Combinatorial CRISPR-Cas9 Metabolic Screens Reveal Critical Redox Control Points Dependent on the KEAP1-NRF2 Regulatory Axis

Graphical Abstract

Highlights

- CRISPR screen reveals gene essentiality and interactions in the metabolic network
- Isozyme families display redundancy and compensation with single-gene knockout
- Integration of CRISPR screen and MFA link genotype to cell-specific phenotype
- KEAP1 mutation alters cancer redox network and essentiality of oxidative PPP genes

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In Brief

Zhao et al. used combinatorial CRISPR screening to elucidate gene essentiality and interactions in the cancer metabolic network. Examination of cell-type-specific essentiality revealed a critical regulation of redox metabolism along the KEAP1-NRF2 signaling axis.
Combinatorial CRISPR-Cas9 Metabolic Screens Reveal Critical Redox Control Points Dependent on the KEAP1-NRF2 Regulatory Axis

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SUMMARY

The metabolic pathways fueling tumor growth have been well characterized, but the specific impact of transforming events on network topology and enzyme essentiality remains poorly understood. To this end, we performed combinatorial CRISPR-Cas9 screens on a set of 51 carbohydrate metabolism genes that represent glycolysis and the pentose phosphate pathway (PPP). This high-throughput methodology enabled systems-level interrogation of metabolic gene dispensability, interactions, and compensation across multiple cell types. The metabolic impact of specific combinatorial knockouts was validated using 13C and 2H isotope tracing, and these assays together revealed key nodes controlling redox homeostasis along the KEAP-NRF2 signaling axis. Specifically, targeting KEAP1 in combination with oxidative PPP genes mitigated the deleterious effects of these knockouts on growth rates. These results demonstrate how our integrated framework, combining genetic, transcriptomic, and flux measurements, can improve elucidation of metabolic network alterations and guide precision targeting of metabolic vulnerabilities based on tumor genetics.

INTRODUCTION

Cancer cells are characterized by unchecked cellular proliferation and the ability to move into distant cellular niches, requiring a rewiring of metabolism to increase biosynthesis and maintain redox homeostasis. This reprogramming of cellular metabolism is now considered an essential hallmark of tumorigenesis (Pavlova and Thompson, 2016). Since the metabolic network is highly redundant at the isozyme and pathway levels, reprogramming is an emergent behavior of the network and manifests itself in non-obvious ways. For instance, a unique metabolic feature of tumor cells is a reliance on aerobic glycolysis to satisfy biosynthetic and ATP demands (Hensley et al., 2016). This metabolic rewiring is coordinated, in part, by the selective expression of distinct isozymes, which may benefit the cell by offering different kinetics or modes of regulation (Chaneton et al., 2012; Christofk et al., 2008; Patra et al., 2013). However, isozyme switching is not solely a consequence of genomic instability and instead can be a coordinated step in tumorigenesis that facilitates cancer cell growth and survival (Castaldo et al., 2000; Guzman et al., 2015). Therefore, understanding which isozymes and pathway branch points are important and how they interact with and compensate for one another is necessary to effectively target metabolism in cancer cells.

In this regard, the advent of CRISPR screening technology now provides a rapid, high-throughput means to functionally characterize large gene sets (Shaem et al., 2014; Wang et al., 2014). This analysis has led to greater annotation of essential genes in human cancers and context-dependent dispensability (Hart et al., 2015; Wang et al., 2015). Correspondingly, single-gene knockout (SKO) CRISPR screens have been able to identify important genes in redox homeostasis and oxidative phosphorylation in conjunction with metabolic perturbations (Arroyo et al., 2016; Birsoy et al., 2015). However, in the context of mammalian metabolism the SKO CRISPR approach comes with limitations, as redundancies and plasticity of the metabolic network may allow the system to remodel around an SKO, thereby confounding analyses of impact on cellular fitness. To overcome this challenge, our group and others recently developed combinatorial gene KO screening approaches that may provide a more suitable platform to study gene dispensability and also systematically map their interactions (Boettcher et al., 2017; Chow et al., 2017; Han et al., 2017; Shen et al., 2017; Wong et al., 2016).
Using this combinatorial CRISPR genetic screening format, coupled with interrogation of metabolic fluxes, we systematically studied the dispensability and interactions within a set of genes encoding enzymes involved in carbohydrate metabolism, including glycolysis and the pentose phosphate pathway (PPP). We illustrated functional relationships between dominant and minor isozymes in various families and discovered multiple genetic interactions within and across glucose catabolic pathways. Aldolase and enzymes in the oxidative PPP (oxPPP) emerged as critical drivers of fitness in two cancer cell lines, HeLa and A549. Distinctions in this dependence are influenced by the KEAP1-NRF2 signaling axis, which coordinates the cellular antioxidant pathway in response to redox stress. We found loss or mutation of KEAP1 E3-ubiquitin ligase upregulates NRF2-mediated transcription of genes involved in glutathione (GSH) synthesis and NADPH regeneration, making the oxPPP less important for NADPH production and less critical for cell growth in these contexts. Thus, mutation status of the KEAP1-NRF2 regulatory axis should be considered when designing therapeutic strategies that target redox pathways in cancer cells.

RESULTS

Combinatorial CRISPR-Cas9 Screening to Probe Metabolic Networks

To systematically study the dispensability and interactions of genes underlying carbohydrate metabolism, we applied a combinatorial CRISPR screening approach (Shen et al., 2017) to interrogate singly and in combination a set of 51 genes, encompassing glycolysis, gluconeogenesis, PPP, and glucose entry into the TCA cycle (Figure 1A). We generated three single guide RNAs (sgRNAs) per gene such that nine unique constructs were present for every gene pair, resulting in a dual-sgRNA library consisting of 459 elements targeting genes individually, as well as 11,475 unique elements targeting two different genes simultaneously (Table S1). The dual-sgRNA constructs were synthesized from oligonucleotide arrays, cloned into a lentiviral vector, and then transduced into HeLa or A549 cells stably expressing Cas9 (Figures 1B, S1A, and S1B). Through sampling of sgRNA frequencies at days 3, 14, 21, and 28 (Figures S1C and S1D), both robust gene-level fitness values ($f_i$) and
interaction scores (\(\tau_{pq}\)) were computed. Finally, impact of SKOs and dual-gene knockouts (DKOs) on cellular growth and metabolic fluxes was validated in a targeted fashion.

**Mapping Metabolic Gene Dependencies in Glucose Catabolism**

Upon analyzing fitness scores for individual gene KOs across the metabolic network (Table S2), we noted that for most (but not all) isozyme families, a dominant gene showed the greatest dispensability (Figures 2A and S2A). Consistent with the notion of a “cancer-specific” isozyme (Hay, 2016), HK2, ALDOA, PGK1, and PKL all showed a fitness defect greater than 2-fold higher as compared to other isozymes. However, not all families exemplified this dynamic, with ENO1/ENO3 and the lactate dehydrogenase (LDH) family showing similar dispensability across gene members (Figures 2A and S2A). The general dispensability of SKOs within the LDH family is notable given the critical role of glycolysis in the maintenance of cancer cell homeostasis and concomitant need to regenerate cytosolic NAD+ when relying on glycolytic flux (Vander Heiden et al., 2009). Importantly, nodes central to the regeneration of equivalents (NADH and NADPH)—GAPDH, G6PD, and PGD—were found to be critical for cellular growth (Figures 2A and S2A).

We hypothesized that gene expression could explain why certain genes were less dispensable and why certain families did not display a dominant member. Indeed, lower fitness score may be associated with higher gene expression (R = −0.461, p = 6.7e−04 and R = −0.429, p = 1.7e−03 for HeLa and A549 cells, respectively). These expression-driven differences also partially explained dynamics within isozyme families. For instance, ALDOA had a much lower fitness score and higher gene expression as compared to ALDOB and ALDOC (Figure 2B). ENO1 and ENO3 both displayed negative fitness scores and both were more highly expressed than ENO2 (Figures 2B and S2C). However, the dispensable isozyme families LDH and PDH (key for maintenance of glycolytic flux and oxidation of pyruvate, respectively) were also found to be highly expressed in both cell types (Figures 2B and S2C). With each family having more than two member enzymes, this result demonstrates that vital functions of cell metabolism can be carried out by multiple genes and show surprising resiliency through isozyme compensation or network behavior.

To this end, SKOs correlated well (R = 0.718, p = 3.1e−09) across both cell lines (Figure 2C). This correlation extended to expression of all enzymes (R = 0.938, p < 2.2e−16). Furthermore, HeLa fitness scores correlated well with previously published HeLa screening data (R = 0.664, p = 1.435e−07) (Hart et al., 2015). However, these results exemplify the challenge in understanding metabolic topology through screening individual genes: few metabolic genes are essential, and essential elements are typically conserved across all cell types.

We subsequently hypothesized that gene interactions could provide information on metabolic network topology and cell-specific adaptations in these pathways. Indeed, a notable number of gene pairs were found to significantly interact (Figures 2D and 2E; Table S3). Specifically, after filtering for genes with reads per kilobase of transcript per million mapped reads (RPKM) < 0.15, we observed 35 interactions (Z score < −3) in the combined HeLa and A549 interaction network (Figure 2E; Table S4), of which 10 (~30%) have been previously reported as protein-protein interactions (Stark et al., 2006). Five gene pair interactions were shared across both cell types.

Notably, the conserved interaction of ENO1/ENO3 demonstrates the possible compensation observed in SKO results (Figure 2A). Previous results have demonstrated that passenger deletion of ENO1 in glioblastoma (GBM) cell lines increases their dependence on ENO2 and generates a GBM synthetic lethality (Muller et al., 2012). As ENO2 is only expressed in neural tissues, our results suggest that ENO1 and ENO3 may compensate for one another in these cell lines. Additionally, redox-associated genes GAPDH and G6PD had many interacting partners, consistent with their negative SKO fitness scores and metabolic functions (Figure 2E). As NAD(P)H is required for both bioenergetics and biosynthetic reactions, alteration of cofactor balance or regeneration fluxes could have large impacts on distal reactions within the network.

**Validation of Significant SKO and DKO Results on Cellular Fitness and Metabolic Fluxes**

Next, to functionally validate the screening results, competition assays and metabolic flux measurements were conducted in the presence of SKO and DKO pairs. Competition assays were performed by mixing control tdTomato+ cells expressing an empty vector, with tdTomato– cells expressing a gRNA of interest (Figure 3A), and relative growth rates were assayed by quantifying the ratio of ± cells in the mixture via flow cytometry (Figure 3B). Dominant family member isozyme fitness was observed in the ALDOA family (Figure 3C), and significant gene interactions over additive SKO effects were observed in multiple gene pairs (Figures 3D and 3E). Correspondingly, perturbations in glycolytic flux were observed through dynamic labeling of metabolites (i.e., pyruvate, lactate, and alanine) from 13C-labeled glucose ([U-13C6]glucose) (Figure 3F). Notably, DKO of ENO1 and ENO3 significantly decreased flux through glycolysis compared to control and SKOs (Figures 3G, S3A, and S3B) and also displayed significantly lower fitness (Figure 3H). Finally, we applied specific 13C and 2H tracers to quantify how the oxPPP contributed to NADPH regeneration (Figure 3I) (Lee et al., 1998; Lewis et al., 2014). SKO of oxPPP enzymes was indeed observed to lower flux (Figures 3J and 3K) and fitness (Figures 3L and S3C) through this pathway.

**Comparison of Metabolic Liabilities across Cell Lines Reveals Key Role of KEAP1-NRF2**

We next focused on differences in the screens of these two cell lines to explore how oncogenic status contributes to metabolic reprogramming. By conducting screens in A549 and HeLa cells and comparing fitness results, we could also gain insights into the impact of SKO results in combination with endogenous mutations. Notably, screening results suggested and we validated that SKO of oxPPP genes (i.e., G6PD and PGD) impacted the growth and survival of HeLa cells more dramatically than A549 cells (Figures 4A, S4A, and S3C) with observed editing rates in each cell line > 95% (Figure S3D). Intriguingly, the expression of G6PD and PGD in these cell lines showed the opposite trend, with A549 cells expressing these genes at significantly higher levels but having a lower dependence on them to maintain
Figure 2. Combinatorial CRISPR Screens Reveal Metabolic Network Dependencies

(A) SKO fitness scores for HeLa cells, plotted as \( f_0 \) (day\(^{-1}\)), with a more negative score representing a loss in fitness with SKO. Plotted as mean ± SD.

(B) Multi-isoform family member fitness scores and gene expression for HeLa (top) and A549 (bottom) cells.

(C) Relative comparison of SKO fitness scores (\( f_0 \)) across both cells.

(D) Relative comparison of genetic interaction scores (\( p_{gg} \)) across both cell lines.

(E) Combined genetic interaction map of both cell lines. Green solid line represents interactions observed in both cell lines. Red and blue lines represent significant genetic interactions in A549 and HeLa cells, respectively. See also Figure S2 and Tables S2, S3, and S4.
Figure 3. Screening Results Validated through Targeted Fitness and Metabolic Flux Measurements

(A) Schematic of cell competition assay used to validate growth. A Cas9-expressing cell is transduced with an sgRNA lentivirus of interest (tdTomato−) and mixed with a control Cas9-expressing cell transduced with a tdTomato lentivirus (tdTomato+). The cells are grown together and the percentage of control (tdTomato+) cells is used to assess relative fitness of SKO.

(B) Non-targeting control (top) is stable for duration of experiment and shows no fitness changes. SKO of ALDOA (bottom) shows decreased fitness over time as control cells take over population.

(C) SKO competition assay of ALDO isozyme family. ALDOA shows greatest loss of fitness.

(D) Growth validation of PFKM/PGD genetic interaction. DKO (green) shows significantly greater decrease in fitness over additive SKO effect (black).

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growth and viability (Figures 4A and S4A). As the oxPPP is critical for maintaining redox homeostasis (i.e., NADPH regeneration) (Kuehne et al., 2015), mutations within control points of redox metabolism could drive this differential sensitivity and further extend the interactions of metabolic genes to known oncogenes or tumor suppressors.

In this regard, A549 non-small-cell lung cancer (NSCLC) cells harbor a loss-of-function mutation in KEAP1 while this regulatory axis is functional in HeLa cells. Loss-of-function mutation of KEAP1 is observed in 20%–50% of NSCLCs (Singh et al., 2006). KEAP1 is a redox-sensitive E3 ubiquitin ligase that targets oxidized NRF2, the master transcriptional regulator of the cellular antioxidant response (DeNicola et al., 2011; Ishii et al., 2000; Thimmulappa et al., 2002), and previous work has demonstrated an ability of NRF2 to alter metabolic fluxes (DeNicola et al., 2015; Mitsuishi et al., 2012; Thimmulappa et al., 2006). Consequently, we hypothesized that the mutational status of this pathway potentially influenced oxPPP sensitivity.

KO of KEAP1 in HeLa cells significantly increased NRF2 levels and expression of oxPPP enzymes G6PD and PGD (Figures S3E and 4B), consistent with the increased expression levels observed in A549 cells (KEAP1-deficient) relative to HeLa cells (KEAP1 WT [wild-type]) (Figure S4A, bottom left). We next determined how oxPPP flux contributed to cytosolic NADPH pools via G6PD measurements, WT KEAP1 increased NRF2 levels as compared to parental cells (Figure S4B), and previous work has demonstrated an ability of NRF2 to alter metabolic fluxes (DeNicola et al., 2015; Mitsuishi et al., 2012; Thimmulappa et al., 2006). Consequently, we hypothesized that the mutational status of this pathway potentially influenced oxPPP sensitivity.

KO of KEAP1 in HeLa cells significantly increased NRF2 levels and expression of oxPPP enzymes G6PD and PGD (Figures S3E and 4B), consistent with the increased expression levels observed in A549 cells (KEAP1-deficient) relative to HeLa cells (KEAP1 WT [wild-type]) (Figure S4A, bottom left). We next determined how oxPPP flux contributed to cytosolic NADPH pools using [3-2H]glucose in KEAP1 KO cells (Lewis et al., 2014). For all sgRNAs we observed a significant decrease in labeling (Figure 4C), which indicates higher pathway flux and loss of label via GSH-mediated H-D exchange (Zhang et al., 2017). This enhanced GSH buffering capacity is consistent with the greater dispensability of oxPPP enzymes observed in A549 cells as compared to HeLa cells (Figure 4A).

We next hypothesized that KEAP1 mutational status could directly alter sensitivity to SKO of oxPPP enzymes and quantified the impact of such SKOs on the fitness and metabolism of an isogenic panel of A549 cells. Ectopic expression of WT KEAP1 decreased NRF2 stabilization as compared to constitutively active C273S mutant KEAP1 (Zhang and Hannink, 2003) (Figure S4B). Interestingly, overexpression of either mutant or WT KEAP1 increased NRF2 levels as compared to parental cells (Figure S4B). Re-expression of WT KEAP1 in A549 cells increased cell sensitivity to PGD KO as compared to C273S KEAP1 mutant cells (Figures 4D and S4C), highlighting the role of KEAP1 in regulating oxPPP enzyme expression and flux. Consistent with these fitness results and the above metabolic measurements, WT KEAP1 expression increased the contribution of PGD to cytosolic NADPH regeneration (Figure 4E) and decreased expression of oxPPP enzymes (Figure 4F).

Finally, we hypothesized that KEAP1 remodels redox metabolism due to its canonical role in the cellular antioxidant response. Indeed, expression of WT KEAP1 was found to decrease expression of both NADPH-regenerating enzymes and those involved in GSH synthesis (Figure 4G). Consistent with decreased expression of GSH synthesis enzymes, intracellular GSH levels were decreased by 45% upon expression of WT KEAP1 (Figure 4H). Presumably, the decreased buffering capacity by GSH and lower expression of other NADPH regenerating contribution to the increased dependence on oxPPP flux observed in cells expressing WT KEAP1. A model therefore emerges from our screening results, whereby KEAP1 mutational status alters the relative importance of the oxPPP by modulating expression of the redox network to drive GSH synthesis and regeneration (Figure 4).

DISCUSSION

While it is clear that cancer cells rely on aerobic glycolysis to maintain biosynthetic fluxes and ATP demands (Hsu and Saba-tini, 2008), how the underlying metabolic network topology changes in response to specific oncogenic events is not fully clear. In this study, we comprehensively interrogated metabolic gene dispensability, interaction, and compensation through a combinatorial CRISPR-Cas9 screening approach. Key nodes within glycolysis were found to significantly interact with one another (e.g., ALDOA and PGD) in an emergent network behavior. Many of these interactions were conserved across cells of different origin, implying such enzyme interaction pairs harbor some function that warrants future interrogation.

Other interactions were demonstrative of metabolic compensation within isozyme families (e.g., ENO1 and ENO3) and consistent with previously described mechanisms of metabolic synthetic lethality (Dey et al., 2017; Muller et al., 2012). These observed network features present a new opportunity through combinatorial (pairwise) screening to understand if/how cells can adapt around loss of a metabolic enzyme. Knowing if a solid tumor of interest is pharmacologically vulnerable to a metabolic inhibitor a priori will allow for future precision medicine applications.

In fact, by comparing relative SKO scores across cell types, we were able to elucidate a paradoxical resistance to targeting the oxPPP along the KEAP1-NRF2 axis. Even though cells

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(E) Growth validation of ALDOA/GAPDH interaction.
(F) Atom transition map depicting glycolysis. Fully labeled [U-13C6]glucose leads to fully labeled pyruvate, lactate, and alanine.
(G) Metabolic validation of DKO interaction in ENO1/ENO3; DKO significantly lowered flux through glycolysis over control or SKOs. * indicates statistical significance (p < 0.05) for all conditions as compared to DKO.
(H) Growth validation of ENO1/ENO3 interaction.
(I) Atom transition map depicting oxPPP tracing. [3-2H]glucose labels cytosolic NADPH through oxPPP. Labeling on glycolytic intermediates from [1,2-13C]glucose is changed by shunting of glucose through oxPPP.
(J) Metabolic validation of PGD SKO at day 4. oxPPP contributes less NADPH with PGD KO. Plotted as mean ± 95% CI. Asterisk (*) indicates statistical significance by non-overlapping confidence intervals.
(K) Metabolic validation of G6PD SKO at day 7. Less glucose is shunted through oxPPP with G6PD KO.
(L) SKO competition assay of oxPPP enzymes.
All experiments were performed in HeLa cells.
(O–E, G, H, K, and L) Data plotted as mean ± SEM. See also Figure S3.
**Figure 4. KEAP1 Mutational Status Alters Redox Metabolism and Impact of oxPPP Gene KOs on Cellular Fitness**

(A) Plot of cell-specific fitness scores for expressed genes. Positive scores are SKOs that are essential in A549s and negative scores are SKOs more essential in HeLa cells. The cell-specific essentiality scores respond to the $Z$ score transformed residuals of linear regression of HeLa and A549 SKO fitness, shown in Figure S4A.

(B) Immunoblot of KEAP1 SKO in HeLa cells.

(C) Contribution of oxPPP to cytosolic NADPH with KEAP1 SKO in HeLa cells. Plotted as mean ± 95% CI. Asterisk (*) indicates statistical significance by non-overlapping confidence intervals.

(D) Relative PGD SKO effect in A549s with KEAP1 mutant panel.

(E) Contribution of oxPPP to cytosolic NADPH in A549s with KEAP1 mutant panel. Plotted as mean ± 95% CI. Asterisk (*) indicates statistical significance by non-overlapping confidence intervals.

(F) Immunoblot of A549s with KEAP1 mutant panel.

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potently upregulated expression of oxPPP enzymes upon loss of KEAP1, cells were less vulnerable to KO of enzymes in this metabolic pathway. In this case, alternate NADPH regeneration pathways and increased antioxidant buffering by GSH pools provide compensation and survival benefits to cells. Our NAPDH tracing data demonstrated that cells lacking functional KEAP1 exhibit higher oxPPP flux, as evidenced by reduced labeling due to increased H-D exchange through GSH-related pathways (Zhang et al., 2017). Indeed, elevated oxPPP enzyme levels and increased GSH pools would specifically increase exchange flux, resulting in the observed decrease in labeling downstream of [3-2H]glucose. The integration of such functional measurements with genetic screening and transcriptional results provides better context to interpret the observed metabolic reprogramming downstream of KEAP1-NRF2.

Our results suggest that KEAP1 mutational status must be considered when targeting the oxPPP therapeutically. In fact, recent work has implicated KEAP1 mutational status as a driver of metabolic reprogramming and potential targeting of glutaminase in pre-clinical models of lung adenocarcinoma (Romero et al., 2017). Consistent with our findings, KEAP1 mutation increases intracellular GSH levels and need for cysteine, causing an increased need for glutamine anaplerosis to support glutamate/cysteine antiporter flux (SLC7A11) (Muir et al., 2017; Romero et al., 2017). Other recent work has also implicated KEAP1 mutational status as a driver of chemotherapeutic resistance in preclinical models of lung cancer and further demonstrates the need for new paradigms connecting oncogenic mutations to cancer cell survival (Kroll et al., 2017).

Moving forward, it will be important to perform such screens across a larger number of cell types to elucidate a more comprehensive picture of metabolic network reprogramming. The high-throughput methodology presented here increases the feasibility of such studies. We note also that comparing the absolute fitness values in screens across cell lines can be confounded by various factors. These include differences in relative cell growth and expression of CRISPR effectors among others, and thus devising new strategies for normalization will be valuable to improve the utility of future screening datasets. We also note the critical importance of sgRNA efficacy and anticipate that continued improvements in sgRNA design (Chari et al., 2015; Doench et al., 2016; Erard et al., 2017) will be critical to improving consistency and signal to noise in the assays. Finally, layering in data from complementary perturbation strategies such as CRISPR activation/inhibition and small molecule inhibition should enable charting of more comprehensive networks underlying cellular function and transformation.

Discovery of the unique metabolic features in transformed cells has spurred much interest in exploiting metabolic vulnerabilities for drug discovery (Vander Heiden, 2011). In fact, metabolic inhibitors have been developed as single-agent therapeutics and combination therapeutics for many different cancer types (Tennant et al., 2010). However, these agents have found varying success in the clinic due an inability to determine proper cancer types in preclinical development. While cancer cells share common hallmarks of metabolic reprogramming, cell-of-origin and tumorigenic drivers uniquely influence the direction and extent of metabolic reprogramming. The new paradigm of incorporating combinatorial CRISPR screening, transcriptomic information, and metabolic flux measurements presented here will provide a new platform to address this limitation. By interrogating metabolism at the network level, new therapeutic targets may be identified, and clinicians may become better equipped at identifying the most responsive patient populations.

**STAR METHODS**

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**SUPPLEMENTAL INFORMATION**

Supplemental Information includes four figures and four tables and can be found with this article online at https://doi.org/10.1016/j.molcel.2018.01.017.

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**AUTHOR CONTRIBUTIONS**

D.Z., M.G.B., C.M.M., and P.M. designed the study. D.Z., M.G.B., J.L., J.H.M., A.B., C.S.A., and P.M. performed experiments and analyzed data. R.S. and T.I. provided technical advice. D.Z., M.G.B., C.M.M., and P.M. wrote the manuscript. All authors read the manuscript.

**DECLARATION OF INTERESTS**

The authors declare no competing interests.

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**REFERENCES**


# STAR★METHODS

## KEY RESOURCES TABLE

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CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Prashant Mali (pmali@ucsd.edu).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Cell Lines and Culture Conditions
HEK293T, A549, HeLa-AAVS-Cas9-Hygro, A549-AAVS-Cas9-Hygro cells were grown in DMEM supplemented with 10% FBS, 2 mM L-glutamine, 100 units/mL of penicillin, 100 μg/mL of streptomycin, and 0.25 μg/mL of Amphotericin B. HeLa-AAVS-Cas9-Hygro and A549-AAVS-Cas9-Hygro cells were purchased from GeneCopoeia.

METHOD DETAILS

Dual-gRNA library design and cloning
A set of 51 genes, encompassing glycolysis, gluconeogenesis, pentose phosphate pathway, and glucose entry into the TCA cycle were selected for this study. Three unique 20-bp sgRNAs were designed for each target gene and three scrambled, non-targeting sequence absent from the genome were used as control. The dual-sgRNA construct library comprised all pairwise gRNA combinations between either two genes or a gene and a scramble, resulting in 11,475 double-gene-knockout constructs and 459 single-gene-knockout constructs. The dual-gRNA library was generated as previously described (Shen et al., 2017).

Lentivirus production
One 15cm dish of HEK293T cells at 60% confluent were transfected with 3 μg PMD2.G, 12 μg of lenti-gag/pol/PCMVR8.2, and 9 μg of lentiviral vector (library or single constructs) using 36 μL of Lipofectamine 2000. Medium containing viral particles were harvested 48 h and 72 h after transfection, then concentrated with Centricon Plus-20 100,000 NMWL centrifugal ultrafilters, divided into aliquots and frozen at −80°C.

CRISPR/Cas9 dual-gRNA screening
CRISPR Cas9 nuclease stable expressing HeLa and A549 cells were obtained from GeneCopoeia and grown in DMEM medium with 10% FBS and Antibiotic-Antimycotic. Hygromycin B was added at the concentrations of 200 μg/ml or 100 μg/ml for HeLa and A549 cells, respectively. For each screen, cells were seeded in three 15cm dishes at a density of 1 × 10^7 per ml and transduced with the lentiviral dual gRNA library at a low MOI of 0.1-0.3. Puromycin was added at 48 h after transduction at a concentration of 5 μg/ml. Then the cells were cultured and passaged for every 3-4 days while 1 × 10^7 cells were sampled at days 3, 14, 21 and 28 and stored at −80°C until extraction of genomic DNA. Two biological replicates of the screens were performed for each cell line.
Quantification of dual gRNAs abundance

Genomic DNA of the cells were purified using QIAGEN DNeasy Blood and Tissue Kits. To amplify the dual gRNAs from each sample, we used 20 μg of genomic DNA as template across ten 50 μL PCR reactions with Kapa Hifi polymerase. By testing the amplification efficiency, we used 22 - 24 cycles at an annealing temperature of 55°C with the following primers:

Forward: ACACTCTTTCCCTACACGACGCTCTTCCGATCTTATATATCTTGTGGAAAGGACGAAACCG;
Reverse: GACTGGAGTTCAGACGTGTGCTCTTCCGATCTCCTTATTTTAACTTGCTATTTCTAGCTCTA.

The amplicons were pooled and purified with Agencourt AMPure XP bead at a double selection of 0.55 and then 0.83. The samples were quantified with Qubit dsDNA High Sensitivity Kit. To attach Illumina sequencing adaptors and indexes, we used 50 ng of purified step I PCR product as template across four 50 μL PCR reactions with Kapa Hifi polymerase using primers of Next Multiplex Oligos for Illumina (New England Biosciences). 7 or 8 PCR cycles were carried out at an annealing temperature of 72°C. The PCR product were purified twice with Agencourt AMPure XP bead at 0.83 ratio, quantified, pooled and sequenced on an Illumina HiSeq rapid-run mode for 75 cycles paired-end runs.

Computation of single and double gene knockout fitness and genetic interaction scores

Analysis was performed with a previously reported software pipeline constructed from Python, R and Jupyter Notebooks (https://github.com/ucsd-ccbb/mali-dual-crispr-pipeline). The following details are adapted from our published paper (Shen et al., 2017). Briefly, the two gRNA sequences were extracted and trimmed to 19bp from 3' end, and then aligned to the known library sequences with one mismatch allowed. We determined a minimum threshold for read counts based on the histograms and masked out pairwise gRNA constructs that have read counts below the threshold. The read counts were used for analysis of fitness and genetic interaction scores as follows:

1. **Estimation of fitness of each pairwise gRNA construct.** The logarithmic transformation of the frequency of each pairwise gRNA construct in the population is:

   \[ X_c = \log_2 \frac{N_c}{\sum N_c} \]

   where \( N_c \) is the number of cells in the population expressing construct \( c \). We assume that each cell subpopulation grows exponentially:

   \[ N_c(t) = N_c(0) \times 2^{f_c t} \]

   where \( t \) is a given time point; \( f_c \) is the fitness of construct \( c \); \( f_0 \) is the absolute fitness of reference cells which don’t express any constructs. Combining these two equations, we get:

   \[ x_c(t) = a_c + f_c t - \log_2 \sum c 2^{x_c - f_c t} \]

   where \( a_c \equiv x_c(0) \) as the initial condition and \( \sum c 2^{x_c} = 1 \) in the whole population. Fitting to this equation from experimental data of frequencies \( X_c(t) \), we minimize the sum of squares:

   \[ E(a_c, f_c) = \sum c \sum t (X_c(t) - x_c(t))^2 \]

   Here \( E \) is invariant under the substitution \( f_c \rightarrow f_c + \delta \), where \( \delta \) is an arbitrary constant, which can be fixed by setting the mean non-targeting gRNA fitness to zero. To resolve this, one should find the minimum of the function:

   \[ E_1 \equiv E - \lambda \left( \sum c 2^{a_c} - 1 \right) \]

   where \( \lambda \) is the Lagrange multiplier. This solution equals:

   \[ \frac{\partial E_1}{\partial a_c} = \frac{\partial E_1}{\partial f_c} = \frac{\partial E_1}{\partial \lambda} = 0 \]

   When the number of constructs is large, \( \sum c 1 \gg 1 \), the approximation solution is:

   \[ f_c = \frac{\text{Cov}(X_c, t)}{\text{Var}(t)} + \delta \]

   and

   \[ a_c = x_c - f_c t + \log_2 \sum c 2^{x_c - f_c t} \]

   where the bars indicate means over time points. The \( a_c \) values do not depend on the choice of \( \delta \).
(2) Estimation of single gRNA fitness and gRNA–gRNA interactions. For each construct containing gRNAs \( g \) and \( g' \), we define:

\[
f_c = f_g + f_{g'} + \pi_{gg'},
\]

where \( \pi_{gg'} \) is the gRNA–gRNA interaction scores. \( f_c \) is calculated from step (1). \( f_g \) values are found by robust fitting of this equation. The gRNA-level \( \pi_{gg'} \) scores are the residuals of the robust fit.

(3) Computation of gene level fitness based on weighted average of gRNA fitness. We ranked the three gRNAs targeting to the same gene as \( r(g) \in (0, 1, 2) \) in ascending order of \( |f_g| \). The gene-level fitness values are calculated as the weighted means of gRNA fitness values with weights given by the squares of gRNA ranks, \( r^2(g) \). The gene-level interaction scores are calculated as the weighted means of gRNA-gRNA interaction scores with weights given by the products of gRNA ranks, \( r(g) r(g') \).

(4) Correction by replicates. As we performed biological replicates for each experiment, we combine replicates for more power rather than looking for two \( f_c \) separately. We fit a single optimal \( f_c \) from all data points excluding those below the threshold, with the assumption that \( f_c \) does not change across experiments while the initial conditions \( a_c \) may be different. The raw \( P \) value associate to each \( f_c \) is:

\[
t_c = \frac{f_c}{SE(f_c)},
\]

where \( SE(f_c) \) is the standard error of \( f_c \):

\[
SE(f_c) = \sqrt{\frac{\sum (X_c(t) - X_c(t'))^2}{(n_c - 2) \sum (t^2 - \bar{t}^2)}}.
\]

The raw \( P \) values then are transformed into posterior probabilities, \( PP_c \), according to the theory of Storey. To scale the genetic interaction scores for comparison across different experiments, we calculated a genetic interaction \( z \) score by dividing the \( f_c \) by the SE of the observed interaction pair in a given experiment. We consider an interaction to be a meaningful candidate if it has an absolute \( z \) score above 3.

(5) Calculation of false discovery rates by numerical Bayesian ensemble of experiments. We assign a fitness value to each construct \( c \) on the basis of change in fitness relative to the standard deviation of repeated-measurements. The assigned value is either 0 with probability \( (1 - PP_c) \), or a random number within \( f_c \pm s.d. \) We perform 1000 permutations and reported gene level \( f_g \) and \( \pi_{gg'} \) for each permutation. The false discovery rate (FDR) of genetic interactions \( \pi_{gg'} \) is calculated as the odds ratio between the observed and permuted results in the null model, which is obtained by mean-centering of the marginal distribution of every \( \pi_{gg'} \).

Single-gRNA construct cloning
The LentiGuide-Puro vector were linearized using restriction enzyme BsmBI at 55 °C for 3 hours. For each individual gRNA, two oligonucleotides containing the spacer sequences were synthesized as listed in Table S1. The two oligos were annealed and extended to make a double stranded DNA fragment using KAPA HiFi polymerase. The fragment was purified and subjected to Gibson assembly (New England Biolabs) with the linearized LentiGuide-Puro vector.

Competitive cell growth assay
We developed a competitive cell growth assay to assess the effect of gene perturbations by mixing control tdTomato+ cells with tdTomato- cells expressing a gRNA of interest (Figure 3A) and sampling relative growth rates through time by flow cytometry. Cas9-expressing cells were transduced with EF1A- tdTomato-T2A-puromycin lentivirus and cultured under puromycin selection for stable expression of tdTomato. To measure the negative impact of a gRNA introduced gene perturbation on the cellular proliferation rate, the Cas9-expressing cells were cultured in 12-well-plate and transduced with gRNA lentivirus at a high MOI (> 5). The day after transduction, the Cas9-expressing cells were re-suspended, counted, mixed with tdTomato+ Cas9-expressing cells, and re-seed into 12-well-plate. The cells were sampled every 3 or 4 days to score the tdTomato+/tdTomato- ratio by longitudinal flow cytometric analysis. By assuming the exponential growth of the cells, from time \( t_1 \) to \( t_2 \), the growth of cells (tdTomato+ or gRNA expressing) in the mixture population fits to the equation:

\[
N(t) = N(t_1) \times 2^{(t - t_1)/\Delta t},
\]

where \( N \) is the cell number of the certain cell subtype, \( \Delta t \) is the absolute fitness of reference cells which in this case is the tdTomato+ cells, and \( \Delta t \) is fitness measurements of the certain cell subtype. For a certain gRNA (or a pair of gRNAs), the \( \Delta t \) is able to be calculated easily according to the equation without measuring the absolutely fitness of reference cells:

\[
\frac{N(t_2)}{N(t_1)} = \frac{N(t_2)}{N(t_1)} \times 2^{(t_2 - t_1)/\Delta t}.
\]
Although the percentage of tdTomato+ cells in the mixtures with the cells expressing non-targeting control gRNAs was stable over time, we normalize the fitness of gRNA of interest to non-targeting control gRNAs for side by side comparisons. The cell viability of a gRNA of interest (non-log transformed fitness) relative to non-targeting controls showed in Figure 3 is as follows:

\[ F_{\text{gRNA}} = \frac{2(\Delta f_{\text{gRNA}}/\Delta f_{0})}{2(\Delta f_{\text{NTC}}/\Delta f_{0})} \times 100\% . \]

The expected cell viability of a pair of gRNAs calculated according to:

\[ F_{\text{gRNA1, gRNA2}} = F_{\text{gRNA1}} \times F_{\text{gRNA2}} . \]

In addition, \( f_0 \) is able to be measured by counting of the absolute cell number over time base on the Equation (1). Then the effects of a gene perturbation (e.g., PGD) relative to non-targeting controls (NTC) in a certain cell subtype (e.g., KEAP1 mutations) are calculable as follows:

\[ R_{\text{PGD, KEAP1}} = \frac{(f_0 + \Delta f_{\text{PGD, KEAP1}}) - (f_0 + \Delta f_{\text{NTC, KEAP1}})}{f_0 + \Delta f_{\text{NTC, KEAP1}}} . \]

**RNA sequencing data analysis**

RNA sequencing data were obtained from the ENCODE project (GEO: GSE30567, sample GSM765402 and GSM758564 for HeLa and A549 cell lines respectively). The results were expressed as the average value of reads per kilobase transcript per million mapped reads (RPKM) across two biological replicates. The average RPKM values were log2 transformed for Pearson correlation analysis.

**Stable isotope tracing**

For isotopic labeling experiments, cells were cultured in glucose- and glutamine-free media (GIBCO) supplemented with 10% dialyzed FBS, 100 U/mL penicillin/streptomycin, 4mM glucose (Sigma), and 20 mM of either [3-2H]glucose (98%, Cambridge Isotope Laboratories), [U-13C6]glucose (99%, Cambridge Isotope Laboratories), or [1,2-13C]glucose (99%, Cambridge Isotope Laboratories).

Cells were rinsed with PBS before addition of tracing media. For glycolytic measurements, basal media was changed 1 hr before addition of tracer media and extracted at indicated time intervals. For measurement of shunting through oxPPP, cells were traced for 4hrs. For estimation of PGD contribution to cytosolic NADPH, cells were traced for 48hrs.

**Metabolite Extraction and GC-MS Analysis**

Cells were rinsed with 0.9% (w/v) saline and 250 μL of –80°C MeOH was added to quench metabolic reactions. 100 μL of ice-cold water supplemented with 10 μg/mL norvaline was then added to each well and cells were collected by scraping. The lysate was moved to a fresh 1.5 mL eppendorf tube and 250 μL of –20°C chloroform supplemented with 4 μg/mL D31 palmitate was added. After vortexing and centrifugation, the top aqueous layer and bottom organic layer were collected and dried under airflow.

Dervatization of aqueous metabolites was performed using the Gerstel MultiPurpose Sampler (MPS 2XL). Methoxime-tBDMS derivatives were formed by addition of 15 μL 2% (w/v) methoxylamine hydrochloride (MP Biomedicals) in pyridine and incubated at 45°C for 60 minutes. Samples were then silylated by addition of 15 μL of N-tert-butyldimethylsilyl-N-methyltrifluoroacetamide (MTBSTFA) with 1% tert-butylimidethylchlorosilane (tBDMS) (Regis Technologies) and incubated at 45°C for 30 minutes. Aqueous metabolites were analyzed by GC-MS using a DB-35MS column (30 m x 0.25mm i.d. x 0.25 μm, Agilent J&W Scientific, Santa Clara, CA) in an Agilent 7890B gas chromatograph (GC) interfaced with a 5977C mass spectrometer (MS). Electron impact ionization was performed with the MS scanning over the range of 100-650 m/z for polar metabolites. For separation of aqueous metabolites the GC oven was held at 100°C for 1 min after injection, increased to 255°C at 3.5°C/min, and finally increased to 320°C at 15°C/min and held for 3 min.

Dried organic fraction was saponified and esterified to form fatty acid methyl esters (FAMEs) by addition of 500 μL of 2% (w/v) H2SO4 in MeOH and incubated at 50°C for 120 minutes. FAMES were then extracted by addition of saturated NaCl and hexane before collection and drying of the inorganic layer. Derivatized fatty acids were analyzed by GC-MS using a select FAME column (100 m x 0.25mm i.d. x 0.25 μm; Agilent J&W Scientific) as above, with the MS scanning over the range 120-400 m/z. For separation the GC oven was held at 80°C for 1 min after injection, increased to 160°C at 20°C/min, increased to 198°C at 1°C/min, and finally increased to 250°C at 5°C/min and held for 15 min.

**Metabolite integration and isotopomer spectral analysis (ISA)**

Mass isotopomer distributions and total abundances were determined by integration of mass fragments (Table S1) and correcting for natural abundances using MATLAB-based algorithm. Glycolytic flux was estimated by normalizing pyruvate, lactate, or alanine abundance by the sum of intracellular branched-chain amino acids abundance and M+3 label. Oxidative PPP shunting was estimated by \( M+1/M+1 + M+2 \) labeling on pyruvate from [1,2-13C]glucose (Lee et al., 1998). Isotopomer spectral analysis (ISA) was performed to estimate contribution of oxPPP to cytosolic NADPH as previously described (Lewis et al., 2014). ISA compares experimental labeling
of fatty acids to simulated labeling using a reaction network where C16:0 is condensation of 14 NADPHs. Parameters for contribution of PGD to lipogenic NADPH (D value) and percentage of newly synthesized fatty acid (g(t) value) and their 95% confidence intervals are then calculated using best-fit model from INCA MFA software (Young, 2014).

**Immunoblotting**
Cultured cells were washed with cold PBS and harvested on ice with mPER (Pierce Biotechnology) with freshly added 1x HALT inhibitor (Thermo Fisher Scientific). Protein concentration was determined by BCA assay and equal amounts of protein were resolved on SDS-PAGE gel and transferred to nitrocellulose membrane. Membrane was blocked with 5% milk in TBST (Tris-buffered saline with 0.1% Tween 20) for 2-3hrs and incubated overnight at 4°C with primary antibody: anti-Vinculin (Abcam), anti-G6PD (Cell Signaling Technologies), anti-PGD (Santa Cruz Biotechnology), anti-KEAP1 (Proteintech), anti-HA (Abcam), or anti-Nrf2 (Cell Signaling Technology). Blots were then incubated with secondary antibody for 1hr at room temp, Anti-Rabbit HRP-conjugate (Cell Signaling Technology) or Anti-Mouse HRP-conjugate (Cell Signaling Technology). Finally blots were incubated with ECL substrate (BioRad) and imaged.

**RT-PCR**
Total mRNA was isolated from cells using RNA isolation kit (RNaseasy Mini Kit; Qiagen). Isolated RNA was reverse transcribed using cDNA synthesis kit (High-capacity cDNA Reverse Transcription Kit; Thermo Fisher Scientific). Real-time PCR was performed using SYBR green reagent (iTaq Universal SYBR Green Supermix; Bio-Rad). Relative expression was determined using Livak (ΔΔCt) method with RPL27 and RPLP0 as housekeeping gene. Primers used were taken from Primerbank (Wang et al., 2012) and tabulated in Table S1. All commercial kits were used per the manufacturer’s protocol.

**Glutathione measurement**
Intracellular glutathione was measured using Glutathione Assay Kit (Sigma) per manufacturer’s protocol. Ten centimeter dishes of cells were assayed in quintuplicate and cells were counted in parallel for normalization.

**QUANTIFICATION AND STATISTICAL ANALYSIS**
Unless indicated, all results shown as mean ± SEM of biological triplicates. P values were calculated using a Student’s two-tailed t test; *, P value between 0.01 and 0.05; **, P value between 0.001 and 0.01; ***, P value less than 0.001.

**DATA AND SOFTWARE AVAILABILITY**
Analysis was performed with a previously reported software pipeline constructed from Python, R and Jupyter Notebooks (https://github.com/ucsd-ccbb/mali-dual-crispr-pipeline; Shen et al., 2017). Information of paired guide RNA designs and raw read counts of screens: Table S1. Single gene fitness: Table S2. Genetic interactions (pi) scores: Table S3. Top hits of genetic interactions: Table S4. Original imaging data have been deposited to Mendeley Data and are available at https://doi.org/10.17632/dnkdmrsc9v.1. The accession numbers for the A549 RNA-seq data and HeLa RNA-seq data are GEO: GSM758564 and GSM765402, respectively.
Supplemental Information

Combinatorial CRISPR-Cas9 Metabolic Screens Reveal Critical Redox Control Points

Dependent on the KEAP1-NRF2 Regulatory Axis

Dongxin Zhao, Mehmet G. Badur, Jens Luebeck, Jose H. Magaña, Amanda Birmingham, Roman Sasik, Christopher S. Ahn, Trey Ideker, Christian M. Metallo, and Prashant Mali
Figure S1. Schematic of dual-gRNA-library construction and quality control of screens; related to Figure 1. (A) Oligonucleotides bearing two sgRNA spacers were synthesized, amplified, and cloned into a lentiviral gRNA cloning vector. Next, a fragment containing a sgRNA scaffold and the mouse U6 promoter was inserted between the two spacers to yield the final dual-gRNA expression construct. A pair of primer matching sites labeled in blue were designed for enrichment of the two spacer regions prior to deep sequencing analysis. (B) Frequency distribution of the metabolism dual-gRNA plasmid library. (C) Principle component analysis (PCA) of the dual-gRNA read count distributions. (D) Cumulative frequency of dual-gRNA constructs by deep sequencing. Shift in the curves at days 14, 21, and 28 represents the depletion of dual-gRNA constructs. Each time point was measured in duplicates.
Figure S2. CRISPR screening results reveal metabolic network dependencies; related to Figure 2. (A) SKO fitness scores for A549 cells, plotted as $f_0$ (day$^{-1}$), with a more negative score representing a decrease in fitness with SKO. Plotted as mean ± SD. (B) Gene pairs with significant genetic interaction scores (z-score < -3) are shown. Conserved interactions cross HeLa and A549 are indicated in blue. Previously reported interactions are indicated in red. Purple indicates the conserved interactions which have been previously reported.
Figure S3.

A

B

C

D

HeLa: gG6PD 100%
HeLa: gPGD 97%
A549: gPGD 96%
A549: gG6PD 95%

E

Non-Targeting 0.084%
gKEAP1-1 100%
gKEAP1-2 100%
gKEAP1-3 100%
Figure S3. Screening results validated through metabolic flux measurements and fitness assays; related to Figure 3. (A, B) Metabolic validation of DKO interaction in ENO1/ENO3. DKO significantly lowered flux through glycolysis over control or SKOs. A, measurement of labeled Lactate. B, measurement of labeled Alanine. † indicates statistical significance (p<0.05) for all conditions as compared to DKO. (C) SKO competition assay of oxPPP genes in HeLa (left) and A549 (right) cells. HeLa data replicated from Figure 3L and log transformed for comparison. (D) Deep sequencing analysis of indels (insertions and deletions) introduced by CRISPR-Cas9 at 10 days after transduction of G6PD or PGD gRNA constructs. (E) Deep sequencing analysis of indels introduced by CRISPR-Cas9 at two weeks after transduction of KEAP1 gRNA constructs in HeLa cells. Ordinate shows the read counts of indels at each corresponding location. Most cells were successfully targeted after transduction of gRNAs, while only a background level of mutagenesis was observed in the cells transduced with non-targeting control gRNAs. These experiments suggest high targeting efficiency in both the A549 and HeLa Cas9-stable cell lines. (A-C) Plotted as mean ± SEM.
**Step 1:** measure the fitness of reference cells

\[ N_e(t_2) = N_e(t_1) \times 2^{(f_0 \times (t_2-t_1))} \]

\[ R^2 = 0.9977 \]
\[ f_0 = 0.8205 \]

**Step 2:** measure the fitness of gRNA 

in Keap mutation cells (slope)

\[ \frac{N_{eRNA}(t_2)}{N_0(t_2)} = \frac{N_e(t_1) \times 2^{(f_0 + \Delta f_{RNA})(t_2-t_1)}}{N_0(t_1) \times 2^{(f_0)(t_2-t_1)}} \]

**Step 3:** calculate the relative effect of PGD knockout

\[ R_{PGD,KEAP1} = \frac{(f_0 + \Delta f_{PGD,KEAP1}) - (f_0 + \Delta f_{NTCKEAP1})}{f_0 + \Delta f_{NTCKEAP1}} \]

**Figure S4.**
Figure S4. *KEAP1* mutational status alters redox metabolism and impact of oxPPP gene knockouts on cellular fitness; related to Figure 4. (A) Scatter plots (left) of SKO fitness and gene expression in HeLa versus A549. Residual plots (right) of linear regressions showing the outliers between HeLa and A549. oxPPP genes (*G6PD* and *PGD*) showed more essentiality in HeLa cells versus A549, while their mRNA expression levels are lower in HeLa cells versus A549. (B) Immunoblot of A549s with *KEAP1* mutant panel. Superfluous lane (negative control) removed from image. (C) Measurement of relative *PGD* perturbation effect in A549 cells across *KEAP1* mutant panel. Growth curve of the reference cells, which is tdtomato+ cells in this case, and its absolute fitness ($f_0$) was extracted by counting average cell numbers in three independent experiments for three days. The fitness of *PGD* perturbation ($\Delta f_{PGD,KEAP1}$) relative to non-targeting controls (NTC) in *KEAP1* mutation cells were measured by competitive assay. Finally, by incorporating also the absolute fitness of reference cells, the relative effects of *PGD* perturbation ($R_{PGD,KEAP1}$) in *KEAP1* mutant cells was calculated.