**Glucose transporter-1 inhibition overcomes imatinib resistance in gastrointestinal stromal tumor cells**

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**Abstract.** Imatinib mesylate (imatinib) is the primary agent of choice used to treat gastrointestinal stromal tumors (GIST). However, drug resistance to imatinib poses a major obstacle to treatment efficacy. In addition, the relationship between imatinib resistance and glycolysis is poorly understood. Glucose transporter (GLUT)-1 is a key component of glycolysis. The present study aimed to assess the potential relationship between components in the glycolytic pathway and the acquisition of imatinib resistance by GIST cells, with particular focus on GLUT-1. An imatinib-resistant GIST cell line was established through the gradual and continuous imatinib treatment of the parental human GIST cell line GIST-T1. The expression of glycolysis-related molecules (GLUT-1, hexokinase 2, pyruvate kinase M2 and lactate dehydrogenase) was assessed in parental and imatinib-resistant cells by western blotting, reverse transcription-quantitative PCR and glucose and lactate measurement kits. In addition, clinical information and transcriptomic data obtained from the gene expression omnibus database (GSE15966) were used to confirm the in vitro results. The potential effects of GLUT-1 inhibition on the expression of proteins in the glycolysis (GLUT-1, hexokinase 2, pyruvate kinase M2 and lactate dehydrogenase) and apoptosis pathways (Bcl-2, cleaved PARP, caspase-3 and caspase-9) in imatinib-resistant cells were then investigated following gene silencing and treatment using the GLUT-1 inhibitor WZB117 by western blotting. For gene silencing, the mature siRNAs for SLC2A1 were used for cell transfection. Annexin V-FITC/PI double-staining followed by flow cytometry was used to measure apoptosis whereas three-dimensional culture experiments were used to create three-dimensional spheroid cells where cell viability and spheroid diameter were measured. Although imatinib treatment downregulated GLUT-1 expression and other glycolysis pathway components hexokinase 2, pyruvate kinase M2, and lactate dehydrogenase in parental GIST-T1 cells even at low concentrations. By contrast, expression of these glycolysis pathway components in imatinib-resistant cells were increased by imatinib treatment. WZB117 administration significantly downregulated AKT phosphorylation and Bcl-2 expression in imatinib-resistant cells, whereas the combined administration of imatinib and WZB117 conferred synergistic growth inhibition effects in apoptosis assay. WZB117 was found to exert additional inhibitory effects by inducing apoptosis in imatinib-resistant cells. Therefore, the present study suggests that GLUT-1 is involved in the acquisition of imatinib resistance by GIST cells, which can be overcome by combined treatment with WZB117 and imatinib.

**Introduction**

Gastrointestinal stromal tumors (GIST) is the most common mesenchymal tumor of the digestive tract (1). The incidence of GIST is between 10 and 15 cases per 1 million worldwide (2). Imatinib mesylate (imatinib) is the primary agent of choice used for GIST treatment, which inhibits the expression of the tyrosine kinase receptor gene (KIT) (3). Imatinib is typically administered to patients with unresectable primary GIST or
metastasized recurrent GIST (3). However, drug resistance to imatinib poses a major concern reducing the efficacy of GIST treatment. Generally, resistance to imatinib develops after a median of 18-24 months of treatment (4,5). Therefore, characterization of the mechanism underlying the acquisition of imatinib resistance in patients with GIST is essential for developing novel treatment strategies.

Cancer cells metabolize glucose through aerobic glycolysis in abundant levels of oxygen (6). Although the quantity of ATP produced using mitochondrial oxidative phosphorylation by tumor cells is lower compared with that produced by healthy cells, they proliferate rapidly by using the pentose phosphate pathway for nucleic acid synthesis (7,8). Cancer cells typically exhibit increased rates of glucose uptake and glycolysis (9). This key feature of cancer cells is called the Warburg effect (10,11). In terms of this phenomenon, a number of studies have reported an association between cancer-specific energy metabolism and cancer progression in non-small cell lung cancer, breast ductal carcinoma and brain tumor (11-13). In addition, several studies have reported that the Warburg effect can facilitate to chemotherapeutic resistance (14,15).

The rate of glycolysis in cancer cells depends on the presence of key glycolytic components, including glucose transporters (GLUT) (16). GLUT is the first rate-limiting factor in the glucose metabolic pathway (17). GLUT-1 is primarily responsible for basal glucose uptake and the maintenance of basal glucose metabolism, where it serves as a key modulator of intracellular ATP production (18,19). High GLUT-1 expression has been previously associated with high tumor aggressiveness and high rates of tumorigenesis in GIST, non-small cell lung cancer, breast ductal carcinoma and gastric adenocarcinoma (13,20,21). Furthermore, high GLUT-1 expression has been linked to chemotherapeutic resistance in colon cancer and in head and neck cancer cell lines (22,23). Although a numerous studies have reported an association between imatinib resistance and increased rates of glycolysis (24-28), the mechanism of action and role of GLUT-1 underlying this entire process in GIST remains unclear.

Evasion of apoptosis is a key characteristic feature of cancer cells (29). Specifically, disequilibrium between the activities of pro-and anti-apoptotic Bcl-2 proteins can determine cancer cell survival (30). Overexpression of anti-apoptotic Bcl-2 proteins has been frequently associated with recurrence, poorer prognosis and resistance to various cancer therapies (31). Furthermore, silencing GLUT-1 expression has been reported to induce apoptosis and inhibit cell proliferation in breast cancer cells (30).

Therefore, the present study aimed to investigate the relationship between the expression of components in the glycolytic pathway and imatinib resistance in GIST cells, with emphasis on GLUT-1. The aim of the present study was to explore a potential treatment strategy for overcoming drug resistance in imatinib-resistant GIST cells.

Materials and methods

Cell culture and cell viability assessment. The human gastrointestinal stromal tumor cell line GIST-T1 (cat. no. PMC-GIST01C) showing a negative result for mycoplasma contamination was purchased from Cosmo Bio Co., Ltd., who performed cell line authentication and characterization. The GIST-T1 cell line has a 57-nucleotide in-frame deletion in the KIT exon, which leads to the production of KIT proteins without the valine 570 and tyrosine 578 residues (32). The GIST-T1 cells were cultured in DMEM supplemented with 10% (v/v) FBS and glutamine without penicillin/streptomycin (HyClone; Cytiva) supplementation at 37°C in a humidified atmosphere containing 5% CO₂.

Cell viability. The GLUT-1 inhibitor WZB117 was obtained from Selleck Chemicals. A total of 5x10⁴ cells/well were seeded into 96-well plates. DMSO (control), imatinib (2, 3.25, 4, 6.5, 8, 10, 13, 20, 26, 50, 52, 100, 200 and 500 nM) and/or WZB117 (2.5, 5, 10, 20, 40 and 80 µM) were administered either alone or in combination, respectively. The treatment of cells was at 37°C in a humidified atmosphere containing 5% CO₂ for 72 h. The number of viable cells was determined by using the MTT assay (Sigma-Aldrich; Merck KGaA) assay to calculate the half-maximal inhibitory concentration (IC₅₀) of imatinib and WZB117 either alone or in combination (33). The formazan crystals were dissolved using DMSO. The wavelengths used for the measurements in each well were 550 and 670 nm. The OD value of the control was set at 100 and the percentage of viable cells was calculated relative to the control. The IC₅₀ values were calculated according to the equation in the Boltzman sigmoidal concentration response curve using nonlinear regression fitting models (Graph Pad Prism Version 8; GraphPad Software, Inc.). The combination index (CI), an indicator of drug-drug interactions in combination chemotherapy based on the median-effect principle of the mass-action law (34), was calculated using the Compusyn software Version 1.0 (https://www.combosyn.com/index.html). The fraction of growth inhibition was calculated from the MTT assay data, where each corresponding concentration was entered into Compusyn. CI values at 50% growth inhibition by drug combination treatment (CI₅₀), which are the most reliable CI values, were then calculated using Compusyn according to previous reports (35,36). CI values <1, 1, and >1 indicate synergistic, additive and antagonistic effects, respectively.

Induction of imatinib-resistance. GIST-T1 cells were treated with 1 nM imatinib (Gleevec®; Adooq Bioscience) to generate imatinib-resistant GIST cells. After treatment the cells were cultured at 37°C in a humidified atmosphere containing 5% CO₂. The cells were passaged to 70-80% confluency. The concentration of imatinib was kept constant until the cells start to proliferate. The concentration was then gradually increased from 1 to 100 nM over a period of ~10 months, which was repeated to obtain the imatinib-resistant GIST-T1 cell line (GIST-T1/IM-R). GIST-T1/IM-R cells obtained were cultured in the continuous presence of 100 nM imatinib before use. After the establishment of GIST-T1/IM-R, Sanger sequencing (Genewiz, Inc.) was performed to confirm mutations in KIT (exons 8, 9, 11, 13, 14, 17 and 18) and in platelet-derived growth factor receptor α (PDGFRα) (exons 12, 14 and 18).

Western blotting. A total of 5x10⁴ cells/well were seeded into 6-well plates. After each treatment with different concentrations of DMSO (control), imatinib (0, 6.5, 13 and 500 nM), gene
silencing or WZB117 (10 µM) either alone or in combination for 72 h at 37°C, whole cells were homogenized in a chilled RIPA assay buffer containing 25 mmol/l Tris-HCl (pH 7.6), 1% NP-40, 0.1% deoxycholic acid, 0.1% SDS, and 150 mmol/l NaCl (Thermo Fisher Scientific, Inc.) with protease inhibitor cocktail 1 (cat. no. 25955-24; Nacalai Tesque, Inc.), protease inhibitor cocktail 2 (cat. no. P5726; Sigma-Aldrich; Merck KGaA), and protease inhibitor cocktail 3 (cat. no. P0044; Sigma-Aldrich; Merck KGaA) and incubated for 15 min on ice. After centrifugation at 13,697 x g for 20 min at 4˚C, the supernatants containing whole-cell protein samples were collected. Protein content was measured using a DC Protein Assay Kit (Bio-Rad Laboratories, Inc.). The protein lysate (7 µg) was extracted using NucleoSpin miRNA (cat. no. U0971B; Takara Bio, Inc.) and the Applied Biosystems 7500 Real-Time PCR System (Thermo Fisher Scientific, Inc.). The reaction conditions were 37°C (15 min), 85°C (5 sec) and then the reaction was stopped at 4°C. Predesigned TaqMan fluorogenic probes and primer sets for the solute carrier family 2 member 2 (SLC2A1; Hs00892681 m1), HK2 (Hs00606086 m1), LDHA (Hs01378790 g1) and GAPDH (Hs03929097 g1) were purchased from Applied Biosystems (Thermo Fisher Scientific, Inc.). A dual-labeled fluorescent (fluorescein amidite and minor groove binder) probe and a primer set for PKM2 (PrimePCR™ Probe Assay: PKM2, Human; qHsaCIP004657) were purchased from Bio-Rad Laboratories, Inc. THUNDERBIRD™ Probe qPCR Mix (cat. no. QPS-101) were purchased from Toyobo Life Science. The thermocycling conditions were 95°C (20 sec) followed by 40 cycles 95°C (3 sec) and 60°C (30 sec). Relative quantification was normalized to GAPDH expression using the 2^ΔΔCt method (39).

Measurement of glucose and lactate concentrations. A total of 5x10^3 cells/well were seeded into 96-well plates. After each treatment with different concentrations of imatinib (0, 6.5 and 13 nM) for 72 h at 37°C, the culture medium was collected and cell viability and glucose and lactate concentrations were measured using the CellTiter-Glo 3D Cell Viability Assay (cat. no. G9681; Promega Corporation), Glucose Uptake-Glo Assay (cat. no. J1341; Promega Corporation) and the Lactate-Glo Assay (cat. no. J5021; Promega Corporation), respectively. All procedures were performed according to the manufacturer’s protocols of each kit. Absorbance was measured using the GloMax Multi plus Detection System (Promega Corporation) to calculate the relative concentrations of lactate and glucose.

Gene expression omnibus (GEO) database. Clinical information and previously published transcriptionomics data were obtained from the GEO database. The data reported by Rink et al. (40) (GSE15966; 18-paired human GIST tissues samples) (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE15966) were used to determine the effect of imatinib administration on the expression of the genes of interest. This gene expression profiling was performed using oligonucleotide microarrays on tumor samples obtained before and after imatinib therapy. The inclusion criteria are that the dataset that it contains the response data of imatinib therapy as well as expression data. There were no specific exclusion criteria. The non-responder group (tumor shrink rate ≤0) and the responder group (tumor shrinkage rate >0) were designated (40). These data were analyzed using the publicly available software R (version 4.0.1; https://stat.ethz.ch/pipermail/r-announce/2020/000655.html) to generate a boxplot and statistical significance of the association between the groups was evaluated using paired t-test.

Gene silencing or inhibition experiments. WZB117 is a small compound that inhibits GLUT-1-mediated sugar transport by binding reversibly at the exofacial sugar-binding site (41). The mature siRNAs for SLC2A1 [HSS109812 (siR-SLC2A1 #1; sense, 5'-GCCCAUGUAUGUGGUGAGUCA-3' and antisense, 5'-UGACACUUCCACCCACAUAGUGGC-3')] and HSS109811 (siR-SLC2A1 #2; sense, 5'-GGCGGAUUUUAGUCGAUUGAG-3' and antisense: 5'-UUCAUC
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AUCAGCAUUUGAAUUCGCC-3′ Human Stealth Select RNAi; Invitrogen; Thermo Fisher Scientific, Inc.) were used for the transfection of the cells.

A total of 5x10^4 cells/well were seeded into 6-well plates 1 day before each treatment. The cells were treated with imatinib (13 nM) and WZB117 (10 µM) either alone or in combination for 72 h at 37˚C. The inhibition level was assessed from 0 to 72 h of treatment. DMSO was used to control for non-specific effects during the inhibition experiments. For gene silencing, a total of 5x10^4 cells/well were seeded into 6-well plates. At 6 h after seeding, transfection was performed using Lipofectamine RNAiMAX (Invitrogen; Thermo Fisher Scientific, Inc.). Silencer negative control siRNA (cat. no. 4390843; Invitrogen; Thermo Fisher Scientific, Inc.) was used as a control for nonspecific effects. In total, six experimental groups were defined based on the treatment of cells with the following for 72 h: i) DMSO (control); ii) imatinib (13 nM); iii) siR-SLC2A1 #1 (10 nM) alone; iv) siR-SLC2A1 #2 (10 nM) alone; v) siR-SLC2A1 #1 in combination with imatinib; and vi) siR-SLC2A1 #2 in combination with imatinib. The cells were assessed 72 h after transfection. Silencer negative control siRNA was administered as a negative control for i) and ii).

Apoptosis assay. Cells were seeded into cell culture flasks and treated with the indicated reagents for 72 h and then harvested for Annexin V-FITC/PI staining on 1x10^6 cells using the Annexin V-FITC Early Apoptosis Detection Kit (cat. no. 6592; Cell Signaling Technology, Inc.). To 1x10^6 cells, 1 µl Annexin V-FITC and 12.5 µl PI were added and the cells were stained for 10 min on ice. The double-stained cells were then analyzed using BD FACSAria II (BD Biosciences). Data were analyzed using the CellQuest™ software (BD Biosciences). In total, ≥10,000 cells were analyzed for each sample using the FlowJo software (FlowJo LLC).

Three-dimensional culture experiment. Cells were seeded into a 96-well plate (EZ-BindShunt® II; AGC Techno Glass Co., Ltd.) at a concentration of 0.3x10^4 cells/well 48 h before each experiment. Imatinib (13 or 26 nM) and WZB117 (10 µM) were administered either alone or in combination. At 72 h after administration, the three-dimensional cell viability was assessed using a CellTiter-Glo 3D Cell Viability Assay (Promega Corporation) according to the manufacturer’s protocols and measured using the GloMax Multi Detection System (Promega Corporation). Two-dimensional images of three-dimensional GIST-T1/IM-R cell spheroids were taken using light microscopy (BZ-X700; Keyence Corporation) at x40 magnification (37). The diameter of the spheroid was measured by BZ-X Analyzer software (version 1.4.0.1; Keyence Corporation).

Statistical analysis. Each experiment was performed ≥ three times. Statistical significance between two groups was determined using the two-sided unpaired t-test. The publicly available software R (version 4.0.1; https://stat.ethz.ch/pipermail/r-r-announce/2020/000665.html) was used to plot the data obtained from the analysis of the glycolysis-related genes in GSE15966. Statistical significance among ≥ three groups was determined using one-way analysis of variance and Tukey’s post hoc test were used for statistical analysis. The values are presented as the mean ± standard deviation. P<0.05 was considered to indicate a statistically significant difference.

Results

Effects of imatinib on GIST-T1 and GIST-T1/IM-R cells. The IC_{50} of imatinib in GIST-T1 and GIST-T1/IM-R cells was calculated to be 6.50 and 127.1 nM, respectively, suggesting that imatinib resistance in GIST-T1/IM-R cells was >19-fold higher compared with that in GIST-T1 cells.
Sequencing analysis revealed that GIST-T1/IM-R cells harbored a secondary PDGFRA mutation on exon 12 (c.1701A>G) in addition to a deletion in KIT exon 11 (Fig. S1). Therefore, for the standard dosing concentrations of imatinib in the present study, 6.5 nM was set (IC_{50} of the parental GIST-T1 cells), whilst 13 nM was also set (double the IC_{50}).

(Fig. 1A). Therefore, for the standard dosing concentrations of imatinib in the present study, 6.5 nM was set (IC_{50} of the parental GIST-T1 cells), whilst 13 nM was also set (double the IC_{50}). In
addition, 500 nM was identified to be at the upper limit of the imatinib dosage, which reduced cell viability of GIST-T1/IM-R cells by ~80%. The effect of imatinib administration on the activity of components in the KIT cascade in GIST-T1 and GIST-T1/IM-R cells was subsequently examined. The phosphorylation of KIT, AKT and ERK 1/2 remained unchanged after imatinib administration even at 13 nM in both cell lines tested (Figs. 1B and S2). Treatment with 500 nM imatinib significantly decreased the phosphorylation of KIT, AKT and ERK1/2 in both GIST-T1 and GIST-T1/IM-R cells compared with that in untreated controls (Figs. 1B and S2). Imatinib administration reduced GIST-T1 cell viability even at low concentrations without affecting the phosphorylation of KIT, AKT and ERK1/2. Therefore, other mechanisms independent of KIT phosphorylation may have mediated this suppression of viability after low-dose imatinib administration.

**Effect of imatinib administration on the glycolysis pathway in GIST-T1 and GIST-T1/IM-R cells.** The effect of imatinib administration on the expression of glycolysis-related genes GLUT-1, HK2, PKM2 and LDHA in both GIST-T1 and GIST-T1/IM-R cells was next examined. Administration of 13 nM imatinib, a concentration that did not affect KIT expression or phosphorylation, decreased the mRNA expression of SLC2A1, HK2, PKM2 and LDHA in parental GIST-T1 cells, but instead increased their expression in GIST-T1/IM-R cells (Fig. 2A). GLUT-1 levels in GIST-T1 cells was decreased in a dose-dependent manner after treatment with a low doses imatinib (6.5 or 13 nM), whereas those in GIST-T1/IM-R cells were increased in a dose-dependent manner (Figs. 2B and S3). The expression and/or phosphorylation profile of glycolysis-related proteins PKM2 and LDHA in both cell lines were similar to that of GLUT-1 expression (Figs. 2B and S3). Results from glucose and lactate assays showed that both glucose uptake and lactate production were decreased in GIST-T1 cells after imatinib treatment but were instead increased in GIST-T1/IM-R cells (Fig. 2C). Furthermore, the effects of imatinib administration on GLUT-1, HK2 and LDHA...
expression in human GIST tissues obtained before and after imatinib therapy were measured by utilizing the GSE19566 dataset. In total, 18 paired samples were categorized into the following two groups: Non-responder group (tumor shrink rate, ≤0) and the responder group (tumor shrink rate, >0). Although the expression of \( \text{GLUT-1} \) was significantly downregulated after imatinib treatment in the responder group, it did not change significantly in the non-responder group (Fig. 2D). These results suggest that alterations in \( \text{GLUT-1} \) expression contributed to imatinib resistance in GIST-T1 cells.

**Effect of \( \text{SLC2A1} \) gene silencing on imatinib-resistant GIST cells.** To determine the effects of imatinib treatment on \( \text{GLUT-1} \) expression in imatinib-resistant cells, \( \text{SLC2A1} \) expression was knocked down in GIST-T1/IM-R cells. Transfection with siR-\( \text{SLC2A1} \) downregulated the expression of \( \text{SLC2A1} \) and \( \text{GLUT-1} \) (Fig. 3A and B). In addition, significant reduction in cell viability was observed in GIST-T1/IM-R cells transfected with siR-\( \text{SLC2A1} \) and/or treated with imatinib compared with that in control (Fig. 3C). Furthermore, combined transfection with siR-\( \text{SLC2A1} \) and treatment with imatinib induced significant reductions in cell viability in GIST-T1/IM-R cells compared with that in cells treated with imatinib alone or transfected with siR-\( \text{SLC2A1} \) alone (Fig. 3C). However, HK2, p-PKM2 and LDHA protein levels did not show significant changes in any of the siR-\( \text{SLC2A1} \)-transfected groups compared to
control or imatinib (13 nM) alone (Figs. 3D and S4). These findings suggest that in GIST-T1/IM-R cells, transfection with siR-SLC2A1 and imatinib treatment significantly reduced cell viability but did not affect the expression of glycolysis-related proteins, implying that other mechanisms may have contributed to the reduced cell viability.

Effect of the GLUT-1 inhibitor WZB117 on GIST-T1/IM-R cell viability and glycolysis. The effect of the GLUT-1 inhibitor WZB117 on GIST-T1/IM-R cells was next assessed. The IC_{50} of WZB117 was calculated to be 15.8 µM in GIST-T1/IM-R cells (Fig. S5). Based on this result, 10 µM was set as the standard administration concentration of WZB117, whereas 13 nM was set as the standard administration concentration of imatinib. GIST-T1/IM-R cells were treated with imatinib (0-104 nM) and/or WZB117 (0-80 µM) at a molar rate of 13:10,000 before cell viability was assessed. The CI_{50} value was calculated to be 0.708 (CI value, <1) for both imatinib and WZB117, indicating a synergistic inhibitory effect on GIST-T1/IM-R cell viability (Fig. 4A). Considerable reduction in cell viability was also observed after combination treatment with 13 nM imatinib and 10 µM WZB117 in GIST-T1/IM-R cells compared with that in control or imatinib (13 nM) alone (Fig. 4B). Changes in the expression of glycolysis-related proteins in WZB117-treated GIST-T1/IM-R cells was next measured. GLUT-1 protein expression was reduced to the greatest extent at 6 h after WZB117 treatment, but the protein levels of p-PKM2 and LDHA at 6 h after WZB117 treatment remained unchanged (Figs. 4C and S6A). The protein expression level of HK2 was significantly increased at 48 and 72 h after WZB treatment compared with that in the control group (Figs. 4C and S6A). Similarly, the protein expression of GLUT-1 was significantly downregulated in GIST-T1/IM-R cells treated with both imatinib and WZB117 compared to control or imatinib alone, but those of the other glycolysis-related proteins remained unchanged (Figs. 4D and S6B). These findings also suggest that in GIST-T1/IM-R cells, although GLUT-1 suppression contributed to the reduction of cell viability, factors other than glycolysis were associated with this effect.

Effect of treatment of imatinib, siR-SLC2A1 transfection and/or WZB117 treatment on apoptosis in GIST-T1/IM-R cells. The effects of imatinib treatment and/or siR-SLC2A1 transfection on apoptosis in GIST-T1/IM-R cells were first
investigated. Both treatment with imatinib and siR-SLC2A1 transfection significantly decreased AKT phosphorylation in GIST-T1/IM-R cells compared to control or imatinib alone. However, the expression of Bcl-2, an apoptosis suppressor, remained unchanged in GIST-T1/IM-R cells (Figs. 5A and S7). Treatment with both imatinib and siR-SLC2A1 transfection increased the expression of cleaved PARP and caspase-9 compared with that in the imatinib alone group. However, no significant changes were observed in the expression levels of cleaved PARP, caspase-3 and caspase-9 in siR-SLC2A1-transfected cells treated with or without imatinib (Figs. 5A and S7). Annexin V-FITC/PI staining showed that siR-SLC2A1
transfection exerted no significant effects on the apoptosis in GIST-T1/IM-R cells when combined with imatinib treatment (Fig. 5B). The phosphorylation of AKT (6 h after treatment) and Bcl-2 (12 h after treatment) in GIST-T1/IM-R cells were both decreased after combined treatment with WZB117 and imatinib compared to control or imatinib alone (Figs. 6A and S8). By contrast, the protein levels of cleaved PARP, caspase-3 and caspase-9 were increased in the WZB117 and imatinib combined group compared with those in the control or imatinib groups alone (Figs. 6A and S8). Annexin V-FITC/PI staining revealed the induction of apoptosis after the combined treatment of GIST-T1/IM-R cells with WZB117 and imatinib compared to control or imatinib alone (Fig. 6B). These findings suggest that the combination of siR-SLC2A1 transfection or WZB117 treatment and imatinib suppressed not only the expression of GLUT-1 protein, but also the phosphorylation of AKT in GIST-T1/IM-R cells. In addition, combined treatment with WZB117 and imatinib resulted in significant increases in apoptosis. This result may have been caused by reductions in Bcl-2 protein expression.

**Effect of combination treatment with imatinib and WZB117 on proliferation of GIST-T1/IM-R cell spheroids.** The effect of combined imatinib and WZB117 treatment on the proliferation of GIST-T1/IM-R cells was next examined. WZB117 treatment-induced disaggregation of the edges of colonies, where treatment with WZB117 alone or in combination with 26 nM imatinib significantly suppressed spheroid diameter and cell viability compared with those in the control group (Fig. 6). Treatment with WZB117 and 26 nM imatinib in combination significantly suppressed spheroid diameter and cell viability compared with those in WZB117 alone (Fig. 6C-E). This suggest that this combined treatment was effective in inhibiting the proliferation of imatinib-resistant GIST cells.

These results suggest that GLUT-1 is involved in the acquisition of imatinib resistance, where combined administration of the GLUT-1 inhibitor WZB117 and imatinib may reverse this imatinib resistance in GIST-T1/IM-R cells (Fig. 7).

**Discussion**

The present study revealed that GLUT-1, a key glycolytic molecule, is involved in the acquisition of imatinib resistance by GIST cells. In addition, combined treatment with the GLUT-1 inhibitor WZB117 and imatinib mediated synergistic inhibitory effects by inducing apoptosis in imatinib-resistant GIST cells.

Imatinib resistance in GIST cells poses an additional obstacle due to limited availability of antitumor agents for unresectable GIST treatment (42-45). Recent studies have reported that imatinib-resistant GIST cells exhibit high glycolysis rates (24,25), which may contribute to imatinib resistance. However, it was previously reported that imatinib-resistant GIST cells exhibited higher oxidative phosphorylation and higher glycolytic rates compared with those in imatinib-sensitive GIST cells (24). However, the exact mechanism underlying the acquisition of imatinib resistance remains unclear, where a diverse range of metabolic modifications may be involved (24,25,46). In addition, it has been reported that the PI3K/AKT/mTOR pathway can be partially activated after secondary imatinib resistance in GIST (27). The possible involvement of GLUT-1 in the acquisition of imatinib resistance in BCR-ABL-positive leukemia cells has also been reported, where combining imatinib with Bcl-2 antisense oligonucleotides has been shown to significantly increase apoptosis in imatinib-resistant cells (26,28). However, the function of GLUT-1 in the acquisition of imatinib resistance in GIST cells remains unclear.

Although imatinib can inhibit KIT, downregulates AKT expression and causes the transient dephosphorylation of ERK1/2, the imatinib concentrations required for inhibiting KIT phosphorylation have not been found to correlate well with the proliferation of GIST cells (47). The present study showed that administration of imatinib at the IC50 dose of did not inhibit KIT, AKT or ERK1/2 phosphorylation in both parental GIST and GIST-T1/IM-R cells. Although the administration of low-dose imatinib did not inhibit KIT phosphorylation, significant suppression of the expression of components in

Figure 7. Schematic model from the present study. Low-dose imatinib treatment suppressed glycolysis in the parental gastrointestinal stromal tumor GIST-T1 cells, whereas glycolysis was enhanced in imatinib-resistant GIST-T1 cells. The glycolytic pathway may serve a pivotal role in the survival of GIST-T1 cells. Alterations in the glycolytic pathway contributes to imatinib resistance. In addition, GLUT-1 inhibitor WZB117 treatment overcame imatinib resistance by inducing intrinsic apoptosis in imatinib-resistant GIST cells. Data from the present study also suggest that WZB117 administration causes growth inhibition in imatinib-resistant GIST cells by suppressing p-AKT and Bcl-2. GLUT-1, glucose transporter 1.
the glycolysis pathway, glucose uptake and lactate production were all observed in parental GIST-T1 cells. These results suggest that glycolysis may be an important pathway for energy production in GIST cells. In addition, low-dose imatinib administration to GIST-T1/IM-R cells resulted in increased glycolytic pathway activation. Therefore, the glycolysis pathway may be essential for GIST cells acquiring imatinib resistance and for cell survival.

The present study next examined the effects of GLUT-1 inhibition by gene silencing or WZB117 administration on the expression of glycolysis components. WZB117 is a small compound that inhibits GLUT-1-mediated sugar transport by binding reversibly at the exofacial sugar-binding site and has been reported to inhibit cell proliferation in a number of cancer cell lines, including such as non-small cell lung cancer, colon cancer and KRAS-mutant cancer (13,41,48,49). In the present study, the protein expression of GLUT-1 in GIST-T1/IM-R cells was suppressed at its greatest extent at 6 h after WZB117 administration, which increased again over time. Therefore, combination treatment of imatinib with WZB117 6 h after administration was performed. A previous report showed that the protein expression of GLUT-1 was temporarily decreased after WZB117 administration but increased again over time (13), suggesting that WZB117 administration does not permanently inhibit GLUT-1 activity. Inhibition of glucose transport by WZB117 likely reduced the supply of glucose into GIST-T1/IM-R cells, thereby necessitating an increase in glucose import and an upregulation of GLUT-1 protein expression. Although expression of the GLUT-1 protein was increased again, cell viability was reduced at 72 h after administration. Therefore, suppression of GLUT-1 protein expression by WZB117 may be involved in the survival of GIST-T1/IM-R cells, albeit transiently. Although WZB117 administration reduced cell viability even when administered alone, the CI₅₀ value, which measures the nature of drug-drug interaction in combination chemotherapy (34-36,50), showed that WZB117 was more effective in combination with imatinib. In siR-SLC2A1-transfected and WZB117-treated cells, administration of a low dose of imatinib considerably reduced cell viability. However, the expression of all proteins in the glycolysis pathway apart from GLUT-1 remained unchanged. These results suggest that a robust compensatory mechanism is operational when GLUT-1 is inhibited, since the glycolytic pathway is crucial for tumor cell function (51).

The effect of GLUT-1 inhibition on the AKT and ERK signaling pathways and apoptosis was next investigated in imatinib-resistant GIST-T1/IM-R cells. Combined administration of GULT-1 inhibition and imatinib suppressed AKT phosphorylation, which may have in turn inhibited GIST-T1/IM-R cell proliferation. However, transfection with siR-SLC2A1 and WZB117 treatment conferred different effects on the induction of apoptosis. SLC2A1 silencing did not significantly increase the apoptotic rate, whereas WZB117 treatment resulted in a considerable increase in apoptosis, especially when administered in combination with imatinib. Although SLC2A1 knockdown has been reported to induce apoptosis and inhibit cell proliferation in breast cancer cells, this phenomenon has not been observed in GIST cells (30). A previous report showed that Bcl-2 expression was downregulated by WZB117 treatment in breast cancer cells (35).

In the present study, unlike SLC2A1 knockdown, WZB117 administration suppressed the expression of Bcl-2, GLUT-1 and the phosphorylation of AKT in GIST-T1/IM-R cells. Furthermore, the WZB117 dose (10 µM) used was lower than its IC₅₀ in GIST-T1/IM-R cells (15.8 µM) or that used in a previous study for breast cancer cells (60 µM) (35). Therefore, the results obtained in the present study are unlikely the result of an off-target effect mediated by WZB117. Bcl-2 downregulation may be a key mechanism for the induction of apoptosis in GIST cells treated with imatinib and a GLUT-1 inhibitor in combination.

The relationship between GLUT-1 and AKT is highly complex. Previous studies have reported that GLUT-1 downregulation by suppressing AKT phosphorylation can inhibit pancreatic cancer and hepatocellular carcinoma growth (52,53). By contrast, SLC2A1 silencing has been shown to inhibit cell proliferation by promoting apoptosis in colorectal cancer cells by inactivating the TGF-β/PI3K/AKT/mTOR signaling pathway (54). Therefore, a possible interaction between GLUT-1 and AKT may exist, such that the suppression of one can suppress the other, where GLUT-1 inhibition may inhibit AKT phosphorylation in GIST cells. However, the possible underlying mechanism is not precisely understood at present and further investigation of these pathways is required. WZB117 administration was found to inhibit cell proliferation more efficiently compared with that mediated by SLC2A1 silencing, which was likely due to WZB117 inducing apoptosis by simultaneously suppressing Bcl-2 expression in GIST-T1/IM-R cells.

The present study has a number of limitations. Only one cell line was used. The GIST-T1/IM-R cells established in the present study had a PDGFRA mutation on exon 12. PDGFRA is one of the two PDGFR subunits, which transduces signals through numerous downstream pathways, including PI3K/AKT, Src kinase, phospholipase C/protein kinase C and Ras/Raf/MEK/ERK, all of which have been documented to serve important roles in tumorigenesis (55,56). The activation of PDGFRA may in turn activate the ERK pathway and lead to increased GLUT-1 expression (57). In the present study, a mutation in PDGFRA may have been involved in promoting GLUT-1 synthesis after imatinib treatment, thereby leading to increased cell survival. However, further experiments are required to clarify the relationship between this mutation and glycolysis.

To validate the in vitro results in the present study, additional experiments were performed using human sample data obtained from the GEO database. GLUT-1 expression was found to be significantly suppressed in the responder group compared with that in the non-responder group after imatinib treatment, though the number of sample specimens used was not high (18 pairs). Although this result is in line with the present finding that GLUT-1 expression is involved in the acquisition of imatinib resistance, HK2 expression was significantly suppressed after imatinib treatment in the non-responder group. It may be the complex metabolic morphology in the actual clinical GIST specimen. Therefore, further investigation involving the accumulation of non-responder clinical samples to imatinib is required. In addition, the possible side effects of WZB117 administration was not validated in vivo in the present study, although
they were attempted by transplanting GIST-T1/IM-R cells into mice. However, transplantation proved to be difficult. Therefore, a three-dimensional model examination of the GIST-T1/IM-R cells was performed instead. In this model, administration of 26 nM imatinib and 10 µM WZB117 exerted a synergistic effect, possibly due to the lower efficiency of imatinib treatment in three-dimensional conditions than that in two-dimensional conditions (38). Further validation of these phenomena is required. In addition to the in vivo validation of the possible therapeutic effects of WZB117, it is important to identify the optimal drug delivery system and any side effects of WZB117. Data obtained in this preclinical study should be used to guide the planning of clinical trials involving patients with imatinib-resistant GIST.

In the present study, a key role of GLUT-1 in the acquisition of imatinib resistance by GIST cells was revealed. Furthermore, combined treatment with the GLUT-1 inhibitor WZB117 and imatinib was shown to overcome this resistance.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors’ contributions

Conception and design of the study: TS and KoT. Experiments: TS (in vitro experiments) and YT (data set analysis). Analysis and interpretation of data: TS, KoT, YT, YI, JA, SWL, KY, KaT and KU. All authors read and approved the final manuscript. TS and KT confirm the authenticity of all the raw data.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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