the unmapped reads were extracted and aligned to the reference genome to identify unique anchor positions within the splice site. Anchored reads that aligned in the reversed orientation (head to tail) indicated circRNA splicing and were subsequently subjected to find_circ (Menczak et al., 2013) to identify circRNAs. The circRNAs were blast searched against circBase (Glazar et al., 2014) for annotation. Those that could not be annotated were considered novel circRNAs. To quantify the circRNAs, back-spliced junction reads were scaled to reads per million mapped reads (RPM) according to the following formula: RPM = 10^6C/N, where C is the number of back-spliced junction reads that uniquely aligned to a circRNA and N is the total number of back-spliced junction reads.

**Clustering Analysis and Principal Component Analysis (PCA)**

The relationships between samples were explored using Unsupervised hierarchical clustering and PCA. Unsupervised hierarchical clustering analyses were performed using gmodels in R^2 and PCA was conducted using ggplot2 in R^2.

**Analysis of Differentially Expressed circRNAs**

The edgeR software package^3 was used to identify differentially expressed circRNAs between samples or groups. CircRNAs with an absolute value of fold change of ≥ 2 and a p-value of < 0.05 were identified as differentially expressed in comparisons between samples or groups.

**Trend Analysis**

To examine patterns of differentially expressed circRNAs, the expression data for each sample were normalized to 0, log2 (v1/v0), and log2 (v2/v0) and clustered using Short Time-series Expression Miner (Ernst and Bar-Joseph, 2006). Clusters with p ≤ 0.05 were considered significant.

**Analysis of circRNA-miRNA-mRNA Interactions**

To predict mRNAs that interact with circRNAs and miRNAs, mirEAP, miranda, and TargetScan were used, and the intersection between the three prediction algorithms was used as the prediction results for the miRNA target genes. Pairs with positively correlated expression were identified as ceRNA pairs. Negative correlations indicated miRNA-mRNA and miRNA-circRNA pairs. Competition between circRNAs and mRNAs for the same miRNA and positive correlations between circRNAs and mRNAs were then screened. circRNA-miRNA-mRNA networks conforming to the ceRNA rules were visualized using Cytoscape v3.7.1. The flowchart of the circRNA-miRNA-mRNA ceRNA network analysis shown in Figure 1.

**Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) Pathway Analyses**

Source genes were mapped to GO terms in the Gene Ontology database^4, gene numbers were calculated for every term, and significantly enriched GO terms for source genes compared to the genome background were defined using a hypergeometric test. Calculated p-values were subjected to false discovery rate (FDR) correction, taking FDR ≤ 0.05 as a threshold for significant enrichment.

1. https://cran.r-project.org/web/packages/ggplot2/index.html
2. https://cran.r-project.org/web/packages/gmodels/index.html
3. http://www.r-project.org/
4. http://www.geneontology.org/
KEGG is a major publicly available pathway-related database (Kanehisa et al., 2008). Pathway enrichment analyses were performed to identify significantly enriched metabolic pathways or signal transduction pathways in source genes compared to the whole genome background. The calculated p-values were subjected to FDR correction, taking FDR ≤ 0.05 as a threshold for significance. Bubble charts were generated using ggplot2 in R.

Quantitative Reverse-Transcription Polymerase Chain Reaction (qRT-PCR)
Reverse transcription was performed using a PrimeScript RT Reagent Kit (Takara, Kusatsu, Japan). Fluorescence qRT-PCR was performed using an ABI 7500 detection system (Thermo Fisher Scientific, Waltham, MA, United States). Each 20 µL reaction mixture contained 2 µL of template cDNA, gene-specific primers (0.4 µM), 10 µL of SYBR Premix Ex Taq II (Tli RNaseH Plus, 2X), 0.4 µL of ROX Reference Dye II (50X), and 6 µL of sterilized RNase-free water. The reaction conditions were as follows: 50°C for 2 min, 95°C for 2 min, followed by 40 cycles of 95°C for 15 s and 60°C for 60 s. The relative expression levels of each circRNA normalized against the β-actin level were evaluated using the 2^{-ΔΔCt} method (Livak and Schmittgen, 2001). qRT-PCR was performed with five biological replicates. Divergent primers were designed to confirm the back-splice junction and were synthesized by Shanghai Sangon Biotech (Supplementary Table 1). The qRT-PCR products were tested by electrophoresis on 2% agarose gels. The amplification products were inserted into a pUCm-T vector for Sanger cloning and sequencing to determine their full length.

Statistical Analysis
Statistical analyses were performed using SPSS 21.0 (IBM SPSS, Armonk, NY, United States). Statistically significant differences were identified using Student’s t-tests (two-tailed). All values are presented as means ± standard deviation. Values of p < 0.05 were considered statistically significant.

RESULTS
Overview of circRNA-Seq
In total, 1,043,148,336 and 1,032,539,926 clean reads were generated in the ARM and control groups, respectively. After removing the low-quality, poly-N-containing, and adapter-containing reads, 915,328,052 and 1,024,161,230 high-quality clean reads were obtained from the ARM and control groups, respectively. After using find_circ for mapping, 18,295 circRNAs were obtained. All identified circRNAs were novel. Six types of circRNAs were identified. From most to least frequent, they were categorized as follows: annot_exon, exon_intronmost, antisense, intergenic, one-exon, and intronic (Figure 2A). Most
of the circRNAs were approximately 200–400 nucleotides (nt) in length (Figure 2B).

**Time Course Analysis of circRNA Expression Profiles in Hindguts of the ARM and Control Group Rats**

An unsupervised hierarchical clustering and PCA were performed to determine the relationships between circRNA expression and spatiotemporal dynamics in the hindguts of normal rat embryos and those with ARMs. The circRNA expression patterns in the two groups formed distinct clusters (Figure 2C). As shown in Figure 2D, there were no significant changes in total circRNA expression levels in the groups over time. The majority of circRNAs annotated in the ARM group were also detected in the normal group (Figure 2E). In total, 10651 circRNAs were co-expressed in both groups, 3857 were expressed exclusively in the ARM
FIGURE 3 | Volcano plots showing up- and downregulated circRNAs in the ARM group and normal group. (A–F) Red dots represent upregulated and green dots represent downregulated circRNAs. A, ARM group; N, normal group.

FIGURE 4 | Overall expression patterns of circRNAs in the ARM group and normal group. (A) Expression patterns of circRNAs in the ARM group. (B) Expression patterns of circRNAs in the normal group. Red indicates significant enrichment ($p < 0.05$). The profile IDs are shown in the upper left corner, and the number of genes in the profile is shown in the lower left corner. (C) Venn diagram showing upregulated circRNAs in both the ARM group and normal group. (D) Venn diagram showing downregulated circRNAs in both the ARM group and normal group.
FIGURE 5 | (A–C) Volcano plots showing the up- and downregulated circRNAs in the ARM group and normal group. Red dots represent upregulated and green dots represent downregulated circRNAs. (D) Venn diagram showing the overlap among these circRNAs between the three time-points. (E) Hierarchical cluster analysis and heat map showing differentially expressed circRNAs between the ARM group and normal group. Colors from blue to red indicate circRNA expression levels from low to high. Each column in the heat map represents a mean of three biological replicates, and each row represents a circRNA. A, ARM group; N, normal group.
To characterize the changes in circRNA expression, we evaluated trends in the highly differentially expressed circRNAs in each group during development. We identified seven main circRNA profiles in the ARM and normal groups, and each showed a characteristic expression pattern (referred to as Profiles 1–7).
Profile 6 was significant in the ARM group (Figure 4A) and Profile 7 was significantly upregulated in the normal group (Figure 4B). We also analyzed the up- and downregulated circRNAs in both groups. We found that the expression levels of 60 circRNAs were higher in the normal group and 42 were higher in the ARM group (Figure 4C), whereas levels of 33 circRNAs were lower in the normal group and 47 were lower in the ARM group (Figure 4D). These findings suggested that the dysregulation of circRNAs plays a role in hindgut development.

Screening of Differentially Expressed circRNAs and Clustering Analysis

The expression profiles in hindgut samples from the normal and ARM groups were compared at key time-points during anus formation (GD14, GD15, and GD16). There were significant differences in the expression levels of 425 circRNAs \(|\text{Fc}| > 2, p < 0.05\). Compared to the normal group, 60 circRNAs were significantly upregulated and 58 were significantly downregulated in the ARM group on GD14,
94 circRNAs were significantly upregulated and 74 were significantly downregulated in the ARM group on GD15, and 65 circRNAs were significantly upregulated and 87 were significantly downregulated in the ARM group on GD16 (Figures 5A–C). The overlap among these circRNAs between the three time-points is shown in Figure 5D. A heat map of significant differential expression is shown in Figure 5E.

**Predicted ceRNA Networks**

We constructed ceRNA networks based on 425 differentially expressed circRNAs. According to StarBase v3.0 and TargetScan v7.2, among differentially expressed circRNAs, 55 (14 up- and 41 downregulated) were predicted to bind to 195 miRNAs and 947 mRNAs. The detailed information of the 55 differentially expressed circRNAs at the three time-points are shown in Table 1. The circRNA-miRNA-mRNA correlations were visualized using Cytoscape 3.7.1 (Figures 6, 7). According to a connectivity analysis, the top five circRNAs were novel_circ_001042, novel_circ_012639, novel_circ_003974, novel_circ_006611, and novel_circ_014878. novel_circ_001042 was predicted to interact with 64 miRNAs and 328 mRNAs. These included the wingless-type MMTV integration site family
(WNT) signaling pathway-related genes Fzd4, Ccnd3, and Dkk2, the transforming growth factor-β (TGF-β) signaling pathway-related gene Bmp2, and the mitogen-activated protein kinase (MAPK) signaling pathway-related genes Flna, Egfr, Mef2c, Dusp3, and Cacng6. This complicated ceRNA network suggested that novel_circ_001042 played regulatory roles in various pathways via miRNAs and their targets during hindgut development.

**Enrichment of Differentially Expressed circRNAs**

Since circRNAs can directly regulate source genes, we conducted an enrichment analysis of source genes of circRNAs. According to a GO analysis, circRNA source genes were enriched for 33 GO terms in the three main categories, biological processes, molecular functions, and cellular components (**Figure 8A**). The top 20 KEGG pathways are listed in **Figure 8B**. The enriched pathways included the WNT pathway and the MAPK pathway, which are closely associated with embryonic development and ARMs.

For non-coding RNAs, functional annotations are determined by the functions of associated mRNAs. We evaluated potential functions of significantly up-regulated and down-regulated circRNAs in the ARM and normal groups by an enrichment analysis of target genes. Based on a GO analysis, the target genes related to the circRNA signature were involved in various biological processes, including single-organism processes, cellular processes, and metabolic process (**Figure 8C**). In a KEGG pathway analysis, endocytosis and PI3K-Akt were identified as the most significant pathways (**Figure 8D**).

**Validation of Sequencing Data**

To validate the circRNAs identified by RNA-Seq, we randomly selected 7 up- and 7 downregulated circRNAs for validation by qRT-PCR. Among the 14 differentially expressed circRNAs, novel_circ_008138, novel_circ_001042, novel_circ_012938, novel_circ_01300, novel_circ_016600, and novel_circ_015194 were differentially expressed on GD14; novel_circ_012938, novel_circ_01300, novel_circ_016600, and novel_circ_015194 were differentially expressed on GD15; novel_circ_011174 was differentially expressed at two time-points (GD14 and GD15) ($p < 0.05$; **Supplementary Figure 1A**). The relative expression levels...
were consistent with the RNA-Seq data. Sanger cloning and sequencing results for novel_circ_002017 and novel_circ_008138 are shown in Supplementary Figures 1B,C. Agarose gels electrophoresis was used to validate that a single product had been amplified are shown in Supplementary Figure 1D.

DISCUSSION

ARMs are complex deformities determined by many factors. Genetic factors have an important influence on the etiology of ARMs (Khanna et al., 2018). The roles of circRNA in embryonic development and congenital malformations of the gastrointestinal system have rarely been reported. Different from most RNAs involved in translation, circRNAs often act at the level of gene expression (Esteller, 2011). The versatility of circRNA expression is mainly reflected in post-transcriptional regulation, which is usually achieved by RNA–RNA interactions (Guil and Esteller, 2015). Studies of circRNAs related to gastrointestinal development are very limited. High-throughput sequencing has confirmed that circRNA expression levels differ among anatomical parts of the digestive tract, and these differences may be closely related to cell proliferation and differentiation during development (Danan et al., 2012; Salzman et al., 2013; Shen et al., 2017). Increasing number of studies have shown that circRNAs, acting as miRNA sponges, inhibit miRNAs and eventually mediate the expression of mRNAs (Hansen et al., 2013). Peng et al. (2017) found that circ-ZNF609 participates in the pathogenesis of congenital megacolon via the mir-150-5P/AKT3 pathway. Zhou et al. performed a sequencing analysis of human congenital megacolon specimens and found that the down-regulation of circRNAs/miRNAs and inhibition of the expression of the mir-1324 target gene PLCB1 (Zhou et al., 2018). However, the precise circRNAs with key regulatory roles in the normal development of the anus and rectum and the mechanism by which these circRNAs affect the pathogenesis of anorectal atresia and rectourethral fistula are unclear. Therefore, further research is required on this topic.

According to previous morphological studies, urorectal fistulas in rat embryos with ARMs can be explained by a failure of the fusion of the urorectal septum with the cloacal membrane during the critical period of anorectal development. Deduced proliferation and massive apoptosis play important roles in the development of ARMs (Qi et al., 2000a,b, 2002). In the present study, 18,295 circRNAs were identified at key time-points (GD14, GD15, and GD16) during anorectal development in the rat hindgut by high-throughput sequencing. We analyzed expression patterns and performed a bioinformatics analysis of circRNAs that may be involved in the pathogenesis of ARMs. We detected aberrantly expressed circRNAs in a comparison of rat fetuses with ARMs and normal rat fetuses. After screening (|Fc| > 2, p < 0.05), we detected 118, 168, and 152 differentially expressed circRNAs at GD14, GD15, and GD16, respectively. We can infer that there is a temporal imbalance in circRNA expression.

Using a bioinformatics approach for target prediction, among 425 differentially expressed circRNAs, 55 had ceRNA network connectivity. Compared to the normal group, 14 genes were upregulated and 41 genes were downregulated, suggesting that reductions in circRNA levels were dominant in the ARM group and might contribute to the development of ARMs. A circRNA-miRNA-mRNA ceRNA network was also constructed. A KEGG pathway enrichment analysis of miRNA-target genes identified the WNT, TGF-β, and MAPK signaling pathways, which are closely related to ARM (Nakamura et al., 2007; Wong et al., 2013; Wang et al., 2015). GO terms of related genes provided compelling evidence that circRNAs can influence metabolic, regulatory, and cellular functions. Our results demonstrated that these differentially expressed circRNAs may play important roles in the development of ETU-induced ARMs.

As a member of the miR-17-92 family, miR-92 plays an important role in the regulation of early embryonic development as well as in tumorigenesis and differentiation in various cancers (Monzo et al., 2008; Jevnaker et al., 2011). Our research group has previously shown that miR-92a-2-5p could have a regulatory role in ARMs. The overexpression of miR-92a-2-5p leads to a decrease in proliferation and an increase in apoptosis in an intestinal cell line (Long et al., 2020). Our results based on RNA-seq data showed that rno-miR-92a-1-5p is related to the regulation of 90 mRNAs. novel_circ_001042, novel_circ_003974, and novel_circ_010342 were associated with rno-miR-92a-1-5p and may function as ceRNAs to regulate the progression of ARMs. Notably, most of the circRNAs identified in this analysis have not been studied previously. Therefore, further research should explore the relationships between circRNAs/miRNAs and ARMs based on our ceRNA-associated network.

CONCLUSION

In conclusion, a high-throughput sequencing approach was used to identify numerous differentially expressed circRNAs between the hindguts of normal rat embryos and those with ARMs on GD14–GD16. Using a bioinformatics approach, a circRNA-miRNA-mRNA network was constructed. The potential target genes of novel_circ_001042 may be closely related to the development of ARMs. circRNAs are candidate biomarkers for disease diagnosis and therapeutic targets owing to their high conservation, stability, and tissue-specificity. Our results provide novel insight into the roles of circRNAs in the development of ARMs and provide a valuable resource for further analyses of molecular mechanisms and signaling networks. The construction of regulatory networks could improve our understanding of the basic molecular mechanisms underlying ARM.

DATA AVAILABILITY STATEMENT

The datasets generated for this study can be found in NCBI GEO accession GSE159306: go to https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE159306.
ETHICS STATEMENT

The animal study was reviewed and approved by the Ethical Committee on Science and New Techniques of Shengjing Hospital of China Medical University.

AUTHOR CONTRIBUTIONS

All authors listed have made a substantial, direct and intellectual contribution to the work, and approved it for publication.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fgene.2021.605015/full#supplementary-material

Supplementary Figure 1 | (A) Fourteen circRNAs in rat embryonic hindgut tissues in the ARM group and normal group were validated by qRT-PCR. *p < 0.05, n = 5. (B,C) Two representative examples [novel_circ_000217 and novel_circ_008138] of qRT-PCR products confirmed by Sanger cloning and sequencing. (D) qRT-PCR products were visualized using agarose gels electrophoresis. GD, gestational day; A, ARM group; N, normal group.

Supplementary Table 1 | The sequences of the primers pairs.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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