Inhibition of Mitochondrial Complex II by the Anticancer Agent Lonidamine*

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The antitumor agent lonidamine (LND); 1-(2,4-dichlorobenzyl)-1H-indazole-3-carboxylic acid is known to interfere with energy-yielding processes in cancer cells. However, the effect of LND on central energy metabolism has never been fully characterized. In this study, we report that a significant amount of succinate is accumulated in LND-treated cells. LND inhibits the formation of fumarate and malate and suppresses succinate-induced respiration of isolated mitochondria. Utilizing biochemical assays, we determined that LND inhibits the succinate-ubiquinone reductase activity of respiratory complex II without fully blocking succinate dehydrogenase activity. LND also induces cellular reactive oxygen species through complex II, which reduced the viability of the DB-1 melanoma cell line. The ability of LND to promote cell death was potentiated by its suppression of the pentose phosphate pathway, which resulted in inhibition of NADPH and glutathione generation. Using stable isotope tracers in combination with isotopologue analysis, we showed that LND increased glutaminolysis but decreased reductive carboxylation of glutamine-derived α-ketoglutarate. Our findings on the previously uncharacterized effects of LND may provide potential combinational therapeutic approaches for targeting cancer metabolism.

Targeting cancer metabolism has received renewed interest as a strategy for anticancer therapy (1–3). Most cancers have altered metabolism and a high demand for nutrients to support their rapid proliferation (4–7). Warburg et al. (8) first reported that malignant cells metabolize glucose via aerobic glycolysis, which is often associated with increased glucose uptake and lactate production. Many other metabolic alterations have been discovered in cancer cells, including a high rate of glutamine consumption, enhanced lipid synthesis, and increased pentose phosphate pathway (PPP) flux (1, 7). Approaches targeting cancer metabolism such as inhibition of glycolysis and restriction of glutamine availability have shown promising results in preclinical and clinical studies (1, 7).

In addition to direct targeting of metabolic enzymes or transporters to restrict nutrient availability, modulation of mitochondrial metabolism that occurs through the tricarboxylic acid (TCA) cycle and electron transport chain (ETC) has been proposed as a novel approach for anticancer drug development (3, 9). The catabolic processing of sugars, glutamine, fatty acids, and amino acids leads to the generation of reducing equivalents in the form of NADH and FADH₂, which are subsequently oxidized. The electrons transferred during this process are shuttled down the ETC, a process that generates an electrochemical gradient across the inner mitochondrial membrane necessary for ATP production. In addition to the crucial role of energy production, the TCA cycle also provides intermediates for lipid and amino acid synthesis. The increased demand for energy production and anabolic building blocks observed in cancer cells makes selectively targeting the TCA cycle and ETC promising approaches to limiting malignancy and proliferation. It is noteworthy that the type II diabetes drug metformin has been reported to inhibit mitochondrial complex I activity and provide beneficial effects on a number of cancer models (10–12). A recent study indicates that metformin blocks gluconeogenesis resulting from the inhibition of mitochondrial glycerol-3-phosphate dehydrogenase, another contributor of electrons to the ETC (13). This newly discovered activity of metformin could also be responsible in part for the reduction in risk of cancer and diminished cancer-related mortality in patients using the drug (12).

Lonidamine (LND); 1-(2,4-dichlorobenzyl)-1H-indazole-3-carboxylic acid has been used in combination with other therapeutic agents to improve efficacy and overall response to can-
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Chemicals and Reagents—Rotenone, 4,4,4-trifluoro-1-(2-thienyl)-1,3-butanedione (TTFA), 3-nitropropionic acid (3-NPA), malonate, 2,6-dichlorophenolindophenol, decylubiquinone, 2-(4,5-dimethyl-2-thiazolyl)-3,5-diphenyl-2H-tetrazolium (MTT), phenazine methosulfate (PMS), carbonyl cyanide 4( trifluoromethoxy)phenylhydrazone (FCCP), roxolium bromide (MTT), phenazine methosulfate (PMS), car

Experimental Procedures

Cell Culture and Treatment—DB-1 cells were human melanoma cells derived from a lymph node metastasis as described previously (29). DB-1 and HepG2 were maintained in minimum Eagle’s medium α and RPMI 1640 medium, respectively. HeLa and HCT116 were cultured in DMEM. All the media were supplemented with 10% fetal bovine serum, 100 units/ml penicillin, and 100 mg/liter streptomycin. LND, TTFA, and 3-NPA were freshly prepared in DMSO. DB-1 cells were grown to 80% confluence and treated for 1 h with vehicle (0.1% DMSO) or the indicated treatment in glutamine-free DMEM containing 5 mM glucose. HePG2, HeLa, and HCT116 were treated with DMSO or LND (150 μM) for 1 h in their respective media. Cell volumes were determined using a Multisizer 3 Coulter Counter (Beckman Coulter).

Mitochondria Labeling by [13C4]Succinate—Mouse liver mitochondria were freshly prepared from adult mice as described previously (30). 60 μg of isolated mitochondria were suspended in 200 μl of reaction buffer (135 mM sucrose, 65 mM KCl, 5 mM KH2PO4, 10 mM Tris/HCL, 20 μM ETGA, 2.5 mM MgCl2, pH 7.4) containing 5 mM [13C4]succinate. Mitochondria samples were pulse-sonicated for 5 s before incubation at 37 °C for 30 min.

Organic Acid Extraction and Derivatization—Cells were washed twice with phosphate-buffered saline (PBS) followed by scraping into 750 μl of ice-cold methanol/water (4:1, v/v) containing 500 ng of internal standard ([13C3]lactate, [13C4]glucose, [13C6]citrate, and [13C4]fumarate). Samples were pulse-sonicated for 30 s and centrifuged at 16,000 × g for 10 min. The supernatant was transferred to a clean tube. For α-ketoglutarate derivatization, methoxyamine HCl (2 mg) was added, and samples were incubated at 37 °C for 1 h. Following incubation, samples were evaporated to dryness under nitrogen and suspended in 100 μl of mobile phase A (400 mM 1,1,1,3,3,3-hexafluoro-2-propanol and 10 mM Dipea in water) prior to LC-MS analysis. For NADPH and NADP+ analysis, the supernatant from methanol/water extracts was diluted 1:4 using 50 mM ammonium carbonate for LC-MS analysis.

For labeled mitochondria, 800 μl of ice-cold methanol containing 500 ng of [13C4]glutamine was added to quench the reaction. For quantifying glutamine in the culturing medium, 5 μl of medium were added to 500 μl of ice-cold methanol/water (4:1, v/v) containing 1.5 μg of [13C5,15N2]glutamine. Samples were pulse-sonicated for 30 s and centrifuged at 16,000 × g for 10 min. The supernatant was transferred to a clean tube and evaporated to dryness under nitrogen. The dried residues were derivatized with 100 μl of DIPA in acetonitrile (0.5:99.5, v/v) and 100 μl of pentafluoro benzyol chloride in acetonitrile (1:4, v/v) at 60 °C for 1 h. Derivatized samples were evaporated to dryness under nitrogen and reconstituted in hexanes/ethanol (95:5, v/v) prior to LC-electron capture atmospheric pressure chemical ionization-MS analysis (31).

LC-Selected Reaction Monitoring/MS Analysis—Organic acids from cell samples were analyzed using an Agilent 1200 series HPLC system coupled to an Agilent 6460 triple quadrupole mass spectrometer equipped with an electrospray ionization source operated in negative ion mode. Analytes were separated by reversed-phase ion-pairing chromatography utilizing a Phenomenex Luna C18 column (250 × 2.00 mm, 3-μm particle size) at a flow rate of 200 μl/min maintained at 45 °C. A two-solvent gradient system was used with solvent A as 400 mM 1,1,1,3,3,3-hexafluoro-2-propanol and 10 mM DIPA in water and solvent B as 300 mM 1,1,1,3,3,3-hexafluoro-2-propanol and 10 mM DIPA in methanol. The linear gradient conditions were as follows: 2% B at 0 min, 2% B at 3 min, 10% B at 28 min, 95% B at 31 min, 95% B at 38 min, 2% B at 39 min followed by a 6-min equilibration. The Agilent 6460 mass spectrometer operating conditions were as follows. The gas temperature was set at 320 °C, and the gas flow was set to 8 liters/min. The sheath gas temperature was 400 °C, and the sheath gas flow was set to 10 liters/min. The capillary voltage was set to 3000 V, and the
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nozzle voltage was set to 1000 V. The isotopic distribution of TCA cycle metabolites was calculated as described previously (32). The analysis of NADPH and NADP⁺ was similar except that an LC method with a 15-min run time was performed on a 150 × 2-mm Phenomenex Luna C₁₈ column. This ensured that all the samples were analyzed within 3 h after the extraction and minimized decomposition of the NADPH and NADP⁺.

[¹³C₄]Fumarate and [¹³C₅,¹⁵N₂]malate generated in mitochondria and glutamine from the medium were converted to pentafluorobenzyl bromide derivatives and analyzed as described previously (33). Briefly, pentafluorobenzyl bromide derivatives were separated using a CHIRALPAK AD-H column (250 × 4.6 mm, 5-µm particle size; Daicel Chemical Industries, Ltd., Tokyo, Japan) at a flow rate of 1 ml/min maintained at 30 °C. A post-column addition (0.75 ml/min) of methanol was used. Solvent A was hexanes, and solvent B was isopropanol/methanol (1:1, v/v). The linear gradient was as follows: 1% B for 3 min, 60% B at 25 min, 60% B at 29 min, 1% at 30 min followed by a 5-min equilibration. MS analysis was performed on a Thermo Quantum Triple Stage Quadrupole mass spectrometer (Thermo Scientific) with an atmospheric pressure chemical ionization source operated in the negative ion mode. The operating conditions were as follows: vaporizer temperature, 350 °C; heated capillary temperature, 300 °C; corona discharge needle, 30 µA. The sheath gas and auxiliary gas pressures were 35 and 10 (arbitrary units), respectively. Cellular glutathione (GSH) levels were quantified as described previously (34).

Mitochondrial Oxygen Consumption Assay—For measurements of oxygen consumption via complex II, 80 µg of isolated mouse liver mitochondria were added to respiration buffer (135 mM sucrose, 65 mM KCl, 5 mM KH₂PO₄, 10 mM Tris/HCl, 20 µM EGTA, 2.5 mM MgCl₂, pH 7.4) containing 5 mM succinate and LND at the indicated concentrations. ADP (100 µM) and FCCP (60 nM) were added sequentially. The measurements were carried out using a Clark electrode (model 949, Strathkelvin Instruments) in a water-jacketed glass reaction vessel (Mitocell MT200) containing 100 µl of stirred, air-saturated respiration buffer at 37 °C.

Enzyme Assays—Succinate-ubiquinone reductase (SQR) activity and succinate dehydrogenase (SDH) activity were measured as described (35, 36) with minor modifications. Briefly, mouse liver mitochondria were resuspended in assay buffer (0.3 M mannitol, 25 mM KH₂PO₄, pH 7.4) at 20 µg/ml and supplemented with 20 mM succinate, 5 µM rotenone, 2 µM antimycin A, and 10 mM NaN₃. Following a 10-min incubation at room temperature, inhibitors were added and incubated for an additional 15 min before initiation of reactions. For SQR activity, reactions were initiated by adding 50 µM decylubiquinone and 50 µM 2,6-dichlorophenolindophenol (extinction coefficient = 21 mM⁻¹ cm⁻¹). Absorbance at 600 nm was monitored every min for 20 min. For the SDH activity of complex II, reactions were initiated by adding 150 µM MTT and 400 µM PMS. The change in absorbance of MTT was monitored at 570 nm for 20 min.

Intracellular Reactive Oxygen Species (ROS) Measurement—Cells were treated with vehicle or drugs for 4 h followed by staining with 2',7'-dichlorofluorescin diacetate (10 µM) in minimum Eagle's medium α for 20 min at 37 °C. Cells were then trypsinized and rinsed twice with PBS. The cellular fluorescence of the oxidized product, 2',7'-dichlorofluorescein, was analyzed by flow cytometry (Accuri C6, BD Biosciences). At least 10,000 data events were collected for each sample.

Propidium Iodide Staining for Cell Viability—After treatment with vehicle or drugs for 24–48 h, cells were trypsinized and rinsed twice with PBS. Cells resuspended in PBS were mixed with PI (final concentration of 1 µg/ml) right before analysis by flow cytometry (Accuri C6). At least 10,000 data events were collected for each sample.

Dynamic Isotopic Labeling of Cells—DB-1 cells were treated with DMSO, LND, or TTFA for 2 h before culturing in glutamine-free DMEM containing 2 mM [¹³C₆]glutamine supplemented with 10% fetal bovine serum and drugs. After 0, 0.5, 1, 2, 4, and 6 h of incubation, medium was aspirated, and 750 µl of ice-cold methanol/water (4:1, v/v) were added to cell culture plates snap frozen on dry ice. The glucose labeling experiment was performed similarly except that cells were labeled with glucose-free DMEM containing [¹³C₆]glucose, 10% fetal bovine serum, and relevant drugs. Cells were harvested after 0, 1, 2, 5, 10, 30, and 60 min of incubation.

Metabolic Flux Analysis Using ¹³C Metabolic Fragmented Mass Isotopologue Distribution Modeling—A two-compartment (extracellular medium, intracellular cellular distribution) metabolic model was used to fit experimental ¹³C dynamic mass isotopologues of labeled citrate and malate to determine metabolic fluxes. The metabolic network included glycolysis, the TCA cycle, extracellular glutamine uptake, cell glutamate production via cytosolic and mitochondrial glutaminase, reductive carboxylation of α-ketoglutarate, anaplerosis through pyruvate carboxylase, glutaminolysis and malic enzyme activity.

The model was expressed mathematically using two types of mass balance equations: 1) mass balance for total metabolite concentration and 2) ¹³C mass isotopologue mass balance for labeled metabolites and their related fragments based on bio-network and atom distribution matrices (fragmented mass isotopologue framework) (37). Mass isotopologue dynamics of the system were formulated as the Cauchy initial value problem for ordinary differential equations using mass isotopologue fractions as the state variables. Mass isotopologue balance equations were derived in a similar manner as equations for bonded cumulative isotopologues as described previously (38). In terms of ordinary differential equations, this model describes the rates of loss and creation of particular labeled and unlabeled metabolite forms (mass isotopologues) after incubations with labeled glutamine in extracellular media. For the [¹³C₆,¹⁵N₂]glutamine experiment, the fitted experimental dynamic mass isotopologues were seven mass isotopologue forms of citrate (unlabeled, M + 1 to M + 6 mass isotopologues) and five forms of malate with a total of 12 dynamic mass isotopologue curves. The extracted absolute fluxes for glutaminolysis and reductive carboxylation were determined. For all metabolite mass isotopologues, the ¹³C natural abundance of ¹³C isotope (1.078%) was taken into account.

Solving a system of non-linear differential equations in terms of whole/fragmented mass isotopologues with the Runge-Kutta fourth order procedure for stiff systems provided time courses
for all possible $^{13}$C mass isotopologues (e.g. citrate and glutamate). Dynamic mass isotopologue values at the experimental time points were taken into account for fitting procedures. The cost function was used to quantify differences between measurements and computational results for labeled dynamic data and to select the corresponding vector of fluxes that minimizes the cost function in $[^{13}$C$_5^{15}$N$_2]$ glutamine experiments. Minimization was performed with simplex or Broyden-Fletcher-Goldfarb-Shanno algorithms. Correct mean square convergence was confirmed by verifying that goodness of fit values were close to expected theoretical values. To overcome potential local minima, several sets of initial random fluxes were used (4).

Reliability of the flux values was evaluated by using Monte-Carlo simulations as described previously (39). The Monte-Carlo simulation procedure was performed by generation of synthetic dynamic mass isotopologues for citrate and malate by solving differential equations. Metabolic fluxes, which were representative for metabolic fluxes in DB-1 melanoma cells, were used as nominal values to generate these synthetic dynamic mass isotopologues. For each Monte-Carlo draw (at least 500 draws total), random Gaussian noise with a mean of zero and standard deviation $\sigma$ of 0.01 was added to dynamic $^{13}$C mass isotopologues. The chosen noise level was slightly above a typical noise level for $in vitro$ $^{13}$C LC-MS cell studies. These synthetic renormalized dynamic mass isotopologues were then fitted using a $^{13}$C metabolic model to obtain best fit values. Different starting values were chosen randomly to ensure that the results were independent of the initial condition of the fit. Resulting parametric probability density functions directly reflect the uncertainty of each fitted parameter. Distributions were characterized by their standard deviations. Other statistics not reported include probability density functions, confidence intervals, and cross-correlation between metabolic fluxes/parameters. All numerical procedures were carried out in Matlab (Mathworks, Natick, MA) as described previously (4).

Results

**LND Treatment Results in the Accumulation of Succinate—**To examine the effect of LND on melanoma cell metabolism, we used LC-selected reaction monitoring/MS to analyze the metabolites in DB-1 melanoma cells treated with LND. As a monocarboxylate transporter inhibitor, LND is known to stimulate lactate accumulation in cells (25, 26). After 1 h of treatment with 150 $\mu$m LND, lactate was elevated nearly 5-fold over control cells (Fig. 1A). The levels of succinate and $a$-ketoglutarate in LND-treated DB-1 cells were found to increase, whereas the levels of citrate, fumarate, and malate decreased (Fig. 1, B and C). A 3–5-fold accumulation of succinate was also observed in a variety of cancer cell lines, including HepG2 (Fig. 1D), HCT116 (Fig. 1E), and HeLa (Fig. 1F).

The accumulation of succinate and decrease in fumarate and malate in DB-1 cells suggested that the conversion of succinate to fumarate was inhibited by LND (Fig. 1C). The oxidation of succinate to fumarate is catalyzed by the succinate dehydrogenase activity of complex II. Therefore, we investigated the effect of LND on succinate accumulation in DB-1 cells at various concentrations and compared its potency with two known complex II inhibitors, 3-NPA and TTFA. At 50 $\mu$m, both LND and 3-NPA induced a 2-fold increase in succinate levels, whereas TTFA treatment resulted in a larger increase (Fig. 1G). Neither 3-NPA nor TTFA affected lactate levels (Fig. 1H). LND failed to induce further succinate or lactate accumulation in DB-1 cells beyond a dose of 150 $\mu$m (Fig. 1, G and H). This observation indicates that cells may not be able to effectively take up LND from the cell culture medium at concentrations beyond 150 $\mu$m.

**LND Inhibits the Formation of Fumarate and Malate in Isolated Mitochondria—**To rule out the possibility that the inhibition of the succinate dehydrogenase activity of complex II by LND was an indirect effect resulting from metabolic alterations in the cytosol, we examined the conversion of succinate to fumarate using isolated mitochondria. Importantly, the utilization of isolated mitochondria reduces interference of cytoplasmic metabolites and enzymes. To interrogate the specific action of complex II, isolated mitochondria were incubated with 5 $\mu$m $[^{13}$C$_4$] fumarate, $[^{13}$C$_4$] malate, and $[^{13}$C$_4$] citrate was quantified to assess succinate metabolism. Within 30 min of incubation, 106 nmol of $[^{13}$C$_4$] fumarate and 232 nmol of $[^{13}$C$_4$] malate were generated from 60 $\mu$g of mitochondria with vehicle treatment. No heavy labeled citrate was detected, most likely due to the lack of external acetyl-CoA supply. The addition of LND inhibited the production of $[^{13}$C$_4$] fumarate and $[^{13}$C$_4$] malate (Fig. 2, A and B). 150 $\mu$m LND inhibited ~40% of fumarate and 50% of malate production. The extent of inhibition caused by LND was similar to that of 3-NPA at various concentrations, whereas TTFA was a more potent inhibitor of complex II (Fig. 2, A and B). Overall, our observations suggest that the inhibition of complex II activity by LND is independent of lactate accumulation or other metabolic changes in the cytosol.

**LND Inhibits Complex II-dependent Respiration—**Floridi and Lehninger (28) have previously explored the ability of LND to inhibit cellular respiration. To examine the effect of LND on complex II-dependent respiration, we included the complex I inhibitor rotenone in the respiration assay utilizing succinate as the metabolic substrate. We measured the mitochondrial oxygen consumption rate at the basal level, in the ADP-stimulated condition, and in the presence of the mitochondrial uncoupler FCCP (Fig. 2C). LND inhibited mitochondrial oxygen consumption rates under all three conditions. Similar to that observed in the previous study (28), the inhibition was much stronger under ADP- and FCCP-stimulated conditions than at basal levels (Fig. 2, C and D). LND did not inhibit the basal respiration up to 100 $\mu$m. Under all conditions, half-maximal inhibition by LND was observed between 100 and 200 $\mu$m.

**Mechanism of Complex II Inhibition by LND—**Complex II contains four subunits: A (flavoprotein), B (iron-sulfur subunit), C (15-kDa integral membrane protein), and D (cytochrome b small subunit) encoded by the SDHA, SDHB, SDHC, and SDHD genes, respectively. Electrons obtained from the oxidation of succinate within SDHA are transferred from the SDHA-bound flavin adenine dinucleotide (FAD) cofactor to the Fe-S clusters of SDHB and finally to the ubiquinone reduction site residing between SDHC and SDHD where ubiquinone is reduced to ubiquinol (Fig. 3A). To determine the specificity of complex II inhibition by LND, we measured the formation of ubiquinol by complex II (SQR activity) using a water-soluble
ubiquinone analogue decylubiquinone in combination with the artificial electron acceptor 2,6-dichlorophenolindophenol. In the SQR activity assay, isolated mouse liver mitochondria were incubated with succinate and decylubiquinone together with rotenone and the complex III inhibitor antimycin A. Our results showed that LND inhibited the formation of ubiquinol by complex II in a dose-dependent manner (Fig. 3B). At 500 μM, around 90% of the SQR activity was inhibited by LND.

**FIGURE 1. LND treatment alters the levels of TCA cycle metabolites.** A and B, DB-1 cells were incubated with 150 μM LND or DMSO for 1 h. Levels of lactate (A) and metabolites in the TCA cycle (B) were measured by LC-MS. The levels of metabolites in LND-treated group were normalized with respect to the relevant metabolites in DMSO controls. C, a scheme of the TCA cycle. The changes observed in LND-treated cells are indicated. HepG2 (D), HCT116 (E), and HeLa (F) cells were incubated with DMSO or LND (150 μM) for 1 h. Levels of lactate and TCA metabolites were quantified by LC-MS. For A, B, and D–H, the means of three samples are shown. Error bars represent S.D. *, p < 0.05; **, p < 0.01; ***, p < 0.001 (Student’s t test).
To further investigate the mechanism of complex II inhibition by LND, we also examined the SDH activity of complex II. SDH activity is the first step of the SQR reaction and can be determined by two artificial electron acceptors, PMS and MTT. PMS and MTT create a bypass around ubiquinone in SDHC and SDHD and directly accept electrons from iron-sulfur clusters in subunit B. The SDH activity was determined by monitoring the color change of MTT (Fig. 3A).

To determine the specificity of the SQR and SDH activity assays, we compared the inhibition by TTFA and malonate in isolated mitochondria. TTFA inhibits complex II activity by primarily binding to the ubiquinone-binding site, whereas malonate competes with succinate for the dehydrogenase-binding site on complex II. TTFA (50 μM) potently inhibited SQR activity (89%) but exhibited only modest inhibition of MTT reduction (36%) as shown in Fig. 3B. In contrast, malonate (5 mM) inhibited SQR by 84% and MTT reduction by 65%.

Increasing doses of LND were used to compare its effect on the SQR and SDH activities of complex II in isolated mitochondria. The inhibition of SQR activity was found to be much greater than inhibition of SDH activity at all LND concentrations tested. LND at 1 mM almost completely inhibited the activity of SQR, whereas the SDH activity was only reduced by 34% (Fig. 3B). This finding indicates that LND failed to fully
block the electron transfer from the iron-sulfur cluster to PMS and MTT. Thus, LND appears to inhibit complex II activity by interfering with the ubiquinone reduction, possibly at the ubiquinone-binding site in SDHC and SDHD.

LND Induces ROS Formation and Cell Death—As part of the ETC, complex II is also a source of ROS. It mainly produces ROS from either the reduced FAD or ubiquinone site when downstream components of the ETC are blocked (40, 41). To examine whether LND induces intracellular ROS generation, we quantified the level of ROS by $2\text{-H}11032\text{-dichlorofluorescein fluorescence (Fig. 4, A–C). Surprisingly, the level of ROS formed in LND-treated cells was even higher than the level in cells treated with TTFA (Fig. 4, A and C).}

To determine whether LND induces ROS through complex II inhibition, we examined ROS generation from a combined treatment with LND and TTFA or LND and 3-NPA. 3-NPA binds to the active site of SDHA and inhibits the electron transfer from succinate to FAD. Thus, 3-NPA alone does not trigger substantial ROS production and can reduce ROS induced by TTFA, which inhibits complex II at the ubiquinone-binding site (34, 36) (Fig. 4C). LND and TTFA treatments showed an additive effect of ROS production. In contrast, the addition of 3-NPA markedly decreased the level of ROS generated in LND-treated cells (Fig. 4, A–C). This result suggests that LND induces ROS at a site within complex II downstream of SDHA. It also supports our finding that LND inhibits complex II by interfering with ubiquinone reduction.

LND also induced substantial cell death after treatment of DB-1 cells for 48 (Fig. 4D) or 24 h (Fig. 4E). TTFA alone was unable to induce large amounts of cell death. However, it significantly promoted cell death triggered by LND (Fig. 4, D and F). The increased cell death caused by the addition of TTFA to LND-treated cells indicates that LND renders cells vulnerable to further complex II inhibition or additional ROS. Incubation with the ROS scavenger N-acetylcysteine reduced the rates of cell death in cells treated with LND alone or with TTFA (Fig. 4, C and E). Thus, ROS is responsible, at least in part, for cell death induction under these conditions.

LND Reduces Cellular Levels of Glutathione and NADPH and the PPP—To investigate why the LND-treated cells were more susceptible to complex II inhibition than TTFA-treated cells, levels of the cellular antioxidant GSH were compared. LND caused a 40% drop in GSH levels at a concentration of 150 μM and above (Fig. 5A). The extent of GSH reduction is similar to that caused by 100 μM diethyl maleate (DEM), a compound known to deplete GSH (Fig. 5A). In contrast, TTFA (50 or 200

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**FIGURE 3. LND inhibits the SQR activity of complex II.** A, a schematic representation of complex II subunits and enzyme activities. SQR activity was determined by measuring the electron transfer from succinate to decylubiquinone and 2,6-dichlorophenolindophenol (DCPIP). SDH of complex II was measured by the electron transfer from succinate to the iron-sulfur cluster and finally to the water-soluble dyes PMS and MTT. Q, ubiquinone; QH2, reduced ubiquinone. B, isolated mouse liver mitochondria were incubated with LND, TTFA, and malonate at the indicated concentrations. SQR and SDH activities were measured as described in A. The percentage of inhibition was determined by comparing the enzyme activities of the drug-treated groups with DMSO-treated controls. The data presented are means of three samples. Error bars represent S.D. The SQR and SDH activities of complex II in all treatment groups were determined on the same batch of freshly prepared mitochondria.
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Figure 4. LND induces cellular ROS release and reduces cell viability. A and B, ROS in DB-1 cells was measured by 2′,7′-dichlorofluorescein (DCF) fluorescence. Shown are the representative flow cytometry histograms of samples treated with DMSO or the indicated drugs. C, mean fluorescence intensities of cells treated with DMSO or the indicated drugs. D and E, PI exclusion method was used to examine the viability of DB-1 cells treated with DMSO or indicated drugs. Shown are the percentages of PI-positive cells. The data presented are means of three samples. Error bars represent S.D. *, p < 0.05; **, p < 0.01; ***, p < 0.001 (Student’s t test). F, representative flow cytometry histograms for PI-stained cells from D. Percentages of PI-positive cells are indicated. Cells were treated with the indicated drugs for 48 (D and F) or 24 h (E). The following drug concentrations were used: LND, 150 μM; TTFA, 150 μM unless indicated otherwise; 3-NPA, 500 μM; N-acetylcysteine (NAC), 10 mM.

μM) caused a modest reduction of 6% (Fig. 5A). Consistent with the reduced GSH levels, the levels of NADPH and the NADPH/NADP⁺ ratio both decreased after LND treatment but not after TTFA treatment (Fig. 5, C and D). The PPP is an important source of NADPH (5) required for the GSH reductase-mediated reduction of glutathione disulfide back to GSH (42). LND has been reported previously to be an inhibitor of hexokinase (Fig. 5B) (23, 43), which catalyzes the first step of glycolysis. In contrast, some studies have shown no effect on hexokinase based on the analysis of glucose 6-phosphate after LND treatment of breast cancer and glioma cell lines (25, 26, 43). However, these latter studies did not examine the effect of LND on
specific PPP metabolites, which could also be down-regulated as a result of hexokinase inhibition (Fig. 5B). In support of this possibility, we found that the concentration of 6-phosphogluconate, an important PPP metabolite, was markedly decreased in LND-treated DB-1 cells (Fig. 5E). In addition, a time course for the incorporation of [13C6]glucose into PPP metabolites (determined by LC-MS) revealed that incorporation into the M + 6 isotopologue [13C6]6-phosphogluconate as well as the glycolysis metabolite [13C6]fructose 1,6-bisphosphate was significantly delayed (Fig. 5, F and G). These data, together with the reduced levels of 6-phosphogluconate, suggest that the flux into the PPP was greatly reduced by LND, most likely through inhibition of hexokinase. Thus, the reduced NADPH levels (Fig. 5C) and NADPH/NADP+ ratios (Fig. 5D) and decreased GSH levels (Fig. 5A) in LND-treated cells resulted, in part, from inhibition of the PPP (Fig. 5B).

**GSH Depletion Sensitizes Melanoma Cells to LND or TTFA treatment**—The reduced PPP and GSH levels caused by LND may render cells more susceptible to damage from ROS. To test this hypothesis, we incubated DB-1 cells with LND or TTFA in the presence or absence of the GSH depletion agent DEM. TTFA induced less ROS production (Fig. 6A) and less cell death.
can be metabolized reductively to citrate to provide carbon atoms for lipid synthesis (Fig. 7A). We performed a dynamic labeling assay to analyze both oxidative and reductive pathways of glutamine metabolism. DB-1 cells were first treated with vehicle, LND, or TTFA and then cultured in medium containing \([^{13}\text{C}_5,^{15}\text{N}_2]\)glutamine. Incorporation of \([^{13}\text{C}]\) carbons into TCA cycle metabolites at different time points was then determined. Over the 6 h of labeling, glutamine contributed to succinate, fumarate, malate, and citrate through oxidative metabolism as shown by their M + 4 isotopologues (Figs. 7 and 8). Most metabolites reached steady state labeling beyond 4 h in control cells (Fig. 8). Treatment of DB-1 cells with LND strikingly increased the labeling in succinate (Fig. 8A), fumarate (Fig. 8B), malate (Fig. 8C), and citrate (Fig. 8D) over the time course of the experiment as well as at steady state (Fig. 7, C–F). Similar effects were also observed in TTFA-treated cells (Figs. 7, C–F, and 8, A–D).

To determine whether LND and TTFA altered flux from glutamine into TCA cycle metabolites, we integrated the kinetic labeling patterns with cellular concentrations of metabolites in a two-compartment metabolic flux model (Table 1 and Fig. 9). The analysis showed that glutaminolysis flux was significantly higher in LND- and TTFA-treated cells (Fig. 7G). Consistently, the M + 6 isotopologue of glutamate was also increased upon drug treatment (Fig. 7H), although its level remained unchanged (Table 1). The enhanced glutamine utilization in LND- and TTFA-treated cells is underscored by their increased glutamine uptake during a 12-h incubation (Fig. 7I). Although the increase in overall glutamine uptake is less than 10% for LND, this percentage was calculated based on the number of cells at the beginning of treatment. Given the arrested cell growth and cell death induced by LND, the actual glutamine uptake may be even higher.

Glutamine-dependent Reductive Carboxylation Is Reduced by LND—DB-1 cells also use glutamine through reductive metabolism (Fig. 7A) as demonstrated by the formation of malate and fumarate M + 3 isotopologues and the citrate M + 5 isotopologue (Fig. 10, A–F). LND, but not TTFA, reduced the labeling of these metabolites resulting from reductive glutamine metabolism over the dynamic labeling or at the steady state (Fig. 10, A–F). LND lowered the flux from reductive carboxylation into citrate by nearly 50%. In contrast, the reductive flux was unchanged in TTFA-treated cells (Fig. 10G).

Discussion

LND is known to interfere with energy metabolism in cancer cells. To improve its efficacy and potentially reduce organ toxicity, recent studies have focused on developing targeted delivery systems for LND (19–22). For example, EGF receptor-targeting nanoparticles have significantly improved the tumor-suppressive effect of LND in combination with the chemotherapeutic agent paclitaxel (20). Utilizing delivery with the targeting nanocarriers, 25 nmol of LND/g of tissue was reached in tumor xenografts (19). It is important to note that the local cellular concentration of LND may be even higher, indicating that the doses used in this study may be physiologically relevant in these settings. Another nanoparticle delivery system that targets LND to mitochondria showed an increase of more than 100-fold in the efficacy of reducing can-
cer cell viability compared with non-targeted LND (21), further highlighting the importance of elucidating unknown mechanisms by which LND exerts antitumor effects.

The effect of LND on central energy metabolism has never been fully characterized. Floridi et al. (23, 28) reported that LND inhibits the respiration in both Ehrlich ascites cells and isolated mitochondria. However, the sites of inhibition were not clearly identified. Through LC-MS analysis of TCA cycle metabolites, we found that LND inhibited the oxidation of succinate to fumarate in various cell lines and in isolated mitochondria (Figs. 1 and 2). Furthermore, we showed that in the presence of artificial electron acceptors LND inhibited ubiquinone reduction, but it did not fully block the transfer of electrons from iron-sulfur clusters in SDHB (Fig. 3). These results indicate that SDH activity is not the target of LND. This notion is further supported by assays for ROS generation upon LND treatment (Fig. 4, A–C). TTFA and 3-NPA had additive and inhibitory effects on LND-triggered ROS, respectively, suggesting that free electrons are released from a location between the SDH active site and ubiquinone-binding site. Because LND effectively inhibits the SQR activity of complex II, we propose that LND inhibits complex II by interfering with the reduction of ubiquinone. However, it is unclear whether the inhibition is caused by direct blockade of the ubiquinone-binding site or through allosteric modifications to the tertiary structure of SDHC and SDHD.

**FIGURE 7. LND and TTFA reduce oxidative glutamine metabolism in DB-1 cells.** A, a scheme showing the $^{13}$C labeling of metabolites by $[^{13}$C$_5$,$^{15}$N$_2$]glutamine in the oxidative and reductive glutamine metabolism. B–H, DB-1 cells were treated with DMSO, LND (300 μM), or TTFA (50 μM) for 2 h. Levels of succinate (B) and other metabolites (Table 1) were quantified in one set of samples, whereas the rest of the cells were further incubated with 2 mM $[^{13}$C$_5$,$^{15}$N$_2$]glutamine and the indicated drugs. After 6 h of incubation, the $^{13}$C labeling from the oxidative glutamine metabolism is shown by the M + 4 isotopic enrichment of succinate (C), fumarate (D), malate (E), and citrate (F). Rates of glutaminolysis as well as the standard deviation and p values were calculated as described under “Experimental Procedures” (G). The glutamate labeling is shown by M + 6 isotopic enrichment (H). I, glutamine uptake was measured by incubating cells with DMSO, LND (300 μM), or TTFA (50 μM) for 12 h. For all the panels, the means of three samples are shown. Error bars represent S.D. *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$. 
Incubation of the DB-1 cells with the classical complex II inhibitor TTFA triggered less ROS release compared with that caused by LND treatment (Fig. 4, A and C). There was a concomitant increase in the amount of cell death in the LND-treated cells when compared with the TTFA-treated cells (Fig. 4, D and F). The increase in ROS production and cell death was amplified by DEM for both TTFA (Fig. 6, A and C) and LND treatments (Fig. 6, B and D). Importantly, TTFA induced a significant amount of additional cell death when combined with LND treatment (Fig. 4, D and F). These results suggest that the compromised antioxidant system in the LND-treated cells resulting from PPP inhibition (reduced NADPH and GSH) contributes to the susceptibility to cell death in response to complex II inhibition and ROS release.

Inhibition of TCA cycle enzymes can have consequences for tumor progression depending upon tumor types and the stage of tumorigenesis. For example, fumarate hydratase deficiency...

**FIGURE 8.** Dynamic labeling through oxidative glutamine metabolism in LND- and TTFA-treated cells. Dynamic labeling of DMSO-, LND- (150 μM), and TTFA (50 μM)-treated cells after switching to [13C5,15N2]glutamine. Data were collected from the same dynamic labeling experiment as described in Fig. 7. Labeling percentages of M + 4 succinate (A), fumarate (B), malate (C), and citrate (D) were plotted over time. The data presented are the means of three samples. Error bars represent S.D.

**TABLE 1**

Concentrations of metabolites quantified after 2 h of drug treatment (means ± S.D.; n = 3)

| Metabolite | Vehicle Mean ± S.D. (n = 3) | LND Mean ± S.D. (n = 3) | TTFA Mean ± S.D. (n = 3) |
|------------|----------------------------|-------------------------|--------------------------|
| Succinate  | 0.48 ± 0.03                | 0.90 ± 0.06             | 0.87 ± 0.2               |
| Fumarate   | 0.35 ± 0.05                | 0.35 ± 0.05             | 0.28 ± 0.023             |
| Malate     | 0.74 ± 0.08                | 0.96 ± 0.04             | 0.71 ± 0.03              |
| Citrate    | 0.89 ± 0.004               | 0.27 ± 0.01             | 0.90 ± 0.06              |
| Glutamate  | 28.08 ± 2.02               | 31.81 ± 2.03            | 27.01 ± 1.45             |

**FIGURE 9.** Flux analysis of glutamine metabolism. A, kinetics of citrate labeling in control cells after switching to [13C5,15N2]glutamine. B, kinetics of malate labeling in DMSO-treated cells after switching to [13C5,15N2]glutamine. Solid lines represent simulated labeling curves that best fit the data points.
and complex II subunit mutations are both associated with the development of cancer (41, 45). Accumulation of succinate resulting from complex II deficiency inhibits prolyl hydroxylases in the cytosol, leading to stabilization and activation of HIF1α (46). Complex II inhibition can also induce ROS production (41, 47). Both of these pathways are involved in tumorigenesis (46–48). Our findings raise a concern that complex II inhibition by LND may be advantageous to certain tumor types. In line with this notion, both LND- and TTFA-treated cells showed increased labeling through oxidative glutamine metabolism (Fig. 7, C–F) as well as higher rates of glutaminolysis (Fig. 7G) and glutamine uptake (Fig. 7I). These effect are reminiscent of the greater reliance on glutamine by many cancer cells (44). Consistent with our findings, previous studies have shown that complex II knockdown or fumarate hydratase deficiency resulted in higher glutamine contribution to the TCA cycle and higher overall glutamine consumption (45, 49). It is still unknown why deficiencies in TCA cycle enzymes would result in a higher consumption of glutamine. Nevertheless, our finding suggests that a combination of LND with a glutaminase inhibitor (50) could have a positive synergistic effect when used as an anticancer therapeutic approach.

Flux analysis was conducted to more fully understand the effect of LND treatment on the metabolism of DB-1 cells (Figs. 7G and 10G). This revealed that LND lowered the flux from reductive carboxylation of glutamine-derived α-ketoglutarate into citrate (Fig. 7A) by nearly 50% (Fig. 10G). In contrast, flux through the reductive carboxylation pathway was unchanged in
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