Phosphatidylinositol 3-kinase inhibition potentiates glucocorticoid response in B-cell acute lymphoblastic leukemia

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Despite remarkable progress in polychemotherapy protocols, pediatric B-cell acute lymphoblastic leukemia (B-ALL) remains fatal in around 20% of cases. Hence, novel targeted therapies are needed for patients with poor prognosis. Glucocorticoids (GCs) are drugs commonly administrated for B-ALL treatment. Activation of the phosphatidylinositol 3-kinase (PI3K)/Akt/mammalian target of rapamycin signaling pathway is frequently observed in B-ALL and contributes to GC-resistance. Here, we analyzed for the first time to our knowledge, the therapeutic potential of pan and isoform-selective PI3K p110 inhibitors, alone or combined with dexamethasone (DEX), in B-ALL leukemia cell lines and patient samples. We found that a pan PI3K p110 inhibitor displayed the most powerful cytotoxic effects in B-ALL cells, by inducing cell cycle arrest and apoptosis. Both a pan PI3K p110 inhibitor and a dual γ/δ PI3K p110 inhibitor sensitized B-ALL cells to DEX by restoring nuclear translocation of the GC receptor and counteracted stroma-induced DEX-resistance. Finally, gene expression analysis documented that, on one hand the combination consisting of a pan PI3K p110 inhibitor and DEX strengthened the DEX-induced up- or down-regulation of several genes involved in apoptosis, while on the other, it rescued the effects of genes that might be involved in GC-resistance. Overall, our findings strongly suggest that PI3K p110 inhibition could be a promising strategy for treating B-ALL patients by improving GC therapeutic effects and/or overcoming GC-resistance.

KEYWORDS
ALL, dexamethasone, PI3K inhibitors, targeted therapy
1 | INTRODUCTION

B-cell acute lymphoblastic leukemia (B-ALL) is the most common malignancy in children, accounting for 80% of pediatric ALL (Graux, 2011; Siegel, Naishadham, & Jemal, 2012). Thank to progress in polychemotherapy protocols, the majority of B-ALL pediatric patients achieve complete remission. However, the prognosis of relapsed and chemoresistant patients remains poor (Harned & Gaynon, 2008). Also, infants and young adults with B-ALL display a much worse prognosis (Guest & Stam, 2017; Harned & Gaynon, 2008). Therefore, the identification of novel targeted therapies to support conventional chemotherapy is urgently required to further improve the outcome of this disorder.

Phosphatidylinositol 3-kinase (PI3K)/Akt/mammalian target of rapamycin (mTOR) (PI3K/Akt/mTOR) signaling pathway is a highly conserved signal transduction axis involved in many cellular processes (Laplante & Sabatini, 2012). It is well established that aberrant activation of this cascade is associated with the pathogenesis of several types of hematologic malignancies (Polak & Buitenhuis, 2012), including B-ALL (Fuka et al., 2012; Tasián, Teachev, & Rheingold, 2014). Therefore, this pathway is an attractive target to efficiently treat B-ALL patients. In particular, PI3K isoforms are now regarded as key targets for the development of innovative therapeutic strategies (Miller, Goulet, & Johnson, 2017). PI3K is a family of enzymes grouped into three classes (I–III) with distinct structures and functions. Class I PI3Ks comprise four heterodimers that phosphorylate phosphatidylinositol-4,5-bisphosphate (PtdIns 4,5P2) to yield PtdIns 3,4,5-trisphosphate (PtdIns 3,4,5P3). PtdIns 3,4,5P3 in turn recruits a number of downstream proteins containing a pleckstrin homology domain, including Akt. Class I PI3Ks, consisting of a p110 catalytic (-α, -β, -γ, -δ) and a regulatory subunit, display different patterns of expression in mammalian tissues. p110α and p110β catalytic subunits are ubiquitously expressed, whereas p110δ and p110ε catalytic subunits are mainly expressed in leukocytes (Chantry et al., 1997). Class I PI3Ks are the most studied class of isozymes and sustain not only carcinogenesis, but also several tumor-promoting aspects of the neoplastic microenvironment (Engelman, Luo, & Cantley, 2006; Hanahan & Weinberg, 2011).

Over the last few years, several pan and isoform-selective class I PI3K p110 inhibitors have been disclosed by pharmaceutical companies. They display favorable drug properties and suppress tumor growth in different preclinical models of cancer. Therefore, some of them have recently entered clinical trials, also for hematological malignancies (Fruman & Rommel, 2014). Glucocorticoids (GCs) represent one of the most effective agents used in chemotherapeutic protocols for B-ALL, due to their strong pro-apoptotic activity to leukemic cells. Early response to GCs is one of the major prognostic factors of B-ALL, while GC-resistance has been associated with a poor outcome (Inaba & Pui, 2010; Kaspers et al., 1998). Despite their clinical significance, the molecular mechanisms underlying GC activity and resistance in lymphoid malignancies remain unclear (Bhadri, Trahair, & Lock, 2012; Norman & Hearing, 2002). In this connection, it is known that interactions between B-ALL cells and bone marrow stromal cells (BMSCs) in the marrow microenvironment can support leukemic cell survival and resistance to several drugs, including GCs (Randhawa et al., 2016).

Therefore, strategies aimed to reverse GC-resistance may considerably improve the outcome of B-ALL patients. It was recently shown that inhibition of the PI3K/Akt/mTOR pathway could reverse GC-resistance in ALL cells, as either MK2206 (an Akt inhibitor) or rapamycin (an mTOR inhibitor) were identified as GC sensitzers (Piovano et al., 2013; Wei et al., 2006).

Moreover, also migration of leukemic cells toward BMSCs is a poor prognostic factor in B-ALL (Aries et al., 2014; Konoplev et al., 2011). Therefore, molecules targeting motility of leukemic cells are interesting candidates for drug development to optimize treatment of B-ALL patients.

With the above in mind, we used preclinical models of B-ALL to test the cytotoxic effects of both pan and isoform-selective PI3K p110 inhibitors. Moreover, using dexamethasone (DEX), a GC frequently used in B-ALL therapy, we investigated the ability of PI3K p110 inhibitors to improve the therapeutic efficacy of GCs and/or their capability to decrease GC-resistance.

2 | MATERIALS AND METHODS

2.1 | Cell lines, primary samples, human bone marrow stromal cells, and reagents

B-ALL cell lines (KOPN8, REH, NALM6) were obtained from Deutsche Sammlung von Mikroorganismen und Zellkulturen GmbH (DSMZ, Braunschweig, Germany). It is worth remembering that KOPN8 and NALM6 cells are representative of infant and pediatric B-ALL, respectively, while NALM6 cells are representative of young adult B-ALL. All cell lines were cultured in RPMI-1640 medium (Life Technologies Italia, Monza, Italy) supplemented with 10% fetal bovine serum (FBS, Life Technologies). Samples from B-ALL pediatric patients were obtained after written and informed consent from the parents. Experimental protocols were in accordance with the guidelines and the regulations of the 1975 Helsinki declaration. The samples analyzed have been obtained from patients enrolled in the Associazione Italiana Ematologia Oncologia Pediatrica 2002/01 clinical trial, which was approved by the institutional review board of the Sant’Orsola Hospital, Bologna, Italy. Samples were isolated using Ficoll-Paque (Amersham Biosciences, Little Chalfont, UK) and grown in RPMI 1640 supplemented with 20% FBS and insulin-transferrin-sodium selenite. Human BMSCs (hBMSCs) were purchased from Creative Bioarray (Shirley, NY) and cultured in MSCGM™ Mesenchymal Stem Cell Growth Medium (Lonza, Williamsport, PA).

The PI3K p110 inhibitors (ZSTK-474, AS-605240, CAL-101, IPI-145), the Akt inhibitor (MK2206) and DEX were from Selleck Chemicals (Houston, TX). Antibodies to PI3K p110δ and to glucocorticoid receptor (GR) were from Santa Cruz Biotechnology (Heidelberg, Germany), whereas the Ser134 p-GR antibody was from Merck Millipore (Darmstadt, Germany). All other primary and secondary antibodies were from Cell Signaling Technology (Danvers, MA).
To test the effects of PI3K p110 inhibitors, B-ALL cell lines were cultured for 24 hr in the presence of the vehicle (0.1 % dimethylsulfoxide [DMSO]) or increasing drug concentrations, and cell viability was determined using the MTT [3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide] cell proliferation kit (Roche Diagnostic, Basel, Switzerland) (Huang et al., 2017; Poli et al., 2017). The combination effects and potential synergism were evaluated by quantitative analysis of dose–effect relationship, based on the Chou and Talalay method (Chou & Talalay, 1984). For each combination experiment, a combination index (CI) was calculated using the CalcuSyn software (Biosoft, Cambridge, UK). This method of analysis generally defines CI values of <0.3 as strongly synergistic, 0.3–0.9 as synergistic, 0.9–1.1 as additive, whereas values >1.1 are considered antagonistic.

2.2 Annexin V-FITC/propidium iodide (PI) staining and cell cycle analysis

Apoptosis and cell cycle analysis were performed as previously described (He, Zhang, & Zhao, 2016; Shrestha et al., 2016). B-ALL cell lines were treated with the different compounds or the vehicle alone. Analyses were performed on a FC500 flow cytometer (Beckman Coulter, Miami, FL) with the appropriate software (CXP, Beckman Coulter).

2.3 Western blotting analysis

Western blotting analysis was performed by standard methods, as previously described (Ramazzotti et al., 2016; Tang et al., 2016). Cells were lysed using M-PER Mammalian Protein Extraction Reagent supplemented with the Protease and Phosphatase Inhibitor Cocktail (Thermo Fisher Scientific, Inc., Rockford, IL). Fifty micrograms of protein was loaded for each lane, unless otherwise specified. An antibody to β-actin was used as a loading control.

2.4 PtdIns 3,4,5P3 quantification by flow cytometry

B-ALL cell lines were treated with the PI3K p110 inhibitors for 2 hr, fixed in 4% paraformaldehyde for 15 min, permeabilized in 0.4% Triton X-100 for 10 min, washed in phosphate-buffered saline (PBS) with 1% bovine serum albumin (BSA) and incubated over night at 4°C with a FITC-conjugated anti-PtdIns 3,4,5P3 antibody (Echelon, Biosciences, Inc., Salt Lake City, UT). Analyses were performed on a FC500 flow cytometer with the appropriate software (CXP).

2.5 Immunofluorescence microscopy

Cells were seeded on electrostatically charged glass slides using a Shandon Cytospin (Thermo Electron Corporation, Pittsburgh, PA) at low acceleration and 200 rpm for 5 min. Slides were fixed in 4% paraformaldehyde at 37°C for 10 min and permeabilized with 0.15% Triton X-100 in PBS for 8 min. Slides were then blocked with PBS containing 5% BSA for 1 hr. Incubation with anti-GR antibody (1:100) was performed overnight at 4°C in blocking medium. Then, a FITC-conjugated anti-mouse IgG antibody (1:200) was used for 1 hr at room temperature. Slides were washed three times for 10 min at room temperature with PBS/Tween 20 and mounted with a 4′,6-diamidino-2-phenylindole (DAPI) anti-fade reagent in glycerol (Molecular Probes, Eugene, OR). Images were taken on a Zeiss Axio ImagerZ1 microscope, equipped with 60X/NA 1.4 optics and Apotome apparatus, coupled to a computer driven Zeiss AxioCam digital camera (MRm), using Zeiss Axio Vision 4.4 software (Carl Zeiss, Oberkochen, Germany). All images were taken at similar exposures within an experiment for each antibody and processed using Adobe Photoshop 7 (Adobe Systems, San Jose, CA).

2.6 Subcellular fractionation

Cells were collected, centrifuged, and washed in PBS. Pellets were resuspended in 1 ml of 10 mM Tris–HCl pH 7.4, 2 mM MgCl2 for 2 min. Then, 0.6% Triton X-100 was added and the cells were passed twice through a syringe fitted with a 22 ½ gauge needle. Next, MgCl2 was added to a final concentration of 5 mM and the samples were centrifuged for 10 min at 800 rpm. Nuclear pellets were washed twice in 10 mM Tris–HCl pH 7.4, 5 mM MgCl2, while the supernatant was transferred to a new vial and used as a cytoplasmic fraction.

2.7 Migration assay and co-culture of B-ALL cells

Migration assay of B-ALL cells was performed using 24-well plates with Transwell® culture inserts (diameter 6.5 mm, pore size 8 μm; Costar Corning, New York, NY). 1 × 10⁶ cells were suspended in 100 μl of medium and added to the upper chamber. Inserts were transferred to bottom wells containing 600 μl of medium with or without 150 ng/ml of CXCL-12 (Biodesign International, Saco, ME). Drugs or solvent were added and incubated for 4 hr at 37°C in 5% humidified CO2 atmosphere. Inserts were then removed and cells transmigrated into the lower chamber were recovered and counted using an inverted microscope. Results are shown as the fold change of the percentage of migration compared with the untreated condition.

For co-culture, hBMSC were grown in the lower chamber of Transwell® 6-well plates containing a 0.4 μm polyester membrane (Corning) at a density of 5,000 to 6,000 cells/cm² (Frolova et al., 2012). Then, NALM6 and REH cells (375,000) were added to the upper chamber and treated with the drugs or solvent at the indicated concentrations. After 48 hr, the viability of drug-treated hBMSCs, NALM6, and REH cells, grown either alone or co-cultured, was evaluated by MTT assays.

2.8 Gene expression analysis

Total RNA was isolated from untreated cells and cells treated for 8 hr using a RNeasy Mini Kit (QIAGEN, Valencia, CA) according to the manufacturer’s instructions. Total RNA was reverse-transcribed into cDNA using the iScriptTM Advanced cDNA Synthesis Kit (Bio-rad, Hercules, CA). Gene expression of GC signaling markers was measured using the PrimePCRTM Assay for real-time (Bio-rad). For each sample,
cDNA was mixed with 2× SsoAdvancedTM universal supermix (25 ng cDNA/reaction) containing SYBR Green (Bio-rad) and aliquoted in equal volumes to each well of the real-time PCR arrays. The quantitative PCR reaction was performed using a 7300 Real-Time PCR system (Applied Biosystems, Foster City, CA).

### 2.9 Statistical analysis

Statistical analyses were performed using Student's t-test or one-way ANOVA (Dunnett's test) at a significance level of $p < 0.05$ (GraphPad Prism Software, La Jolla, CA).

# 3 RESULTS

### 3.1 PI3K p110 inhibitors display cytotoxic effects in B-ALL cell lines

We investigated the efficacy of PI3K p110 inhibitors in suppressing leukemic cell proliferation and survival in in vitro models of B-ALL. We first evaluated the expression of PI3K p110 isoforms in a panel of B-ALL cell lines. All PI3K p110 isoforms were expressed in B-ALL cell lines, although protein levels varied (Figure 1a). Then, the phosphorylation status of Akt (Ser473 and Thr308) and PTEN (Ser380) was studied. It is well established that, when phosphorylated at Ser380 by CK2, PTEN is post-translationally inactivated with a consequent PI3K/Akt/mTOR pathway activation in B-ALL cells (Gomes et al., 2014). Accordingly, all cell lines displayed activated (phosphorylated) Akt with a concomitant PTEN phosphorylation at Ser380 (Figure 1a).

Next, we evaluated the viability of B-cell lines upon treatment with the pan PI3K p110 inhibitor ZSTK-474. As both p110γ and p110δ are highly expressed in B-lymphocytes, we also evaluated the isoform-selective inhibitors AS-605240 (selective to PI3K p110γ; Camps et al., 2005) and CAL-101 (selective to PI3K p110δ; Herman et al., 2010), as well as the γ/δ dual inhibitor IPI-145 (Duvelisib®) and the combination AS-605240 + CAL-101. IPI-145 antagonizes intrinsic and extrinsic survival signals in chronic lymphocytic leukemia (CLL) cells with an overactive PI3K/Akt/mTOR pathway (Dong et al., 2014).

Cells were treated for 24 hr with increasing concentrations of the drugs and the effects of PI3K p110 inhibitors on cell viability were analyzed by MTT assay (Figure 1b). The pan PI3K p110 inhibitor ZSTK-474 displayed marked cytotoxic effects on all B-ALL cell lines (IC50 between 0.1 and 4.9 μM), whereas isoform-selective inhibitors were less cytotoxic and showed differences among different cell lines (Table 1). Thereafter, the combination of p110δ (CAL-101) and p110γ (AS-605240) inhibitors at a fixed ratio (1:1) was evaluated. Interestingly, dual p110γ/δ inhibition (using either IPI-145 or CAL-101 along with AS-605240) displayed stronger cytotoxic effects, especially in REH cells, compared to isoform-selective inhibitors (Figure 1b and Table 1). In general, REH cells were the most sensitive to all PI3K inhibitors, whereas KOPN8 and NALM6 were more resistant, except to ZSTK-474.

Based on these results, in most of the following experiments we mainly focused on the two PI3K p110 inhibitors that showed the most potent cytotoxic effects on B-ALL cells, that is, ZSTK-474 and IPI-145.

### 3.2 ZSTK-474 affects cell cycle progression, induces apoptosis, and decreases migration of B-ALL cell lines

To determine whether treatment of B-ALL cell lines with PI3K p110 inhibitors could affect cell cycle progression, cells were incubated for 24 hr with the drugs and the cell cycle was studied by means of flow cytometry.

#### TABLE 1

| Drug       | IC50 (μM) |
|------------|-----------|
| KOPN8      | 4.9       |
| NALM6      | 0.1       |
| REH        | 0.1       |
| AS-605240  | N.D.      |
| CAL-101    | >10       |
| AS + CAL   | >10       |
| IPI-145    | N.D.      |

**N.D., not determined. AS-605240 + CAL-101: AS + CAL.**
FIGURE 2 PI3K p110 inhibitors affect cycle progression, induce apoptosis, and down-regulate migration of B-ALL cell lines. (a) B-ALL cell lines were treated for 24 hr with 5 \( \mu \)M of the following drugs: ZSTK-474 (ZSTK), AS-605240 (AS), CAL-101 (CAL), AS-605240 + CAL-101 (AS + CAL), IPI-145 (IPI). Cell cycle analysis was then performed by flow cytometry. CTR: untreated cells. *\( p < 0.05 \) relative to CTR. (b) B-ALL cell lines were treated for 24 hr with 5 \( \mu \)M of the following drugs: ZSTK-474 (ZSTK), AS-605240 (AS), CAL-101 (CAL), AS-605240 + CAL-101 (AS + CAL), IPI-145 (IPI). Then, cells were stained with Annexin V-FITC/PI and analyzed by flow cytometry. CTR: untreated cells. *\( p < 0.05 \) relative to CTR. (c) REH cells were assayed for apoptosis by Annexin V-FITC/PI staining upon treatment for 24 hr with 5 \( \mu \)M of the following drugs ZSTK-474 (ZSTK), AS-605240 (AS), CAL-101 (CAL), AS-605240 + CAL-101 (AS + CAL), IPI-145 (IPI). Results from a representative experiment are shown. CTR: untreated cells. (d) Western blotting analysis documenting caspase 3 and PARP cleavage in response to PI3K inhibitor treatment (5 \( \mu \)M for 24 hr). CTR: untreated cells. (e) B-ALL cell lines were treated with 5 \( \mu \)M of either ZSTK-474 (ZSTK) or IPI-145. Migration toward the chemoattractant agent CXCL12 (200 ng/ml) was tested using a Transwell® system as described in “section 2.” After 4 hr of drug treatment, transmigrated cells were recovered and counted under the microscope. CTR: untreated cells. *\( p < 0.05 \) relative to CTR.
cytometric analysis of PI-stained samples (Figure 2a). Only ZSTK-474 induced a statistically significant G0/G1 block and a concomitant decrease in both S and G2/M phases of the cell cycle in all the cell lines tested.

Then, apoptosis induction was investigated by cytofluorimetric (Figures 2b and 2c) and Western blotting (Figure 2d) analyses. Cells treated with ZSTK-474 underwent apoptosis after 24 hr of treatment, as demonstrated by the increased number of Annexin-V+/PI− and Annexin-V+/PI+ cells which correspond to early and late apoptosis, respectively (Figures 2b and 2c).

Caspase-dependent apoptosis was further examined in B-ALL cells by Western blotting analysis. A marked cleavage of caspase 3 as well as poly ADP-ribose polymerase (PARP) was induced by all the drugs, with the exception of AS-605240, in REH cells after 24 hr of treatment (Figure 2d). On the contrary, NALM6 and KOPN8 cell lines resulted less sensitive to apoptosis, as documented by fainter or absent bands corresponding to cleaved-caspase 3 and PARP.

It is well established that CXCL12/CXCR4 axis is involved in many physiological and pathological processes, including tumor progression, angiogenesis, metastasis, and survival (Teicher & Fricker, 2010). CXCL12 is secreted by BMSCs and plays an important role in homing, migration, and survival of both hematopoietic progenitor cells and leukemic cells (Burger & Peled, 2009), including B-ALL cells (Bradstock et al., 2000). Indeed, the G-protein-coupled transmembrane receptor CXCR4 is highly expressed in B-ALL, where it is involved in survival of leukemic cells (Bradstock et al., 2000). Moreover, high levels of CXCR4 are linked to poor patient outcome in B-ALL patients (Konoplev et al., 2011; van den Berk et al., 2014).

Therefore, we investigated whether PI3K inhibitors could affect migration of B-ALL cells. KOPN8, NALM6, and REH cells were treated with either ZSTK-474 or IPI-145 for 4 hr and migration toward CXCL12 was measured using Transwell assays. Both PI3K inhibitors strongly reduced (approximately 80%) migration of REH cells compared to controls, whereas reduction of transmigration was observed to a lower extent in both NALM6 and KOPN8 cells (Figure 2e).

3.3 | PI3K p110 inhibitors down-modulate PI3K/Akt/mTOR signaling

To confirm that ZSTK-474 and IPI-145 inhibited PI3K activity at the concentrations we used, PtdIns 3,4,5P3 levels were measured in B-ALL cells. Cells were treated with either drug and flow cytometric analysis was performed. ZSTK-474 and, to a lower extent, IPI-145 were both able to significantly reduce the percentage of PtdIns3,4,5P3-positive in all the three cell lines (Figure 3a). Therefore, we examined the effects of PI3K p110 inhibitors on PI3K/Akt/mTOR signaling pathway effectors through Western blot analysis (Figure 3b). After 4 hr of drug treatment, we observed that ZSTK-474 negatively affected PI3K

![Figure 3](image-url)
downstream targets in all the cell lines analyzed, as indicated by a marked dephosphorylation of Akt (both at Ser473 and at Thr308) and S6 ribosomal protein (S6RP). Similar effects were obtained in NALM6 and REH cells treated with IPI-145 (Figure 3b).

3.4 | PI3K p110 inhibitors synergize with GCs

To further evaluate the clinical potential of PI3K inhibitors in B-ALL treatment, we investigated whether these drugs could synergize with DEX, a GC commonly used for the treatment of pediatric B-ALL patients. Firstly, cells were incubated for 24 hr with increasing concentrations of DEX and cell viability was measured via MTT assays. As shown in Figure 4a, DEX reduced viability of REH, NALM6 and, to a lower extent, KOPN8 cell lines.

Then, the existence of possible synergisms of PI3K inhibitors (ZSTK-474 and IPI-145) with DEX was analyzed (Figure 4b). Cells were treated for 24 hr with one of the above-mentioned drugs, alone or combined to DEX at a fixed ratio (DEX:PI3K inhibitors, 1:50) and CsIs were calculated. A strong synergism of DEX with IPI-145 and, less frequently, with ZSTK-474, was observed in the analyzed cell lines, including KOPN8 cells, the most resistant cells both to PI3K p110 inhibitors and to DEX (Table 2). Synergisms induced by the combination of DEX with AS-605240, CAL-101 and AS-605240 + CAL-101 were in general less strong or undetectable in all the three cell lines analyzed (Table 2).

We further tested the pro-apoptotic effects of combination treatments. Cells were stained with Annexin V-FITC/PI and flow cytometric analysis was performed. The combination of DEX with either ZSTK-474 or IPI-145 was more effective in inducing apoptosis in all B-ALL cell lines analyzed when compared to single drug treatment, thus confirming that PI3K p110 inhibitors sensitized B-ALL cells to GCs (Figure 5a). Moreover, apoptosis induction was confirmed by caspase 3 and PARP cleavage in KOPN8, NALM6 and REH cells, as documented by Western blotting analysis in samples treated with the drug combinations (Figure 5b).

The Bcl-2 family proteins are critical in mediating GC-induced apoptosis and could be involved in GC resistance (Bachmann, Gorman, Mackenzie, Lutze-Mann, & Lock, 2005). In particular, it has been observed that reduction of pro-apoptotic Bim expression (Jiang et al., 2011) as well as increase of anti-apoptotic Mcl-1 levels (Holleman et al., 2004) are related to GC resistance in pediatric ALL. We, therefore, analyzed the effects of treatment with DEX, ZSTK-474, or IPI-145, alone or in combination, on Bim and Mcl-1 expression in KOPN8, NALM6, and REH cells. Western blotting analysis documented a decrease of Mcl-1 expression in response to DEX plus PI3K inhibitor treatment and a concomitant increase of Bim in all cell lines studied (Figure 5c).

3.5 | PI3K p110 inhibition counteracts stroma-induced DEX resistance

As human hBMSCs are known to be responsible for DEX-resistance in vitro (Randhawa et al., 2016), we investigated whether PI3K p110 inhibitors could reverse DEX-resistance in NALM6 and REH cells co-cultured with hBMSCs in a Transwell® system.

At first, we demonstrated that DEX and PI3K inhibitors did not affect hBMSC viability, as documented by MTT assays (Figure 6a). Secondly, we observed that either ZSTK-474 or IPI-145 were able to partially counteract the increase in both NALM6 and REH leukemic cell viability in response to DEX treatment caused by co-culturing with hBMSCs (Figure 6b).

3.6 | PI3K p110 inhibitors induce GR nuclear translocation

GC cytotoxicity is mediated by binding of GCs to their intracellular receptor, GR, which is a ligand-inducible transcription factor
Indeed, when DEX binds to GR, the receptor is activated and translocates to the nucleus, thus activating the transcription of target genes that induce cell cycle arrest and apoptosis (Inaba & Pui, 2010).

In order to determine whether resistance to DEX correlated with GR down-regulation, basal levels of GR protein were compared among B-ALL cell lines by Western blotting analysis. However, GR was expressed at approximately equivalent levels in all cell lines (Figure 7a). The phosphorylation status of GR was then investigated. It has been documented that GR phosphorylation at Ser134 plays an important role in the modulation of GC-driven apoptosis (Galliher-Beckley, Williams, & Cidlowski, 2011; Piovan et al., 2013). Indeed, when GR is phosphorylated at Ser134, a nuclear translocation impairment occurs and this interferes with the patterns of GC-induced genes. Of note, this mechanism is regulated by Akt in T-ALL cells (Piovan et al., 2013). In order to investigate whether PI3K/Akt signaling was involved in Ser134 phosphorylation and in nuclear translocation also in B-ALL cells, KOPN8 and NALM6 cells were treated with DEX, IPI-145, or ZSTK-474, and Western blotting analysis for Ser134 p-GR was carried out. PI3K inhibitor treatment caused a marked dephosphorylation of GR at Ser134 (Figure 7b), whereas DEX alone did not. A similar inhibitory effect was caused by MK2206, an allosteric Akt inhibitor (Hirai et al., 2010), thus confirming the crucial role played by Akt in this phenomenon.

As the GC-induced apoptotic response depends on translocation of the receptor-ligand complex to the nucleus, GR nuclear translocation was investigated by immunofluorescence and Western blotting analyses following a 4 hr treatment with DEX in less sensitive cell lines, namely NALM6 and KOPN8 cells. As shown in Figure 7c, GR was predominantly cytoplasmic in untreated and drug-treated (ZSTK-474 or IPI-145) samples, while DEX caused a partial re-localization of GR to the nucleus. On the contrary, the combination of DEX with either ZSTK-474 or IPI-145 caused a striking nuclear translocation of GR. Similar results regarding GR localization were obtained by Western blotting analysis in NALM6 cells (Figure 7d).

### 3.7 Gene expression analysis

To study in more depth, the synergisms between PI3K inhibitors and GCs in B-ALL cells, the GC signaling pathway was analyzed in NALM6 cells using a quantitative real-time PCR array with 82 genes (Supplementary Table S1). A 8 hr treatment with the drugs (DEX or ZSTK-474), either used alone or in combination, was chosen in order to identify genes that were modulated in early phases of GC activation (Jing et al., 2015). Only differentially expressed genes (fold-change ≥ 2) were considered (Figures 8a and 8b).

Thirty-one genes, of which four were induced and 27 repressed, were modulated in ZSTK-474 + DEX versus CTR samples (Figure 8a and Supplementary Table S2). De-regulation of transcripts involved in signal transduction (e.g., PI3KR1, RGS2, SESN1, SPHK1, and USP2), nucleotide metabolism (e.g., AMPD3), glucose and fatty acid metabolism (e.g., ASPH, GOT1, and H6PD), cytokines and chemokines (e.g., EDN1 and TNF), cell surface receptors (PDGFRB and VLDLR) as well as transcription factors (e.g., ARID5B, FOSL2, TSC22D3, and BCL6) was observed. In particular, ZSTK-474 combined with DEX induced a remarkable up-regulation of TSC22D3, a gene that plays a pivotal role in the anti-inflammatory and immunosuppressive effects of glucocorticoids (Bruscoli et al., 2015) as well as a down-regulation of several genes associated with cellular immunity.

| Table 2: Analysis of the effects of DEX treatment combined with PI3K p110 inhibitors in B-ALL cell lines |
| --- |
| **CI** |
| **[μM]** | ZSTK + DEX | AS + DEX | CAL + DEX | AS + CAL + DEX | IPI + DEX |
| KOPN8 | 0.0065/0.3125 | 0.28 | 0.68 | 0.69 | 0.15 | 0.05 |
| | 0.025/1.25 | 0.26 | 0.79 | 0.77 | 0.20 | 0.01 |
| | 0.1/5 | 0.39 | 1.77 | 0.82 | 0.47 | 0.02 |
| | 0.2/10 | 0.44 | 1.16 | 0.71 | 0.58 | 0.04 |
| NALM6 | 0.0065/0.3125 | 0.43 | 1.33 | 0.35 | 0.88 | 0.41 |
| | 0.025/1.25 | 0.39 | 0.76 | 0.33 | 0.27 | 0.19 |
| | 0.1/5 | 0.55 | 0.48 | 0.39 | 0.26 | 0.22 |
| | 0.2/10 | 0.43 | 0.71 | 0.49 | 0.33 | 0.24 |
| REH | 0.0065/0.3125 | 0.43 | 0.93 | 0.52 | 1.01 | 0.57 |
| | 0.025/1.25 | 0.16 | 1.24 | 0.70 | 0.70 | 0.25 |
| | 0.1/5 | 0.17 | 0.82 | 0.65 | 0.87 | 0.21 |
| | 0.2/10 | 0.19 | 1.10 | 0.93 | 0.85 | 0.22 |

PI3K inhibitor:DEX ratio was 50:1. Combination index (CI) values, calculated with the CalcuSyn software, are presented. ZSTK-474: ZSTK; AS-605240: AS; CAL-101: CAL; AS-605240 + CAL-101: AS + CAL; IPI-145: IPI.

(Whitfield, Jurutka, Haussler, & Haussler, 1999). Indeed, when DEX binds to GR, the receptor is activated and translocates to the nucleus, thus activating the transcription of target genes that induce cell cycle arrest and apoptosis (Inaba & Pui, 2010).
proliferation and survival such as BLC6, CREB3, EDN1, PIK3R1, SPHK1, STAT5B, and USP2. In order to gain further insight into the mechanisms that might influence the synergism between GCs and PI3K inhibitors, we also analyzed genes differentially expressed in ZSTK-474 + DEX versus DEX alone samples (Figure 8b and Supplementary Table S3). ZSTK-474 rescued the effects of genes that might be involved in GC resistance given that their expression goes in an opposite direction to GC-cytotoxic effects. For instance, DEX alone increased the expression of genes involved in cell survival such as AQP1, CTGF, DDIT4, FKBP5, and IL10, while the addition of ZSTK-474 counteracted this effect, restoring the balance toward a cell death fate. In particular, DDIT4 blocks cell growth, proliferation and survival through negative regulation of mTOR (Sofer, Lei, Johannessen, & Ellisen, 2005) and FKBP5 functions as a tumor suppressor of PI3K/Akt/mTOR signaling when it is hyper-activated (Li, Lou, & Wang, 2011). Finally, ZSTK-474 was able to counteract the increase in AQP1 and CTGF mRNA, a water channel protein and a growth factor, respectively, both involved in

**FIGURE 5** DEX enhances apoptosis induced by PI3K p110 inhibitors in B-ALL cells. (a) B-ALL cell lines were treated for 24 hr with DEX (0.1 μM), ZSTK-474 (ZSTK, 5 μM), AS-605240 (AS, 5 μM), CAL-101 (CAL, 5 μM), AS-605240 + CAL-101 (AS + CAL, both at 5 μM), IPI-145 (IPI, 5 μM), alone, or with a combination of the PI3K p110 inhibitors with DEX at the same concentrations as above. Then, Annexin V-FITC/PI staining analysis was performed by flow cytometry. CTR: untreated cells. *p < 0.05 relative to CTR. (b) B-ALL cell lines were treated for 24 hr with DEX (0.1 μM), IPI-145 (IPI, 5 μM), ZSTK-474 (ZSTK, 5 μM) alone, with a combination of the PI3K p110 inhibitors with DEX at the same concentrations as above. Then, Western blot analysis for caspase 3 and PARP cleavage was performed. CTR: untreated cells. (c) Western blotting analysis documenting Mcl-1 and Bim expression after treatment for 24 hr with DEX (0.1 μM), IPI-145 (IPI, 5 μM) or ZSTK-145 (ZSTK, 5 μM) or their combinations at the same concentrations. CTR: untreated cells. Band intensities were quantified by NIH Image J densitometry analysis. The values represent the amount of protein present in treated samples relative to untreated cells after normalizing to β-actin density. CTR: untreated cells.
lymphoblasts synergistic cytotoxic effects in B-ALL patient analyzed by MTT assays, 48 hr after treatment with DEX (0.1 μM), IPI-145 (IPI, 5 μM) or ZSTK-474 (ZSTK, 5 μM). CTR: untreated cells. Cell viability was then measured by MTT assays. The results are the mean of three different experiments ± sd. (b) PI3K p110 inhibitors sensitize B-ALL cell lines to DEX. Cell viability was analyzed by MTT assays, 48 hr after treatment with DEX (0.1 μM) in either the absence or the presence of hBMSCs, using a Transwell® system. IPI-145 (IPI) and ZSTK-474 (ZSTK) were used at 5 μM. CTR: untreated cells. The results are the mean of three different experiments ± sd. Asterisks indicate statistically significant differences with respect to samples cultured without hBMSC (p < 0.05).

FIGURE 6 PI3K p110 inhibitors lower hBMSC-induced DEX-resistance in B-ALL cell lines. (a) hBMSC were treated for 48 hr with DEX (0.1 μM), IPI-145 (IPI, 5 μM) or ZSTK-474 (ZSTK, 5 μM). CTR: untreated cells. Cell viability was then measured by MTT assays. The results are the mean of three different experiments ± sd. Asterisks indicate statistically significant differences with respect to samples cultured without hBMSC (p < 0.05).

3.8 | PI3K p110 inhibitors and DEX display synergistic cytotoxic effects in B-ALL patient lymphoblasts

To better evaluate the efficacy of PI3K p110 inhibitors combined with DEX as potential therapeutic agents, we analyzed four pediatric B-ALL patient samples, isolated from peripheral blood. The effects of PI3K p110 inhibitors were first evaluated by performing MTT assays (Figure 9a). Cells were incubated with either single drugs (DEX, IPI-145, ZSTK-474) or their combinations at a fixed ratio (DEX:PI3K inhibitor ratio, 1:50). A marked reduction of cell viability was detected, thus documenting that the combined treatments were effective also in primary samples. The strongest effects were observed with the ZSTK + DEX combination.

Then, we analyzed apoptosis induction following combined treatment using Annexin V-FITC/PI staining. Flow cytometric analysis showed that both drug combinations increased apoptosis when compared to single drug treatments (Figure 9b). Finally, the effects on the PI3K/Akt/mTOR pathway were assessed by Western blotting after 24 hr of treatment with ZSTK-474 or IPI-145, used either alone or combined with DEX in a representative patient sample (Figure 9c). High levels of p-Akt, p-S6RP, and p-4EBP1 were observed in untreated lymphoblasts which indicated a constitutive activation of the PI3K/Akt/mTOR signaling axis. Drug treatments caused a marked decrease in the levels the phosphoproteins, documenting an inhibition of this signaling pathway.

4 | DISCUSSION

Over the last 40 years, the survival rate for pediatric B-ALL has increased from 30% to 80% (Pui & Evans, 2006). Protocols used to treat B-ALL include GCs, vincristine and asparaginase, with or without anthracyclines. However, relapsing B-ALL pediatric patients persist and this remains a leading cause of childhood morbidity and mortality. This therapy failure is mostly due to GC-resistance that is widely recognized as a determinant of poor prognosis (Den Boer et al., 2003). GCs are cytotoxic not only to neoplastic lymphoid cells, but also to healthy lymphocytes, thereby increasing the risk of severe infections. Furthermore, other adverse effects of GC therapy include hyperglycemia, osteonecrosis, neuropsychological alterations, and myopathy (Inaba & Pui, 2010). In light of this, it is urgent to develop novel therapeutic strategies against B-ALL cells aimed to further optimize therapeutic regimens and to overcome GC-resistance.

The PI3K/Akt/mTOR pathway is aberrantly activated in hematological malignancies, including T-ALL and B-ALL (Khwaja, 2010) and it has been demonstrated that up-regulation of this signaling network is associated with bad prognosis and GC-resistance, both in T-ALL and B-ALL (Morishita et al., 2012; Silva et al., 2008). For instance, it is known that Akt activation underlies GC-resistance in T-ALL via a direct phosphorylation of GR at the Ser134 residue (Piovan et al., 2013) and PI3K p110 inhibitors sensitized resistant MLL-rearranged ALL cells to prednisolone (Spijkers-Hagelstein, Pinhancos, Schneider, Pieters, & Stam, 2014). Therefore, PI3K p110 inhibitors are regarded as innovative drugs for the treatment of these types of blood cancer.

Although in T-ALL it has been investigated whether either pan or isoform-selective PI3K p110 inhibitors are more cytotoxic to leukemic cells (Lonetti et al., 2015; Stengel, Jenner, Meja, Mayekar, & Khwaja, 2013; Subramaniam et al., 2012), similar studies have not been performed at all in B-ALL.

Starting from these premises, the aim of this study was to investigate the therapeutic potential of PI3K p110 inhibitors, alone and in combination with DEX, using a panel of B-ALL cell lines and primary samples. We tested pan, isoform-selective (γ or δ) and dual γ/δ PI3K p110 inhibitors, and evaluated their cytotoxic effects. To the best of our knowledge, this is the first time that the effects of these drugs were analyzed in B-ALL cells.

We showed that the strongest cytotoxic effects against B-ALL cells were displayed by ZSTK-474, a novel pan PI3K p110 inhibitor that has entered clinical trials in patients with advanced solid
malignancies (NCT01682473 and NCT01280487). ZSTK-474 reduced cell viability in a concentration-dependent manner, as demonstrated by MTT assays. Flow cytometric and Western blotting analyses documented that ZSTK-474 cytotoxic effects were associated with both cell cycle arrest and apoptosis induction. ZSTK-474 induced a dephosphorylation of downstream targets of PI3K, namely Akt and S6RP. A decrease in PtdIns 3,4,5P3 levels following treatment with ZSTK-474 was also detected. White blood cells are enriched in PI3K p110γ and δ catalytic isoforms (Chantry et al., 1997). Therefore, the use of either selective (γ or δ) or dual γ/δ PI3K p110 inhibitors is considered as an innovative, rationale-based therapeutic strategy for hematological malignancies. Indeed, the use of these inhibitors should reduce toxicity and adverse effects in patients when compared to pan PI3K p110 inhibitors. For instance,
safety and efficacy of the dual γ/δ PI3K p110 inhibitor IPI-145 have been already documented, as this drug entered clinical trials for treating non-Hodgkin lymphoma (Curran & Smith, 2014) and CLL (Balakrishnan et al., 2015) patients. However, both selective and dual PI3K p110 inhibitors showed weaker effects on B-ALL cell viability when compared with ZSTK-474. In our study, only IPI-145 displayed anti-leukemic effects when administrated alone, but not in all tested cell lines. In particular, only REH cells were sensitive to IPI-145, displaying an IC50 of 2.6 μM. No significant synergy between AS-605240 and CAL-101 was seen in B-ALL cells as far as Akt dephoshorylation was concerned. This suggests that PI3K p110δ is likely the most important isoform involved in Akt phosphorylation (but not in S6 ribosomal protein phosphorylation) in B-ALL cell lines. Nevertheless, the cytotoxic effects of either CAL-101 or IPI-145 were generally inferior to ZSTK-474, supporting the concept that also in B-ALL inhibition of all PI3K p110 catalytic subunit is required to attain the strongest cytotoxic effects. This is in agreement with the results previously reported by our group for T-ALL (Lonetti et al., 2015; Stengel et al., 2013).

Intriguingly, we observed that the REH cell line is sensitive to both pan and selective or dual PI3K p110 inhibitors and this might be due to the presence of gain-of-function PI3K mutations in this cell line. In particular, REH cells carry mutations in PI3KR1, PIK3CG, and PIK3CD genes (encoding the regulatory p85α, p110γ, and p110δ catalytic subunits, respectively) according to the COSMIC database (http://cancer.sanger.ac.uk/cell_lines/sample/overview?id=909696).

We have also shown that ZSTK-474 and IPI-145 could inhibit the in vitro motility of B-ALL cells. It has been demonstrated that migration mediated by CXCL12/CXCR4 axis is associated with relapses and a bad patient outcome in B-ALL patients (Bruggemann, Raff, & Kneba, 2012). Moreover, chemokines, such as CXCL12, secreted by stromal cells in the bone marrow microenvironment, increase leukemic survival and GC resistance (Randhawa et al., 2016) and our results, obtained by co-culturing NALM6 and REH cells with hBMSCs, are consistent with these recent observations.

As GCs are used for the treatment of B-ALL, we wanted to analyze the effects of PI3K p110 inhibitors on DEX response. In particular, both ZSTK-474 and IPI-145 strongly synergized with DEX in all tested cell lines and in samples treated with PI3K p110 inhibitors + DEX combination an increased GC-induced apoptosis was observed in both GC-sensitive and -resistant cell lines.

GC-induced cell death depends on the interactions between the hormone and the cytosolic GR that mediates apoptosis induction.
Multiple mechanisms are responsible for GC resistance, including decreased expression of GR, altered affinity of GR for its ligand and dysregulation of GR nuclear translocation (Azher et al., 2016; Bhadri et al., 2012). In particular, it seems that this latter process is controlled by PI3K/Akt/mTOR signaling. Indeed, phosphorylation of GR at Ser134 by Akt causes an impairment of GR translocation in T-ALL cells (Piovan et al., 2013) and this phenomenon could be particularly relevant in cells which display a hyper-activated PI3K/Akt/mTOR cascade, such as B-ALL cells.

We demonstrated that single treatment with either PI3K p110 inhibitors (ZSTK-474 and IPI-145) or DEX, did not lead to GR nuclear translocation in KOPN8 and NALM6 cells, however, translocation could be induced by treating cells with drug combinations. Moreover, we demonstrated that both ZSTK-474 and IPI-145 caused a de-phosphorylation of Ser134 p-GR and a rescue of GCs response, in agreement with what was documented by Piovan et al. (2013) in T-ALL cells.

An association between the modulation of specific anti-apoptotic and pro-apoptotic proteins, such as Mcl-1 and Bim, and GCs resistance has been proposed in pediatric B-ALL (Holleman et al., 2006; Jiang et al., 2011). In addition, it has been demonstrated that rapamycin, an mTOR inhibitor, induced GC sensitivity in primary ALL samples via Mcl-1 modulation (Wei et al., 2006). Interestingly, Wei et al. (2006)
demonstrated that even a weak decrease in Mcl-1 expression could have a significant effect on regulation of apoptotic intrinsic pathway. We demonstrated that DEX + IPI-145 and DEX + ZSTK-474 combinations modulated Mcl-1 and Bim expression levels, suggesting that this could be another mechanism through which PI3K p110 inhibitors may restore GC sensitivity.

The synergism between ZSTK-474 and DEX was examined in more depth by means of gene expression analysis in NALM6 cells.

Remarkably, ZSTK-474 potentiated the DEX-induced up- and down-regulation of genes that are involved in the complex process of GC-dependent apoptosis. In agreement with the literature data, DEX induced a rapid and marked increase in TSC22D3 expression, a master regulator of GC anti-proliferative, pro-apoptotic, and anti-inflammatory effects (Bruscoli et al., 2015), and this increase was strengthened by the addition of ZSTK-474 (a further 2.5-fold increase in TSC22D3 mRNA compared to samples treated with DEX alone).

Interestingly, on one hand ZSTK-474 seemed to reinforce the ability of DEX to down-regulate genes which are involved in proliferation, survival, and anti-apoptotic mechanisms such as BCL6, CEBPB, COL4A2, CREB3, PIK3R1, POU2F2, SPHK1, STAT5B, USP2, on the other, ZSTK-474 rescued the effects of some genes involved in tumor progression and most likely also in GC-resistance such as AQP1, CTGF, DDIT4, FKBP5, and IL10.

Indeed, the immunomodulatory cytokine interleukin-10 (IL10) is frequently found up-regulated in various types of cancer and can be produced by tumor cells, thus contributing to tolerance and tumor-escape processes (Mannino et al., 2015). AQP1 and CTGF, a water channel protein and a growth factor, respectively, participate in cell growth and migration processes (Chu et al., 2008; Papadopoulos et al., 2008). Finally, ZSTK-474 caused an increase in RGS2 and SESN1 expression, which are negative regulators of MAP kinase and mTOR signaling, respectively (Chantranupong et al., 2014; Nguyen et al., 2009).

However, it is essential to consider that the combined treatment also up-regulated some anti-apoptotic and/or pro-inflammatory genes, such as IL6, RASA3, and SGK1. Although these findings have been obtained using only a single cell line and need to be validated in primary samples, we think that they are important as they highlighted some genes that could blunt the efficacy of a targeted therapy based on PI3K inhibitors in B-ALL cells.

As to the toxicity of PI3K p110 inhibitors on CD19+ peripheral blood B-lymphocytes from healthy donors, our unpublished data documented that ZSTK-474, even when employed at the maximal concentration of 10 μM, was less cytotoxic to normal than to neoplastic B-cells, as about 70% of B-lymphocytes were still viable. Regarding IPI-145, several lines of evidence indicate that this drug does not affect the viability of normal B-cells (Balakrishnan et al., 2015; Dong et al., 2014; Gockeritz et al., 2015).

Others inhibitors of the PI3K/Akt/mTOR pathway have been tested in combination with GCs in preclinical models of ALL, including rapamycin, an allosteric mTORC1 inhibitor, (Zhang et al., 2012), and MK2206, an allosteric Akt inhibitor, (Piovan et al., 2013). However, clinical trials, performed with these inhibitors in acute leukemia patients, have provided quite disappointing results (Konopleva et al., 2014; Perl et al., 2009). Therefore, new drugs targeting this pathway, including PI3K p110 inhibitors, should be tested.

In conclusion, our findings strongly advocate the use of PI3K p110 inhibitors for improving the clinical outcome of pediatric B-ALL patients. A pan PI3K p110 inhibitor displayed the most powerful cytotoxic effects and synergized with DEX. However, also a dual γ/δ PI3K p110 inhibitor was effective in combination with DEX, even when used on DEX-resistant cells. Therefore, IPI-145 could have a clinical relevance not only for B-ALL patients that respond to GCs by decreasing the drug dosage needed and adverse effects, but also for patients who do not respond to GCs. Moreover, the use of a dual inhibitor could spare the patients the side effects elicited by the inhibition of all the four PI3K p110 catalytic subunits (Greenwell, Flowers, Blum, & Cohen, 2017).

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest with respect to the authorship and/or publication of this article.

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