Inhibition of cell proliferation by wortmannin in T98G cells involved induced inhibition of NF-κB transcriptional activity

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Abstract

Wortmannin is an important regulator of Phosphoinositide 3-kinase (PI(3)K) signaling pathway. Changes in expression and activity of PI3-kinase and PDGF are major positive and negative regulators, respectively, of the PI3-kinase pathway, which regulates growth, survival, and proliferation. Here we have shown that cells dosed with platelet-derived-growth-factor (PDGF) and /or wortmannin, an inhibitor of PI3 kinase, proliferated at expected rates with respect to cells deprived of any additions. Cells with added platelet-derived-growth-factor (PDGF) multiplied substantially faster than naturally growing cells-some thirty percent. As anticipated, cells given only wortmannin divided over forty percent slower than cells without any dosage. Additionally, cells transfected with a luciferase reporter carrying a consensus sequences of the nuclear factor NF-κB binding site and treated with wortmannin inhibited the activation of luciferase in T98G cells. However, this inhibition was not affected by the treatment of PDGF. Our data indicate that Wortmannin and PDGF play different role in the control of expression of Phosphoinositide 3-kinase in glioma T98G cell line.

Introduction

Wortmannin is a cell-permeable, fungal metabolite that acts as a potent, selective and irreversible inhibitor of phosphatidylinositol 3-kinase (PI3K) [1-3]. This kinase Phosphoinositide-3-kinase phosphorylates PtdIns[4,5] P2 (Phosphatidylinositol 4,5-bisphosphate) to generate phosphatidylinositol 3,4,5-trisphosphate (PIP3). PIP3 plays a key role by recruiting PH domain-containing proteins to the membrane, including AKT1 and PDPK1, activating signaling cascades involved in cell growth, survival, proliferation, motility and morphology [3,4]. Involved in immune, inflammatory and allergic responses. PI3Ks have been studied intensively since the discovery of a PI3K activity associated with two viral oncoproteins: polyoma middle T (mT) antigen and pp60v-src [5] having an important role not only in growth regulation but also in various other cellular responses [6,7]. In addition, following recent findings PI3K activation prevents cell death [8], that PI3K is a retrovirus-encoded oncogene [9], and that PI3K mutations increase lifespan in Caenorhabditis elegans [10]. Three mammalian PI3Ks sharing 42–58% amino acid sequence identity has been cloned and designated p110α, p110β, and p110δ [11]. In addition, Wortmannin, also potently inhibits mammalian polo-like kinase [12]. The fungal metabolite wortmannin is a potent inhibitor of the lipid and protein kinase activities of class I PI3Ks [1]. The 50% inhibitory concentration (IC50) values for inhibition of the isolated enzymes are all in the range of 1–10 nM. Wortmannin irreversibly inhibits p110α by reacting covalently with lysine-802 [13,14], a residue required for catalytic activity that is conserved in all phosphoinositide kinases (and in protein kinases). Wortmannin have been used extensively to study the physiological role of class I PI3Ks in various cellular responses. On the other, considerable circumstantial evidence has accumulated which strongly implicates platelet derived growth factor (PDGF) as an autocrine and/or paracrine agent in the development of numerous human tumor types [15]. PDGF and its corresponding receptor (PDGFR) have important roles in the regulation of human physiological functions [16]. It has been shown that PDGF plays a role in blood vessel formation, healing regulation and maintenance of interstitial fluid pressure [17,18]. The receptors of PDGF exists in two forms of strong structural homology: PDGFRα and PDGFRβ. It is normally expressed by the cells of the supporting tissue; it plays a role in embryonic development and tissue scarring [19]. Its role in the tumorigenesis of certain cancers is currently well documented. In dermatofibrosarcoma protuberans (DFSP), PDGFR is constituatively activated inducing intracellular oncogenic signals [20]. Similarly, glioblastomas are characterized by autocrine activation of PDGFR [21]. The PDGF family consists of four protein chains that form five biologically active dimers (PDGF-AA, -AB, -BB, -CC, and -DD) [22]. Ligand binding induces dimerization and autophosphorylation of PDGFRs. Phosphorylated tyrosine residues function as docking sites for signal transduction proteins with SH2 domains [23]. Proteins interacting with PDGFRs

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include SHP2, PI 3-kinase, Src family kinases, Grb2, and Shc [24]. In addition, PDGF stimulates cell proliferation and has also been shown to be a chemoattractant for several cell types including fibroblasts, smooth muscle cells, neutrophils, and monocytes [25].

Materials and methods

Cell culture

The T98G brain tumor cell line, established from human glioblastoma, was obtained from the American Type Culture Collection (ATCC, Manassas, VA, USA). The cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) (Invitrogen, Carlsbad, CA, USA) supplemented with 10% fetal bovine serum (FBS), 1 mM sodium pyruvate, 100 U/ml penicillin G, 100 μg/ml, 1 streptomycin, 2 mM glutamine, 1 mM MEM non-essential amino acids and 1 μM 2-mercaptoethanol in a 5% CO2 incubator at 37°C. The cells were subcultured once in 3–5 days.

Reagents

Tris-borate-EDTA and acrylamide:biacrylamide (29:1) were obtained from Bio-Rad (Richmond, CA, USA). Lipofectamine was obtained from Life Technologies, Inc., USA. Complete Mini EDTA-free protease inhibitor cocktail tablets and Annexin V-Fluor were purchased from Roche Diagnostics GmbH (Mannheim, Germany). Phorbol 12-myristate-13-acetate (TPA) (50 ng/ml) and wortmannin (10 nM) and PDGF (100 ng/ml) were purchased from Stratagene Inc. (La Jolla, CA, USA). Luciferase assay reagent, lysis buffer and the pGL-2 luciferase vector were obtained from Promega (Madison, WI). Recombinant human TNF-α was obtained from Sigma-Aldrich (St. Louis, MO).

DNA synthesis assay

DNA replication rate was measured by a (3H)-Thymidine incorporation assay. Cells were seeded in 96-well tissue culture plates (1,000 cells/well) and treated with wortmannin (10 nM) or PDGF (100 ng/ml) (see Treatment of Cells with wortmannin and PDGF). Twenty four, 48, 72, 96 and 120 hours after treatment with wortmannin and/or PDGF (3H)-Thymidine (0.5 μCi/well) was added for 3 hours. Cells were harvested with a PhD-200A cell harvester (Cambridge Technologies, Cambridge, MA), which transferred labeled lysates to paper spots. These were subsequently washed, and the amount of radioactive DNA was quantitated by scintillation counting using Biosafe II scintillation liquid.

Preparation of cell lysates

T98G human glioblastoma cells were washed once with PBS and suspended in lysis buffer (40 mM HEPES, pH 7.4, with 10% glycerol, 1% Triton X-100, 0.5% Nonidet P-40 (NP-40), 150 mM NaCl, 50 mM NaF, 20 mM β-glycerophosphate, 1 mM EDTA, 1 mM EGTA, 1 mM phenylmethylsulfonyl fluoride and 0.1 mM vanadate) containing a protease inhibitor mixture (1 mg/ml aprotinin, leupeptin and pepstatin). Cells lysates were cleared by centrifugation at 15,000 rpm for 30 min, collected and stored -80°C.

Luciferase assay

The NF-kB reporter plasmid driven by the rat prolactin minimal promoter (-36 to +37) under the control of the two copies of the NF-kB binding site of the human Ig κ light chain enhancer 5′-GGGACCTTCC-3′ was kindly provided by M. Rincón and R.A. Flavell (Section of Immunobiology, Howard Hughes Medical Institute, Yale University School of Medicine, New Haven, CT). To assay for luciferase activity, transfected cells in duplicate wells were cultured for 24 h before being stimulated with or without wortmannin (100 nM) or PDGF (100 ng/ml) for a defined length of time. Cells were harvested, washed twice in PBS and treated with lysis buffer (Luciferase Assay, Promega) for 5-10 min on ice. Lysates were spun down for 1 min, and the total supernatants were analyzed using Luciferase Reagent (Promega) and measured in a luminometer (MicroLumat LB-96P, Berthold) for 5 sec. Background measurement was subtracted from each duplicate, and experimental values are expressed either as recorded light units, luciferase activity or as relative activity compared to extracts from unstimulated cells [26,27].

Results

Cells with added wortmannin (10 nM) divided slower than cells without any dosage (Figures 1)

As showed in figure 1 and as we anticipated, cells given only wortmannin, an inhibitor of PI3 kinase, proliferated at less expected rates with respect to cells deprived of any additions (control). The treated cells divided over twenty eight percent slower than cells without any dosage. The cells were incubated by 24, 48, 72, 96 and 120 hours. After treatment with wortmannin and/or PDGF (3H)-Thymidine (0.5 μCi/well) was added for 3 hours.

Cells with added platelet-derived-growth-factor (PDGF) multiplied substantially faster than naturally growing cells (Figures 1).

In contrast to the effect of wortmannin, T98G glioblastoma cells treated only with PDGF, proliferate at more expected rates with respect to cells deprived of any additions (Control) Using the averages on the last counting day, we see that the addition of wortmannin to cells treated with PDGF accounts for a proliferation decrease of about 15888 (82876-66987), a number which also equals the difference in averages between control cells (58654) and cells dosed with wortmannin only (41678). From this data it can be inferred that wortmannin inhibits cells boosted with PDGF by the same amount as it inhibits control cells.

Wortmannin treatment causes inhibition of the Luc-reporter-NF-kB (Luc-2x GGGACCTTCC) in TPA activated T98G cells (Figure 2)

In order to test whether wortmannin is an effective inhibitor of NF-kB transcriptional activity in T98G cells, transient transfection assay was performed. One Luciferase- base reporter constructs regulated by binding of NF-kB transcription complex to multiple copies of classical NF-kB site (Luc-2x GGGACCTTCC) was examined: pGL-2-Luc-kB). Thus, specific inhibition of the activated PI3K pathway is predicted to inhibit the NF-kB dependent reporter. The reporter showed enhanced expression up to 10 fold and 7 respectively when activated with TPA an activator of NF-kB or with PDGF. As control, parallel transfection was carried out with a control reporter vector, pGL-2 Luc, without kB. Cells treated with wortmannin and transfected with de reporter vector Luc-NF-kB, but not the untreated cells, led to a substantial inhibition (approx. 75%) of the reporter activated by TPA in the transiently transfected NF-kB in T98G glioblastoma cells (Figure 2). The observation that expression of wortmannin inhibit TPA activated NF-kB-Luc reporter construct is consistent with the role of NF-kB-dependent system in T98G cells. However, cells treated with PDGF led to a substantial increase in luciferase reporter gene expression (Figure 2).
In the present study we investigated the effect of PDGF and wortmannin in T98G glioblastoma human cells. PDGF is a multifunctional peptide, with a biologically active form that is a dimer composed of A, B, C, and D chains. The activity of PDGF depends on the dimer formed: AA, AB, BB, CC, or DD (19, 20, 21). On the other hand, Wortmannin, a steroid metabolite of the fungi Penicillium funiculosum, Talaromyces wortmanni, is a non-specific, covalent inhibitor of phosphoinositide 3-kinases (PI3Ks) (1, 2). In summary, in this study we showed that cells treated with platelet-derived-growth-factor (PDGF) and/or wortmannin, an inhibitor of PI3 kinase proliferated at expected rates with respect to cells control (deprived of any additions). Cells with added platelet-derived-growth-factor (PDGF) or wortmannin proliferated at expected rates with respect to cells control (deprived in this study we showed that cells treated with platelet-derived-growth-factor (PDGF) and wortmannin inhibited cells boosted with PDGF by the same amount as it inhibits control cells. PDGF, then increases cell growth by means of the same pathway which wortmannin inhibits the basal pathway, likely through down-regulation of PI3K signaling and NF-kappa B protein expression.

The observation that expression of wortmannin inhibit TPA activated NF-kB-Luc reporter construct is consistent with the role of NF-kB-dependent system in T98G cells. The assay showed that cells treated with PDGF led to a substantial increase in luciferase reporter gene expression while wortmannin decrease de expression of the Luciferase reporter gene. From this data it can be inferred that wortmannin inhibits cells boosted with PDGF by the same amount as it inhibits control cells.

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