| タイトル | Title |
|---------|-------|
| Glucose metabolism-targeted therapy and withaferin A are effective for epidermal growth factor receptor tyrosine kinase inhibitor-induced drug-tolerant persisters |

| 著者 Author(s) |
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| Kunimasa, Kei / Nagano, Tatsuya / Shimono, Yohei / Dokuni, Ryota / Kiriu, Tatsunori / Tokunaga, Shuntaro / Tamura, Daisuke / Yamamoto, Masatsugu / Tachihara, Motoko / Kobayashi, Kazuyuki / Satouchi, Miyako / Nishimura, Yoshihiro |

| 掲載誌・巻号・ページ Citation |
|-----------------------------|
| Cancer Science, 108(7):1368-1377 |

| 刊行日 Issue date |
|-------------------|
| 2017-07 |

| 資源タイプ Resource Type |
|------------------------|
| Journal Article / 学術雑誌論文 |

| 版区分 Resource Version |
|------------------------|
| publisher |

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|------------|
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| DOI |
|-----|
| 10.1111/cas.13266 |

| JaLCDOI |
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|         |

| URL |
|-----|
| http://www.lib.kobe-u.ac.jp/handle_kernel/90004368 |

PDF issue: 2020-05-05
Glucose metabolism-targeted therapy and withaferin A are effective for epidermal growth factor receptor tyrosine kinase inhibitor-induced drug-tolerant persisters

Kei Kunimasa, Tatsuya Nagano, Yohei Shimono, Ryota Dokuni, Tatsunori Kiriu, Shuntaro Tokunaga, Daisuke Tamura, Masatsugu Yamamoto, Motoko Tachihara, Kazuyuki Kobayashi, Miyako Satouchi and Yoshihiro Nishimura

1Division of Respiratory Medicine; 2Division of Medical Oncology/Hematology, Department of Internal Medicine, Kobe University Graduate School of Medicine, Kobe; 3Department of Thoracic Oncology, Hyogo Cancer Center, Akashi, Japan

Key words
Cell interaction, drug resistance, EGFR-TKI, senescent cells, stem-like cells

Correspondence
Tatsuya Nagano, Division of Respiratory Medicine, Department of Internal Medicine, Kobe University Graduate School of Medicine, 7-5-1 Kusunoki-cho, Chuo-ku, Kobe 650-0017, Japan.
Tel: +81-78-382-5660; Fax: +81-78-382-5661; E-mail: tnagano@med.kobe-u.ac.jp

Funding Information
Eli Lilly Japan; AstraZeneca; Novartis Pharma; the Japan Society for the Promotion of Science; Kobe University School of Medicine Alumni Association Shinryokukai General Incorporated Association; Hyogo Medical Association.

Received September 25, 2016; Revised April 19, 2017; Accepted April 20, 2017

Cancer Sci 108 (2017) 1368–1377
doi: 10.1111/cas.13266

In pathway-targeted cancer drug therapies, the relatively rapid emergence of drug-tolerant persisters (DTPs) substantially limits the overall therapeutic benefit. However, little is known about the roles of DTPs in drug resistance. In this study, we investigated the features of epidermal growth factor receptor–tyrosine kinase inhibitor-induced DTPs and explored a new treatment strategy to overcome the emergence of these DTPs. We used two EGFR-mutated lung adenocarcinoma cell lines, PC9 and II-18. They were treated with 2 μM gefitinib for 6, 12, or 24 days or 6 months. We analyzed the mRNA expression of the stem cell–related markers by quantitative RT-PCR and the expression of the cellular senescence-associated proteins. Then we sorted DTPs according to the expression pattern of CD133 and analyzed the features of sorted cells. Finally, we tried to ablate DTPs by glucose metabolism-targeting therapies and a stem-like cell targeting drug, withaferin A. Drug-tolerant persisters were composed of at least two types of cells, one with the properties of cancer stem-like cells (CSCs) and the other with the properties of therapy-induced senescent (TIS) cells. The CD133high cell population had CSC properties and the CD133low cell population had TIS properties. The CD133low cell population containing TIS cells showed a senescence-associated secretory phenotype that supported the emergence of the CD133high cell population containing CSCs. Glucose metabolism inhibitors effectively eliminated the CD133low cell population. Withaferin A effectively eliminated the CD133high cell population. The combination of phloretin and withaferin A effectively suppressed gefitinib-resistant tumor growth.
mutations. The EGFR-TKI-induced DTPs show >100-fold reduced drug sensitivity and maintain viability through engagement of IGFR-1R signaling and an altered chromatin state that requires the HDAC RBP2/KDM5A/Jarid1A. Although an HDAC inhibitor and an IGFR-1R inhibitor did prevent the emergence of DTPs in vitro, the combination of an HDAC inhibitor (vornostat) and erlotinib did not show a meaningful clinical benefit in EGFR-mutated NSCLC patients. In addition, a phase III trial (ADWIGO 1018) that examined the effects of tiglitumub, a fully human anti-IGFR-1R G2 mAb, in combination with erlotinib as a second/third-line treatment in previously treated NSCLC patients, failed and was discontinued.

A major contributor to EGFR-TKI treatment failure is the formation and emergence of EGFR-TKI-induced DTPs. Conventional anticancer drug treatments are of minor benefit to overcome and prevent EGFR-TKI-induced DTPs. In this context, we investigated the features of EGFR-TKI-induced DTPs and explored a new treatment strategy to overcome the emergence of these DTPs.

Materials and Methods

Cells lines and reagents. Human EGFR-mutant PC9 (exon19del E746-A750; Sigma-Aldrich, St. Louis, MO, USA), HCCS87 (exon19del E746-A750; ATCC, Manassas, VA, USA) and II-18 (L858R; RIKEN Cell Bank, Riken, Tsukuba, Japan) cells were purchased. All of the cell lines were cultured in RPMI-1640 media supplemented with 10% FBS and 1% BSA (Sigma-Aldrich), 1 µg/mL insulin (Sigma-Aldrich), 1× B27 supplement (Life Technologies), and 0.4% BSA (Sigma-Aldrich). The cells were cultured under 5% CO2 at 37°C for 1 week.

Cell surface marker analysis using flow cytometry. Cells (1 × 106) were resuspended in 100 µL Hanks’ balanced salt solution supplemented with 2% BSA. Cells were incubated with 5 µL polyclonal mouse anti-human CD133-FITC conjugated antibody (1:20; Miltenyi Biotec, Bergisch Gladbach, Germany) and were analyzed using a BD FACSARia III (BD Biosciences, Franklin Lakes, NJ, USA). In this study, we sorted DTPs into CD133high and CD133low cell fractions based on CD133 expression level using the FACSARia III cell sorter.

Tumorigenicity in nude mice. Female BALB/c nude mice (6 weeks old) were purchased from SLC (Shizuoka, Japan). Mice were inoculated s.c. in the flank with 5.0 × 104 cells/100 µL suspension of parental PC9 cells and CD133high and CD133low-sorted PC9 DTPs. This study was approved by the Institutional Animal Care and Use Committee (permit number: 150108) and was carried out according to the Animal Experimentation Regulations of Kobe University (Kobe, Japan).

In vivo tumor growth inhibition assay. Suspensions of PC9 cells (5 × 105) were injected s.c. into the backs of female BALB/c nude mice (6 weeks old; SLC). The length (x) and width (y) of the tumor masses were measured twice a week, and tumor volume (TV) was calculated using the formula TV = (x × y2)/2. Relative tumor volume on day n was calculated using the formula: relative tumor volume = TVn / TV0, where TVn is the tumor volume on day n and TV0 is the tumor volume on day 0. When TV reached 100 mm3, mice were divided into five groups consisting of four mice per group (day 0) and were then allocated into one vehicle group and four gefitinib (6.25 mg/kg daily) groups. When the TV re-reached 100 mm3 after it was transiently reduced by gefitinib, the four gefitinib groups were re-allocated as follows: (i) a gefitinib (6.25 mg/kg daily) treatment group; (ii) gefitinib (6.25 mg/kg daily) and phloretin (10 mg/kg three times per week) combination treatment group; (iii) a gefitinib (6.25 mg/kg daily) and WFA (2 mg/kg three times per week) combination treatment group; and (iv) gefitinib (6.25 mg/kg daily), phloretin (10 mg/kg three times per week), and WFA (2 mg/kg three times per week) combination treatment group. Gefitinib was dissolved in DMSO and in glucose solution for the in vivo study. Phloretin and WFA were dissolved in methanol and in glucose solution for the in vivo study.

Results

Treatment with EGFR-TKI induces TIS cells and CSC marker expression. Drug-tolerant persisters were generated within a few days of exposure of PC9 and II-18 cells to a concentration (2 µM) of an EGFR-TKI (gefitinib) that was higher than the IC50 value. Their drug-tolerant phenotype was confirmed using flow cytometric and pathological analyses as described by Sharma et al. (data not shown). Because the genotoxic stress induced by anticancer drugs, including gefitinib, is known to induce cellular senescence (TIS), we investigated the characteristics of these gefitinib-induced DTPs, focusing on cellular senescence. β-Galactosidase staining revealed the induction of senescence in PC9 cells following gefitinib treatment (Fig. 1a). Similarly, Western blot analyses showed time-course-dependent upregulation of senescence-associated proteins (pRb, p21, and γH2AX) (Fig. 1b). Cellular
Senescence is considered to be a state of stable exit from the cell cycle and senescent cells are assumed to be permanently arrested. However, response to gefitinib treatment in EGFR-mutated lung cancer is often short-lived and has failed to result in cures. Hence, the acquisition of resistance and recurrence to gefitinib treatment is thought to result from a feature of DTPs other than the cellular senescence of cells. We therefore further explored whether DTPs have CSC properties. The upregulation of stem cell-related markers such as Oct3/4, Nanog, Sox2, Myc, CD133, CD44, ALDH1A1, ABCB, and ABCG following gefitinib treatment of PC9 and II-18 cells was confirmed using qRT-PCR (Fig. 1c). These data suggested that gefitinib induced DTPs that have both cellular senescent and CSC properties.

Gefitinib treatment increases the fraction of CD133high cells in gefitinib-induced DTPs. Next we determined if it was possible to discriminate senescent cells from CSCs of gefitinib-induced DTPs. Recent studies have indicated that NSCLC contains cells that express the glycoprotein prominin-1 (CD133), which is a CSC marker that is essential for tumor cell propagation and metastasis. We therefore examined the expression of CD133 in gefitinib-induced DTPs. We exposed PC9 cells to gefitinib (2 μM) and followed the CD133high cell population using flow cytometric analysis. We removed apoptotic PC9 cells (17%–31% of total cells) before flow cytometric analysis because such cells lose the CD133 membrane marker and could therefore bias the analysis. A significant increase in the CD133high cell population was observed 12 days after gefitinib treatment (Fig. 2). We could separate DTPs into two cell fractions based on CD133 expression level using the FACSAria III cell sorter.

CD133high cell population has CSC properties and CD133low cell population has senescent cell properties. To investigate the differences between CD133high and CD133low cell populations,
we first carried out Western blotting for senescence-associated proteins (Figs 3a,S1). The expression of these proteins was increased in the CD133<sup>high</sup> cell population compared to the CD133<sup>low</sup> population. The expression of stem cell-related markers was then evaluated using qRT-PCR (Figs 3b,S2) and relatively higher expression of these markers was observed in the CD133<sup>high</sup> cell population compared to the CD133<sup>low</sup> cell population. We next undertook an <em>in vitro</em> sphere forming assay (Fig. 3c) to explore CSC properties of the CD133<sup>high</sup> cell population. Compared to the parental and the CD133<sup>low</sup> cell population, the CD133<sup>high</sup> cell population gradually formed significantly more sphere colonies of various sizes and irregular shapes after approximately 1 week of culture. Furthermore, we injected the CD133<sup>high</sup> cell population s.c. into BALB/c nude mice. After approximately 10 days, the injected CD133<sup>high</sup> cells started to gradually form tumors (Fig. 3d). These results indicated that the CD133<sup>high</sup> cell population possessed strong tumorigenicity. Collectively, these data suggested that the CD133<sup>low</sup> cell population has senescent cell properties and the CD133<sup>high</sup> cell population has CSC properties.

**Senescence-associated secretory phenotype of CD133<sup>low</sup> cell population drives the emergence of CD133<sup>high</sup> cell population.** We then investigated the relationship between the CD133<sup>high</sup> and the CD133<sup>low</sup> cell populations. We first undertook <em>in vitro</em> assays that focused on the assay of soluble factors from the CD133<sup>low</sup> cell population. Recent increasing evidence indicates that senescent cells actively communicate with neighboring cells through a plethora of secretory factors including inflammatory cytokines, chemokines, and growth factors. This senescence-related phenotype is called the SASP. We evaluated the SASP of gefitinib-treated PC9 and II-18 cells using qRT-PCR (Fig. 4a). Gefitinib-induced SASP was enhanced in a time-course-dependent manner. Interestingly, the CD133<sup>low</sup> cell population containing senescent cells showed a stronger SASP than the CD133<sup>high</sup> cell population (Fig. 4b). To understand how the SASP of senescent cells might contribute to an increase in the proportion of CSCs (the CD133<sup>high</sup> cell population), we compared the effect of incubation of PC9 cells with gefitinib plus the conditioned medium of gefitinib-treated (12 days) cells with incubation of PC9 cells with gefitinib plus conditioned medium. When PC9 cells were cultured in the conditioned medium plus gefitinib, there was a greater induction of the CD133<sup>high</sup> cell population than when the PC9 cells were cultured in the standard, non-conditioned medium containing gefitinib (Fig. 4c). In contrast, neutralizing anti-bodies against some SASP effectively suppressed the induction of the CD133<sup>high</sup> cell population (Fig. S3). These results suggest that the SASP regulates an increase in the proportion of cancer stem cells. Rapamycin, an mTOR inhibitor and autophagy inducer, evokes cellular senescence but represses the SASP. We investigated whether rapamycin could suppress the SASP induced by gefitinib treatment. Quantitative RT-PCR analyses of the SASP showed that the addition of rapamycin suppressed the gefitinib-induced SASP (Fig. 4d). Furthermore, flow cytometric analysis indicated that the addition of rapamycin suppressed the gefitinib-induced increase in the proportion of the CD133<sup>high</sup> cell population (Fig. 4e). These data suggested that the gefitinib-induced senescent cells (the CD133<sup>low</sup> cell population) contributed to an increase in the proportion of CSCs (CD133<sup>high</sup> cell population) through the SASP.

**Glucose metabolism-targeting therapies and CSC targeting DTPs.** We determined whether conventional anticancer drugs including cisplatin and pemetrexed were effective against gefitinib-induced DTPs. We generated gefitinib-induced DTPs by exposing PC9 cells to gefitinib (2 μM) for 7 days. The DTPs were examined for their response to conventional anticancer drugs by using a growth inhibition assay (Fig. 5a). Conventional anticancer drugs were not effective against these DTPs. Recently, therapy of induced senescent cells has been reported to be dependent on senescence-related metabolic reprogramming, which is composed of enhanced glycolysis and a hypermetabolic phenotype. We therefore investigated whether gefitinib-induced senescent cells were dependent on glucose metabolism. Compared to the CD133<sup>high</sup> cell population, the CD133<sup>low</sup> cell population showed higher phosphorylation reflecting activation of AMPK (Figs 5b,S4) and qRT-PCR analysis showed upregulation of the glucose transporters Glut1.

Fig. 2. Epidermal growth factor receptor tyrosine kinase inhibitor treatment induces CD133<sup>high</sup> cells in drug-tolerant persisters. (a) PC9 cells treated with gefitinib (2 μM) for 12 days were stained with CD133 and analyzed using flow cytometry. Gating was used to identify CD133<sup>high</sup> cells. (b) CD133<sup>high</sup> cells were quantified using flow cytometry. Results are the means ± SD of three independent experiments. *<em>P</em> < 0.001.
Glut3, and the glycolytic enzyme hexokinase 2 (Hk2) in the CD133low cell population (Fig. 5c). These results suggested that the CD133low cell population that contained senescent cells was more dependent on glucose metabolism than the CD133high cell population. Thus, glucose metabolism targeting therapies and WFA, which are novel therapeutic agents for CSCs, are promising strategies for combating gefitinib-induced DTPs. To evaluate their efficacy, we assayed the

Fig. 3. Comparison of the features of CD133high and CD133low non-small-cell lung cancer cells. (a) CD133high and CD133low cells were sorted from PC9 drug-tolerant persisters 12 days after gefitinib (2 μM) treatment. Whole-cell lysates were prepared and analyzed by Western blotting as indicated. (b) mRNA expressions of stem cell-related markers in CD133high and CD133low cells sorted from PC9 drug-tolerant persisters 12 days after gefitinib (2 μM) treatment was analyzed using quantitative RT-PCR. Relative expression levels were obtained by dividing the mRNA level in CD133high by that in CD133low cells. Results are the means ± SD of three independent experiments. (c) Sphere-forming assay of parental, CD133high, and CD133low cells. Left, phase contrast images. Scale bar = 50 μm. Right, quantification of sphere number. CD133high cells showed an enhanced capacity for self-renewal compared with parental and CD133low cells. Results are the means ± SD of at least three independent experiments. *P < 0.01. (d) Xenograft tumor volume was measured on the indicated days following parental, CD133high, and CD133low cell injection (left panel). Results are means ± SD (four mice). Images show the CD133high tumors of four mice 18 days after injection (right panel).

Fig. 4. Senescence-associated secretory phenotype (SASP) induced by epidermal growth factor receptor–tyrosine kinase inhibitor treatment contributes to the emergence of CD133high cells in drug-tolerant persisters. (a) PC9 and II-18 cells were treated with gefitinib (2 μM) over the indicated time. An SASP was determined by analysis of the cellular mRNA levels of the indicated proteins using quantitative (q)RT-PCR. Results are means ± SD of three independent experiments. (b) mRNA expressions of SASP-associated proteins in CD133high and CD133low cells sorted from PC9 drug-tolerant persisters 12 days after gefitinib (2 μM) treatment was analyzed using qRT-PCR. Relative expression levels were obtained by dividing the mRNA level in CD133high by that in CD133low cells. Results are the means ± SD of three independent experiments. (c) Sphere-forming assay of parental, CD133high, and CD133low cells. Left, phase contrast images. Scale bar = 50 μm. Right, quantification of sphere number. CD133high cells showed an enhanced capacity for self-renewal compared with parental and CD133low cells. Results are the means ± SD of at least three independent experiments. *P < 0.01. (d) Xenograft tumor volume was measured on the indicated days following parental, CD133high, and CD133low cell injection (left panel). Results are means ± SD (four mice). Images show the CD133high tumors of four mice 18 days after injection (right panel).
effect of glucose metabolism-targeting therapies including 2DG and phloretin, and WFA on gefitinib-induced DTPs (Fig. 5a). Interestingly, these drugs had an antitumor activity against the parent cells as well as the DTPs. To assess the efficacy of phloretin and WFA in a separate in vivo model, we treated mice bearing xenografts of PC9 cells. To generate gefitinib-induced DTPs in this in vivo model, we first treated PC9 cell xenograft models with gefitinib (6.25 mg/kg daily) and, after the tumors had progressed on gefitinib treatment, we added phloretin, WFA, or their combination, to gefitinib. These additional drugs had a dramatic inhibitory effect on progressed tumor progress (Fig. 5d). Thus, glucose metabolism-targeting drugs and the CSC targeting drug, WFA, are promising drugs to overcome gefitinib-induced DTPs and to prevent cancer recurrence.

Discussion

In this study, we analyzed the features of gefitinib-induced DTPs. Our analysis suggested that gefitinib-induced DTPs are composed of at least two types of cells: one has the properties of CSCs, and the other has the properties of TIS cells. The FACS analysis of CD133 (Prominin-1) expression, a marker for CSCs, divided the DTPs into CD133high and CD133low cell populations. The CD133high cell population has CSC properties and the CD133low cell population has senescent cell properties. Numerous studies have found that high levels of CD133 expression confers chemotherapy resistance and cancer stemness on various types of cancer cells. (30–33) In glioblastoma, CD133 expression induces senescence by increasing expression of MDR1 through the PI3K-Akt signal pathway. (34) A high level of expression of CD133 therefore suggested that the CD133high cell population in gefitinib-induced DTPs possesses cancer stem-like properties and chemoresistance.

Whether DTPs emerge de novo during gefitinib treatment or exist in the parental cell population before treatment is an intriguing and significant question. Recent studies have reported that NSCLC cells before treatment contained CD133 high expressing population with strong self-renewal and multipotent differentiation capacities. (35,36) The ClonTracer barcode system tracking of more than 1 million individual cells in cultured cancer models revealed that chemoresistant clones are present before treatment. Gefitinib treatment abolishes chemosensitive parental clones. However, CD133-expressing parental chemoresistant clones would survive this treatment and proliferate, resulting in the emergence of CD133-expressing DTPs. The CD133low cell population would be induced in response to gefitinib-induced genotoxic stress. In the current study (Fig. 1b), senescence-associated protein expression was identified in parental cells before treatment but initially disappeared after gefitinib treatment. These proteins were re-expressed on day 12 after gefitinib treatment. The senescence-associated protein bands observed by Western blotting before treatment (day 0) imply oncogene-induced senescence, which is characterized as a proliferative arrest elicited by oncogenic RAS and RAF through induction of p16ink4a. (38,39) However, the senescence-associated protein bands that emerged after treatment imply TIS, which is thought to be an irreversible cell cycle arrest induced by anticancer drug treatment, including gefitinib, through genotoxic stress of the cancer cells. (24) Based on the above, it was considered that parental oncogene-induced senescence cells abolished by gefitinib treatment and that chemoresistant TIS cells emerged de novo.

The TIS CD133low cell population showed an SASP (Fig. 4a,b). Recently, increasing evidence from sophisticated mouse models indicates that senescent cells actively communicate with neighboring cells through an SASP. (26,27) Our analysis showed that the CD133low cell population has higher potential for secreting various inflammatory cytokines and chemokines including interleukin-6, transforming growth factor-β, tumor necrosis factor-α than the CD133high population. Some studies have found that these pro-inflammatory cytokines and survival signals, such as transforming growth factor-β and interleukin-6 can induce functional changes in some cell populations that not only increase their drug resistance but also contribute to the preservation of stemness in certain conditions. (40,41) To analyze the correlation between the SASP of the CD133low cell population after gefitinib treatment and the increase in the proportion of the CD133high cell population, we used rapamycin, which effectively suppressed the SASP. (28) Addition of rapamycin to gefitinib treatment effectively suppressed the TIS-induced SASP and simultaneously inhibited the increase in the proportion of the CD133high cell population. These results suggested that the CD133low cell population forms a CSC niche that produces an SASP, and that blocking the SASP with rapamycin is an effective strategy for prevention of gefitinib induced DTP emergence, although we need to examine the adequate amount of SASP before the CD133high cell population started to increase before the upregulation of SASP on day 6 (Figs 1c,4a). In support of this possibility, the combination of BGT226, a novel PI3K/mTOR dual inhibitor and gefitinib showed supra-additive growth inhibitory effects on EGFR-mutated NSCLC cells in vitro and in vivo. (42) Furthermore, mTOR inhibitors temsirolimus and everolimus overcome hepatocyte growth factor-dependent resistance to EGFR-TKIs in EGFR-mutated NSCLC cells. (43) Although recent clinical studies of mTOR inhibitors combined with EGFR-TKIs have failed to show clinical benefit in unselected NSCLC patients, (44–46) mTOR inhibitor and EGFR-TKI combination therapy should be re-evaluated in EGFR-mutated NSCLC patients after acquired resistance to EGFR-TKIs.

To investigate other anticancer drugs that might show promise for overcoming DTPs, and that might act as an alternative to mTOR inhibitors, we examined the sensitivity of gefitinib-induced DTPs to various types of drugs, including conventional chemotherapeutic agents (cisplatin, pemetrexed, and docetaxel) and EGFR-TKI. However, the DTPs showed significant resistance to conventional chemotherapeutic agents and gefitinib (Fig. 5a). The recent IMPRESS trial (47) is the only randomized phase III trial to compare continuation of gefitinib in combination with conventional cytotoxic chemotherapy versus chemotherapy alone in patients with advanced EGFR-mutation-positive NSCLC with resistance to first-line gefitinib. This trial showed that the additional conventional cytotoxic chemotherapy does not contribute to improvement in progression-free survival. This result means that the combination of existing chemotherapeutic agents is not a promising strategy for overcoming gefitinib-induced DTPs.

We therefore considered new strategies to eliminate gefitinib-induced DTPs. Anticancer chemotherapy has recently been reported to induce not only TIS but also senescence-related metabolic reprogramming in cancer cells. (24) Therapy-induced senescence resulted in increased glycolytic activity in cancer cells and enforced the Warburg effect. The enhanced glycolytic activity induces insensitivity to TKIs used for chronic myeloid leukemia. (48) Our analysis showed that TIS cells of the CD133low cell population increased expression

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levels of the glucose transporters Glut1 and Glut3, and the glycolytic enzyme hexokinase 2 (HK2), and enhanced AMPK activity compared to the CD133<sup>high</sup> cell population (Figs 5b,c, S4, S5). The CD133<sup>low</sup> cell population was susceptible to a glucose transporter inhibitor (phloretin) and a glycolytic blocking agent (2DG) <em>in vitro</em> or phloretin <em>in vivo</em>. To our knowledge, there is no report that 2DG has been used <em>in vivo</em> analyses. Further functional <em>in vitro</em> analysis is necessary to link between glucose metabolism and the resistance to TKIs. The CSC-containing CD133<sup>high</sup> cell population was also resistant to conventional chemotherapeutic agents. In order to overcome this treatment limitation, novel therapeutic strategies to eliminate CSCs have been developed.<sup>[49–51]</sup> Withaferin A, a bioactive compound isolated from the plant <em>Withania somnifera</em>, is a potent inhibitor of CSCs.<sup>[29]</sup> To the best of our knowledge, we are the first to show that WFA has a promising antitumor effect on lung cancer cells both <em>in vitro</em> and <em>in vivo</em>, and that WFA is a potent therapeutic agent to overcome gefitinib-
induced DTPs. Unfortunately, we could not culture CD133\(^{\text{low}}\) cell and CD133\(^{\text{high}}\) cell populations separately (data not shown). After isolation of the CD133\(^{\text{high}}\) cell population from gefitinib-induced DTPs, we attempted to culture the CD133\(^{\text{high}}\) cell population alone. However, after a few days of culture, a CD133\(^{\text{low}}\) cell population emerged in the culture dish. Addition of some cytokines in the supernatant failed to maintain the CD133\(^{\text{high}}\) cell population. These results meant that we could not evaluate the drug effects of phloretin, 2DG, and WFA on individual CD133\(^{\text{low}}\) and CD133\(^{\text{high}}\) cell populations.

Based on our results, we conclude that gefitinib-induced DTPs are composed of TIS cells containing a CD133\(^{\text{low}}\) cell population and CSCs containing a CD133\(^{\text{high}}\) cell population. Glucose metabolism-targeting therapeutic agents have effective antitumor activity against the CD133\(^{\text{low}}\) cell population; the CSC-targeting agent, WFA, has effective antitumor activity against the CD133\(^{\text{high}}\) cell population, as summarized in Figure 5(e). Although these glucose-targeting therapeutic agents and WFA are not available in a clinical setting at present, they are potent agents for overcoming gefitinib-induced DTPs and are worth evaluating in prospective clinical trials. The combination of these agents and gefitinib will be a promising therapeutic strategy to overcome EGFR-TKI resistance.

**Acknowledgments**

The authors thank Dr. Keisuke Nishimura for kindly supporting the flow cytometry techniques, Dr. Koji Yamamoto for technical assistance in taking images for immunochemical staining, and Dr. Michiru Nishita for technical assistance in Western blot analyses. This work was supported by research grants from the Japan Society for the Promotion of Science (KAKENHI no. 25860643 to D. Tamura), a Kobe University School of Medicine Alumni Association (Shinryokukai General Incorporated Association) grant (to D. Tamura), and the Medical Research Fund of Hyogo Medical Association (to K. Kunimasa).

**Disclosure Statement**

Y. Nishimura received research grants from Eli Lilly Japan, AstraZeneca, and Novartis Pharma. The other authors have no conflict of interest.

**Abbreviations**

| Abbreviation | Definition |
|--------------|------------|
| 2DG          | 2-deoxy-D-glucose |
| Akt          | protein kinase B |
| AMPK         | adenosine monophosphate-activated protein kinase |
| CSC          | cancer stem-like cell |
| DTP          | drug-tolerant persister |
| EGFR         | epidermal growth factor receptor |
| HDAC         | histone demethylase |
| IGF-1R       | insulin-like growth factor 1 receptor |
| mTOR         | mammalian target of rapamycin |
| NSCLC        | non-small-cell lung cancer |
| PI3K         | phosphatidylinositol 3-kinase |
| qRT-PCR      | quantitative RT-PCR |
| Rb           | retinoblastoma protein |
| SASP         | senescence-associated secretory phenotype |
| TIS          | therapy-induced senescence |
| TKI          | tyrosine kinase inhibitor |
| WFA          | withaferin A |

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Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

Fig S1. Cellular senescence-associated proteins production from parental cells and drug-tolerant persisters. Whole-cell lysates were prepared and analyzed by Western blotting as indicated.

Fig S2. The CD133high cell population showed relatively higher expression of stem cell-related markers compared to the CD133low cell population. mRNA expressions of stem cell-related markers in parental, drug-tolerant persisters (DTP), CD133high, and CD133low cells sorted from PC9 cells 12 days after medium or gefitinib (2 μM) treatment was analyzed using quantitative RT-PCR. Results are the means ± SD of three independent experiments.

Fig S3. Neutralizing antibodies against the senescence-associated secretory phenotype suppressed the induction of the CD133high cell population. PC9 cells were cultured either in medium or in medium containing gefitinib (2 μM) for 12 days. Drug-tolerant persisters were incubated with gefitinib (2 μM), gefitinib plus anti-transforming growth factor-β (TGF-β) antibody (1 mg/mL; R&D Systems, Minneapolis, MN, USA), gefitinib plus anti interleukin (IL)-6 antibody (1 mg/mL; R&D Systems), or gefitinib plus anti-CCL5 antibody (1 mg/mL; R&D Systems) for the last 3 days. The cells were stained with anti-CD133 antibody after 3 days of culture and were analyzed using flow cytometry. The results are the means ± SD of at least three independent experiments. The CD133high cell population showed relatively higher expression of stem cell-related markers compared to the CD133low cell population. The mRNA expressions of stem cell-related markers in parental, drug-tolerant persisters, and CD133high and CD133low cells sorted from PC9 cells 12 days after medium or gefitinib (2 μM) treatment was analyzed using quantitative RT-PCR. Results are the means ± SD of three independent experiments.

Fig S4. Adenosine monophosphate-activated protein kinase (AMPK) activity was enhanced in drug-tolerant persisters compared to parental cells. Whole-cell lysates were prepared and analyzed by Western blotting as indicated.

Fig S5. Drug-tolerant persisters showed relatively higher expression of glucose transporters (GLUT1 and GLUT3) and the glycolytic enzyme hexokinase2 (HK2) compared to the parental cells. The mRNA expressions of GLUT1, GLUT3, and HK2 in parental cells and drug-tolerant persisters 12 days after medium or gefitinib (2 μM) treatment was analyzed using quantitative RT-PCR. Results are the means ± SD of three independent experiments.

Table S1. Primer sequences used for RT-PCR.