Integration of microRNA–mRNA profiles and pathway analysis of plant isoquinoline alkaloid berberine in SGC-7901 gastric cancers cells

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Purpose: Berberine (BBR) is a traditional Chinese medicine normally used for gastroenteritis, and recent research found that it could fight against tumors. In this study, we focused on integrating miRNA sequencing and RNA sequencing of SGC-7901 gastric cancer cells treated by BBR to elucidate their underlying mechanisms.

Materials and methods: WST-1 assay and flow cytometry were used to check the effects of BBR on SGC-7901. miRNA sequencing and RNA sequencing were used to establish the miRNA and mRNA profiles of BBR-treated SGC-7901.

Results: The results showed that BBR could inhibit the proliferation of SGC-7901 cells and induce G1 arrest in cell cycle phase and apoptosis. A total of 1,960 upregulated genes and 4,837 downregulated genes were identified in the RNA sequencing and 347 upregulated and 93 downregulated miRNAs in the miRNA sequencing. A total of 78 novel miRNAs were also found. Gene Ontology and Kyoto Encyclopedia of Genes and Genomes analysis showed that the genes were related to pathways in cancer and metabolism. We also analyzed the miRNA–mRNA network of genes grouped into cell cycle, apoptosis, inflammation, metabolism, cell junction, acetylation process, TGF- β pathway, and Wnt signaling pathway.

Conclusion: BBR could inhibit the proliferation of SGC-7901 cells and induce apoptosis. Integrated analysis of microRNA–mRNA profiles is a promising approach to validate gene expression patterns associated with malignant phenotype and study the mechanisms of anticancer.

Keywords: berberine, gastric cancer, RNA sequencing, microRNA sequencing, integrated analysis

Introduction

Berberine (BBR) is a bioactive alkaloid that can be isolated from several Chinese herbs, such as Huang lian (*Rhizoma Coptidis*).¹ BBR is often used for the treatment of diarrhea and gastroenteritis, but many other pharmacological and biochemical effects of the alkaloid have been found recently. It has been reported that BBR is effective against a number of diseases including hyperlipidemia, diabetes, metabolic syndrome, and Alzheimer's disease,^{2–5} and a growing number of reports show that BBR has anticancer effects.^{6,7}

However, the diversity of BBR targets that mediate these pharmacological activities has not been well understood yet. It has been reported that BBR directly regulates gene expression by targeting TATA boxes in transcriptional regulatory regions as well as the poly(A) tail at the mRNA terminus, and the final outcome of gene regulation by BBR depends on the structure of the individual gene.⁸ Recent researches have shown that BBR could also regulate the expression of miRNAs in several diseases

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including cancers and diabetes.^{9,10} It was reported that treatment of 3T3-L1 cells with BBR inhibited cell viability and differentiation in a dose- and time-dependent manner, and significantly increased the expression levels of miRNA-27a and miRNA-27b.¹¹ In human hepatocellular carcinoma cells treated by BBR, miR-23a was upregulated and the upregulation of miR-23a by BBR is p53 dependent.¹² It was also reported that treatment of human U266 multiple myeloma cell line with BBR suppressed cell proliferation and IL-6 secretion in a dose- and time-dependent manner. Meanwhile, BBR dose dependently induced ROS generation, G2/M phase arrest, and apoptosis in U266 cells, and decreased the levels of miR-21 and Bcl-2.¹³

For the effects of BBR on gastric cancer, Zhang et al reported that BBR or D-limonene alone could inhibit the growth of MGC803 cells in a dose- and time-dependent manner, and the combination of BBR and D-limonene showed more remarkable effects on MGC803 cells by cell cycle arrest, ROS production, and apoptosis induction through the mitochondria-mediated intrinsic pathway compared with drugs used singly.¹⁴ You et al reported that BBR treatment significantly reversed cisplatin sensitivity and induced caspase-dependent apoptosis in cisplatin-resistant SGC-7901 and BGC-823 cells by modulating the miR-203/Bcl-w apoptotic axis.¹⁵

Although BBR has anticancer effects on gastric cancer cells, the mechanisms are not clear yet. More advanced tools with a large scale of data are required for the deep interpretation of the problems. Here, in this study, we performed an integrative analysis of the miRNA-mRNA expression profiles in BBR-treated SGC-7901 gastric cancer cells by high-throughput miRNA sequencing and mRNA sequencing technologies. We have integrated miRNA and mRNA expression profiles and regular laboratory analysis to understand the effects of BBR on gastric cancer and the mechanisms. Although BBR is an “old” medicine, it may become a novel agent for the treatment of gastric cancer.

Materials and methods

Cell culture

The human gastric cancer cell line SGC-7901 was purchased from American Type Culture Collection. Cells were cultured in Dulbecco's Modified Eagle's Medium (Gibco; Thermo Fisher Scientific Waltham, MA, USA), supplemented with 10% fetal bovine serum (Hyclone, Logan, UT, USA) at 37°C in a humidified atmosphere containing 5% CO₂. When needed, cells were treated with BBR (Beyotime Biotechnology Company, Shanghai, People's Republic of China,

molecular weight 371.82, purified by high-performance liquid chromatography) at different concentrations.

WST-1 assay

SGC-7901 cells were plated into a 96-well plate at a density of 4×10^3 /well and treated with BBR for 12 and 24 h at different concentrations. Then, cells were treated with WST-1 and culture was continued for 2.5 h according to the protocol of the WST-1 Assay Kit (Beyotime Biotechnology Company). Finally, the observation density was read at 450 nm by a plate reader (BioTek Epoch, Winooski, VT, USA). The percentage of proliferation inhibition was calculated using the following formula: $(1 - \text{the OD value of the BBR-treated group} / \text{the OD value of the control group}) \times 100\%$. The experiment was repeated at least three times.

Cell cycle and apoptosis detection by flow cytometry

SGC-7901 cells were seeded in six-well plates at a density of 1×10^5 /well and incubated with BBR at different concentrations for 24 h. Cells were harvested and washed with phosphate-buffered saline, and fixed with 70% ethanol at 4°C overnight. After staining with propidium iodide (PI) solution for 30 min, cells were collected on a FACScan flow cytometer (Accuri™ C6 Flow cytometer; BD Biosciences, San Jose, CA, USA) equipped with a 488 nm argon laser and analyzed using ModFit LT. Cells for detection of apoptosis were treated with BBR at different concentrations for 24 h, and stained with Annexin-V and PI (BD Pharmingen™; BD Biosciences) according to the manufacturer's instruction. Flow cytometry analysis was also performed by FACScan flow cytometer and analyzed by FlowJo 7.6.

Library construction and miRNA sequencing analysis

MiRNA isolation, library construction, and miRNA sequencing were carried out by Gene Denovo Biotechnology Company (Guangzhou, People's Republic of China). Total RNA was extracted by TRIzol reagent, and the RNA molecules in a size range of 18–30 nt were enriched by polyacrylamide gel electrophoresis. Then the 3' adapters were added and the 36–44 nt RNAs were enriched. The 5' adapters were then ligated to the RNAs as well. The ligation products were reverse transcribed by PCR amplification, and the 140–160 bp size PCR products were enriched to generate a cDNA library. The cDNA library was sequenced by Illumina HiSeq™2500, (San Diego, CA, USA) following the vendor's recommended protocol.

Library construction and RNA sequencing analysis

RNA isolation, library construction, and RNA sequencing were carried out by Gene Denovo Biotechnology Company. After total RNA was extracted, mRNA was enriched by Oligo (dT) beads, and the enriched mRNA was fragmented into short fragments and reverse transcribed into cDNA with random primers. Then the cDNA fragments were purified with QiaQuick PCR extraction kit, poly(A) added and ligated to Illumina sequencing adapters. The ligation products were size selected by agarose gel electrophoresis, PCR amplified, and sequenced using Illumina HiSeq™2500.

Gene Ontology and Kyoto Encyclopedia of Genes and Genomes pathway analysis

Enrichment analysis of the genes was conducted with Gene Ontology (GO) (<http://www.geneontology.org/>) and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway (<http://www.genome.jp/kegg/>). A heatmap chart was drawn by transforming the normalized data clustering using the gplots heatmap.2 of R program.

Statistical analysis

Statistical differences were determined by using SPSS 23.0 software. A *P*-value <0.05 was considered to be significant.

Results

Effect of BBR on the proliferation of SGC-7901 cells detected by WST-I assay

BBR at concentrations of 2.5, 5, 10, 20, and 30 μ M was used to treat SGC-7901 cells for 12 and 24 h, respectively, and the results showed that BBR could inhibit the proliferation

of SGC-7901 cells (Figure 1). BBR could still inhibit the proliferation of SGC-7901 cells at 48 h (Figure S1).

Effect of BBR on the cell cycle and apoptosis of SGC-7901 cells detected by flow cytometry

SGC-7901 cells were treated with BBR at different concentrations for 24 h, and the effect of BBR on the cell cycle and apoptosis of SGC-7901 cells was detected by flow cytometry. It was shown that the cells were accumulated in G1 phase, and less cells entered S phase (Figure 2). BBR could also induce the apoptosis of SGC-7901 cells, and the apoptosis was more obvious with the higher concentration (Figure 3). When the time of SGC-7901 cells treated by BBR was prolonged to 48 h, the number of dead cells was significantly increased in BBR-treated SGC-7901 cells compared to the control group (Figure S2), but there was no difference in apoptosis between the BBR groups and the control group.

Analysis of gene expression profile of RNA sequencing

According to the sequence data, 28,214,592 and 26,977,858 clean reads were identified in the control group and the BBR-treated group, respectively. In the control group, high-quality reads were 27,371,360 (97.01% in total) and in the BBR-treated group, high-quality reads were 26,319,510 (97.56% in total). Here, 1,960 upregulated genes and 4,837 downregulated genes were identified in the BBR-treated SGC-7901 cells compared to the control group (Figure 4).

To understand the functions of differentially expressed genes, all the differentially expressed genes were mapped to terms in the GO database and compared with the background

Figure 1 Effect of BBR on the proliferation of SGC-7901 cells by WST-I assay.

Note: The proliferation of SGC-7901 cells was inhibited by different concentrations of BBR treated for (A) 12 and (B) 24 h, respectively.

Abbreviation: BBR, berberine.

Figure 2 Effect of BBR at different concentrations on the cell cycle of SGC-7901 cells as detected by flow cytometry.

Notes: (A) Control; (B) 5 $\mu\text{mol/L}$; (C) 10 $\mu\text{mol/L}$; (D) 20 $\mu\text{mol/L}$. (E) The histogram showed that the cells treated by BBR at different concentrations were accumulated in G1 phase, and less cells entered S phase. * $P < 0.05$, ** $P < 0.01$.

Abbreviation: BBR, berberine.

(Figure 5). The GO has 3 ontologies, which are “biological process”, “cellular component”, and “molecular function”. In the “biological process” part, most genes were related to “cellular process”, “metabolic process”, and “single-organism process”, but few genes were related to “cell killing” and

“reproductive process”. In the “cellular component” part, most genes were related to the “cell part” and “organelle part”, and few genes were related with “cell junction” and “extracellular matrix”. In the “molecular function” part, most genes were related with “binding” and “catalytic activity”.

Figure 3 Effect of BBR at different concentrations on apoptosis of SGC-7901 cells as detected by flow cytometry.

Notes: (A) Control; (B) 5 μmol/L; (C) 10 μmol/L; (D) 20 μmol/L. (E) The histogram showed that BBR could induce the apoptosis of SGC-7901, and the apoptosis was more obvious with the higher concentration. * $P < 0.01$.

Abbreviation: BBR, berberine.

The KEGG assignments were used to classify functional annotations of the identified differentially expressed genes to further understand the biological functions (Figure 6). Top 20 pathways were shown in the result, including regulation of autophagy, mTOR signaling pathway, MAPK (mitogen-activated protein kinase) signaling pathway, insulin resistance, microRNAs in cancer, AMPK (AMP-activated protein kinase) pathway, pathways in cancer, and pathways related to metabolism.

Analysis of miRNA profile of miRNA sequencing

The result of the miRNA sequencing showed that 11,551,248 and 10,360,439 clean reads were identified in the control

group and the BBR-treated group, respectively. In the control group, high-quality reads were 11,135,724 (96.40% in total), and in the BBR-treated group, high-quality reads were 9,987,143 (96.40% in total). Here, 347 upregulated and 93 downregulated miRNAs were identified in the BBR-treated SGC-7901 cells compared to the control group (Figure 7).

The target genes of the upregulated or downregulated miRNAs were mapped to terms in the GO database (Figure 8). In the “biological process” part, most genes were related to “cellular process”, “metabolic process”, and “single-organism process”. In the “cellular component” part, most genes were related to the “cell”, “cell part”, and “organelle part”. In the “molecular function” part, most genes were related with “binding” and “catalytic activity”.

Figure 4 The scatter plot of the mRNA expression profile of SGC-7901 cells treated by BBR.

Note: A total of 1,960 upregulated genes and 4,837 downregulated genes were identified.

Abbreviation: BBR, berberine.

The top 20 pathways of the KEGG assignments of the target genes of differentially expressed miRNAs are shown in Figure 9, including Ras signaling pathway, Jak-STAT signaling pathway, P13K-Akt signaling pathway, Rap1 signaling

pathway, insulin signaling pathway, AMPK signaling pathway, cAMP signaling pathway, and Oxytocin signaling pathways.

Novel miRNAs and target genes

According to the results of miRNA sequencing, 78 novel miRNAs were found, 71 found in the control group and 55 found in the BBR-treated group. KEGG showed that the pathways of the target genes of differentially expressed miRNAs mainly focused to “calcium pathway”, “PI3K-Akt signaling pathway”, “Ras signaling pathway”, “cAMP signaling pathway”, and “pathways in cancer” (Figure 10).

Analysis of miRNA–mRNA network

Based on the result that BBR could inhibit the SGC-7901 cells, the gene expression profile was further analyzed in detail. Genes related to cell cycle, apoptosis, inflammation, and metabolism were picked up, and their expression level and miRNA–RNA network were further analyzed. TGF- β pathway and Wnt signaling pathway were also analyzed. In addition, genes related to cell junction and acetylation were also differentially expressed in the BBR-treated group. The merged heatmap is shown in Figure 11, and the merged miRNA–RNA relation network is shown in Figure 12. Several novel anticancer pathways were also found in BBR-treated SGC-7901 cells, such as Hippo pathway, Notch pathway, and FoxO pathway. The merged heatmap and miRNA–RNA relation network of the three pathways are shown in Figures S3 and S4, respectively.

Figure 5 The presentation of GO classification of the mRNA expression profile.

Abbreviations: GO, Gene Ontology; BBR, berberine.

Figure 6 The presentation of KEGG assignments of differentially expressed genes.
Abbreviation: KEGG, Kyoto Encyclopedia of Genes and Genomes.

Discussion

Gastric cancer is the fourth most common cancer and the second leading cause of cancer deaths worldwide. Although its incidence and mortality have been declining due to the decline in the prevalence of *Helicobacter pylori* infection and the improvements in food preservation and diet, the burden

of gastric cancer remains very high in several countries in Asia, including China.¹⁶ Most of the chemotherapy drugs for gastric cancer are cytotoxic, which have serious adverse reactions and directly affect the curative effect of gastric cancer. It is one of the effective ways to treat gastric cancer by looking for drugs from natural plants with obvious anti-tumor activity but little side effects.

Our data displayed in this paper demonstrated that BBR could inhibit the proliferation of SGC-7901 gastric cancer cells and induce G1 phase arrest and apoptosis. The KEGG analysis of the RNA sequencing showed that BBR was involved in autophagy and mTOR and MAPK signaling pathways. Pathway analysis of the miRNA target genes included cell cycle, Ras signaling pathway, Jak-STAT signaling pathway, and P13K-Akt signaling pathway. In the gene expression profiling, we also found genes related to cell cycle and apoptosis, and genes in TGF- β pathway and Wnt signaling pathway. These data of miRNA sequencing and RNA sequencing supported our results that BBR could inhibit the proliferation and induce apoptosis of SGC-7901 cells, and provided us the clues that BBR may exert its anti-tumor activities through these pathways. Li et al reported that inactivation of p38 MAPK, extracellular signal-regulated

Figure 7 The miRNA expression profile of SGC-7901 cells treated by BBR.
Note: A total of 347 upregulated and 93 downregulated miRNAs were identified.
Abbreviation: BBR, berberine.

Figure 8 The presentation of GO classification of the upregulated or downregulated miRNA target genes.

Abbreviations: GO, Gene Ontology; BBR, berberine.

Figure 9 The presentation of KEGG assignments of the upregulated or downregulated miRNA target genes.

Abbreviation: KEGG, Kyoto Encyclopedia of Genes and Genomes.

Figure 10 The presentation of KEGG assignments of the upregulated or downregulated novel miRNA target genes.

Abbreviation: KEGG, Kyoto Encyclopedia of Genes and Genomes.

kinase 1/2, and c-Jun N-terminal kinase by BBR contributed to the decreased proliferation and tumorigenesis of MGC-803 gastric cancer cells.¹⁷ Yi et al demonstrated that Akt-related mitochondrial pathway may contribute to the BBR-induced apoptosis in human gastric cancer cells and might represent an important molecular basis for BBR to act as an anticancer agent.¹⁸ Another research demonstrated that BBR could modulate PI3K-AKT and MAPK signaling pathways in thyroid carcinoma cells, which leads to mitochondrial apoptosis, G0/G1 cell cycle arrest and suppressive migration.¹⁹ Okubo et al showed that BBR promptly localized to the nucleus after treatment in HL-60 human promyelocytic leukemia cells and induced apoptotic cell death by activation of caspase-3 and caspase-8.²⁰ It has been reported that BBR inhibits colon tumor formation through inhibition of Wnt/ β -catenin signaling and might be a promising drug for the prevention of

colon cancer.²¹ All the conclusions from these papers have shown that it is an effective method to study the mechanisms of BBR through high-throughput miRNA sequencing and RNA sequencing technologies.

The results of KEGG analysis also demonstrated that BBR is also involved in insulin resistance and metabolism pathways, inflammation pathways, cell junction, and acetylation. Wang et al reported that BBR was able to significantly reduce the alcohol-induced gastrointestinal injury, inhibit increases of alcohol-induced TNF α and IL-1 β expression in gastrointestinal mucosa as well as their upstream signals TLR2 and TLR4, and regulate cytokines that modulate tight junctions.²² Zhao et al demonstrated that BBR and nandinine attenuated insulin resistance in adipocytes by inhibiting inflammation in an AMP-activated protein in a kinase-dependent manner, and BBR and nandinine may be

Figure 11 The merged heatmap of genes related to cell cycle, apoptosis, inflammation, metabolism, TGF- β pathway, Wnt signaling pathway, cell junction, and acetylation.

used as dietary supplements for obesity treatment.²³ It has also been reported that ischemia/reperfusion-induced intestinal tight junction dysfunction could be improved by BBR, thereby demonstrating the therapeutic potential of BBR for intestinal ischemia/reperfusion injury.²⁴

Among the epigenetic changes, histone acetylation has been recognized as a fundamental process that strongly affects gene expression regulation, and disruption of this phenomenon has been linked to carcinogenesis.²⁵ Seldom research is about the effects of BBR on acetylation and

well as the mechanisms involved in cell cycle, apoptosis, inflammation, metabolism, cell junction, acetylation process, TGF- β pathway, and Wnt signaling pathway. Based on the results of the mRNA expression profile, genes related to cell cycle, apoptosis, inflammation, metabolism, TGF- β pathway, Wnt signaling pathway, cell junction, and acetylation were further analyzed in the miRNA-mRNA network. We also found some novel miRNAs.

Conclusion

We have got the miRNA and mRNA expression profiles of SGC-7901 gastric cancer cells treated with BBR as

Figure 12 (Continued)

Figure 12 The merged miRNA-RNA relation network of genes related to cell cycle, apoptosis, inflammation, metabolism, TGF- β pathway, Wnt signaling pathway, cell junction, and acetylation.

of BBR on gastric cancer by high-throughput sequencing technology with network pathway analysis. Our findings have illustrated the power of high-throughput sequencing for large-scale biological data sets, which may be a promising method to provide insight into the action mechanisms of BBR and help to guide its predictions.

Data availability

The RNA sequencing, miRNA sequencing, and the novel miRNAs data have been uploaded to figshare (<https://figshare.com/>), doi:10.6084/m9.figshare.5738577.

Anyone who is interested in the study of miRNAs, BBR, cancers, and related fields can download the data for free.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

Figure S1 Inhibition on the proliferation of SGC-7901 cells treated with BBR for 48 h as checked by WST-I assay.

Note: The proliferation of SGC-7901 cells was inhibited by different concentrations of BBR treatment for 48 h.

Abbreviation: BBR, berberine.

Figure S2 Effect of BBR treatment for 48 h on the apoptosis of SGC-7901 cells as detected by flow cytometry.

Notes: (A) Control; (B) 5 μmol/L; (C) 10 μmol/L; (D) 20 μmol/L. (E) The histogram showed that the number of dead cells was significantly increased in BBR-treated SGC-7901 cells compared to the control group. * $P < 0.01$. The number of dead cells was significantly increased in BBR-treated SGC-7901 cells compared to the control group, but there was no difference in apoptosis between the BBR group and the control group.

Abbreviation: BBR, berberine.

Figure S3 The merged heatmap of genes related to Hippo pathway, Notch pathway, and FOXO pathway.

Figure S4 The merged miRNA–RNA relation network of genes related to Hippo pathway, Notch pathway, and FOXO pathway.

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