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MYCN mediates cysteine addiction and sensitizes neuroblastoma to ferroptosis

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Aberrant expression of MYC transcription factor family members predicts poor clinical outcome in many human cancers. Oncogenic MYC profoundly alters metabolism and mediates an antioxidant response to maintain redox balance. Here we show that MYCN induces massive lipid peroxidation on depletion of cysteine, the rate-limiting amino acid for glutathione (GSH) biosynthesis, and sensitizes cells to ferroptosis, an oxidative, non-apoptotic and iron-dependent type of cell death. The high cysteine demand of MYCN-amplified childhood neuroblastoma is met by uptake and transsulfuration. When uptake is limited, cysteine usage for protein synthesis is maintained at the expense of GSH triggering ferroptosis and potentially contributing to spontaneous tumor regression in low-risk neuroblastomas. Pharmacological inhibition of both cystine uptake and transsulfuration combined with GPX4 inactivation resulted in tumor remission in an orthotopic MYCN-amplified neuroblastoma model. These findings provide a proof of concept of combining multiple ferroptosis targets as a promising therapeutic strategy for aggressive MYCN-amplified tumors.

Many human cancers rely on aberrant expression of MYC transcription factor family members to allow unhindered growth and proliferation; high expression levels are predictive of poor clinical outcome¹. Aberrant MYC oncoprotein levels lead to gross transcriptional changes with hundreds, if not thousands, of upregulated and downregulated genes, which together drive various hallmark features of malignant cells. Pharmacological approaches to target aberrant MYC have largely failed. Therefore, MYC synthetic lethal interactions have been exploited for the development of therapeutic concepts to specifically target MYC-driven

tumors, yet with limited success². Remarkably, in view of MYC's oncogenic activity, its potential to promote cell death apart from cell proliferation is paradoxical. Transgenic mouse models support a role for MYC for both tumor development but also spontaneous cell death depending on tissue type and context³. Both may ultimately be related to MYC's profound influence on various aspects of cellular metabolism, with their interdependencies still poorly understood.

Childhood neuroblastoma, an embryonic tumor derived from progenitors of the sympathetic nervous system, is a paradigmatic

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model for MYC-driven cancers⁴. Amplified *MYCN* identifies a highly aggressive subtype associated with malignant progression and poor outcome despite intensive multimodal treatments. However, a substantial proportion of low-risk neuroblastomas with elevated *MYCN* expressed from a normal *MYCN* locus, particularly those arising in children younger than 18 months, regress spontaneously (stage 4S disease) by unknown mechanisms even when the disease is metastatic⁵. High-risk neuroblastomas lacking amplified *MYCN* harbor rearrangements of other *MYC* gene family members, *TERT* or alternative mechanisms of telomere lengthening (ALT) often associated with *ATRX* gene mutations, the latter subtype being incompatible with high *MYCN* or *MYC* expression^{6,7} (later referred to as MYC(N)). Beside these alterations linked to telomere maintenance mechanisms, mutations in *ALK* or other developmental genes lead to stalled differentiation and tumors composed of heterogeneous cell types resembling different states of the normal neuroendocrine differentiation trajectories. The spectrum of cell types ranges from differentiated over undifferentiated adrenergic-to-mesenchymal cell types (triggered by adrenergic-to-mesenchymal transition), where malignant progression, therapy resistance and disease relapse are strongly associated with undifferentiated cell types.^{8,9}

MYCN-amplified neuroblastoma cells, like other MYC-driven cancer cells, have been found to be addicted to the amino acid glutamine (Gln), the absence of which causes growth arrest or apoptosis¹⁰. More recently, reports showed that neuroblastoma cells are also addicted to iron and are sensitized to ferroptosis^{11,12}, a new iron-dependent oxidative form of cell death associated with lipid peroxidation and insufficient capacity to eliminate lipid peroxides¹³. Whereas the apoptosis pathway of regulated cell death is often genetically or epigenetically impaired in primary high-risk and relapsed neuroblastomas¹⁴, there is only limited knowledge of how metabolic rewiring as a consequence of aberrant *MYCN* controls the liabilities of fast-proliferating malignant cells that are confronted with accumulating reactive oxygen species (ROS).

In this study, by performing single amino acid deprivations in high *MYCN* and low *MYCN* neuroblastoma cells, we discovered strong dependency of high *MYCN* cells on the amino acid cysteine. Using functional *MYCN* synthetic lethal metabolic and genetic screens, we further identified cyst(e)ine deprivation and glutathione peroxidase 4 (GPX4) inhibition as selective liabilities in *MYCN*-amplified neuroblastomas. Combined targeting of cystine uptake, cysteine synthesis via transsulfuration and GPX4 in an orthotopic neuroblastoma model strongly reduced tumor growth in vivo. Multi-omics profiling identified multiple cell type-specific and *MYCN*-regulated mechanisms inhibiting ferroptosis in adrenergic and mesenchymal neuroblastoma cells. Taken together, our study uncovered mechanisms crucial to ferroptosis escape in *MYCN*-amplified neuroblastomas; simultaneous inhibition of those mechanisms led to tumor regression in vivo.

Results

Cystine deprivation induces MYCN-dependent ferroptosis. First, we analyzed the interplay of oncogenic *MYCN* activity with amino acid metabolism. Downregulating *MYCN* in the *MYCN*-amplified IMR5/75 neuroblastoma cell model¹⁵ (approximately 65% reduction; Fig. 1a) slowed cell proliferation without inducing cell death (Extended Data Fig. 1a,b) and reduced the intracellular pools of all amino acids (Fig. 1b). Most prominently, cysteine was reduced nearly tenfold (Extended Data Fig. 1c). Inhibiting *MYCN* binding to Myc-associated factor X (MAX) using 10058-F4 (ref. 16) yielded similar results (Fig. 1b and Extended Data Fig. 1d). These data show that high *MYCN* levels are associated with high levels of cellular cysteine, probably mediated by increased synthesis and/or uptake from the microenvironment. Systematic depletion of individual amino acids from the growth medium impaired cell viability in both high *MYCN* and low *MYCN* cells in most cases (Fig. 1c).

However, in line with recent reports^{11,12}, cells with high *MYCN* expression exhibited stronger dependency on cystine imported by cystine/glutamate-exchange transporter x_c^- and readily reduced to two cysteine molecules intracellularly. Cystine deprivation caused robust cell death in high *MYCN* cells, which was largely prevented by downregulation of *MYCN* expression (Fig. 1c) or inhibition of *MYCN*-MAX binding (Fig. 1d). Overexpressing *MYCN* in *MYCN* diploid cells (Tet21N neuroblastoma cell model¹⁷), rendered these cells highly vulnerable to cystine deprivation (Fig. 1e). Neuroblastoma cell lines with intermediate *MYCN* or *MYC* levels caused by gene translocations are known to exist. We inferred MYC(N) activity by target gene expression score⁵, which agrees with transcript and protein MYC(N) levels (Extended Data Fig. 1e,f). Cell death after cystine deprivation increased with MYC(N) activity score, being virtually absent in cell lines immortalized by alternative telomere lengthening and lacking MYC(N) aberrations and peaking in *MYCN*-amplified cell lines with the highest activity scores (Fig. 1f). These data demonstrate that oncogenic MYC(N) expression is associated with cysteine addiction, with cysteine reduction resulting in massive cell death in an MYC(N)-dependent manner.

Cell death in the cystine-deprived high *MYCN* cells was not abrogated by the inhibition of (1) caspases to prevent apoptosis, (2) lysosomal function to downregulate autophagy or (3) receptor-interacting serine/threonine-protein kinase 1 (RIPK1) to prevent necroptosis (Extended Data Fig. 1g). Availability of cystine/cysteine is the rate-limiting step in GSH synthesis counteracting ROS¹⁸, which prevents the execution of ferroptosis, an oxidative, non-apoptotic and iron-dependent form of regulated cell death caused by ROS-mediated lipid peroxidation^{13,19,20}. Indeed, ferrostatin-1 (Fer-1), a specific inhibitor of ferroptosis, or a lipophilic antioxidant or an intracellular iron chelator such as ciclopirox olamine (CPX) averted death in cystine-deprived high *MYCN* neuroblastoma cells (Fig. 1g and Extended Data Fig. 1h). Monitoring ROS formation by flow cytometry using the lipid peroxidation sensor, C11-BODIPY, showed that cystine deprivation dramatically increased cellular lipid peroxidation selectively in high *MYCN* cells (Fig. 1h). Inhibiting ferroptosis with lipophilic antioxidants or an intracellular iron chelator inhibited lipid peroxidation in cystine-deprived high *MYCN* cells (Fig. 1i). Hence, when deprived of cystine, high *MYCN* neuroblastoma cells exhibited lipid peroxidation and died via ferroptosis.

Addition of GSH to the cystine-free medium also prevented lipid peroxidation in *MYCN*-amplified cells similar to Fer-1 (Fig. 1i) and rescued cell viability similar to Fer-1 (Fig. 1j and Extended Data Fig. 1i), suggesting that high GSH levels protect against ferroptosis. Consistent with their higher sensitivity to ferroptosis, *MYCN*-amplified neuroblastoma cells had lower baseline GSH and cysteine levels compared to cell lines with lower MYC(N) activity scores (Extended Data Fig. 1j,k). However, on downregulation of *MYCN* in *MYCN*-amplified cells, intracellular GSH levels were reduced threefold, the reduced-to-oxidized GSH ratio was halved and intracellular ROS levels increased (Fig. 1k), suggesting that high *MYCN* expression increases GSH synthesis and ROS clearance. Overall, these results indicate that although oncogenic *MYCN* activates the production of GSH, it is maintained at a low steady state level due to its rapid consumption in fast-proliferating cells.

The only other amino acid showing a selective dependency on *MYCN* was Gln, confirming previous reports^{10,21}, although this was significantly less pronounced (Fig. 1c). Interestingly, cysteine and Gln (when converted to glutamate) are two important GSH precursors. Depleting Gln in addition to cystine partially restored GSH levels probably because of less lipid peroxidation²² and rescued cell viability (Fig. 2a–c). Gln/cystine deprivation prevented the induction of ferroptosis-promoting genes (*CHAC1*, *HMOX1*) and reduced transcriptional activation of genes protecting against cellular stress (*ATF4*, *ATF3*) shown to be activated on cystine depletion²³.

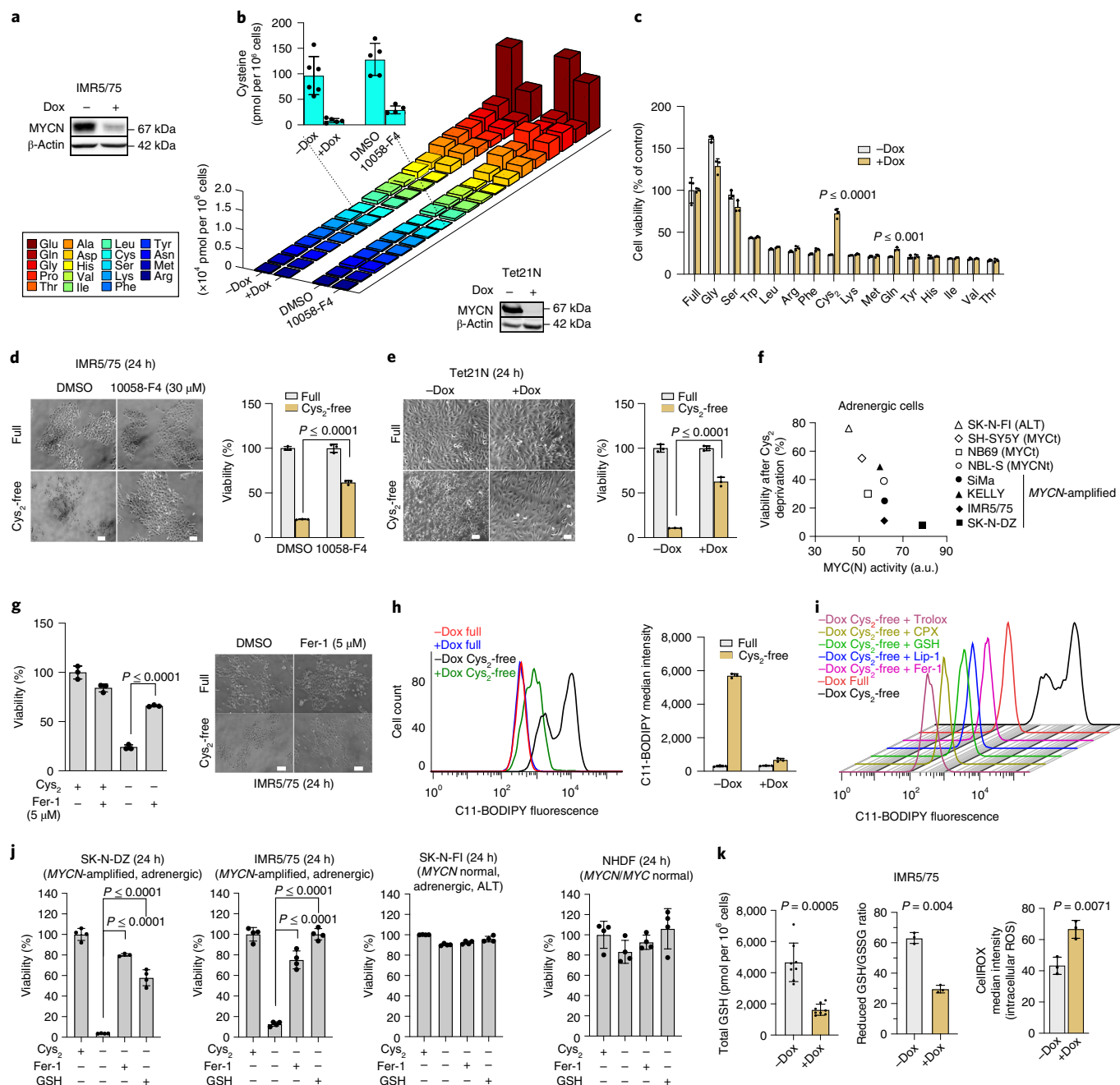


Fig. 1 | Cystine addiction in MYCN-expressing neuroblastoma cells. **a**, Representative western blot of IMR5/75 neuroblastoma cells on MYCN knockdown using Dox (expression: -Dox, high; +Dox, low); the experiment was replicated three times. **b**, Intracellular amino acid quantification after MYCN inhibition for 96 h (+Dox, $n = 5$ samples and -Dox, $n = 6$ samples, or 10058-F4, inhibiting MYCN-MAX binding, $n = 4$ samples and dimethyl sulfoxide (DMSO)-treated, $n = 5$ samples). Data represent the mean \pm s.e.m. The experiment was replicated three times. **c**, Standardized viability of IMR5/75 after single amino acid depletions (48 h). Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. **d,e**, Cellular responses to Cys₂ deprivation in high MYCN (-Dox) and low MYCN (+Dox) state in IMR5/75 (**d**) and Tet21N (**e**) cells; the mean viability of cells was standardized to untreated (full medium) and representative western blot of neuroblastoma Tet21N cells. Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. Images of the cells are shown on the left. Scale bar, 50 μm. **f**, Sensitivity to Cys₂ deprivation versus level of MYC(N) activity in a panel of neuroblastoma cell lines. **g**, Relative viability (survival of compound-treated cells divided by survival of vehicle-treated cells) of IMR5/75 cells after Cys₂ deprivation for 72 h in the presence or absence of Fer-1. Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. Images of the cells are shown on the left. Scale bar, 50 μm. **h**, Analysis of lipid peroxidation in Cys₂-deprived high or low MYCN IMR5/75 cells ($n = 3$ samples; the experiment was replicated 3 times). **i**, Analysis of lipid peroxidation in Cys₂-deprived high or low MYCN IMR5/75 cells in the presence or absence of Trolox, CPX, Lip-1, GSH or Fer-1. The experiment was replicated three times. **j**, Relative viability of SK-N-DZ, IMR5/75, SK-N-FI and normal human dermal fibroblasts (NHDFs) after Cys₂ deprivation in the presence or absence of GSH and Fer-1. $n = 4$ samples. The experiment was replicated three times. **k**, Quantification of total intracellular GSH ($n = 8$ samples) levels and the reduced GSH/GSH disulfide (GSSG) ratio in IMR5/75 cells ($n = 3$ samples). Analysis of intracellular ROS levels using CellROX staining and flow cytometry in IMR5/75 cells in high MYCN (-Dox) and low MYCN (+Dox) state ($n = 3$ samples). Data represent the mean \pm s.e.m. The experiment was replicated three times. Statistical analysis was performed using a two-tailed Student's *t*-test.

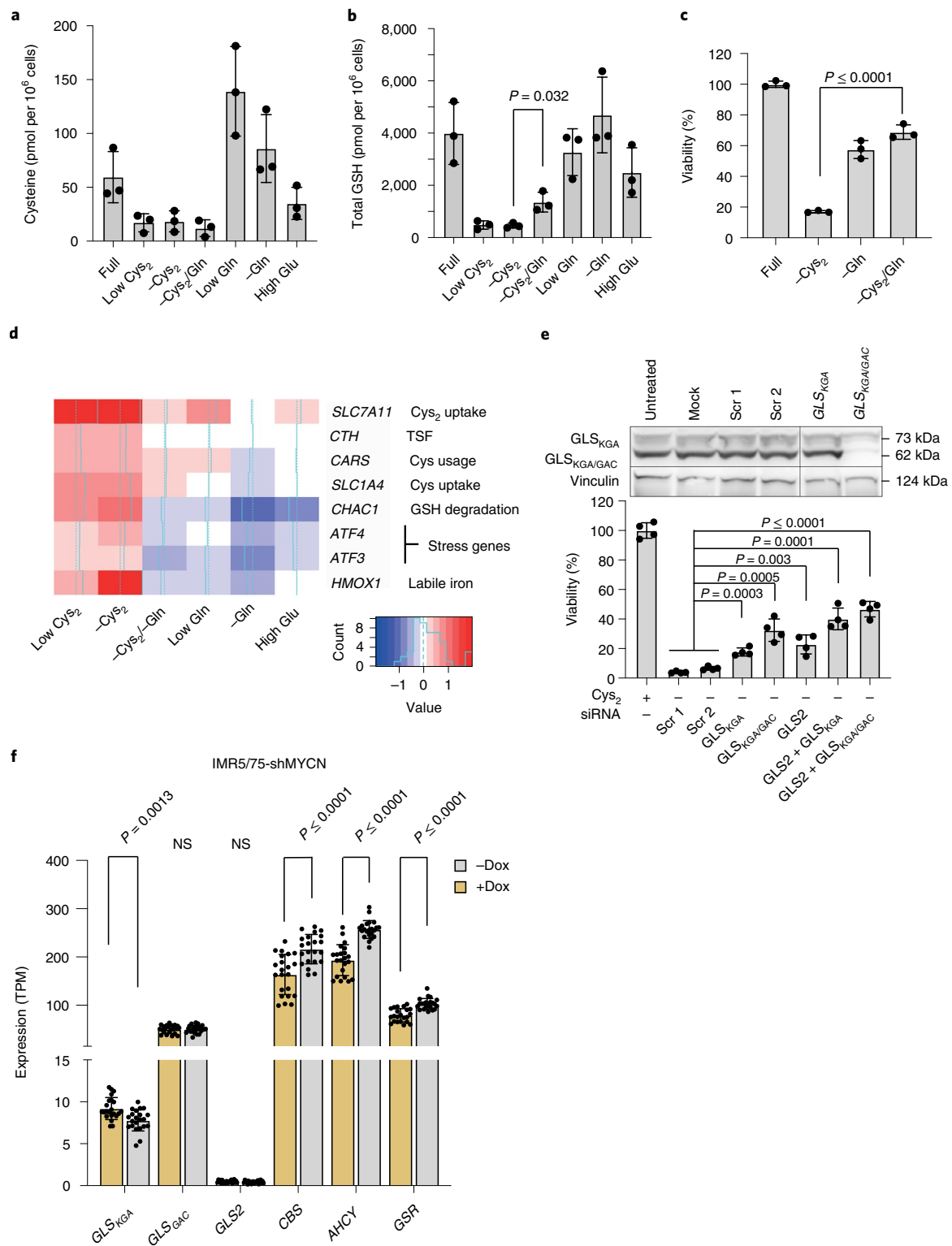


Fig. 2 | Gln is required for ferroptosis in high MYCN cells. **a–c**, Cysteine (**a**), total GSH levels (**b**) and cell viability (**c**) on Cys₂ and Gln deprivation. Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. **d**, RNA-seq of IMR5/75 cells on treatment with the conditions indicated in the figure. $n = 3$ samples. **e**, siRNA-mediated *GLS_{KGA/GAC}* knockdown (72 h) on Cys₂ deprivation (24 h) in IMR5/75 cells. Representative western blots of each isoform. Data represent the mean \pm s.e.m.; $n = 4$ samples. The experiment was replicated three times. **f**, Expression of glutaminolysis genes (mitochondrial *GLS_{GAC}*; cytosolic *GLS_{KGA}* and *GLS2*) compared with MYCN-regulated genes (*CBS*, *AHCY*, *GSR*) in high MYCN and low MYCN IMR5/75 cells. TPM, transcripts per million. Data represent the mean \pm s.e.m. The statistical analysis was performed using a two-tailed Student's *t*-test.

(Fig. 2d). Besides Gln depletion, inhibiting glutaminolysis should prevent ferroptosis²⁴. Indeed, knockdown of either glutaminase isoform, $GLS_{KGA/GAC}$ averted ferroptosis confirming that glutaminolysis is required to induce ferroptosis in cystine-deprived high *MYCN* cells (Fig. 2e). However, since GLS_{KGA} is repressed by *MYCN* (Fig. 2f), GLS_{GAC} appears to be the main glutaminase isoform involved in these cells.

Identification of ferroptosis genes in *MYCN*-amplified cells.

To systematically identify the cellular vulnerabilities associated with high *MYCN* expression, we performed an unbiased synthetic lethal small interfering (siRNA) screen using high/low *MYCN* IMR5/75 cells (Fig. 3a). Among the top hits that reduced the survival of high *MYCN* cells were enzymes involved in GSH metabolism that detoxify lipid peroxides (Fig. 3b,c, Extended Data Fig. 2a and Supplementary Table 1). Namely, inhibition of *GSR*, *GPX4*, *GPX6* or *GSTM1*, *GSTM5* and *GSTK1* (Fig. 3b and Extended Data Fig. 2a) caused a selective reduction in viability in high *MYCN* cells. Knockdown of either of the two enzymes catalyzing GSH biosynthesis, glutamate–cysteine ligase catalytic subunit (*GCLC*) and glutathione synthetase (*GSS*) also showed synthetic lethality with high *MYCN* state (Fig. 3b and Extended Data Fig. 2a).

One of the screening hits, *GPX4*, is an anti-ferroptotic selenoprotein that protects against lipid peroxidation using GSH^{25,26}. siRNA-mediated *GPX4* knockdown induced ferroptosis in high *MYCN* cells, which was rescued by Fer-1 (Fig. 3d and Extended Data Fig. 2b). Similarly, we observed reduction of viability using the *GPX4* inhibitor RSL3, enhanced by supplementing iron and rescued by Fer-1 or GSH (Fig. 3e,f) or simultaneous MYC(N) activity inhibition using 10058-F4 (Extended Data Fig. 2c). The dependency of cell viability on *GPX4* activity increased with MYC(N) activity across our broader cell line panel (Fig. 3g,h and Extended Data Fig. 2d,e) unlike *GCLC* or *GSR*, which exhibited no dependency (Extended Data Fig. 2f). Next, we studied whether the effect of *GPX4* knockdown was linked to cystine uptake and metabolic gene expression in 348 cancer cell lines²⁷. Cells dependent on *GPX4* showed low expression levels of *SLC7A11* of the x_c^- system, importing cystine in exchange for glutamate (Extended Data Fig. 2g). Neuroblastoma cell lines were, after ovarian cancer cell lines, the second most dependent on *GPX4* (Extended Data Fig. 2h).

To therapeutically investigate the role of *GPX4* inhibition in vivo, we used an inducible CRISPR–Cas9 system to eliminate *GPX4* (*GPX4* knockout). In the *MYCN*-amplified SK-N-DZ three-dimensional

(3D) cell culture model, *GPX4* knockout induced ferroptotic cell death. This was rescued by simultaneously treating cells with the 10058-F4 MYC–MAX inhibitor (Fig. 3i). We then generated an orthotopic mouse model for human neuroblastoma by transplanting these SK-N-DZ neuroblastoma cells orthotopically into the adrenal gland and allow the cells expressing native *GPX4* levels to develop into small tumors (Fig. 3j). Subsequently, the CRISPR–Cas9 *GPX4* knockout was induced by doxycycline (Dox) treatment 7 d after transplantation. Although this resulted in a significant reduction in tumor weight compared to controls (Fig. 3k), it was not sufficient for tumor eradication as also shown by others²⁸. Ferroptosis markers such as *CHAC1* and *TFRC*²⁹ increased despite only partial reduction of *GPX4* transcripts, suggesting activation of ferroptosis in vivo (Fig. 3l). In summary, our data show that *GPX4* partially protects high *MYCN* neuroblastoma cells from ferroptosis in vitro and in an orthotopic neuroblastoma mouse model.

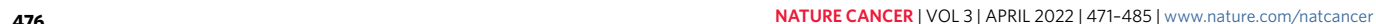
Both neuroblastoma cell lines and primary neuroblastomas had the lowest *SLC7A11* expression compared to other tumor entities (Extended Data Fig. 2i,j). This suggests that the sensitivity of neuroblastoma to cystine depletion and *GPX4* inhibition is due, at least in part, to reduced uptake of cystine¹⁸. *SLC7A11* knockdown or inhibition by sulfasalazine moderately reduced viability regardless of *MYCN* level (Fig. 4a–c) as shown recently¹². Erastin, a more potent *SLC7A11* inhibitor²³, selectively induced ferroptosis in the high *MYCN* state (Fig. 4d), which was partially rescued by additional inhibition of MYC(N) activity (Fig. 4e). Erastin-induced cell death increased with MYC(N) activity score in neuroblastoma cell lines (Fig. 4f,g and Extended Data Fig. 2k), was associated with decreased GSH and reduction of the GSH reduced-to-oxidized ratio (Fig. 4h) and could be prevented by providing either GSH or Fer-1 (Fig. 4i). Taken together, cystine uptake by *SLC7A11* maintains part of the necessary cysteine for GSH production to protect against ferroptosis.

MYCN induces transsulfuration and prevents ferroptosis.

Cellular cysteine can also be produced in some cell types by transsulfuration^{30,31}. In this process, homocysteine (Hcy), an intermediate of the methionine cycle, and serine are combined to form cystathionine (Cysta), which is further converted to cysteine³² (Fig. 5a). *GPX4* dependency in cancer cell lines was associated with enhanced expression of cystathionine beta-synthase (*CBS*), the rate-limiting enzyme for transsulfuration (Extended Data Fig. 2g), with neuroblastoma being among the cancer entities with the

Fig. 3 | Inhibition of *GPX4* is synthetic lethal with high *MYCN*. **a**, *MYCN* synthetic lethal druggable genome-wide siRNA screening approach in IMR5/75 cells. **b**, Effects of individual siRNAs (gray dots): high *MYCN* versus low *MYCN*, including key players (median of two or three siRNAs) of the top *MYCN* synthetic lethal hits (single star symbol) of GSH metabolism (black) and biosynthesis (green). **c**, *MYCN* effects on lipid peroxide formation and intracellular amino acid levels (fold changes shown in red), the x_c^- system (Cys₂ uptake), the two-step biosynthesis of GSH and GSH metabolism; the single star symbol marks the top *MYCN* synthetic lethal hits of GSH metabolism, with an FDR of 0.2. The action of ferroptosis inhibitors (CPX, Fer-1, Lip-1, Trolox and 10058-F4), class I (erastin, IKE, sulfasalazine), class II ferroptosis inducers (RSL3, ML-210) and the GSH biosynthesis inhibitor buthionine sulfoximine as indicated. **d**, siRNA *GPX4* knockdown in the presence or absence of Fer-1. Data represent the mean \pm s.e.m.; $n = 4$ samples. The experiment was replicated three times. **e**, Relative viability (survival of compound-treated cells divided by survival of vehicle-treated cells) of the KELLY cell line after cotreatment with iron sucrose (25 μ g ml⁻¹) and RSL3. Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. **f**, Relative viability of IMR5/75 cells treated with RSL3 in the presence or absence of Fer-1 or GSH. $n = 4$ samples. The experiment was replicated three times. **g**, DRIVE database²⁷ (RSA values, Pearson's correlation, $P = 0.01$; filled circle, *MYCN*-amplified; white circle, *MYCN*-non-amplified). **h**, Cellular responses of neuroblastoma cell lines to 72 h of RSL3 treatment: cells with *MYCN* amplification (black symbols), moderate *MYCN* expression (white circle) and lack thereof (white triangle). Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. **i**, Dox-inducible *GPX4* CRISPR–Cas9 knockout in a 3D model with *MYCN*-amplified SK-N-DZ cells in the presence or absence of MYC–MAX inhibition. Data represent the mean \pm s.e.m. Right: representative western blot; $n = 4$ samples. The experiment was replicated three times. **j**, Orthotopic mouse neuroblastoma model allowing CRISPR–Cas9-mediated *GPX4* deletion. Panel created with BioRender. **k**, Tumor weight after *GPX4* knockout (+Dox) ($n = 5$ mice per group). A representative western blot for CRISPR–Cas9-mediated *GPX4* deletion is shown. **l**, Elevated messenger RNA expression of the ferroptosis markers *CHAC1* and *TFRC*. Data represent the mean \pm s.e.m.; $n = 4$ samples from each group. Statistical analysis was performed using a one-tailed Student's *t*-test for the in vivo experiments and a two-tailed Student's *t*-test for the in vitro experiments. Box plots: the center line indicates the median value, the lower and upper hinges represent either the 25th and 75th percentiles or the minimum and maximum points and the whiskers denote 1.5 \times the interquartile range (IQR). Each dot corresponds to one sample; one-sided Student's *t*-test; *P* values as indicated.

and S-adenosyl-L-homocysteine hydrolase (AHCY), synthesizing Hcy for transsulfuration, show synthetic lethality with high *MYCN* (Fig. 5b,c and Extended Data Fig. 3a), as are two methyltransferases that feed into Hcy production (Extended Data Fig. 3b).



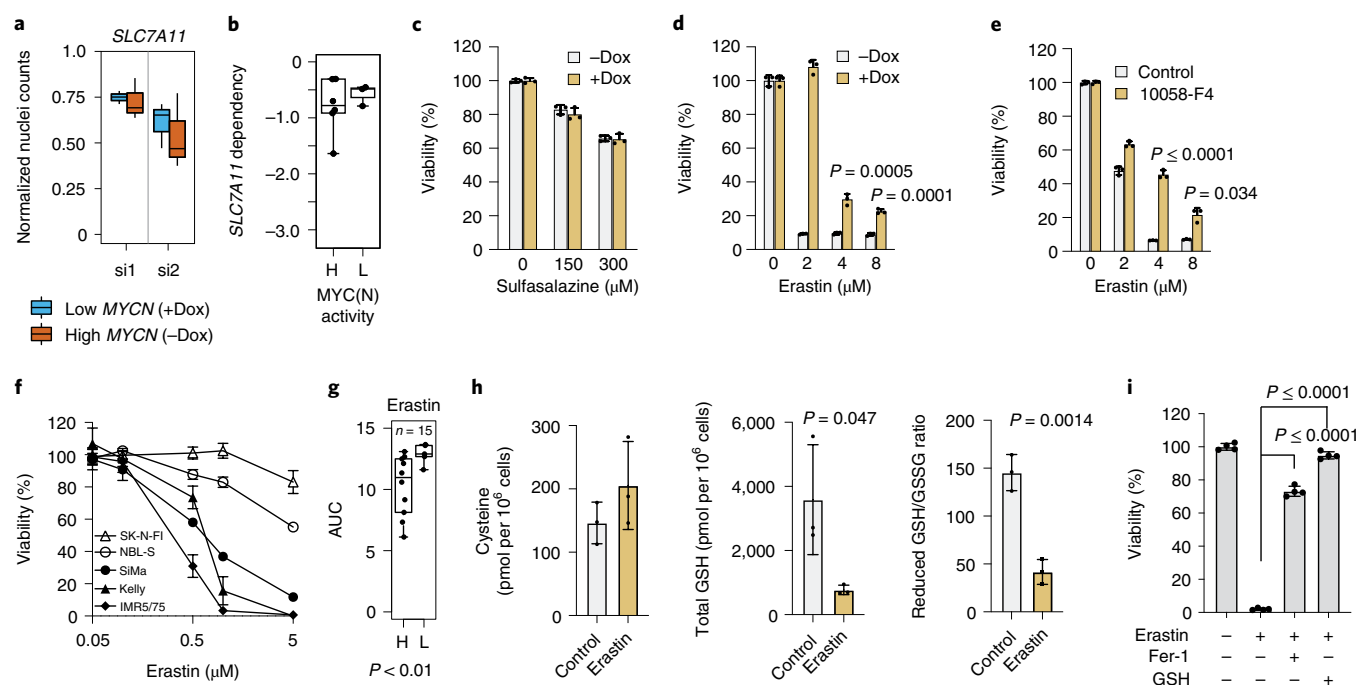


Fig. 4 | Inhibition of cystine uptake by SLC7A11 induces ferroptosis in high MYCN cells. **a**, SLC7A11 knockdown effect in the IMR5/75 cell line. **b**, Correlation between MYC(N) activity and sensitivity to SLC7A11 repression (DRIVE²⁷). **c**, Relative viability (survival of compound-treated cells divided by survival of vehicle-treated cells) of both high and low MYCN IMR5/75 cells after treatment with sulfasalazine. **d,e**, Treatment with erastin for 72 h with MYCN knockdown (**d**) and MYC/MAX inhibitor (**e**). Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. **f,g**, Cellular responses of neuroblastoma cell lines to 72 h of erastin treatment. Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times (**f**) and confirmed by the CTRPv2 (ref. ⁵⁷) data (**g**). AUC, area under the curve. **h**, Cysteine and GSH levels and reduced GSH/GSSG ratio in IMR5/75 cells treated with erastin. Data represent the mean \pm s.e.m.; $n = 3$ samples. The experiment was replicated three times. **i**, Relative viability of IMR5/75 cells treated with erastin in the presence or absence of Fer-1 or GSH (72 h). Data represent the mean \pm s.e.m.; $n = 4$ samples. The experiment was replicated three times. Box plots: the center line indicates the median value, the lower and upper hinges represent the 25th and 75th percentiles, respectively and the whiskers denote 1.5 \times the IQR. Each dot corresponds to one sample; Wilcoxon rank-sum test, P values as indicated.

Supplementing cystine-deprived cells with either Hcy or Cys₂ prevented ferroptosis in all adrenergic neuroblastoma cell lines tested with high or intermediate oncogenic MYC(N) expression, but not in the less common mesenchymal neuroblastoma lines (Fig. 5d and Extended Data Fig. 3c). Pharmacologically inhibiting CTH using propargylglycine (PPG)³³ sensitized adrenergic, but not mesenchymal, high MYCN cell lines to either erastin- or imidazole ketone erastin (IKE)-induced cell death (Fig. 5e). Knockdown of AHCY in adrenergic high MYCN but not mesenchymal neuroblastoma cells impaired colony formation, which was associated with reduced GSH levels and reduction of GSH reduced-to-oxidized ratios (Fig. 5f, g). In summary, these data indicate that transsulfuration provides an internal cysteine source for GSH biosynthesis protecting high MYCN adrenergic neuroblastoma cells from ferroptosis.

Intracellular cysteine is required for two rate-limiting cellular processes: GSH-mediated ROS clearance and production of building blocks in the synthesis of proteins, nucleotides and lipids³⁴ (Fig. 5a). In line with this, protein synthesis inhibition with cycloheximide increased cysteine and total GSH levels (Extended Data Fig. 3d). Notably, reduction/deprivation of cystine (Cys₂) in the medium drastically reduced intracellular cysteine and GSH in the high MYCN state before ferroptosis (Fig. 2a,b) but did not affect cell cycle progression and only moderately reduced protein synthesis (Extended Data Fig. 3e,f). MYCN protein levels were not affected on cystine deprivation compared to methionine- or Gln-deprived cells (Extended Data Fig. 3g). These data suggest robust cysteine channeling into protein synthesis under limited cysteine supply when GSH production is already diminished.

To investigate this further, we performed transcriptomic analysis of IMR5/75 cells cultured in cystine-free medium or inhibiting the x_c^- system with erastin. This revealed activation of a stress response³⁵ before ferroptosis that channeled cysteine into protein synthesis by inducing CARS³¹ and cysteine recycling from GSH (Fig. 2d and Extended Data Fig. 3h–j). *HMOX1* was also activated in these cells, suggesting that the free iron pool is increased before inducing ferroptosis³⁶ (Fig. 2d and Extended Data Fig. 3h–j). Inhibiting CARS prevented ferroptosis in the cystine-deprived high MYCN state, highlighting the competition for intracellularly synthesized cysteine between protein synthesis and redox balance (Fig. 5h). Taken together, transsulfuration supplies cysteine for both protein and GSH synthesis in adrenergic MYCN-amplified neuroblastoma cells but prioritizes cysteine for protein synthesis at the expense of GSH and redox balance when cystine uptake is limited, thus triggering ferroptosis.

Next, we asked how cysteine metabolism and redox homeostasis are affected by oncogenic MYC(N) activity in the adrenergic (active transsulfuration) or mesenchymal (inactive transsulfuration) neuroblastoma subtypes⁸. Adrenergic cells upregulated three key enzymes in transsulfuration, *CBS*, *AHCY* and D-3-phosphoglycerate dehydrogenase (*PHGDH*), in the high MYCN state, while the x_c^- system (*SLC7A11*) controlling cystine uptake was unaffected by changes in MYCN (Extended Data Fig. 4a)³⁷. In a panel of 32 neuroblastoma cell lines, MYCN or MYC amplification or translocation in adrenergic subtypes was accompanied by upregulated *CBS* expression, while MYC translocation/activation in mesenchymal cell lines was not (Fig. 5i). Ectopically expressing MYCN in mesenchymal Tet21N cells left transsulfuration via its rate-limiting enzyme *CBS*

unchanged but induced *SLC7A11*, *GSR*, *GCLC* and thioredoxin reductase 1 (*TXNRD1*), the last one being significantly expressed in *MYCN*-amplified primary neuroblastomas (Extended Data Fig. 4b,c)³⁷. At the gene regulatory level, we found that the *CBS* locus harbored both activating (H3K27ac, H3K4me3) and silencing (H3K27me3) histone modifications, with the former being increased and the latter decreased in the presence of amplified *MYCN* in adrenergic cells compared to non-amplified or mesenchymal cells (Fig. 5j and Extended Data Fig. 5). In primary neuroblastomas, differences in *CBS* expression correlated with histone modifications and methylation of intragenic CpGs dependent on genomic *MYCN* status (Fig. 5k and Extended Data Fig. 6). Together, this suggests that transsulfuration is active in the adrenergic state and regulated at the epigenetic level in high *MYCN* cells.

We found that the levels of *CBS* and *AHCY* were associated with poor patient survival (Fig. 6a). Global gene expression profiles from 498 primary neuroblastomas³⁸ confirmed elevated *AHCY* and *CBS* in *MYCN*-amplified neuroblastomas (Fig. 6b,c). Higher *HMOX1* and lower *SLC3A2* and *TXNRD1* expression were found in stage 4S tumors (Fig. 6d), which tend to regress spontaneously. *CBS*, *AHCY* and *PHGDH* expression was also elevated in mass spectrometry-based global proteomes from *MYCN*-amplified neuroblastoma tumors (Fig. 6e). *SLC7A11* antiporter expression did not correlate with *MYCN* amplification or other risk factors for poor patient outcomes (Fig. 6c). In addition, genes involved in Gln (that is, *SLC38A5*, *SLC1A5*), methionine (that is, *SLC7A5*) and iron uptake (*TFRC*) had higher expression in *MYCN*-amplified or *MYCN*-overexpressed neuroblastomas; *TFRC* was recently associated with ferroptosis in neuroblastomas¹² (Fig. 6c,d and Extended Data Fig. 4c). Genes associated with GSH synthesis/metabolism (that is, *GCLC*, *GSR*) were also upregulated both at the transcript and protein level (Fig. 6c,e,f). Higher expression of the mitochondrial glutaminase (*GLS_{GAC}*)^{24,39} and lower expression of the cytoplasmic glutaminase (*GLS_{KGA}*) isoforms were observed in *MYCN*-amplified tumor transcriptomes and proteomes (Fig. 6c,e), suggesting dependence on mitochondrial glutaminolysis in these neuroblastomas. Together, our results show that *MYCN*-amplified neuroblastomas increase transsulfuration activity, iron import, glutaminolysis and GSH production through coordinated changes in gene expression thereby increasing the susceptibility to ferroptosis.

Cyst(e)ine and GPX4 inhibition as therapeutic targets. To exploit our findings in vivo, we first tested simultaneous inhibition of cysteine uptake and transsulfuration in vitro using IKE, an erastin analog with acceptable pharmacokinetic properties^{40,41}, and PPG³³ (Fig. 7a). We observed a strong synergistic effect with

the two drugs in only *MYCN*-amplified adrenergic cells (Fig. 5e). We then injected mice orthotopically with SK-N-DZ neuroblastoma cells and treated them for two 5-d cycles with a combination of IKE 45 mg kg d⁻¹ and PPG 45 mg kg d⁻¹ (Fig. 7b). When combining these two drugs, we observed a 60% reduction in tumor growth in *MYCN*-amplified SK-N-DZ-driven tumors (Fig. 7c). These data suggest that *MYCN*-amplified neuroblastoma cells with very high levels of *MYCN* are sensitive to a reduction of the intracellular cysteine pool by simultaneous inhibition of cystine import and cysteine synthesis via transsulfuration.

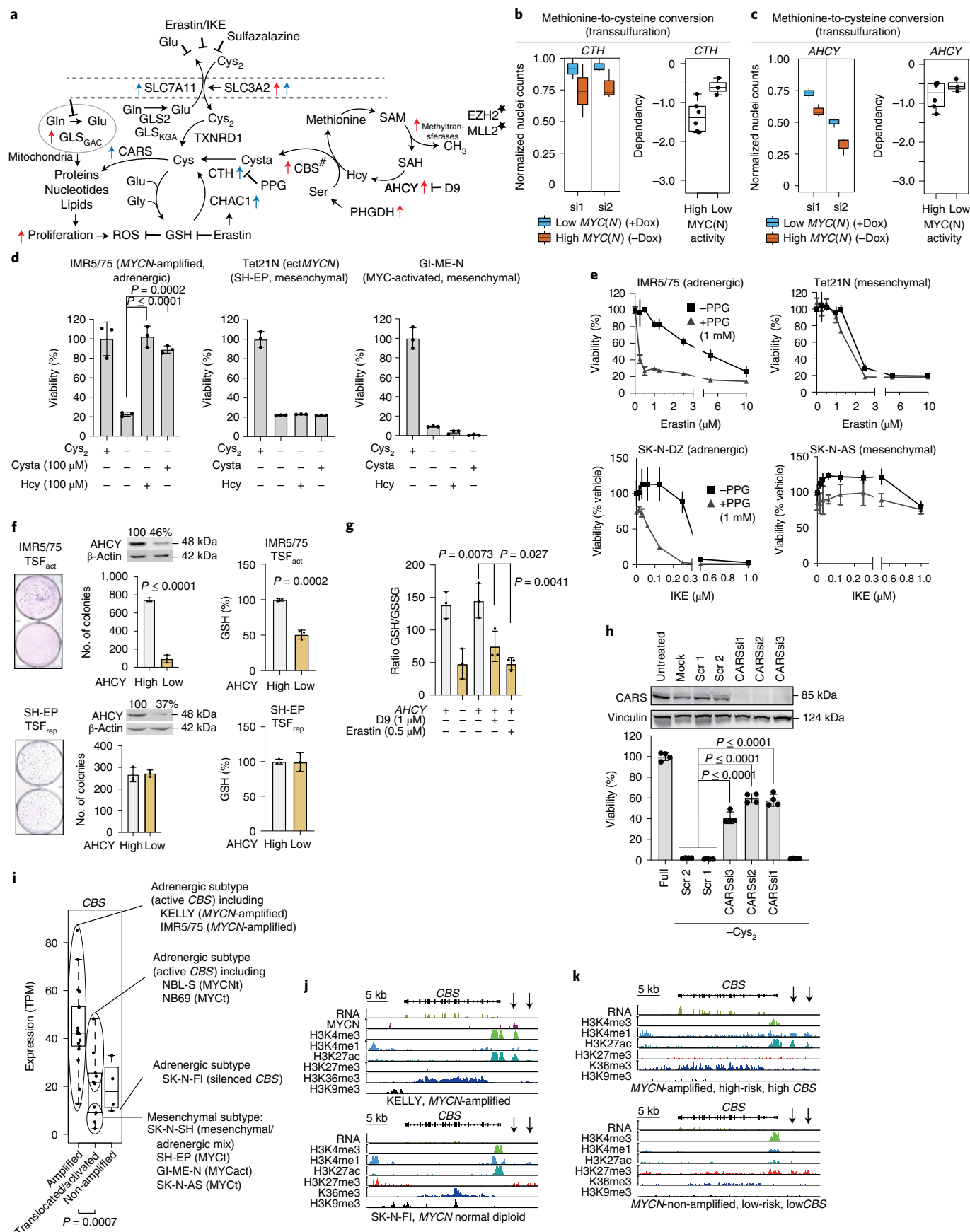
Next, we combined this protocol for reducing cellular cysteine with genetic targeting of GPX4 activity (Fig. 7b). We observed a robust effect with complete remission in most animals (Fig. 7d,e). Transcriptional profiling of residual small tumors revealed induction of ferroptosis markers after combined inhibition of cystine uptake/cysteine synthesis and GPX4 compared to tumors treated with vehicle control (Fig. 7f). These data provide strong in vivo evidence that concomitant reduction of cystine uptake, transsulfuration and GPX4 activity can be utilized as a new therapeutic strategy for high-risk, *MYCN*-amplified neuroblastomas.

Discussion

We demonstrated that oncogenic *MYCN* sensitizes neuroblastoma cells to ferroptosis when intracellular cysteine availability for GSH synthesis and the cystine/cysteine redox cycle are limited. A high *MYCN* state in neuroblastoma cells sensitizes them to lipid peroxidation, which in combination with acute intracellular cysteine reduction triggers massive ferroptotic cell death. Our study shows transsulfuration and GSH redox activity to be crucial to escape ferroptosis, whereas Gln import and glutaminolysis are required for ferroptosis in neuroblastoma cells with oncogenic *MYCN* (Fig. 7a). Expression levels of key genes for these processes are correlated with *MYCN* expression in high-risk tumors. The recently identified adrenergic and mesenchymal subtypes⁸ also appear to determine how cysteine is maintained in neuroblastoma cells, with transsulfuration only being activated by oncogenic MYC(N) in adrenergic cells, where *CBS* is accessible and not epigenetically repressed as in mesenchymal cells.

In this study, we describe metabolic rewiring in *MYCN*-amplified adrenergic neuroblastoma cells, where high consumption of cysteine used for the synthesis of cellular building blocks at the expense of GSH synthesis and ROS clearance creates a new *MYCN*-dependent liability. To prove that this liability can be exploited as a new therapeutic concept, we established and optimized an orthotopic model for neuroblastoma using intra-adrenal gland tumor cell transplantation. This model allowed robust in vivo testing of

Fig. 5 | MYCN activates transsulfuration genes controlling methionine-to-cysteine conversion in MYCN-amplified cells. **a**, Illustration of Cys₂ uptake/metabolism, glutaminolysis, methionine cycle and transsulfuration (#). The single star indicates the top hits from the MYCN siRNA screen; the red arrows indicate genes transcriptionally activated by MYCN in IMR5/75 (FDR = 0.001, using likelihood ratio testing); the blue arrows indicate genes transcriptionally activated by Cys₂ deprivation in IMR5/75 cells. **b,c**, siRNA screen in the IMR5/75 cell line and MYCN synthetic lethality with knockdown of key transsulfuration genes (two individual siRNAs) and MYC(N) activity correlation with *CTH* (**b**) and *AHCY* (**c**). **d**, Relative viability of IMR5/75, Tet21N and GI-ME-N cells after Cys₂ deprivation for 48 h in the presence or absence of Hcy or Cysta. Data represent the mean ± s.e.m.; n = 3 samples. The experiment was replicated three times. **e**, Relative viability (survival of compound-treated cells divided by survival of vehicle-treated cells) of IMR5/75, SK-N-DZ and MYCN-expressing mesenchymal Tet21N and MYC-expressing SK-N-AS cells after cotreatment with PPG and erastin or IKE. Data represent the mean ± s.e.m.; n = 3 samples. The experiment was replicated three times. **f,g**, Colony formation, GSH synthesis (**f**) and reduced GSH/GSSG ratio (**g**) in IMR5/75 and SH-EP cells on *AHCY* inhibition (by knockdown or using small molecule D9). Representative western blots are shown in the figure. Data represent the mean ± s.e.m.; n = 3 samples. The experiment was replicated three times. **h**, Relative viability of IMR5/75 cells after siRNA-mediated *CARS* knockdown (72 h) and Cys₂ deprivation (24 h). Representative western blots are shown in the figure. Data represent the mean ± s.e.m.; n = 4 samples. The experiment was replicated three times. **i**, *CBS* expression in 32 neuroblastoma cell lines (RNA-seq) categorized using *MYCN*-amplified, *MYCN*/*MYC*-translocated/activated cell lines and *MYC(N)* non-expressors. MYCNt/*MYC*t, MYCN or MYC translocation; MYCact, activated MYC due to unknown mechanism. **j,k**, Bimodal *CBS* expression for *MYCN*/*MYC*-translocated/activated cell lines along with adrenergic and mesenchymal cell type mRNA expression (RNA-seq) and input-normalized read counts of histone modifications and MYCN (ChIP-seq) at the *CBS* locus in representative *MYCN*-amplified and *MYCN*-non-amplified cells (**j**) and tumors (**k**). Box plots: the center line indicates the median value, the lower and upper hinges represent the 25th and 75th percentiles, respectively and the whiskers denote 1.5× the IQR. Each dot corresponds to one sample; Wilcoxon rank-sum test; P values as indicated.



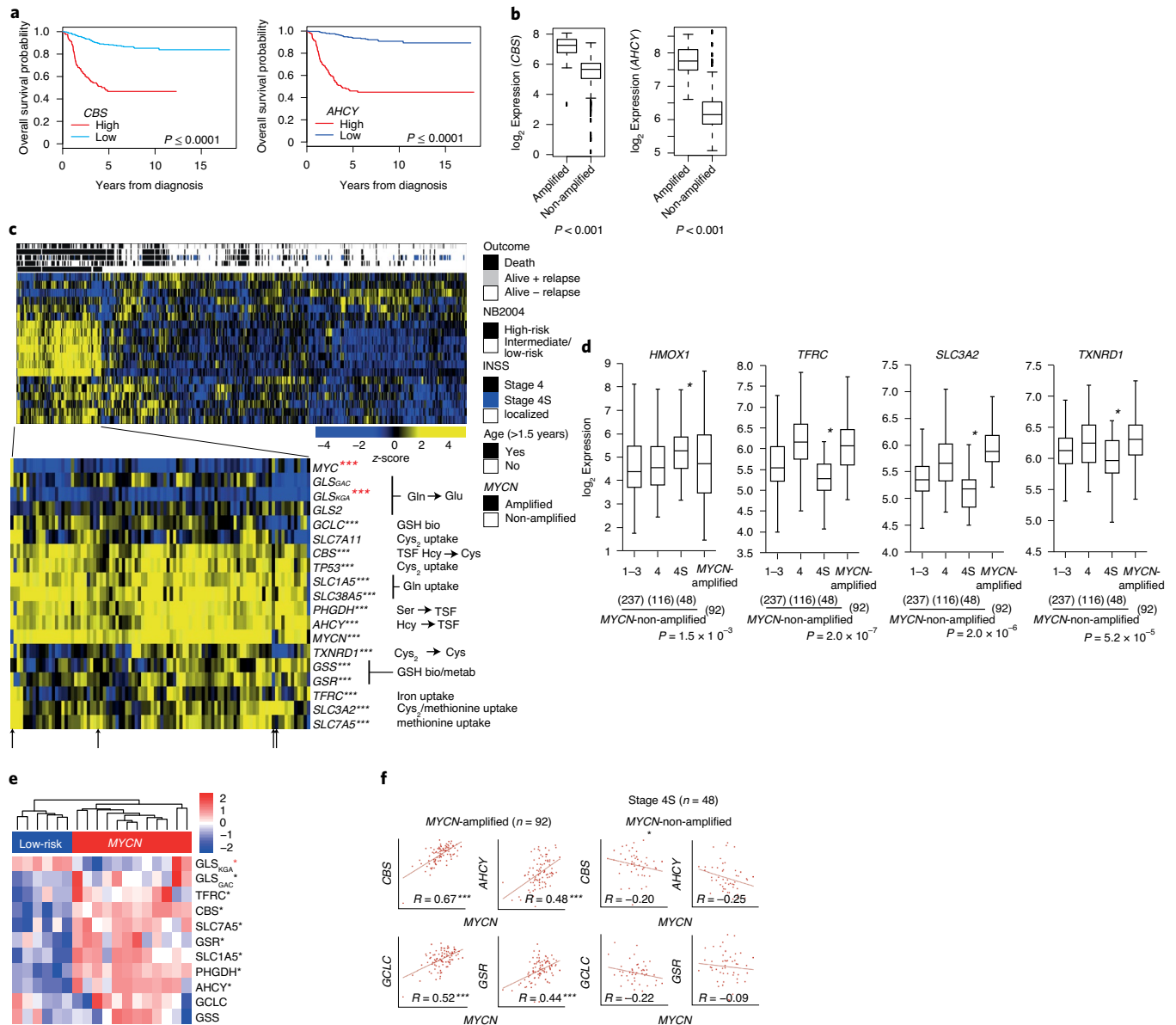


Fig. 6 | Gene and protein markers indicating ferroptosis sensitivity in high- and low-risk neuroblastomas. **a**, Kaplan-Meier overall survival estimates for subgroups defined by CBS expression, high CBS ($n=123$) and low CBS ($n=375$). Kaplan-Meier overall survival estimates for subgroups defined by AHCY expression. The cutoff values for the dichotomization of AHCY expression were estimated by maximally selected log-rank statistics, high AHCY expression ($n=165$) and low AHCY expression ($n=333$). **b**, CBS and AHCY expression in MYCN status-dependent expression in 498 primary neuroblastomas (92 MYCN-amplified, 406 MYCN non-amplified tumors, RNA-seq); Wilcoxon rank-sum test. **c**, Gene expression heatmap of subgroups hierarchically clustered with average linkage and non-centralized correlation distance function. Row: transcript; column: sample, according to the NB2004 study. *** $P < 0.001$ (black) higher and **** $P < 0.001$ (red) lower gene expression. Wilcoxon rank-sum test; the arrows mark MYCN-non-amplified tumors. **d**, Differential expression of cysteine uptake and iron-regulating genes in primary neuroblastomas dependent on the stage of the disease and MYCN amplification status. Differential expression in stage 4S versus all other neuroblastoma subtypes tested by Wilcoxon rank-sum test. **e**, Differential expression of proteins mediating GSH synthesis/metabolism and GSH-relevant amino acid uptake/metabolism compared between MYCN-amplified and low-risk, normal MYCN tumors. * $P \leq 0.01$; higher (black) and lower (red) protein expression. P values were calculated using a two-sided Welch's t -test. Exact P values are given in the source data. Intensities are given as z-scores. See the legend for color-coding. **f**, Pearson's correlation analysis of MYCN expression and GSH synthesis/metabolism and transsulfuration genes in MYCN-amplified ($n=92$) and MYCN-non-amplified, low-risk stage 4S ($n=48$) tumors. Box plots: the center line indicates the median value, the lower and upper hinges represent the 25th and 75th percentiles, respectively and the whiskers denote the 1.5x the IQR. Each dot corresponds to one sample; Wilcoxon rank-sum test; P values as indicated.

ferroptosis induction: we obtained remarkable tumor remission in high MYCN neuroblastoma by combining inhibition of (1) cystine import using IKE and (2) transsulfuration using PPG together with (3) CRISPR-Cas9-mediated GPX4 deletion. Among the different parameters tested, this was the most effective strategy

revealing almost complete tumor remission. Currently there are no GPX4 inhibitors for in vivo use, hence future improvements of this therapeutic strategy would involve the development of potent GPX4 inhibitors with optimal pharmacokinetics and pharmacodynamics.

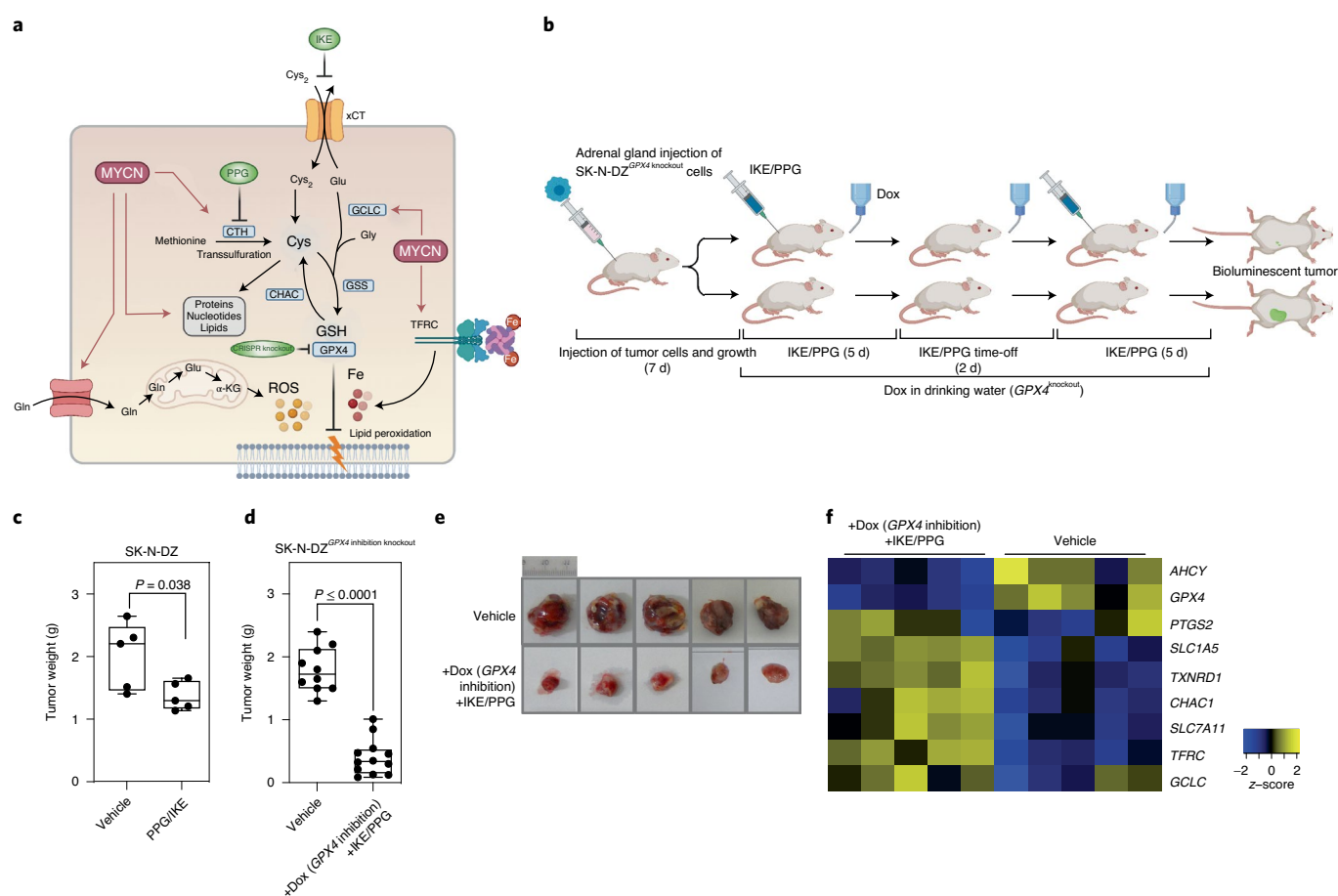


Fig. 7 | Triple combination inhibiting GPX4, cystine uptake and transsulfuration eliminates MYCN-amplified tumors in vivo. **a**, Regulation of pro- and anti-ferroptotic players by oncogenic MYCN in MYCN-amplified neuroblastoma cells that may trigger ferroptosis when both cysteine and GSH availability is limited. Therapeutic intervention points are indicated in green. **b**, Dosage scheme applied with either double (IKE/PPG) or triple combination (GPX4 knockout, IKE and PPG). **c,d**, Tumor weight after combination treatment with IKE and PPG ($n=5$ mice per group; **c**) and the triple combination of GPX4 reduction plus IKE/PPG treatment ($n=9$ in the control group and $n=12$ mice in the treated group; **d**). **e**, Representative photographs of fully grown tumors in the vehicle group versus residual tumors after triple combination. **f**, Transcriptional changes of ferroptosis markers after GPX4 reduction plus IKE/PPG in residual tumor tissue ($n=5$ tumor samples from each group). Statistical analysis was performed using a one-tailed Student's *t*-test for the in vivo experiments. Box plots: the center line indicates the median value and the lower and upper hinges represent the minimum and maximum points. Each dot corresponds to one sample; *P* values as indicated. Panels **a** and **b** created with BioRender.

Recent studies^{11,12} highlighted the role of iron in MYCN-dependent neuroblastoma and ferroptosis. The *TFRC* gene, involved in iron uptake, is activated by MYC in several cell types enhancing cellular proliferation⁴². In line with these studies, we showed a synergistic effect of iron in drug-induced ferroptosis in MYCN-amplified cells. In addition, we demonstrated that the iron chelator CPX rescues ferroptotic cell death under cystine deprivation conditions (Extended Data Fig. 1h). We further observed high expression of *TFRC* in MYCN-amplified cell lines and tumors (Fig. 6c and Extended Data Figs. 3j and 4a) reflected on the protein level (Fig. 6e). Similarly, we observed *HMOX1* upregulation, involved in increasing the labile iron pool, in cystine-deprived neuroblastoma cells before ferroptosis induction. Together, this highlights a prominent role for iron metabolism in promoting ferroptosis in neuroblastoma cells.

We have described a detailed mechanism that explains how cystine deprivation triggers ferroptosis in MYCN-amplified tumors, which depends on the (epi)genetic context of adrenergic or mesenchymal subtypes, the later having transsulfuration silenced. These results differ from Floros et al.¹¹, who suggested that high MYCN mediates upregulation of *SLC7A11* in neuroblastoma cells. In contrast, our results indicate that in most, if not all, adrenergic

neuroblastoma cells (>95% of neuroblastomas) high MYCN fails to boost cystine uptake via x_c^- . Instead, MYCN strongly activates transsulfuration, the intracellular conversion of methionine to cysteine. In general, neuroblastoma cells have very low *SLC7A11* expression compared to other cancer types, which may explain the remarkable sensitivity of neuroblastoma cells to ferroptosis stimuli. Our results demonstrate that the uptake of cystine via x_c^- is not regulated on tumor progression in adrenergic MYCN-amplified cells. Upregulation of *SLC7A11* by MYC oncoproteins was only found in cells of the rare mesenchymal subtype, having epigenetically silenced transsulfuration. Therefore, upregulation of *SLC7A11* may become relevant in the relapse scenario, which is associated with elevated proportions of mesenchymal cells.

Intriguingly, our data may also explain spontaneous neuroblastoma regression, the mechanisms of which have remained elusive. In low-risk, metastatic neuroblastomas (stage 4S), which are likely to regress spontaneously⁵, we identified multiple markers predisposing them to ferroptosis (Fig. 7a), including: (1) failure to upregulate enzymes involved in GSH biosynthesis (*GCLC*) and redox activity (*GSR*) and transsulfuration (*AHCY*, *CBS*) with increasing MYCN activity (Fig. 6f); (2) *CBS* silencing by intragenic CpG methylation

(Extended Data Fig. 6a); (3) *HMOX1* upregulation to increase free iron pools; and (4) *SLC3A2* and *TXNRD1* downregulation (Fig. 6d). Spontaneous regression may be the physiological resolution of this cellular state sensitive to ferroptosis. Unlike low-risk tumors, *MYCN*-amplified neuroblastomas appear to metabolically adapt to survive events that deplete intracellular cysteine, such as high systemic glutamate present in the first two years of life⁴³. The cysteine requirement of cancers dependent on oncogenic *MYCN* activity creates a previously unknown Achilles' heel that could be exploited to selectively induce ferroptosis for treatment. Our findings identify cysteine uptake, transsulfuration and the lipid peroxidation-specific scavenging system as vulnerabilities in cancer cells driven by oncogenic *MYC(N)* activity such as *MYCN*-amplified neuroblastomas.

Methods

Ethics. All patients with neuroblastoma were enrolled in the German Neuroblastoma Trial (NB97, NB2004, NB 2016) and approved by the ethics committee of the University of Cologne. All studies involving mice and experimental protocols were conducted in compliance with German Cancer Research Center guidelines and approved by the governmental review board of the state of Baden-Württemberg.

Experimental in vitro procedures. Cell culture. Human neuroblastoma cells (IMR5/75, KELLY, SiMa, NBL-S, SK-N-FI, SH-SY5Y, NB69, SK-N-DZ, SH-EP, GI-ME-N) were cultivated at 37 °C with 5% CO₂ in Roswell Park Memorial Institute (RPMI) 1640 medium (Gibco) supplemented with 10% fetal calf dialyzed serum (Gibco) and penicillin/streptomycin (AppliChem). The KELLY and SiMa cell lines were purchased from Deutsche Sammlung von Mikroorganismen und Zellkulturen. SK-N-FI cells were purchased from ATCC. NBL-S and Tet21N (SH-EP) cells were provided by G.M. Brodeur and W. Lutz, respectively. Tunable cell lines, IMR5/75 *MYCN* short hairpin RNA (shRNA) and SH-EP *MYCN* transgene (Tet21N) cells were generated and cultured as described previously^{15,17}. Cell line identity/unique single-nucleotide polymorphism profiles were confirmed by the Multiplexion Multiplex Cell Authentication service as described recently⁴⁴. The purity of cell lines was validated using the Multiplex Cell Contamination Test (Multiplexion) as described recently⁴⁴. No *Mycoplasma*, squirrel monkey retrovirus or interspecies contamination was detected. To assess the effects of amino acid deprivation, cells were cultivated using modified amino acid-free DMEM powder (PAN-Biotech) supplemented with individual amino acids (Sigma-Aldrich) as indicated, at final concentrations used in standard DMEM. For the 3D culture experiments, we used the hanging drop method. Twenty thousand cells were placed in hanging drop culture and incubated under physiological conditions until they form 3D spheroids.

Analysis of cell viability and proliferation. The impact of various treatments on cellular proliferation/viability was assessed using a sulforhodamine B (SRB)⁴⁵ or CellTiter-Blue (Promega Corporation) cell assay. To determine changes in cellular proliferation, approximately 2 × 10⁴ cells were seeded per well (48-well format for the SRB assay and 96-well format for the CellTiter-Blue assay) in full medium. After 24 h, cells were washed with PBS, fed with the chosen medium and treated as indicated. Cell viability was analyzed in full or Cys₂-free medium cotreated with 10058-F4 (30 μM, catalog no. F3680; Sigma-Aldrich), Z-VAD-FMK (30 μM; catalog no. sc-3067; Santa Cruz Biotechnology), bafilomycin A1 (30 nM, catalog no. sc-201550; Santa Cruz Biotechnology), necrostatin-1 (2 μM, catalog no. N9037; Sigma-Aldrich), Fer-1 (5 μM, catalog no. SML0583; Sigma-Aldrich), Trolox (100 μM, catalog no. 238813; Sigma-Aldrich), aminooxyacetate (500 μM, Sigma-Aldrich), dimethylloxalylglycine, N-(methoxyoxoacetyl)-glycine methyl ester (5 mM, Sigma-Aldrich), CPX (1 μM, catalog no. sc-204688; Santa Cruz Biotechnology), D9 (synthesized and provided by Bayer Pharma AG), erastin (Cay17754; Biomol), IKE (MedChemExpress), 2-mercaptoethanol (50 μM; Sigma-Aldrich), glutaminase inhibitors BPTES (catalog no. SML0601; Sigma-Aldrich) and compound 968 (catalog no. 352010; Merck Millipore), RSL3 (MedchemExpress), sulfasalazine (catalog no. 599-79-1; Sigma-Aldrich), PPG (1 mM, catalog no. P7888; Sigma-Aldrich). DNA content analysis was performed by fixing cells with 4% paraformaldehyde and staining with FxCycle Violet Stain (Thermo Fisher Scientific) followed by fluorescence-activated cell sorting (FACS) using a MACSQuant Flow Cytometer (Miltenyi Biotec). To assess the effects of Cys₂ deprivation on cell viability, cells were washed and fed with Cys₂-free DMEM supplemented with 10% dialyzed FCS, 200 μM L-methionine (catalog no. 63-68-3; Sigma-Aldrich) and 4 mM Gln (catalog no. 25030081; Gibco) 48 h after seeding. To determine the cell death rescue potential, Cys₂-deprived cells were also cotreated with Hcy (catalog no. 69453; Sigma-Aldrich), Cysta (catalog no. C7505; Sigma-Aldrich) or GSH (2 mM, catalog no. G4251; Sigma-Aldrich). We further compared effects on cell viability after Cys₂ and Gln depletion or Gln excess using the following conditions: full RPMI (208 μM Cys₂, 2.055 mM Gln, 0.136 μM Glu); low Cys₂ (5 μM); low Gln (0.2 mM); and high Gln (2.5 mM). Fluorescence was read (540/580 nm) 24 h and 48 h after deprivation/treatment. Doubling times

were calculated using two different methods: (1) impedance measurements were assessed using the RTCA system (Roche) by seeding at different cell densities and registering impedance signals every 20 min; (2) standard growth curves were generated by counting cells by FACS at the indicated time points and excluding propidium iodide-positive cells as necrotic. In both methods, exponential curves were fitted and doubling times calculated.

Inducible stable AHCY knockdown cell cultures. Stable IMR5/75 and SH-EP cell lines expressing shRNA against AHCY under control of the Tet repressor were generated stepwise as described previously¹⁵ using the following oligonucleotide sequence to target AHCY: forward: gatccccGGATCACTACCGCTACTGAttaaga-gaTCAGTAGCGGTAGTGATCctttttgaaa; reverse: gcttttccaaaaaGGATCACTACC GCTACTGAtctcttgaaTCAGTAGCGGTAGTGATCCggg.

IMR5/75 (5,000 cells per well) and SH-EP-AHCYsh (1,000 cells per well) were seeded in 6-well plates and simultaneously treated with Dox (1 μg ml⁻¹) to induce the AHCY-targeting shRNA. Cells were fixed (11% glutaraldehyde; Sigma-Aldrich) and Giemsa-stained five (SH-EP-AHCYsh) or 7 d (IMR5/75-AHCYsh) later. Colony counting was performed using a GelDoc Documentation System and Quantity One software version 4.6.6 (Bio-Rad Laboratories) and quantification using Microsoft Excel 2016.

Quantification of amino acids. Pellets of 2 × 10⁶ cells were extracted with 0.1 ml ice-cold 0.1 M HCl. Non-thiol-containing amino acids were quantified after specific fluorescent labeling with AccQ-TagTM (Waters) as described previously⁴⁶. Cysteine and GSH levels were determined after labeling with monobromobimane (Calbiochem) as described previously⁴⁷.

Western blot analysis of proteins in cell extracts. Whole cell lysates were prepared and protein expression was visualized as described previously⁴⁸. Protein lysates (20 μg per lane) were separated on 12.5% SDS-polyacrylamide gel electrophoresis. Blots were probed with antibodies directed against MYCN (1:1,000 dilution, catalog no. sc-53993; Santa Cruz Biotechnology), c-MYC (1:1,000 dilution, catalog no. ab32072; Abcam), CTH (1:1,000 dilution, catalog no. ab54573; Abcam), SAHH (A-11) (AHCY antibody) (1:1,000 dilution, catalog no. sc-271389; Santa Cruz Biotechnology), GPX4 (1:1,000 dilution, catalog no. ab41787; Abcam), CARS (1:1,000 dilution, catalog no. ab126714; Abcam), glutaminase 1 (1:40,000 dilution, catalog no. ab156876; Abcam), vinculin (1:1,000 dilution, catalog no. sc-73614; Santa Cruz Biotechnology) or horseradish peroxidase-conjugated anti-β-actin (1:5,000, catalog no. ab20272; Abcam). Peroxidase-AffiniPure goat anti-mouse IgG (H+L) (1:1,000 dilution, catalog no. 115-035-003; Dianova) or peroxidase AffiniPure goat anti-rabbit IgG (H+L) (1:1,000 dilution, catalog no. 111-035-144; Dianova) antibodies were used as secondary antibodies. Proteins were visualized using enhanced chemiluminescence detection reagents (GE Healthcare) and a chemiluminescence reader (VILBER). Protein quantification was performed with ImageJ (<https://imagej.net>).

Flow cytometry. Analysis of intracellular ROS levels and lipid peroxidation. Low MYCN populations were established by incubating cells with 1 μg ml⁻¹ Dox at least 48 h before further treatment. Cells were then fed either with full or cystine-free medium and cotreated with Fer-1 (5 μM), liproxstatin-1 (Lip-1) (1 μM; catalog no. SML1414; Sigma-Aldrich), CPX (1 μM), Trolox (100 μM) or GSH (2 mM, catalog no. G4251; Sigma-Aldrich) for 20 h. Lipid peroxidation was analyzed with the C11-BODIPY BD FACSARIA III cell sorter. Total intracellular ROS levels were determined using CellROX (Thermo Fisher Scientific). Gating strategy is shown in Extended Data Fig. 7.

MYCN synthetic lethal screen. Large-scale druggable genome siRNA screen. For high-throughput screening, a Silencer Select siRNA custom library (catalog no. 4404034; Ambion) was used encompassing 31,242 unpooled siRNAs targeting 10,414 genes (3 siRNAs per gene). Lipofectamine RNAiMax Transfection Reagent (Thermo Fisher Scientific) only and ON-TARGETplus Non-targeting siRNA no. 1 (Dharmacon) served as negative transfection controls; PLK1 (Silencer Select siRNA no. 1; Ambion) served as positive control. Liquid reverse transfection was performed in 384-well plates (2,100 cells per well) using a Freedom EVO 200 robotic platform. Two treatment conditions were screened in triplicate: (1) culture medium only (IMR5/75 high MYCN); and (2) plus Dox (1 μg ml⁻¹ final concentration) to induce the shRNA targeting MYCN (IMR5/75 low MYCN); 96 h after transfection, cells were fixed with 11% glutaraldehyde and subsequently Hoechst-stained (10 mg ml⁻¹ stock in 1 × PBS, 1:2,500; Invitrogen). The number of Hoechst-positive cell nuclei was determined using an OPERA fluorescence microscope based on nine sites per well and a BHC in-house program. We applied redundant siRNA activity (RSA) to the ratio between high and low MYCN data to select top-ranked hits and a false discovery rate (FDR) of 0.2.

Transient siRNA-mediated gene knockdown. IMR5/75 cells were seeded in 96-well plates (3,000 cells per well); 24 h later, they were transiently transfected with a mix of RNAiMax (0.04 μl per well; Thermo Fisher Scientific) and 0.01 μM per well of siRNA according to the manufacturer's instructions. siRNA sequences are listed in Supplementary Table 2.

Generation of inducible Cas9 neuroblastoma cell lines. *Molecular cloning.* The CRISPR–Cas9-mediated inducible knockout experiments were performed using the lentiviral pCW-Cas9-EGFP plasmid. The plasmid was obtained as follows: the lentiviral pCW-Cas9 plasmid (plasmid no. 50661; Addgene) was subcloned by cutting it with the restriction enzymes HincII and XbaI (New England Biolabs). Next, a gBlock DNA fragment (Integrated DNA Technologies) encoding enhanced green fluorescent protein (eGFP) with complementary overhangs was cloned into the plasmid backbone thus replacing the puromycin resistance gene and generating the pCW-Cas9-EGFP plasmid. Guide RNAs (gRNAs) for the CRISPR knockout experiments were designed with Benchling (<https://www.benchling.com>) using the human reference genome GRCh38. Oligonucleotides were ordered (Sigma-Aldrich) with complementary overhangs to the lentiviral delivery plasmid backbone pLKO5.sgRNA.EFS.tRFP6572 and listed in Supplementary Table 2.

Lentivirus production. Lentivirus production was performed using a second-generation lentiviral system and a Calcium Phosphate Transfection Kit (Invitrogen) in HEK 293T cells. Briefly, early passaged HEK 293T cells were cotransfected with the lentiviral transfer plasmid, a packaging plasmid (psPAX2; plasmid no. 12260; Addgene), and with a plasmid coding for the VSV-G envelope (pMD2.G; plasmid no. 12259; Addgene). All experimental procedures for lentivirus production were performed in a biosafety level 2 laboratory.

The SK-N-DZ cell line was previously transduced with lentiviral particles carrying the luciferase reporter pLX-puroR-Luc at a multiplicity of infection (MOI) of 0.3. After recovery, resistant cells were transduced with pCW-Cas9-EGFP lentiviral particles at an MOI of 0.3. Polyclonal cell lines were maintained in DMEM supplemented with 10% tetracycline-free FCS (Clontech Laboratories). Next, cells expressing eGFP and the luciferase reporter were transduced with lentiviral particles carrying the gRNA targeting *GPX4* at an MOI of 0.3. To generate monoclonal CRISPR cell lines, cells were individualized based on eGFP and RFP657 expression using FACS. *In vitro* Cas9 expression was induced supplementing the culture medium with $1 \mu\text{g ml}^{-1}$ Dox (Sigma-Aldrich).

In vivo orthotopic mouse experiments. All studies involving mice and experimental protocols were conducted in compliance with German Cancer Research Center guidelines and approved by the governmental review board of the state of Baden-Württemberg, Karlsruhe District Council, under authorization no. G-176/19, according to German legal regulations. The mouse strains used in the study were NOD.Cg-Prkdc^{scid}Il2rgtm1^{WJ}/SzJ (stock no. 005557; The Jackson Laboratory). Female mice (3–4 months old) were used for the experiments. Mice were housed in individually ventilated cages under temperature and humidity control. Cages contained an enriched environment with bedding material. To generate orthotopic mouse models for neuroblastoma, 2×10^5 SK-N-DZ cells were transplanted into the right adrenal gland after the surgical site was prepared⁴⁹. Cells were resuspended in a 1:1 (vol/vol) mix of growth factor-reduced Matrigel (Corning) and PBS. Overall, 20 μl of this cell suspension was injected into the right adrenal gland of anesthetized mice. After tumor cell transplantation, we monitored mice for evidence of tumor development by bioluminescent signal using an IVIS Spectrum Xenogen device (Caliper Life Sciences). We observed a clear signal from the tumors 1 week after the injection of 2×10^5 SK-N-DZ cells. IKE and PPG were used at a concentration of 45 mg kg^{-1} through intraperitoneal injection. Animal health was monitored daily and mice were euthanized as soon as they reached the termination criteria defined in the procedure. Sample size was calculated with the help of a biostatistician using R v3.4.0. Assumptions for the power analysis were as follows: Alpha error, 5%; Beta error, 20%. Mice were randomized into treatment groups before treatment. In case animals had to be euthanized before the predefined end point (due to weight loss or other termination criteria), they were excluded from any downstream analyses. All animal experiments (apart from animal treatment) were blinded during the experiments and follow-up assessment.

Transcript profiling of tumors and cell lines. *Microarray analysis.* RNA isolated from the remaining tumors (five vehicle-treated versus five triple combination of *GPX4* knockout + IKE + PPG) was analyzed using the Affymetrix GeneChip Clariom S Assay. Microarray data (GSE192976) were processed using a modification of a pipeline described previously⁵⁰. Briefly, raw CEL files were robust multichip average-normalized and two-group comparisons were performed using the limma package v.3.46 (Bioconductor) with an empirical Bayes test for differential expression. An FDR < 0.1 was regarded as statistically significant.

RNA sequencing. Total RNA was isolated using the miRNeasy Mini Kit (QIAGEN) according to the manufacturer's protocols. RNA libraries were prepared using the NEBNext Ultra Directional RNA Library Prep Kit for Illumina (New England Biolabs). Raw RNA sequencing (RNA-seq) sequences of neuroblastoma cell lines and model systems were mapped to the UCSC hg19 genome using STAR⁵¹ v.2.5.3a with default parameters. RNA-seq sequences from the IMR5/75 high and low *MYCN* cells were processed as described previously³⁷. Differentially expressed genes were identified using the edgeR v.3.20.9 generalized linear model (glmFit) approach in R v3.4.3. The expression profiles of the high *MYCN* depletion/rescue experiments were visualized in R v4.3.0 using the gplots v.3.1.1 function heatmap.2, after filtering out lowly expressed genes (count per million ≤ 1) and normalizing for library size using trimmed mean of *M*-values normalization as implemented in

edgeR⁵² v.3.20.9. RNA-seq expression profiles from 498 primary neuroblastomas³⁸ (GSE49711) were analyzed. The Wilcoxon rank-sum test was used to test the association between candidate gene expression and amplified *MYCN* oncogene. Maximally selected log-rank statistics were used to describe the relationship between patient survival and candidate gene expression; the resulting expression cutoff points were used for dichotomization. Survival curves were estimated using the Kaplan–Meier method. Pearson correlation coefficients were calculated to estimate linear dependence between the expression values of candidate genes.

Quantitative PCR with reverse transcription. Complementary DNA was synthesized using the SuperScript IV Reverse Transcriptase Kit (catalog no. 18090-200; Invitrogen) according to the manufacturer's instructions. Quantitative PCR with reverse transcription (RT-qPCR) was performed for the genes of interest and two housekeeping genes using the Platinum SYBR Green qPCR Supermix-UDG kit (catalog nos. 11733038/11733-046; Invitrogen). Primers are listed in Supplementary Table 2.

Epigenetic characterization of tumors and cell lines. *Chromatin immunoprecipitation followed by sequencing analysis of histone modifications in neuroblastoma primary tumors and cell lines.* Formaldehyde cross-linking of cells, cell lysis, sonication, chromatin immunoprecipitation and library preparation were performed as described previously⁵³, starting with approximately 4×10^6 cells (1×10^6 cells per individual immunoprecipitation). Direct cell lysis for each sample was achieved by 30 min incubation on ice in 950 μl radioimmunoprecipitation assay I buffer using approximately 30 mg of fresh-frozen tumor tissue per individual chromatin immunoprecipitation followed by sequencing (ChIP-seq) experiment. Library preparation was performed using the NEBNext Ultra DNA Library Prep Kit (New England Biolabs) according to the manufacturer's protocol. Samples were mixed in equal molar ratios and sequenced on an Illumina sequencing platform.

ChIPmentation of MYCN transcription factor in neuroblastoma cell lines.

Formaldehyde cross-linking, cell lysis, sonication and chromatin immunoprecipitation were performed as described previously⁶, adding the ChIPmentation module by Schmidl et al.⁵⁴ with the following changes: a Bioruptor Plus with automated cooling (4 °C) was used for high-intensity sonication (20–30 min each with 30 s on and 30 s off intervals) and 10 μg MYCN antibody and 10^6 cells for ChIP. The tagmentation reaction (Illumina Nextera DNA library Prep Kit) was performed at 37 °C for 1 min with the bead-bound chromatin sample or 5 ng purified input DNA for normalization. After de-cross-linking, purified samples were amplified using the dual index barcodes of the Nextera Index Kit and 13 PCR cycles. Enriched libraries were purified and pooled. ChIPmentation libraries were sequenced (50 single-end bases) on the Illumina sequencing platform (German Cancer Research Center Core Facility).

Data analysis of ChIP-seq and ChIPmentation. Single-end reads were aligned to the hg19 genome using Bowtie2 v.2.3.0, keeping uniquely aligned reads only. The BAM files of aligned reads were further processed using the deepTools suite v.3.0 (ref. 55). Input files were subtracted from the treatment files using the bamCompare tool v.3.0, applying the simple exponential smoothing method to normalize signal to noise. The resulting signals were normalized to an average 1 \times coverage to produce signal (bigWig) files. Peaks were called using the MACS2 v.2.1 tool using default parameters. Data are available at the Gene Expression Omnibus (GEO) under accession no. GSE189174.

DNA methylation analysis. DNA methylation and gene expression data from 105 primary neuroblastomas assessed by Infinium HumanMethylation450 BeadChips and 44K Agilent oligonucleotide microarrays³⁶ (GEO accession no. GSE73518) were analyzed for candidate loci. The R2: Genomics Analysis and Visualization Platform (<http://r2.amc.nl>) was used to visualize the expression/methylation of selected gene–CpG pairs.

Tumor proteome analysis. Tumor proteome data were generated as part of a previous study⁷ and was reanalyzed for this study. Briefly, tumor samples were lysed in SDS, homogenized, split into replicates, reduced, alkylated and purified by Wessel–Flügge precipitation. Samples were then digested by LysC and trypsin and fractionated by strong cation exchange before being measured by reversed phase liquid chromatography–mass spectrometry (LC–MS) on Q Exactive Plus instruments (Thermo Fisher Scientific). Proteins quantified in less than 50% of *MYCN* high- or low-risk cases were excluded. Data were imputed by random draw from a normal distribution with default parameters: 0.3 width and 1.8 downshift. A two-sided Welch's *t*-test was used to calculate the *P* values for the differential protein expression analysis; multiple testing correction by the Benjamini–Hochberg method was applied.

Statistics and reproducibility. No statistical method was used to predetermine sample size. No data were explicitly excluded from the analyses unless they were of poor quality as determined by standard sequencing quality control metrics. The experiments were not randomized and the investigators were not blinded to allocation during the experiments and outcome assessment. Data are presented as the mean \pm s.e.m. Statistical analyses were performed using Prism 7

(GraphPad Software). A two-tailed unpaired or paired Student's *t*-test was used for comparisons between two groups. For the animal experiments, sample size was calculated with the help of a biostatistician using R v.3.4.0. Assumptions for the power analysis were as follows: Alpha error, 5%; Beta error, 20%. Mice were randomized into treatment groups before treatment. In case animals had to be euthanized before the predefined end point (due to weight loss or other termination criteria), they were excluded from any downstream analyses. All animal experiments (apart from animal treatment) were blinded during the experiments and follow-up assessment.

Reporting Summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

Proteome data of neuroblastoma tumors were previously published by Hartlieb et al.⁷. All data were deposited with the European Genome-phenome Archive (as dataset no. EGAD00001006737) as part of study no. EGAS00001004349. Data are available upon request by contacting F.W. The RNA-seq data of 498 primary neuroblastoma patients were previously published³⁸ and are available at the GEO under the accession no. GSE49711. The DNA methylation data of primary neuroblastoma tumors were previously published³⁸ and are available at the GEO under the accession no. GSE73518. Time course RNA-seq profiling of IMR5/75 high MYCN and low MYCN cells was previously published¹¹ and can be accessed at the GEO under the accession no. GSE97774. ChIP-seq data were deposited at the GEO under the accession no. GSE189174. The aligned BAM files of the RNA expression profiles of the high MYCN depletion/rescue experiments were submitted to the European Nucleotide Archive and can be found under accession no. PRJEB25184. The data of the MYCN synthetic lethal siRNA screen has been added to this article as supplementary data. The microarray data from tumor samples were deposited with the GEO under accession no. GSE192976. The remaining data are available within the article, supplementary information or from the corresponding authors upon reasonable request. Source data are provided with this paper.

Code availability

The custom scripts used to analyze the RNA-seq, microarray and ChIP-seq data are available upon request.

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Author contributions

A.F.F. and H.A. found the link between MYCN and ferroptosis. A.F.F., H.A. and F.W. designed the experiments. The metabolic assays and viability studies were conducted and analyzed by H.A., A.F.F., L.M.B., M. Shaikhkarami, M.L.-P. E.M.W., J.K., K.G. and S.K. U.Y., H.A. and R.W. generated/validated the CRISPR knockout cell lines. H.A., A.F.F. and A.T. designed and analyzed the *in vivo* experiments. H.A. performed the *in vivo* orthotopic experiments with support from C.K., J.M. and P.Z. H.A. and A.F.F. performed the flow cytometry assays. S.K. performed the MYCN synthetic lethal siRNA screen. M.G. helped with the MYCN synthetic lethal screen optimization. C. Shao, S.A.S., A.F.F., D.I.O., N.I., M. Schlesner and C.H. assisted with the bioinformatic analyses. M.G. and F.P. performed and analyzed the ChIP-seq and ChIPmentation experiments. M.N.-H., M.Z., S.A.S. and M. Selbach provided the tumor proteomic data. E.B., M.G., E.M.W. and J.K. conducted the RNA-seq experiments. RT-qPCR was performed by E.M.W. Regulatable cell models were established and colony formation assays performed by J.K. The western blot was performed by S.K., E.M.W. and J.K. G.P., J.B. and M.B. implemented and analyzed the LC-MS measurements. K.-O.H. contributed to the primary tumor RNA expression and DNA methylation analyses. B.N., C. Stresemann, J.H.R., M.F., I.A. and S.W. contributed reagents, materials and analysis tools. The manuscript was prepared by F.W., T.H., A.T., H.A., A.F.F., L.M.B. and S.K. with support from K.-O.H. H.A., A.F.F., S.K. and L.M.B. contributed equally to the study. All authors approved the final version of the manuscript.

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Competing interests

The authors declare no competing interests.

Additional information

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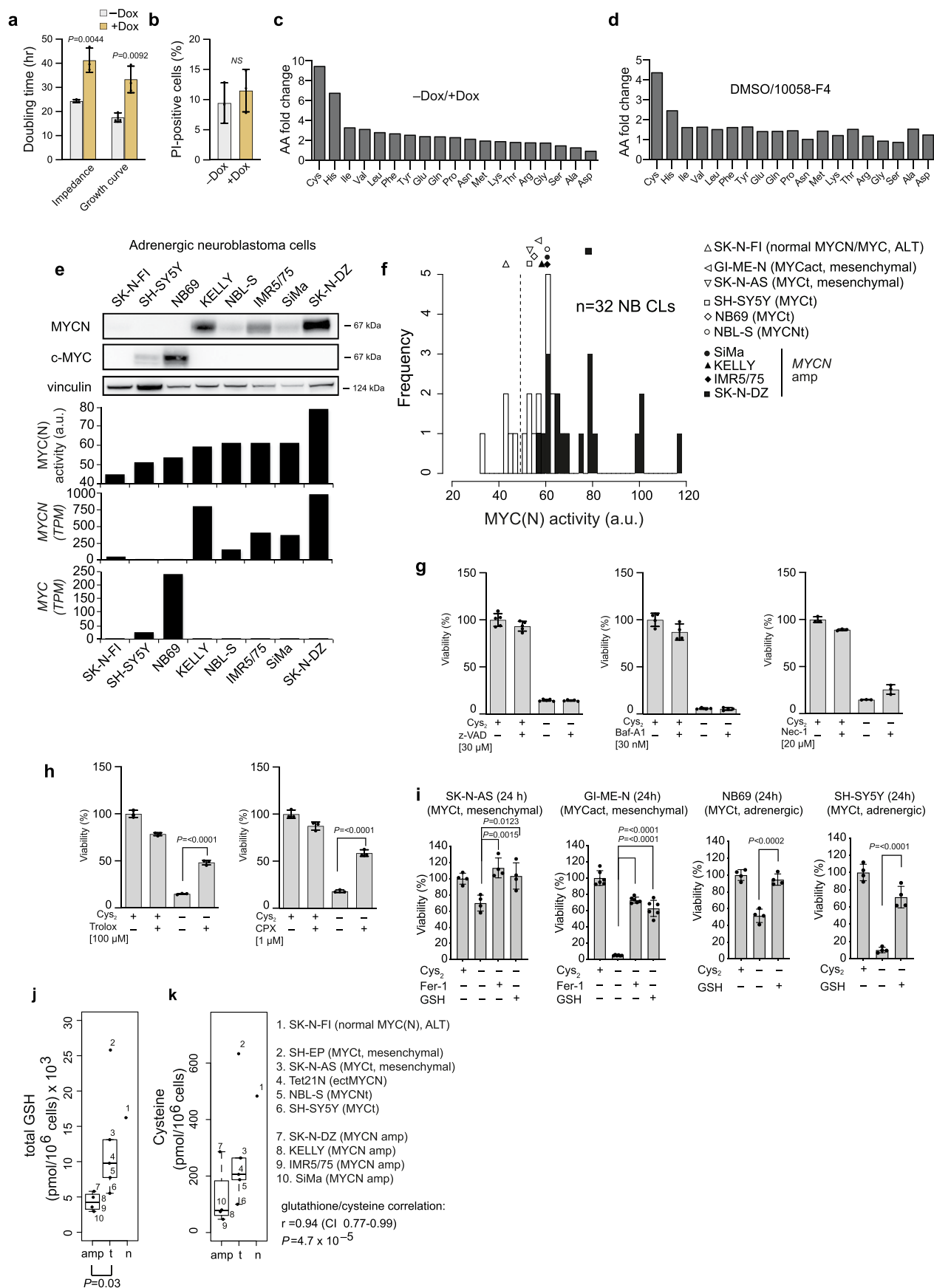
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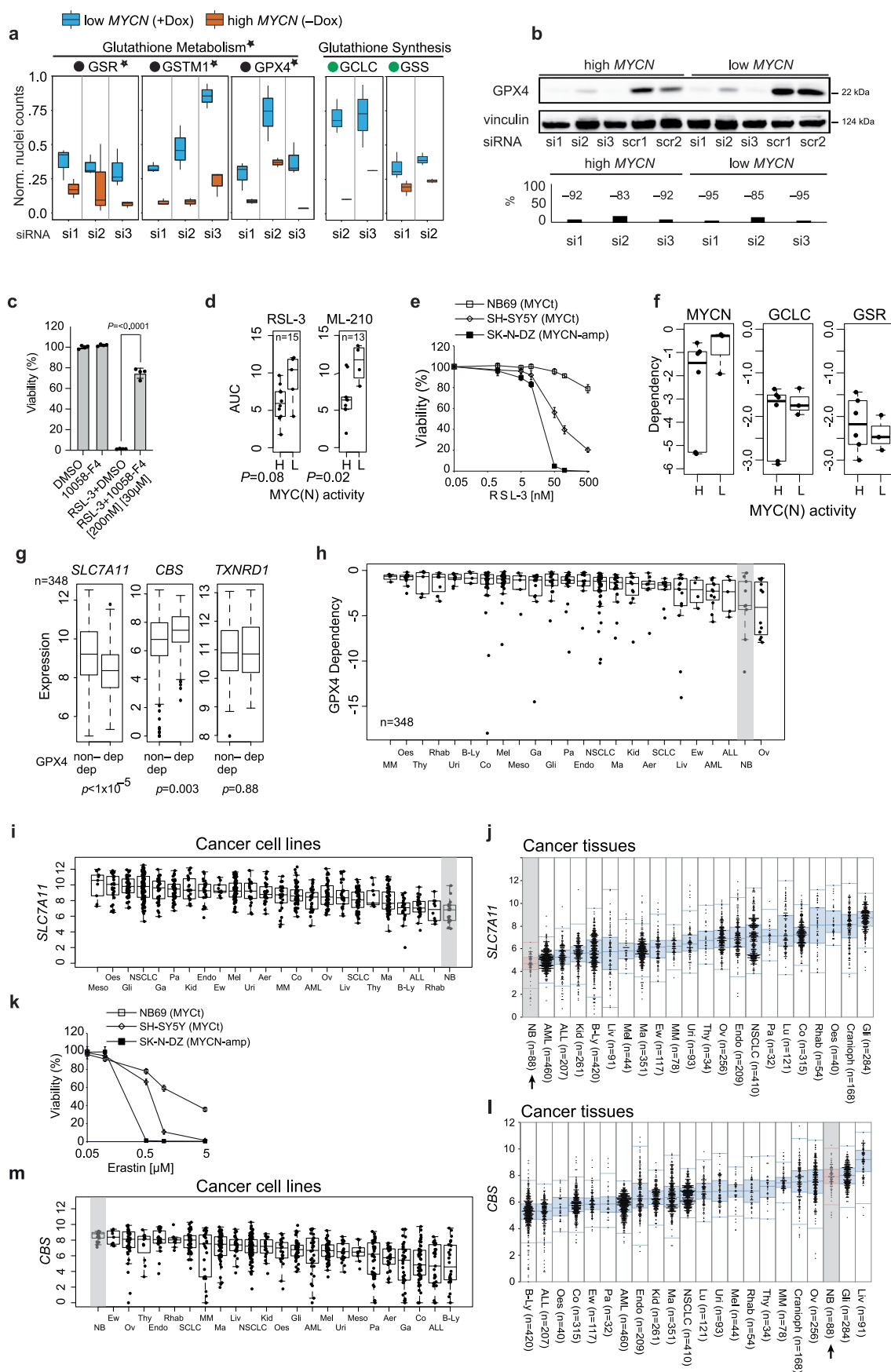
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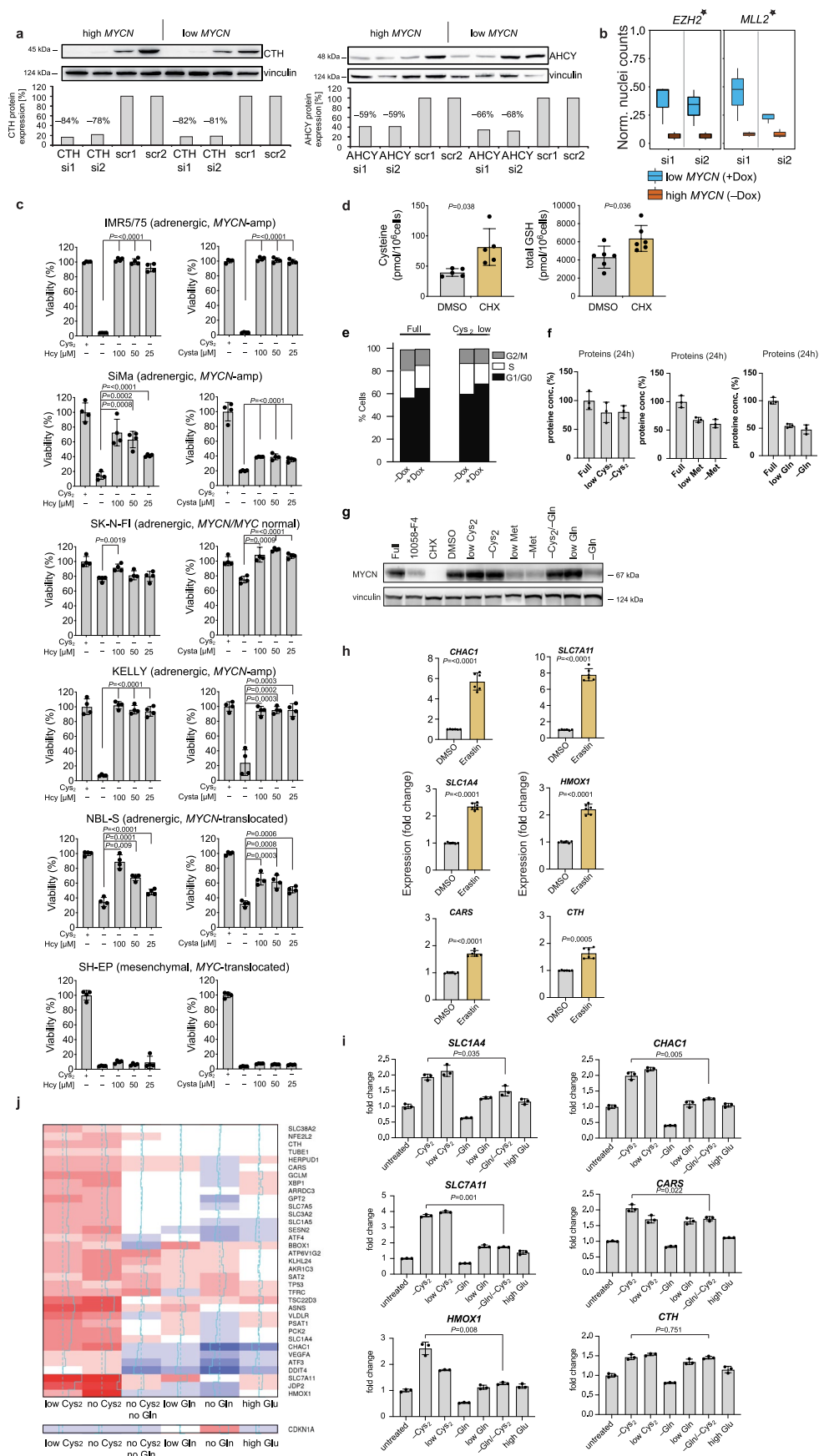
Extended Data Fig. 1 | See next page for caption.

Extended Data Fig. 1 | Cell lines phenotypic data. **a**, Doubling time calculation from exponential growth curve quantified by FACS and impedance measured using the RCTA xCELLigence system for high-MYCN (-Dox) and low-MYCN (+Dox) IMR5/75 cells. Cell proliferation curves of exponentially growing cells ($n=3$ samples, experiment replicated 3 times). **b**, Viability quantification using FACS and propidium iodide staining in exponentially growing cells: high MYCN (-Dox) vs. low MYCN (+Dox) ($n=3$ samples, experiment replicated 3 times). **c**, **d**, Fold changes of intracellular amino acid levels after MYCN inhibition (+Dox or 10058-F4, inhibiting MYCN/MAX binding, 96 h). **e**, MYC(N) RNA, protein expression (experiment replicated 3 times) and **f**, 'MYC(N) activity' score³⁹ for a panel of adrenergic neuroblastoma cell lines. MYC(N) activity score of 32 neuroblastoma (NB) cell lines (CLs) having different MYC(N) genetic status (t = translocation; amp = amplification; act = activated due to unknown mechanism). **g**, Relative viability of IMR5/75 cells after Cys₂ deprivation in the presence or absence of Inhibitors of apoptosis (z-VAD FMK, $n=5$ samples), autophagy (Bafilomycin A1, Baf-A1, $n=4$ samples,) and necroptosis (RIPK1 inhibitor, Necrostatin-1, Nec-1, $n=3$ samples) and **h**, Trolox, or intracellular iron chelator ciclopirox olamine (CPX); $n=3$ samples, experiment replicated 3 times. **i**, Relative viability of neuroblastoma cell lines after Cys₂ deprivation in the presence or absence of ferrostatin-1 (Fer-1) or glutathione (GSH); $n=4$ samples for SK-N-AS, NB69 and SH-SY5Y cell lines and $n=6$ samples for GI-ME-N cell line; experiment replicated 3 times. **j**, Total GSH and **k**, cysteine (Cys) levels in neuroblastoma cells with amplified MYCN, MYC(N) translocations or normal MYC(N) status; Wilcoxon rank-sum test; center line indicates the median value, lower and upper hinges represent the 25th and 75th percentiles, respectively, and whiskers denote 1.5x interquartile range. Each dot corresponds to one sample; p -values as indicated.



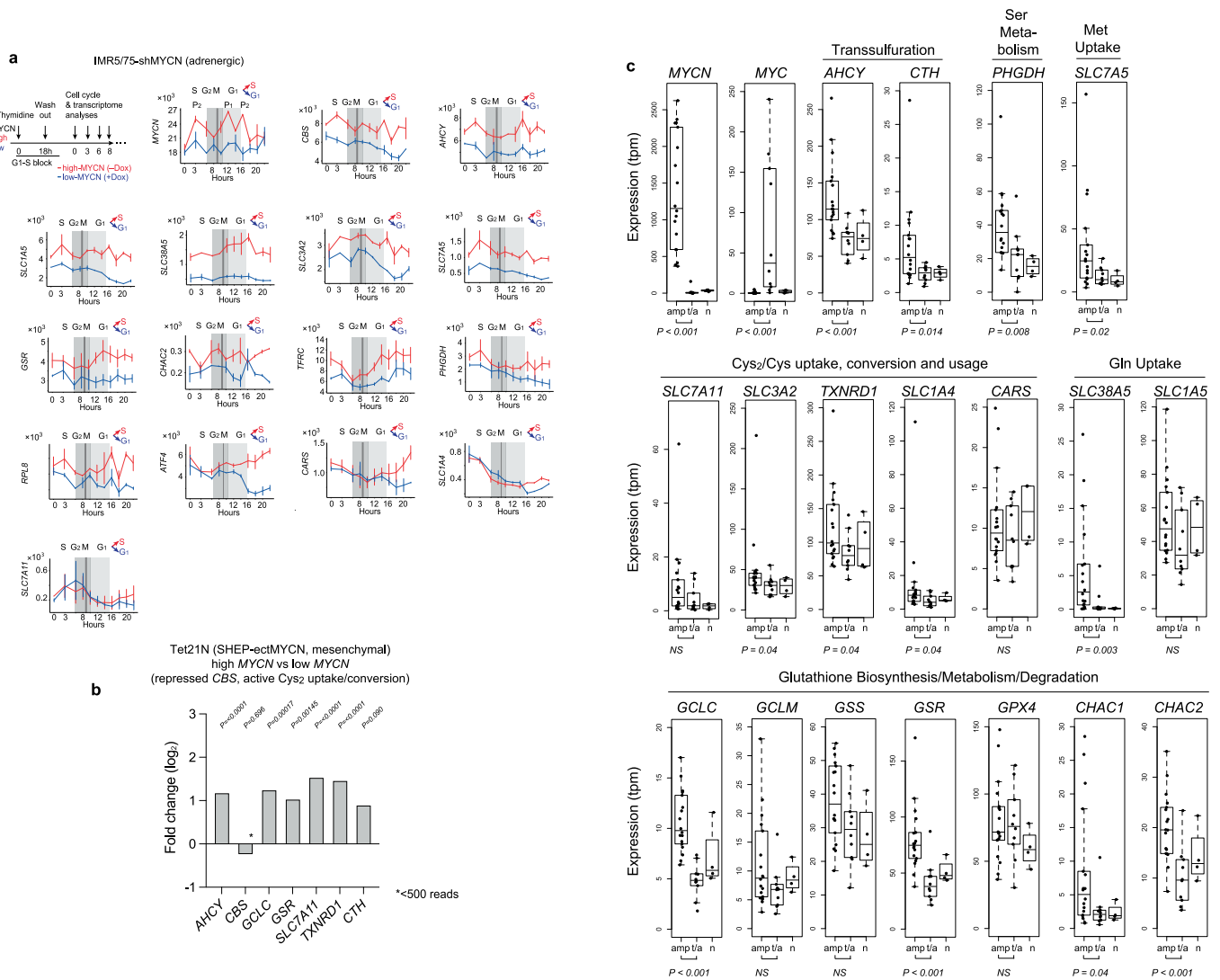
Extended Data Fig. 2 | See next page for caption.

Extended Data Fig. 2 | Additional screen targets and expression data. **a**, MYCN synthetic lethal screen hits. MYCN effects on the two-step biosynthesis of GSH and GSH metabolism (*top MYCN synthetic lethal hits of GSH metabolism, false discovery rate of 0.2) (median of 2-3 siRNAs). **b**, Western Blot and quantification of siRNA-mediated *GPX4* knockdown (96 h) in Tet21N cells with doxycycline-regulatable MYCN (experiment replicated 3 times). **c**, Relative viability (survival of compound-treated cells divided by survival of vehicle-treated cells) of IMR5/75 cell line after treatment with RSL-3 in the presence or absence of MYC inhibitor (10058-F4) ($n = 4$ samples, experiment replicated 3 times) and **d**, Response to GPX4 inhibitors (RSL-3, ML-210) (CTRPv2 database⁵⁴); H = high and L = low MYC(N) activity; Wilcoxon rank-sum test. **e**, Sensitivity to RSL-3-induced ferroptosis in MYCN-amplified neuroblastoma cell lines ($n = 4$ samples, experiment replicated 3 times). **f**, MYC(N) activity correlation with GCLC or GSR dependency in nine neuroblastoma cell lines (DRIVE database³⁹). **g**, Low expression of *SLC7A11* and high expression of *CBS* associated with GPX4 dependency in 384 cancer cell lines (DRIVE database³⁹). *TXNRD1* expression dependent on GPX4 knockdown sensitivity in 384 cancer cells lines (Wilcoxon rank-sum test). **h**, GPX4 dependency in 348 cancer cell lines grouped according to cancer entities (DRIVE database³⁹). **k**, Relative viability of NB69, SH-SY5Y and SK-N-DZ cell lines after treatment with erastin. $n = 3$ samples, experiment replicated 3 times. **i, j, l, m**, Expression of *SLC7A11* or *CBS* in cell lines (CCLE database, $n = 917^{55}$) or tissues (R2 database⁵⁶). Box plots: center line indicates the median value, lower and upper hinges represent the 25th and 75th percentiles, respectively, and whiskers denote 1.5x interquartile range. Each dot corresponds to one sample; Wilcoxon rank-sum test; p -values as indicated.

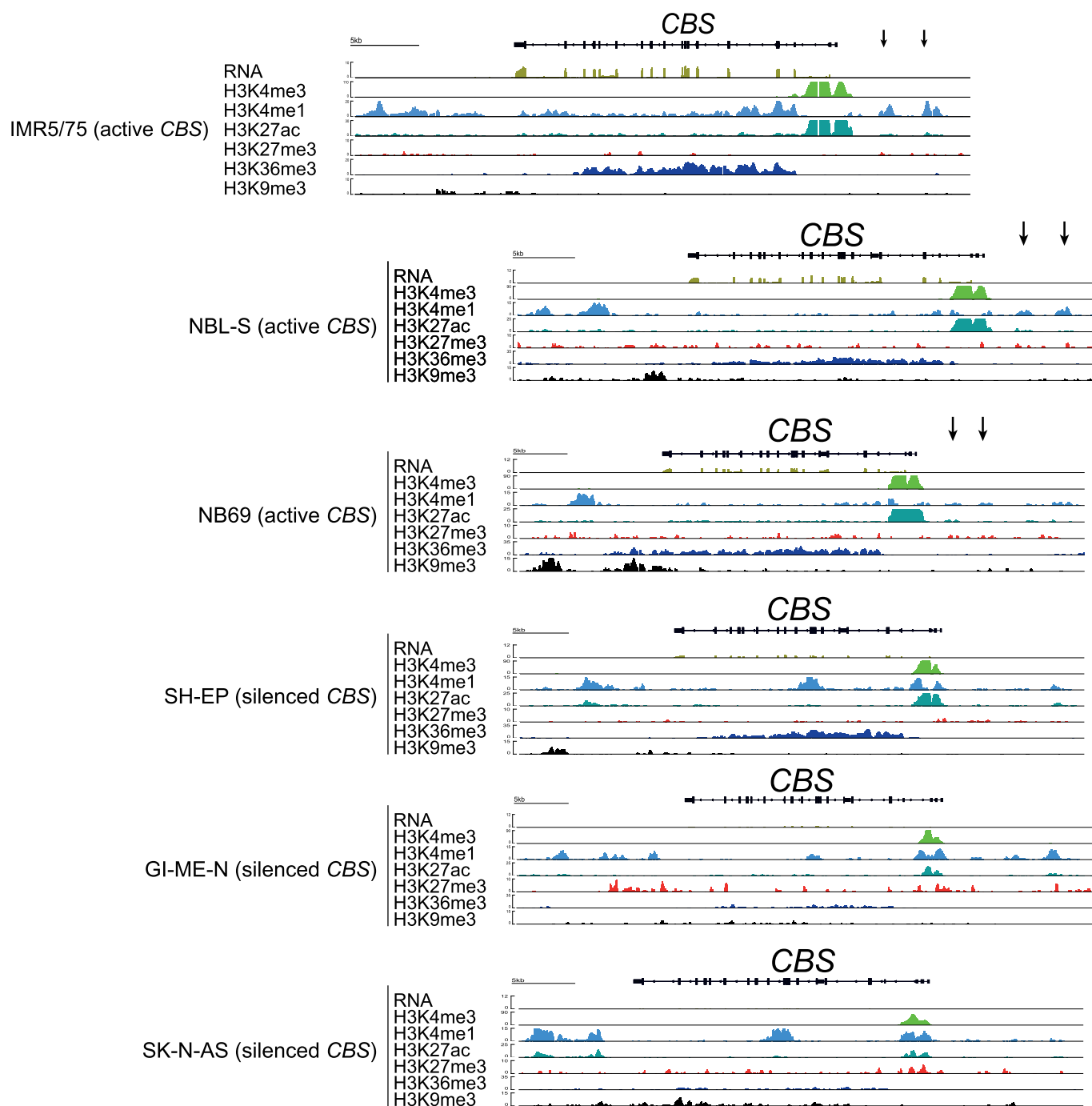


Extended Data Fig. 3 | See next page for caption.

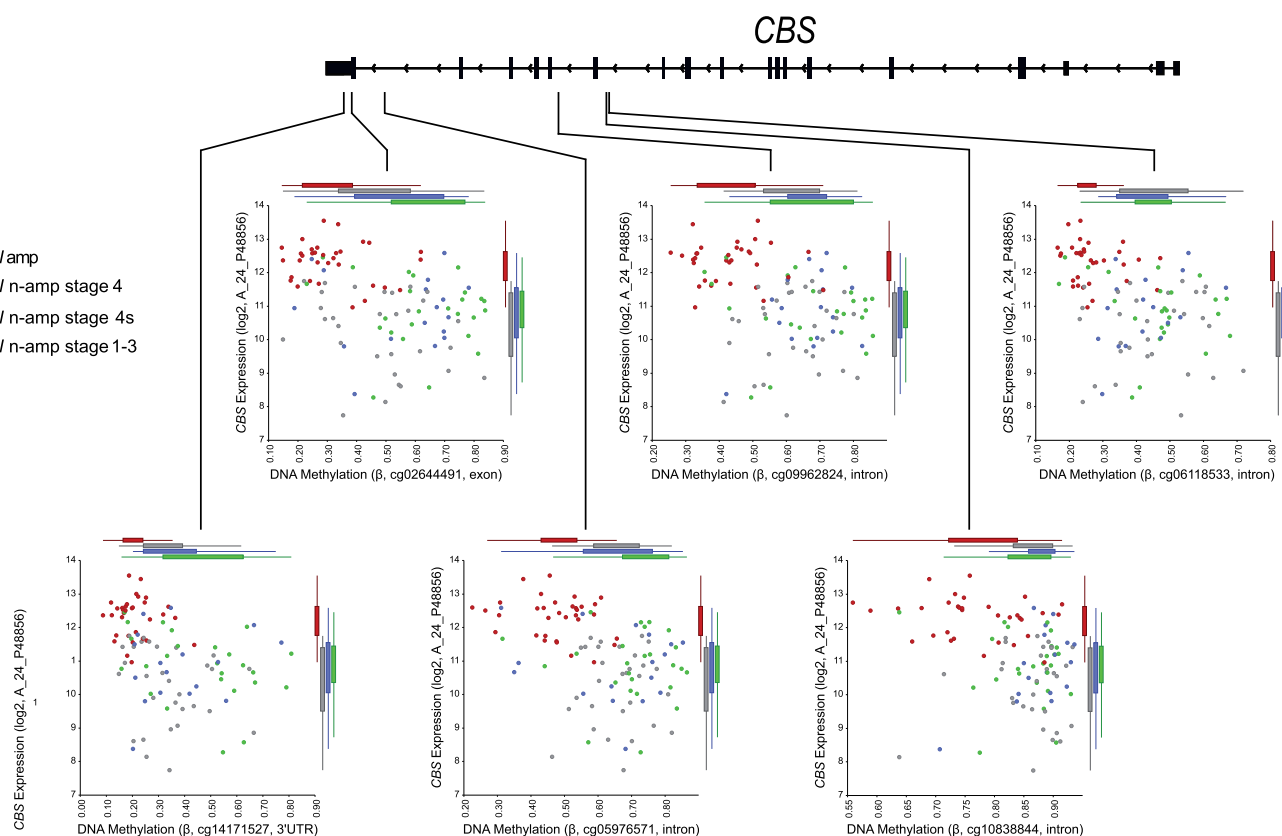
Extended Data Fig. 3 | Quantification of transsulfuration activity. **a**, Western Blot and quantification of CTH or AHCY protein expression upon siRNA-mediated knockdown in IMR5/75 high *MYCN* or low *MYCN* cells. Experiment replicated 3 times. **b**, Two methyltransferases that may indirectly increase Hcy levels are synthetic lethal with high *MYCN* (*MYCN* siRNA screen; *false discovery rate of 0.2); center line indicates the median value of 2-3 siRNAs, lower and upper hinges represent the 25th and 75th percentiles, respectively, and whiskers denote 1.5x interquartile range. **c**, Relative viability (survival of compound-treated cells divided by survival of vehicle-treated cells) of adrenergic and mesenchymal neuroblastoma cell lines after Cys₂ deprivation in the presence or absence of Cysta or Hcy. *n* = 4 samples, experiment replicated 3 times. **d**, Impact of protein synthesis inhibition by cycloheximide (CHX) on cysteine (*n* = 5 samples) and glutathione level (*n* = 6 samples and CHX, 100 ng/ml, 24 h and experiment replicated 3 times). **e**, Cell cycle analysis of *MYCN*-amplified IMR5/75 cells (24 h) after Cys₂ depletion (representative measurement of 3 replicates). **f**, Analysis of protein synthesis in *MYCN*-amplified IMR5/75 cells (24 h) after low levels or complete depletion of Cys₂, methionine (Met) or glutamine (Gln) in the culture medium (*n* = 3 samples, experiment replicated 3 times). **g**, *MYCN* protein expression (by western blot) upon *MYCN*-MAX inhibition (10058-F4), protein synthesis inhibition by cycloheximide (CHX, 100 ng/ml) and low levels or complete depletion of Cys₂, methionine (Met) or glutamine (Gln) in the culture medium after 8 h. Experiment replicated 3 times. **h**, **i**, **j**, RNA-seq and RT-qPCR of IMR5/75 cells 24 h following Cys, Gln depletion and Glu excess or erastin treatment and regulation of ferroptosis-related genes (*n* = 6 samples in **h**, *n* = 3 samples in **i**; experiment replicated 3 times). Statistical analysis was performed using Student's two-tailed t-test.



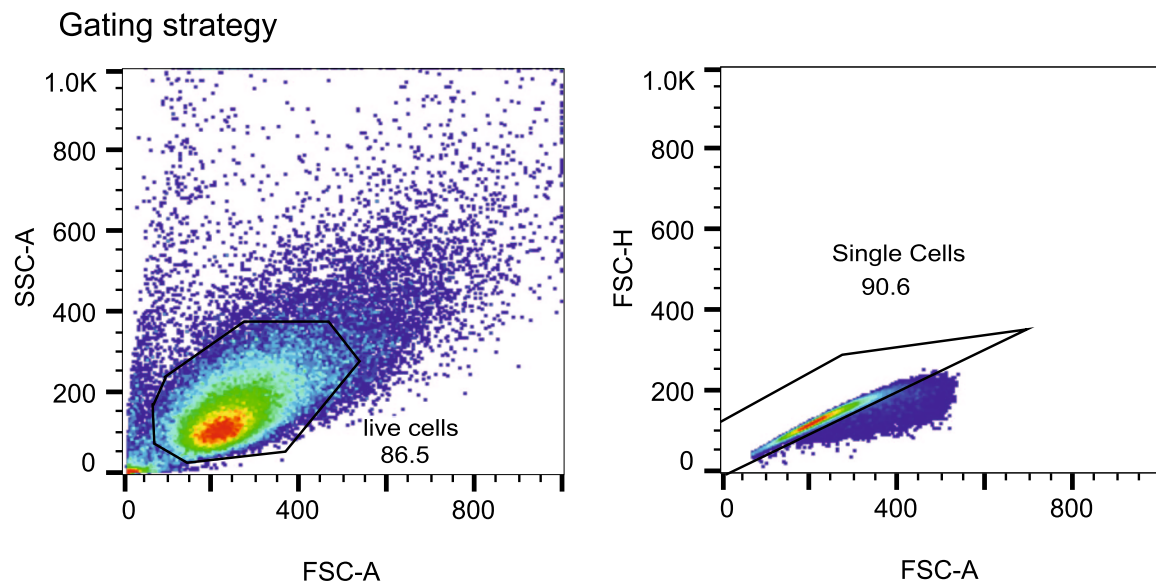
Extended Data Fig. 4 | Expression profiles of neuroblastoma cell lines. a, Time-resolved gene expression profiles during cell cycle progression and cell cycle analysis distinguish pervasive MYCN functions from indirect effects related to cell proliferation rate in high MYCN and low MYCN IMR5/75 cells: MYCN gene expression profile reveals two cell cycle-related peaks (P1 before G1-S transition and P2 before S-G2/M). **b**, Fold change (\log_2) of gene expression in mesenchymal Tet21N (SH-EP-ectMYCN) cells harboring an inducible MYCN transgene upon MYCN induction. Tet21N and parental SH-EP cells have low CBS expression and inactive transsulfuration. MYCN induction has a significant effect on AHCY, GCLC, GSR, SLC7A11 and TXNRD1 but not on CBS expression; experiment replicated 3 times. **c**, Expression of transsulfuration, amino acid uptake/metabolism and glutathione biosynthesis/metabolism/degradation genes in 32 neuroblastoma cell lines (RNA-seq): MYCN-amplified (amp) vs. MYCN/MYC-translocated/activated (t/a) cell lines, MYCN(N)-non-expressors (n) (Wilcoxon rank-sum test; NS = not significant). Center line indicates the median value, lower and upper hinges represent the 25th and 75th percentiles, respectively, and whiskers denote 1.5x interquartile range. Each dot corresponds to one sample; *p*-values as indicated.



Extended Data Fig. 5 | ChIP-seq data for histone modifications. RNA-seq normalized reads and input normalized read counts of ChIP-seq experiments for histone modifications at the *CBS* gene locus in adrenergic *MYCN*-amplified IMR5/75, *MYCN*-translocated NBL-S, and *MYC*-translocated NB69 cells with high *CBS* expression and active transsulfuration, mesenchymal *MYCN*-normal diploid SH-EP, GI-ME-N, and SK-N-AS cells with elevated *MYC* expression due to translocations or unknown mechanism (GI-ME-N) and very low/absent *CBS* expression and inactive transsulfuration.



Extended Data Fig. 6 | Methylation profiles and CBS expression in tumor samples. Genomic position of CBS-annotated CpGs whose methylation is significantly associated with CBS expression and patient risk ($p < 0.01$, Wilcoxon rank-sum statistics and $p < 0.05$, Fisher exact test). DNA methylation assessed by Infinium Human Methylation 450 BeadChips, and CBS expression assessed by 44k customized Agilent oligonucleotide microarrays in 105 primary neuroblastomas (GEO accession GSE73518¹⁸). For all CBS CpGs whose methylation significantly correlated with CBS expression and patient risk, hypomethylation was associated with CBS upregulation and high-risk disease. R2 Genomics Analysis and Visualization Platform (<http://r2.amc.nl>) was used for data visualization.



Extended Data Fig. 7 | Gating strategy for flow cytometry data. For all samples, an initial manual gate in SSC-A by FSC-A was set to identify live cells and exclude debris. From the live cells, a rectangular gate was set on FSC-H by FSC-A to exclude doublets, meaning cells off the diagonal. If DNA staining was carried out, an additional gate was set on DNA (Violet channel)-W by DNA-A to exclude additional cell debris or DNA doublets.

Reporting Summary

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Statistics

For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.

n/a Confirmed

- ☐ ☒ The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement
- ☐ ☒ A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly
- ☐ ☒ The statistical test(s) used AND whether they are one- or two-sided
Only common tests should be described solely by name; describe more complex techniques in the Methods section.
- ☐ ☒ A description of all covariates tested
- ☐ ☒ A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons
- ☐ ☒ A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)
- ☐ ☒ For null hypothesis testing, the test statistic (e.g. F , t , r) with confidence intervals, effect sizes, degrees of freedom and P value noted
Give P values as exact values whenever suitable.
- ☒ ☐ For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
- ☐ ☒ For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes
- ☐ ☒ Estimates of effect sizes (e.g. Cohen's d , Pearson's r), indicating how they were calculated

Our web collection on [statistics for biologists](#) contains articles on many of the points above.

Software and code

Policy information about [availability of computer code](#)

Data collection

NA

Data analysis

Software used in the study:
 R (version 3.4.0)
 Image J (version 1.51)
 FlowJo version 10.6.0
 Microarray analysis (as described in Klaus & Reisenauer, F1000Res 2016)
 Bioconductor - limma (version 3.46)
 BHC in-house software (developed and owned by BAYER Health care, intellectual property)

RNAseq:
 STAR (version 2.5.3a)
 R (version 4.3.0 & 3.4.3)
 R - edge R (version 3.20.9)
 R - gplots (version 3.1.1)

ChIPseq:
 trimalore (version 0.4.3)
 bowtie2 (version 2.3)
 deepTools (version 3.0)
 macs2 (version 2.1)

For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Portfolio [guidelines for submitting code & software](#) for further information.

Data

Policy information about [availability of data](#)

All manuscripts must include a [data availability statement](#). This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A description of any restrictions on data availability
- For clinical datasets or third party data, please ensure that the statement adheres to our [policy](#)

All information about data availability is provided in the data availability statement of the article:

Proteome data of neuroblastoma tumors was previously published by (Hartlieb, S. A. et al. Nat Commun 2021), and all data is deposited at the European Genome-Phenome Archive (EGA) as dataset EGAD00001006737 as part of the study EGAS00001004349. Data is available upon request by contacting Frank Westermann. RNA-seq data of 498 primary neuroblastoma patients was previously published (Lutz, W. et al. 1996) and is available at Gene Expression Omnibus (GEO) under the accession number GSE62564. DNA methylation data of primary neuroblastoma tumors was previously published (Cassago, A. et al. 2012) and is available at GEO under the accession number GSE73518. Time-course RNAseq profiling of IMR5/75 MYCN-high and MYCN-low cells was previously published (Muth, D. et al. 2010), and can be accessed at GEO under the accession number GSE97774. ChIP-seq data is deposited at GEO under the accession number GSE189174. The aligned .bam files of RNA expression profiles of the high-MYCN depletion/rescue experiments were submitted to the European Nucleotide Archive (ENA) and can be found under the accession number PRJEB25184. Data of the MYCN synthetic lethal siRNA screen is added to this article as Supplementary Data. Microarray data from tumor samples is deposited in GEO under the accession number GSE192976. Source data are provided with this paper. The remaining data are available within the Article, Supplementary Information or available from the authors upon reasonable request.

Field-specific reporting

Please select the one below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.

☒ Life sciences ☐ Behavioural & social sciences ☐ Ecological, evolutionary & environmental sciences

For a reference copy of the document with all sections, see [nature.com/documents/nr-reporting-summary-flat.pdf](https://www.nature.com/documents/nr-reporting-summary-flat.pdf)

Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	For in vivo studies, sample size was calculated with the help of a biostatistician using R version 3.4.0. Assumptions for power analysis were as follows: α error, 5%; β error, 20%. Values for standard deviations and differences between experimental groups were based on previous experiments (whenever a similar data type was available). for in vitro studies we used at least 3 biological replicates for each experiments.
Data exclusions	no data were excluded
Replication	in vitro experiments were repeated at least three times with similar results.
Randomization	For in vivo experiments mice were randomized into treatment groups prior to treatment. In case animals had to be sacrificed before the pre-defined endpoint (due to weight loss or other termination criteria), they were excluded from any downstream analyses. For the in vitro experiments, all samples were analyzed equally with no subsampling; therefore, there was no requirement for randomization
Blinding	For in vivo experiments all animal experiments (except during animal treatment) were blinded during entire experiments or follow up assessment. For in vitro studies, data collection and analysis were not performed blinded to the conditions of the experiments.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

n/a	Involved in the study
<input type="checkbox"/>	<input checked="" type="checkbox"/> Antibodies
<input type="checkbox"/>	<input checked="" type="checkbox"/> Eukaryotic cell lines
<input checked="" type="checkbox"/>	<input type="checkbox"/> Palaeontology and archaeology
<input type="checkbox"/>	<input checked="" type="checkbox"/> Animals and other organisms
<input type="checkbox"/>	<input checked="" type="checkbox"/> Human research participants
<input checked="" type="checkbox"/>	<input type="checkbox"/> Clinical data
<input checked="" type="checkbox"/>	<input type="checkbox"/> Dual use research of concern

Methods

n/a	Involved in the study
<input type="checkbox"/>	<input checked="" type="checkbox"/> ChIP-seq
<input type="checkbox"/>	<input checked="" type="checkbox"/> Flow cytometry
<input checked="" type="checkbox"/>	<input type="checkbox"/> MRI-based neuroimaging

Antibodies

Antibodies used

MYCN (clone B8.4.B, sc-53993, Santa Cruz, Lot. B2316, 1:1000)
 c-MYC (clone Y69, ab32072, Abcam, Lot. 625369, 1:1000)
 CTH (ab54573, Abcam, Lot. GR3260298-2, 1:1000)
 SAHH (A-11) (AHCY antibody, clone A-11, sc-271389, Santa Cruz, Lot. C2111, 1:1000),
 GPX4 (ab41787, Abcam, Lot. GR56784-1, 1:1000),
 CARS (clone EPR7121, ab126714, Abcam, Lot. Y1081819DS, 1:1000)
 Glutaminase 1 (clone EP7212, ab156876, Abcam, Lot. GR249636-29, 1:40,000)
 Loading control: vinculin (clone 749, sc-73614, Santa Cruz, Lot. A2319, 1:1000) or β -actin-conjugated (ab20272, Abcam, Lot. GR3418697-1, 1:5000)
 Secondary AB: HSR–peroxidase labeled anti-mouse (115-035-003, Dianova, 1:1000) or anti-rabbit (111-035-144, Dianova, 1:1000).

ChIP-seq: For all ChIP-seq experiments 3 μ g of antibody were used per ChIP.
 H3K27me3; rabbit polyclonal; Active Motif 39155; Lot.31014017
 H3K36me3; rabbit polyclonal; Abcam ab9050; Lot.GR273250-1
 H3K9me3; rabbit polyclonal; Abcam; ab8898; Lot.GR148830-2
 H3K27ac; rabbit polyclonal; Abcam; ab4729; Lot.GR183919-2

Validation

All antibodies used in this study were validated by manufactures for the species and specific application. Relevant validation results can be found in the website of each manufacture. A protein size marker was run on every western blot and the size of the assessed bands was compared to the manufactures information (See source data)

Eukaryotic cell lines

Policy information about [cell lines](#)

Cell line source(s)

Human neuroblastoma cell lines: IMR5/75, KELLY, SiMa, NBL-S, SK-N-FI, SH-SY5Y, NB69, SKNDZ, SH-EP, GI-ME-N.

SK-N-FI, SK-N-DZ, SH-SY5Y cells were purchased from ATCC.

KELLY, SiMa and GI-ME-N were purchased from DSMZ .

NB69 were kindly provided by Larissa Savelyeva

NBL-S and TET21N (SH-EP) were provided by G.M. Brodeur and W. Lutz, respectively.

Tunable cell lines, IMR5/75 MYCN shRNA, and SH-EP MYCN transgene (Tet21N) were generated and cultured as described previously (please see Methods section)

Authentication

Cell line identity/unique SNP profiles were confirmed by the Multiplexion Multiplex Cell Authentication service (Heidelberg, Germany). The purity of cell lines was validated using the Multiplex cell Contamination Test by Multiplexion (Heidelberg, Germany).

Mycoplasma contamination

All cell lines tested negative for mycoplasma contamination as noted in 'Materials and Methods'

Commonly misidentified lines
(See [ICLAC](#) register)

no commonly misidentified cell lines were used here

Animals and other organisms

Policy information about [studies involving animals](#); [ARRIVE guidelines](#) recommended for reporting animal research

Laboratory animals

Mouse strains used in the study: NOD.Cg-Prkdcscidll2rgtm1Wjl/SzJ (NSG, JAX stock #005557). Female mice, 3 – 4 months of age, were used for experiments. Mice were housed in individually ventilated cages under temperature and humidity control. Cages contained an enriched environment with bedding material.

Wild animals	No wild animals were used in the study.
Field-collected samples	No wild animals were used in the study.
Ethics oversight	All studies involving mice and experimental protocols were conducted in compliance with German Cancer Center Institute guidelines and approved by the governmental review board of the state of Baden-Wuerttemberg, Regierungspraesidium Karlsruhe, under the authorization number G-176/19, followed the German legal regulations. Animals health were monitored daily and mice were euthanized as soon as they reached abortion criteria defined in the procedure

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Human research participants

Policy information about [studies involving human research participants](#)

Population characteristics	Tumor samples are from patients enrolled in the German Neuroblastoma Trials of the GPOH (NB97, NB2004, NB2016). Detailed patient specific information (Age, Status, Gender etc) is provided in the previous publication by Gartlgruber et al, Nature Cancer 2020.
Recruitment	Almost all childhood neuroblastoma in Germany (>99%) are enrolled in a clinical trial, informed consent is given for the use of tumor material for research purposes.
Ethics oversight	All neuroblastoma patients were enrolled in the German Neuroblastoma Trial (NB97, NB2004, NB 2016) approved by the Ethics Committee of the University of Cologne and informed written consent was obtained from the patients' parents.

Note that full information on the approval of the study protocol must also be provided in the manuscript.

Clinical data

Policy information about [clinical studies](#)

All manuscripts should comply with the ICMJE [guidelines for publication of clinical research](#) and a completed [CONSORT checklist](#) must be included with all submissions.

Clinical trial registration	not applicable
Study protocol	not applicable
Data collection	not applicable
Outcomes	not applicable

Dual use research of concern

Policy information about [dual use research of concern](#)

Hazards

Could the accidental, deliberate or reckless misuse of agents or technologies generated in the work, or the application of information presented in the manuscript, pose a threat to:

No	Yes	
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<input checked="" type="checkbox"/>	<input type="checkbox"/>	National security
<input checked="" type="checkbox"/>	<input type="checkbox"/>	Crops and/or livestock
<input checked="" type="checkbox"/>	<input type="checkbox"/>	Ecosystems
<input checked="" type="checkbox"/>	<input type="checkbox"/>	Any other significant area

Experiments of concern

Does the work involve any of these experiments of concern:

- | No | Yes | |
|-------------------------------------|--------------------------|---|
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| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Confer resistance to therapeutically useful antibiotics or antiviral agents |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Enhance the virulence of a pathogen or render a nonpathogen virulent |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Increase transmissibility of a pathogen |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Alter the host range of a pathogen |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Enable evasion of diagnostic/detection modalities |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Enable the weaponization of a biological agent or toxin |
| <input checked="" type="checkbox"/> | <input type="checkbox"/> | Any other potentially harmful combination of experiments and agents |

ChIP-seq

Data deposition

- ☒ Confirm that both raw and final processed data have been deposited in a public database such as [GEO](#).
- ☒ Confirm that you have deposited or provided access to graph files (e.g. BED files) for the called peaks.

Data access links

May remain private before publication.

ChIP-seq data has been uploaded to GEO under the accessions GSE189174.

Files in database submission

GSM5695680	GI-ME-N H3K27ac
GSM5695681	GI-ME-N H3K27me3
GSM5695682	GI-ME-N H3K36me3
GSM5695683	GI-ME-N H3K4me1
GSM5695684	GI-ME-N H3K4me3
GSM5695685	GI-ME-N H3K9me3
GSM5695686	GI-ME-N input DNA
GSM5695687	IMR575 H3K27ac
GSM5695688	IMR575 H3K27me3
GSM5695689	IMR575 H3K36me3
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GSM5695694	Kelly H3K27ac
GSM5695695	Kelly H3K27me3
GSM5695696	Kelly H3K36me3
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GSM5695706	NB69 H3K9me3
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GSM5695713	NBL-S H3K9me3
GSM5695714	NBL-S input DNA
GSM5695715	SH-EP H3K27ac
GSM5695716	SH-EP H3K27me3
GSM5695717	SH-EP H3K36me3
GSM5695718	SH-EP H3K4me1
GSM5695719	SH-EP H3K4me3
GSM5695720	SH-EP H3K9me3
GSM5695721	SH-EP input DNA
GSM5695722	SK-N-FI H3K27ac
GSM5695723	SK-N-FI H3K27me3
GSM5695724	SK-N-FI H3K36me3

GSM5695725 SK-N-FI H3K4me1
 GSM5695726 SK-N-FI H3K4me3
 GSM5695727 SK-N-FI H3K9me3
 GSM5695728 SK-N-FI input DNA
 GSM5695729 SKNDZ H3K27ac
 GSM5695730 SKNDZ H3K27me3
 GSM5695731 SKNDZ H3K36me3
 GSM5695732 SKNDZ H3K4me1
 GSM5695733 SKNDZ H3K4me3
 GSM5695734 SKNDZ H3K9me3
 GSM5695735 SKNDZ input DNA

Genome browser session
 (e.g. [UCSC](#))

No longer applicable.

Methodology

Replicates	For each tumor and cell line one biological replicate was done due to the limited amount of material.
Sequencing depth	ChIP-sequencing was done using Illumina HiSeq2000 50 bp single end sequencing.
Antibodies	H3K27me3; rabbit polyclonal; Active Motif 39155; Lot.31014017 H3K36me3; rabbit polyclonal; Abcam ab9050; Lot.GR273250-1 H3K9me3; rabbit polyclonal; Abcam; ab8898; Lot.GR148830-2 H3K27ac; rabbit polyclonal; Abcam; ab4729; Lot.GR183919-2
Peak calling parameters	Peak calling was performed with the MACS2 algorithm, both with the narrowPeak and broadPeak parameters; the FDR cutoff for peak calling was set to 1%.
Data quality	Only uniquely aligned reads are considered, and duplicates are removed. Data quality was ensured both by visual inspection of reference region in the genome browser. Moreover, we assessed the enrichment signal using the fingerprint method developed as part of the DeepTools package (Diaz et al.), and using criteria recommended by the ENCODE consortium such as the PCR bottleneck coefficient and the FRiP (fraction of reads in peaks).
Software	The ChIP-seq processing was performed using a custom pipeline written in Snakemake; the steps involve (1) read trimming using TrimGalore, (2) alignment using Bowtie2 on the hg19 genome with standard parameters, (3) merginf of replicates if available, (4) peak calling using MACS2, (5) QC using the fingerprint and FRiP method, (6) SES normalization and bigwig generation by subtracting normalized input from IP.

Flow Cytometry

Plots

Confirm that:

- ☒ The axis labels state the marker and fluorochrome used (e.g. CD4-FITC).
- ☒ The axis scales are clearly visible. Include numbers along axes only for bottom left plot of group (a 'group' is an analysis of identical markers).
- ☒ All plots are contour plots with outliers or pseudocolor plots.
- ☒ A numerical value for number of cells or percentage (with statistics) is provided.

Methodology

Sample preparation	MYCN-low populations were established by incubating cells with 1 µg/ml doxycycline at least 48 h prior to further treatment. Cells were harvested and lipid peroxidation was analyzed, using C11-BODIPY BD of 4 µM final concentration in Hanks' Balanced Salt Solution (HBSS). Cells were incubated at 37°C for 15 min and signal intensity was measured. Total intracellular ROS levels were determined using CellROX® (according to Thermo Fisher Scientific instruction).
Instrument	MACSquant VYB, model number: 130-096-116, SN:3050 BD FACS Aria™ cell sorter IIu, model number: 355119, SN: P0087
Software	FlowJo software version 10.6.0 (commercial standard software for analysis of flow cytometric data).
Cell population abundance	In this manuscript, only one pure population at the time was analyzed (neuroblastoma cell lines).
Gating strategy	For all samples, an initial manual gate in SSC-A by FSC-A was set to identify live cells and exclude debris. From the live cells, a rectangular gate was set on FSC-H by FSC-A to exclude doublets, meaning cells off the diagonal. If DNA staining was carried out, an additional gate was set on DNA (Violet channel)-W by DNA-A to exclude additional cell debris or DNA doublets (Extended Data Figure 7))

- ☒ Tick this box to confirm that a figure exemplifying the gating strategy is provided in the Supplementary Information.