IRE-1 Dependent Expression of Phosphoribosyl Pyrophosphate Synthetase Genes in U87 Glioma Cells: Effect of Glucose or Glutamine Deprivation

Dmytro O Minchenko1,2, Iana A Garmash1, Yulia M Bashta1, Ganna S Kustkova1, Yulianna D Zalesna3, Andreas Bikfalvi4 and Oleksandr H Minchenko1*

1Department of Molecular Biology, Palladin Institute of Biochemistry, National Academy of Sciences of Ukraine, Kyiv 01601, Ukraine
2Departments of Pediatrics, National O.O. Bogomolets Medical University, Kyiv 01601, Ukraine
3Department of Neurology and Reflexotherapy, Shupik National Medical Academy of Post-Graduate Education, Kyiv 04112, Ukraine
4INSERM U1029, Angiogenesis and Cancer Microenvironment Laboratory, University Bordeaux 1, Talence 33405, France

Abstract

Nutrient deprivation conditions are capable to induce the endoplasmic reticulum stress and are responsible for regulation of the expression of numerous growth factors which control the cell proliferation and angiogenesis in malignant tumors, including glioma. The ERN1 mediated endoplasmic reticulum stress response-signalling pathway is tightly linked to the proliferation because the blockade of ERN1 function suppresses the tumor growth via specific changes in the cell transcriptome. Activation of the pentose phosphate pathway is an important factor of enhanced cell proliferation and tumor growth because it controls the synthesis of nucleotides and respectively nucleic acids. We studied the effect of blockade the ERN1 signaling enzyme function in glioma cells on the expression of phosphoribosyl pyrophosphate synthetase (PRPS), an important enzyme of pentose phosphate pathway, as well as its dependence of glucose and glutamine deprivation.

The expression of different genes of PRPS enzyme in glioma cells with complete suppression of ERN1 enzyme activity was measured by qPCR. For glucose and glutamine deprivation conditions the cells were exposure in the glucose or glutamine free medium for 16 hrs.

The expression of PRPS1 and PRPS2 genes as well as phosphoribosyl pyrophosphate synthetase associated protein 1(PRPSAP1) is significantly increased in U87 glioma cells with suppressed function of ERN1 gene, but the expression PRPSAP2 gene is decreased. It was also shown that glucose or glutamine deprivation leads to suppression of PRPS1 and PRPS2 gene expressions in U87 glioma cells; however, the blockade of ERN1 function in glioma cells modifies the effect of glutamine deprivation. Moreover, the expression of PRPSAP1 and PRPSAP2 genes was resistant to glucose deprivation condition in control glioma cells, but increased in glioma cells with ERN1 loss of function.

Results of this investigation demonstrated that the blockade of ERN1 signaling enzyme function affects the expression of genes encoding PRPS enzyme subunits in different ways and that dysregulation of the expression of important regulatory subunits of PRPS (PRPSAP1 and PRPSAP2) possibly changes the activity of this enzyme and participates in suppression of glioma growth via pentose phosphate pathway.

Keywords: Gene expression; Pentose phosphate pathway; Phosphoribosyl pyrophosphate synthetase; ERN1; Glucose deprivation; Glioma cells

Abbreviations: PRPS: Phosphoribosyl Pyrophosphate Synthetase; PRPSAP: Phosphoribosyl Pyrophosphate Synthetase Associated Protein; ERN1: Endoplasmic Reticulum-Nuclei-1

Introduction

Malignant gliomas are highly aggressive tumors and are characterized by marked angiogenesis and extensive tumor cell invasion into the normal brain parenchyma and to date there is no efficient treatment available. Pentose phosphate pathway is necessary for purine metabolism as well as nucleotide biosynthesis and plays an important role in the regulation of proliferative processes and possibly participates in endoplasmic reticulum stress, which is an important factor of tumor growth and many other diseases [1-5]. The endoplasmic reticulum is a key organelle in the cellular response to ischemia, hypoxia, and some chemicals which activate a complex set of signaling pathways named the unfolded protein response. This adaptive response is activated upon the accumulation of misfolded proteins in the endoplasmic reticulum and is mainly mediated by endoplasmic reticulum to nuclei-1 signaling enzyme (ERN1); also named by inositol requiring enzyme-1alpha, IRE1). ERN1 is the dominant signaling pathway of the unfolded protein response to the accumulation of misfolded proteins and represents a key regulator of life and death processes [6-9]. As such, it participates in the early cellular response to the accumulation of misfolded proteins in the lumen of the endoplasmic reticulum [10,11].

Two distinct catalytic domains of the bifunctional signaling enzyme ERN1 were identified: a serine/threonine kinase and an endoribonuclease which contribute to ERN1 signalling. The ERN1-
The endoplasmic reticulum stress is recognized as an important determinant of cancer and contributes to the expression profile of many regulatory genes resulting in proliferation and apoptosis [3,9,13]. Moreover, the malignant tumors use the endoplasmic reticulum stress and ischemia for activation of proliferation, because the endoplasmic reticulum stress response-signalling IRE-1 pathway is linked to the apoptosis as well as cell death processes and suppression of IRE-1 gene function significantly decreases the tumor growth [15,16].

The pentose phosphate pathway has an important role in the regulation of glycolysis and proliferation during different stress conditions and participates in control of apoptotic cell death. Phosphoribosyl pyrophosphate synthetase synthetase plays a central role in tumor growth because it catalyzes the phosphoribosylation of ribose-5-phosphate to 5-phosphoribosyl-1-pyrophosphate which is necessary for purine metabolism and nucleotide biosynthesis [4].

The aim of this study was the investigation of the expression of different phosphoribosyl pyrophosphate synthetase genes in U87 glioma cells with ERN1 deficiency and decreased proliferative activity to find some molecular mechanisms between suppression of ERN1 function, glioma growth from these cells and the activity of pentose phosphate pathway.

Materials and Methods

The glioma cell line U87 was obtained from ATCC (USA) and grown in high glucose (4.5 g/l) Dulbecco’s modified Eagle’s minimum essential medium (DMEM; Gibco, Invitrogen, USA) supplemented with glutamine (2 mM), 10% fetal bovine serum (Equitech-Bio, Inc., USA), penicillin (100 units/ml; Gibco) and streptomycin (0.1 mg/ml; Gibco) at 37°C in a 5% CO2 incubator. In this work we used two sublines of this glioma cell line. One subline was obtained by selection of stable transfected clones with overexpression of pcDNA 3.1 vector, which was used for creation of dominant-negative ERN1 constructs (dnERN1). This untreated subline of glioma cells (control glioma cells) was used as control 1 in the study of glucose and glutamine deprivation conditions on the expression level of PRPS1, PRPS2, PRPSAP1, and PRPSAP2 genes. Second subline was obtained by selection of stable transfected clones with overexpression of dnERN1 and has suppressed both protein kinase and endoribonuclease activities of this bifunctional sensing and signaling enzyme of endoplasmic reticulum stress. These cells were obtained from Prof. M. Moenner (France) [15,16].

The expression level of studied genes in these cells was compared with cells, transfected by vector (control 1), but subline which overexpress dnERN1 was also used as control 2 for investigation the effect of glucose and glutamine deprivation conditions on the expression level of these genes under blockade ERN1 function. For creation of these deprivation conditions we changed the growing medium in culture plates with DMEM without glucose or glutamine and exposed to these conditions for 16 hrs.

The enzymatic activity of ERN1 in glioma cells that overexpress dnERN1 construct was previously analyzed by detection of XBP1 alternative splice variant (XBP1s), a key transcription factor in ERN1 signaling, and the level of ERN1 phosphorylated isoform in cells treated by tunicamycin (0.01 mg/ml for 2 hours) [17].

Total RNA was extracted from glioma cells using Trizol reagent according to manufacturer protocol (Invitrogen, USA). Cells were extracted with 2 ml of Trizol reagent and 0.4 ml of chloroform was added to cell lysate, mix and spin down. RNA was precipitated with an equal volume of 2-propanol. RNA pellets were washed with 75% ethanol and dissolved in nuclease-free water. For additional purification the RNA samples were re-precipitated with ethanol and dissolved again in nuclease-free water.

Quanti Tect Reverse Transcription Kit (QIAGEN, Germany) was used for cDNA synthesis according to manufacturer protocol. For this aim 1 µg of RNA was incubated for 2 min at 42°C with gDNA Wipeout buffer, and then placed immediately on ice. Add reverse-transcription master mix, containing QuantiScript Reverse Transcriptase, Quantscript RT Buffer and RT Primer Mix, incubated for 15 min at 42°C for synthesis of cDNA, and then incubated for 3 min at 95°C to inactivate transcriptase.

The expression level of PRPS1, PRPS2, PRPSAP1, and PRPSAP2mRNA were measured in glioma cell line U87 and its subline (clone 1C5) by real-time quantitative polymerase chain reaction using “Mx 3000PQPCR” (Stratagene,USA) and Absolute qPCR SYBR GreenMix (THERMO Fisher Scientific, A bgene House, UK). For qPCR analysis initial step was incubation for 10 min at 95°C, and then 40 cycles included incubation for 0.5 min at 95°C, for 0.5 min at 55°C and for 0.5 min at 72°C. At the end of qPCR was final step for detection of dissociation curve. Polymerase chain reaction was performed in triplicate.

For amplification of phosphoribosyl pyrophosphate synthetase synthetase 1 (PRPS1) cDNA we used next primers: forward (5’-AAAGACGGAGAAGGCAAT-3’) and reverse (5’-GACCGGAAGATTCCATGA-3’) primers. The nucleotide sequences of these primers correspond to sequences 786-805 and 971-952 of human PRPS1 cDNA (GenBank accession number NM_002764). The size of amplified fragment is 186 bp.

Two other primers were used for real time RCR analysis of phosphoribosyl pyrophosphate synthetase synthetase 2 (PRPS2) cDNA: forward (5’-AAACATTTGGCGAGTGGAAG-3’) and reverse (5’-TGACGAGACGAGCTCAAGG-3’) primers. The nucleotide sequences of these primers correspond to sequences 613-632 and 957-948 of human PRPS2 cDNA (GenBank accession number NM_002765). The size of amplified fragment is 345 bp.

For amplification of phosphoribosyl pyrophosphate synthetase synthetase-associated protein 1 (PRPSAP1) cDNA we used next primers: forward (5’-GTCCTATGCGGAGACTGC-3’) and reverse (5’-TGG CGCCTCCTGCTTTCAG-3’). These oligonucleotides correspond to sequences 1088-1107 and 1366-1347 of human PRPSAP1 cDNA (GenBank accession number NM_002766). The size of amplified fragment is 279 bp.

For amplification of phosphoribosyl pyrophosphate synthetase synthetase-associated protein 2 (PRPSAP2) cDNA we used next primers: forward (5’-CACCCATGTCGAGAAGTGTG-3’) and reverse 5’. TTAATCTTGGGGCAGTGGAG-3’. These oligonucleotides correspond to sequences 698-717 and 980-961 of human PRPSAP2
cDNA (GenBank accession number NM_002767). The size of amplified fragment is 329 bp.

The amplification of beta-actin (ACTB) cDNA was performed using forward-5' - GAGCTCGAGCGAGATGG-3' and reverse-5' - AGCATCTGTTGGGCGTACAG-3' primers. These primers nucleotide sequences correspond to 747-766 and 980-961 of human ACTB cDNA (GenBank accession number NM_001101). The size of amplified fragment is 234 bp. The expression of beta-actin mRNA was used as control of analyzed RNA quantity. The primers were received from “Sigma” (USA).

An analysis of quantitative PCR was performed using special computer program “Differential expression calculator” and statistical analysis-in Excel program. The values of PRPS1, PRPS2, PRPSAP1, and PRPSAP2mRNA expressions were normalized to the expression of beta-actin mRNA and represent as percent of control (100%). All values are expressed as mean ± SEM from triplicate measurements performed in 4 independent experiments. The statistical significance of differences was calculated by Students t-test. p<0.05 was considered to be a significant difference.

Results

As shown in Figure 1, the expression level of phosphoribosyl pyrophosphate synthase genes PRPS1, PRPS2, and PRPSAP1 significantly increased in glioma U87 cells with knockdown of ERN1, a major sensor and signaling enzyme of endoplasmic reticulum stress, being more significant for PRPS1 gene. At the same time, the expression of PRPSAP2 gene is decreased in these glioma cells.

It was also shown that exposure of U87 glioma cells under glucose or glutamine deprivation conditions led to suppression of PRPS1 gene expression both in control glioma cells and cells with ERN1 knockdown, but more strong effect of glutamine deprivation condition was observed in glioma cells without ERN1 knockdown (Figure 1). As shown in Figure 2, the similar changes were observed in the expression level of PRPS2 gene in control glioma cells as well as in cells with suppressed function of ERN1. For this gene was also shown more strong effect of glutamine deprivation condition in glioma cells with ERN1 knockdown.

At the same time, the expression of PRPSAP1 gene was resistant to glucose and glutamine deprivation conditions in control glioma cells but suppression of ERN1 signaling enzyme function is modified the sensitivity of this gene expression to both glucose and glutamine deprivation conditions (Figure 4). Thus, the expression of PRPSAP1 gene in glioma cells with ERN1 loss of function is increased in glucose deprivation condition and decreased in glutamine deprivation condition. It was also showed that PRPSAP2 gene expression in control glioma cells was suppressed in glutamine deprivation condition but resistant to glucose deprivation condition (Figure 5). However, in glioma cells with ERN1 knockdown the expression of PRPSAP2 was increased both in glucose and glutamine deprivation conditions.

Discussion

We have studied the expression of four different genes of one enzyme, phosphoribosyl pyrophosphate synthase, encoding two catalytic and two regulatory subunits of associated proteins, in U87 glioma cells with ERN1 deficiency and decreased proliferative activity...
with the goal to identify some molecular mechanisms between suppression of ERN1 function, glioma growth from these cells and the activity of pentose phosphate pathway. However, little is known about the molecular mechanisms of the regulation of different PRPS genes, in particular, its participation in ERN1 signalling pathway of endoplasmic reticulum stress [5,18]. Results of this investigation demonstrate that expression of different subunits of phosphoribosyl pyrophosphate synthase in glioma cells significantly depends from endoplasmic reticulum stress, in particular, its ERN1 signalling system and that the blockade of ERN1 enzyme function affects the expression of different subunits of PRPS enzyme.

Moreover, we have shown that the glucose deprivation condition also affects the expression of PRPS1 and PRPS2 genes in control as well as ERN1 knockdown U87 glioma cells and that this effect of glucose deprivation on the expression of both PRPS1 and PRPS2 gene was similar. At the same time, the expression of PRPS associated protein genes (PRPSAP1 and PRPSAP2) is resistant to glucose deprivation condition in control glioma cells but is increased in glioma cells with ERN1 loss of function. Thus, the glucose deprivation condition affects the expression of different phosphoribosyl pyrophosphate synthase genes except PRPSAP1 in control glioma cells; however, ERN1 knockdown significantly modifies the effect of glucose deprivation on the expression of these phosphoribosyl pyrophosphate synthase genes. At the same time, the exposure cells to glucose deprivation condition leads to suppression of PRPSAP1 gene expression only in U87 glioma cells without ERN1 enzyme function.

Moreover, the blockade of ERN1 gene function leads to up-regulation of PRPSAP2 gene expression in U87 glioma cells under glucose deprivation condition. This data conforms to down-regulation of this gene expression in glioma cells without ERN1 enzyme function. Thus, the effect of glucose deprivation condition on the expression of different phosphoribosyl pyrophosphate synthase genes in U87 glioma cells depends from ERN1 function. These results are consistent with data of other investigation where was shown the dependence of different gene expressions under nutrient deprivation condition as well as hypoxia from ERN1 signalling enzyme function [17,19-22]. There are different effects of ERN1 blockade on hypoxic or nutrient deprivation regulations of gene expression, enhance, reduce or eliminate these regulations.

Results of this investigation clearly demonstrated that the blockade of ERN1 signalling enzyme function affects the expression of genes encoding PRPS enzyme subunits in different ways and that dysregulation of the important regulatory subunits of PRPS (PRPSAP1 and PRPSAP2) possibly participates in suppression of glioma growth also via pentose phosphate pathway. It is possible that...
these important regulatory subunits of PRPS enzyme could be used as targets for developing of new approaches for suppression of glioma growth. However, the detailed molecular mechanisms for PRPS gene expressions under endoplasmic reticulum stress condition are complex and the biological significance remains to be determined.

References

22. Minchenko DO, Kharkova AP, Hubenia OV, Minchenko OH (2013) Insulin receptor, IRS1, IRS2, INSIG1, INSIG2, RRA D, and BAIAP2 gene expressions under endoplasmic reticulum stress condition are complex and the biological significance remains to be determined.