Temporal Molecular and Biological Assessment of an Erlotinib-Resistant Lung Adenocarcinoma Model Reveals Markers of Tumor Progression and Treatment Response

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Abstract

Patients with lung cancer with activating mutations in the EGF receptor (EGFR) kinase, who are treated long-term with tyrosine kinase inhibitors (TKI), often develop secondary mutations in EGFR associated with resistance. Mice engineered to develop lung adenocarcinomas driven by the human EGFR T790M resistance mutation are similarly resistant to the EGFR TKI erlotinib. By tumor volume endpoint analysis, these mouse tumors respond to BIBW 2992 (an irreversible EGFR/HER2 TKI) and rapamycin combination therapy. To correlate EGFR-driven changes in the lung with response to drug treatment, we conducted an integrative analysis of global transcriptome and metabolite profiling compared with quantitative imaging and histopathology at several time points during tumor progression and treatment. Responses to single-drug treatments were temporary, whereas combination therapy elicited a sustained response. During tumor development, metabolomic signatures indicated a shift to high anabolic activity and suppression of antitumor programs with 11 metabolites consistently present in both lung tissue and blood. Combination drug treatment reversed many of the molecular changes found in tumors lung. Data integration linking cancer signaling networks with metabolic activity identified key pathways such as glutamine and glutathione metabolism that signaled response to single or dual treatments. Results from combination drug treatment suggest that metabolic transcriptional control through C-MYC and SREBP, as well as ELK1, NRF1, and NRF2, depends on both EGFR and mTORC1 signaling. Our findings establish the importance of kinetic therapeutic studies in preclinical assessment and provide in vivo evidence that TKI-mediated antiproliferative effects also manifest in specific metabolic regulation. Cancer Res; 72(22); 5921–33. ©2012 AACR.

Introduction

Lung cancer is the leading cause of cancer-related deaths (1). Current screening methods for early disease are ineffective at improving the 5-year survival rate, which is less than 4% for advanced non–small cell lung cancer (NSCLC; ref. 2). For most cancers, including NSCLC, traditional cytotoxic chemotherapy has limited efficacy, and patients develop resistance to targeted inhibitors, indicating the need for identification of molecular networks operative in disease and treatment response. In particular, effective preclinical systems are required for guidance on patient stratification, therapeutic directives, and response biomarkers. Such needs are exemplified by the current status of NSCLC clinical management. Activating mutations in EGFR, such as the L858R mutation, are effective predictors of response to EGF receptor (EGFR) tyrosine kinase inhibitor (TKI) therapy (3). While most patients initially respond to EGFR TKIs, the majority relapse within 6 to 12 months, underscoring the importance of identifying therapies that overcome or mitigate such resistance. Strikingly, a single secondary missense mutation in the EGFR TK domain, T790M, is associated with more than 50% of acquired resistance cases (4).

Recent advances in the ability to globally assess molecular networks responsive to biologic and chemical perturbations in cancer have facilitated unbiased discovery in disease...
mechanisms (5). Genetically engineered mouse (GEM) models can facilitate both assessment of disease etiology from initiation through progression as well as treatment responses within the organs/cells of interest (6). Mice engineered to develop erlotinib-resistant lung adenocarcinomas driven by human EGFR L858R-T790M mutations (EGFR-TL; refs. 7, 8) respond to BIBW 2992 (afatinib, an irreversible EGFR/HER2 TKI currently in clinical trial) and rapamycin combination therapy, based on tumor volume endpoints (9). BIBW 2992 or rapamycin alone minimally inhibit tumor growth. To further assess responses to single and combination therapies and to gain insight into the optimal strategy for preclinical evaluation in robust cancer models, we examined both transcriptional and metabolic profiles in disease induction and therapeutic response with time. These studies underscore the value of temporal well-powered evaluations and offer an opportunity to identify surrogate markers for disease and treatment responses and for tailoring therapeutic regimens.

Materials and Methods

Tumor induction and sampling
Mice (see Supplementary Materials and Methods) were fed 200 mg/kg doxycycline feed from Research Diets. Blood was taken from the mandibular vein of bisnisgenic EGFR-L858R-T790M/CCSP-rtTA mice (E+r+) or mice carrying the CCSP-rtTA driver, but not the EGFR transgene (E−r−) at 2 weeks (n = 6 for both E−r− and E+r+), and when tumor growth required euthanasia of the mice (terminal at 6–8 weeks; n = 8 for both E+r+ and E−r−), and snap-frozen in an equal volume of 9 mmol/L EDTA in phosphate buffered saline. Lung tissue was harvested and snap-frozen at 2 weeks and terminal stages and age-matched control mice were taken at the same time points postinduction. For treatment studies, blood and lung tissue were harvested after 1 (n = 19), 2 (n = 19), or 4 (n = 40) weeks of daily treatment with drug vehicle, BIBW 2992 (20 mg/kg/d, oral gavage), rapamycin (2 mg/kg/d, intraperitoneally), or the combination of both compounds starting at 4 weeks postinduction.

Histology/pathology
Mice were euthanized at the indicated times and right lungs were inflated with formalin and processed for immunohisto-pathologic analysis. Immunohistochemistry experiments are described in the Supplementary Data. Hematoxylin and eosin slides of lungs from drug- and vehicle-treated tumors were scored blindly by a board certified veterinary pathologist (P.L. Martin) according to a grading scheme that was developed for this study (see Supplementary Data for details). All tumors (4–10) of each experimental group (vehicle- or drug-treated) were analyzed.

Automated image acquisition and quantification
Immunohistochemistry for Ki-67 was conducted on lung sections from each mouse and slides were scanned using the Arioil image scanning and analysis system (Applied Imaging Corp.) at × 20. As the severity of lung disease was heterogeneous between lung lobes, the percentage of Ki-67–positive cells was quantified in the lung lobe that was most representative of the overall histopathologic score. The Ki67 image analysis module was used to count the total number of nuclei in the lung section as well as the total number of Ki-67–positive (Brown-DAB) nuclei and Ki-67–negative (blue-hematoxylin) nuclei. For each lung lobe, approximately, 30,000 to 125,000 nuclei were counted.

Gene expression profiling
Preparation of RNA is described in the Supplementary Data. Hybridization to Mouse Gene 1.0 ST chips was conducted following Affymetrix protocols (Affymetrix, Inc.). Hierarchical clustering analysis was conducted using pairwise complete linkage method with the Pearson correlation distance measure using the GenePattern genomic analysis platform. "Single Sample" Gene Set Enrichment Analysis (10, 11) was used to determine the degree of absolute enrichment of gene ontology (GO) biologic processes gene sets (12) in each sample within the gene expression data set. Signature values for each sample were normalized using the entire sample set. Standard gene set enrichment analysis (GSEA) was used to determine the enrichment of transcription factor binding sites between classes. Gene sets sharing a cis-regulatory motif that is conserved across the human, mouse, rat, and dog genomes (13), representing known or predicted regulatory elements in promoters and 3′-UTRs, were obtained from MSigDB (see Supplementary Data for details).

Results

Temporal evaluation of tumor growth and treatment response reveals distinct dynamics of single versus combination therapy
To examine responses elicited during human EGFR L858R-T790M–driven lung adenocarcinoma, as well as experimental treatment-induced regression, we measured changes with time in lung disease severity and tumor burden. Bisnisgenic EGFR-L858R-T790M/CCSP-rtTA mice (designated E+r+) are engineered to develop lung adenocarcinoma upon EGFR transgene induction by doxycycline. Mice carrying the CCSP-rtTA driver, but not the EGFR transgene (E−r−), served as controls. E+r+ mice developed lung tumors when continuously exposed to doxycycline, and consistent with previous studies (7), adenomas were detectable by magnetic resonance imaging (MRI) and histology within 2 to 3 weeks postinduction, progressing to adenocarcinomas that often involved entire lobes at terminal stages. Induced E+r+ mice were treated with BIBW 2992, rapamycin, or the combination daily for up to 4 weeks (Fig. 1A). To quantify histopathology response to drug treatment (or lack thereof) in the tumors, severity of disease was scored based on extent and morphology ranging from 0 (no lesions) to 5 (multifocal to coalescing adenocarcinoma; Supplementary Data and Supplementary Fig. S1). The histopathologic severity grade increased over time in the vehicle-treated group, and while each individual drug had some effect in suppressing tumor growth, the combination treatment was most effective (score of 1.9 compared with 4.3 for vehicle-treated after 4 weeks, P = 0.002) (Fig. 1B). Thus, the present expanded study agrees well with previous findings that used smaller cohorts.
Figure 1. Quantification of tumor progression in CCSP-rTA; EGFR-L858R-T790M mice and response to treatment with BIBW 2992 and/or rapamycin. A, schematic of dosing regimen, blood sampling, and imaging for the tumor progression and treatment study. Note that necropsies before the treatment start date are for the tumor progression component only. Blood collection for the treatment study was 24 hours after last dose. B, histopathological scoring on H&E-stained lung tumor samples from treated mice. The average score is plotted for each treatment after 1, 2, or 4 weeks of treatment. C, automated Ki-67 quantification of vehicle versus drug-treated lungs at 1, 2, and 4 weeks. The percentage of Ki-67–positive nuclei was quantified using the Ariol Scanning System for the most severely affected lung lobe. 30,000 to 120,000 cells per lobe were counted. D, MRI quantification was performed as described in Materials and Methods. Each bar represents the percentage change of posttreatment versus baseline lung scan intensity. B–D, the asterisk indicates P < 0.05 compared with the average change in vehicle-treated lungs. Error bars, SE of the means. E, examples of MRI scans of lungs from doxycycline-induced control mice as well as baseline and posttreatment MRI for vehicle-treated and combination BIBW 2992 + rapamycin–treated mice. Pretreatment, 3 weeks of induction time; posttreatment, 4 weeks of treatment time postinduction. White areas indicate tumor obstruction of the airways, which increases over time in vehicle-treated mice.

(9). We also detected decreased cellular proliferation (by Ki-67 staining) and decrease in overall tumor burden (by hematoxylin and eosin and confirmed by total EGFR staining) in both single and combination treatments. Combination treatment resulted in the greatest decrease in total EGFR, phospho-EGFR, and Ki-67 staining, even after only 1 week of treatment (Supplementary Fig. S2). Automated quantification of Ki-67 staining supports the hypothesis that the strongest effect on proliferation is from BIBW 2992 and rapamycin dual treatment at all time points (Fig. 1C); however, the Ki-67 data indicates an increase in proliferation after 4 weeks even in the combination-treated mice. Immunoblotting confirmed phospho-EGFR or phospho-S6 reduction in lungs of BIBW 2992– or rapamycin-treated mice (respectively) at 4, 8, and 24 hours after single dose, as well as reduction of both in lungs of combination-treated mice (Supplementary Fig. S3). By 24 hours, reduction of phospho-EGFR was more variable, even in the combination-treated mice, indicating the need for continued treatment to achieve target suppression.

To attain another quantitative measure of treatment effects, baseline MRI measurements were compared with posttreatment tumors (Fig. 1D and E and Supplementary Data). In this model, signals on MRI reflect not only tumor burden, but also associated inflammation. After 1 week of treatment, BIBW 2992 alone resulted in tumor regression...
compared with baseline, whereas in the cohort treated for 2 weeks, tumor growth was inhibited by BIBW 2992 relative to vehicle, but regression was not observed. However, BIBW 2992 treatment together with rapamycin resulted in substantial tumor regression compared with baseline, evidenced by both 2 and 4 week posttreatment times.

Taken together, these results indicate that although BIBW 2992 successfully inhibits its primary target in vitro and elicits a short-term tumor regression response, it does not produce a sustained pathway and growth response unless combined with rapamycin. Importantly, maximal response for either single agent occurs early, with rebound apparent within a few weeks despite continuous treatment. The kinetics of such responses are thus critical in evaluating therapeutic efficacy. During the course of treatment, some tumor regions responded to drug and thus overall tumor burden decreased, which is reflected in the MRI measurements (note that Fig. 1E is an example of the maximum regression response observed). However, as discussed above, at the histopathology level small areas of the affected lungs (below the resolution level of the MRI scan) were observed that were either refractory to treatment or developed resistance as detected by pathology score and Ki-67 data. Endpoint tumor size in this case measures the sum of treatment responses and emerging resistant growth, underscoring the importance of temporal assessment in preclinical models for accurate response measurements.

**Gene expression reflects changes in oncogenic signaling in response to therapy over time**

To further explore the mechanism(s) of synergy in combination treatment, lungs from control E−R+, tumor-bearing E+R+ induced mice, and drug-treated E+R+ mice were analyzed for transcriptome changes. Gene expression data was subjected to supervised hierarchical clustering to compare profiles of untreated tumor-bearing mice to drug-treated for 1 or 4 weeks (Fig. 2A). As expected, vehicle-treated E+R+ samples had expression profiles similar to those of untreated E+R+ lung samples. The 1-week treated samples showed an almost universal suppression of gene expression compared with untreated or vehicle-treated E+R+ animals that did not distinguish between responders and nonresponders by pathology score (Fig. 2A). However, after 4 weeks there were clearly several samples in which the profile more closely matched the normal gene expression profile (E−R+ control). These lungs also had lower (less severe) pathology scores than the comparable vehicle-treated or untreated samples. Although most of the combination drug-treated tumors responded at the phenotypic level as seen by MRI scans and histopathology, not all of the samples seemed to be responders at the transcriptome level, indicating some heterogeneity within the tumors as discussed above. These data also point to the variability within treatments due to the mixed tumor cell population.

A single-sample GSEA of GO terms was conducted as a first step to reveal the biologic functions that differentiate normal (E−R+) from tumor-bearing (E+R+) lung and vehicle- from drug-treated tumors. Among others, GO terms involved in cell metabolism, mitosis, and DNA replication were differentially affected in EGFR tumors (Fig. 2B). We chose 4 genes in the signature (BUB1, TOP2A, CCNB2, and PLK1) for validation by quantitative PCR (qPCR; Supplementary Fig. S4). Of note, in a study of gene expression markers in human NSCLC, CCNB2 and TOP2A were part of a 187-gene signature set distinguishing normal tissues from malignant NSCLCs (14). All 4 genes were downregulated after treatment with BIBW 2992 or rapamycin, and the drug combination caused a further decrease in gene expression levels. Fold-change levels, compared with vehicle-treated controls, were consistently greater after 1 week compared with 2 weeks of drug treatment, again indicating a greater early treatment response on average. The 4-week gene expression data was a more accurate indicator of which individual tumors exhibited a sustained response. We conclude that multiple time points during the course of treatment are crucial to profiling the in vivo activity of cancer drugs.

**Tumor induction and drug response result in contrasting metabolomic alterations**

To evaluate additional biologic changes accompanying tumor induction and treatment that may correlate to gene expression responses, we conducted metabolic profiling of blood and lung tissue from induced E−R+ and E+R+ mice at various time points and from E+R+ mice treated with the described drug regimen for 4 weeks. Using a high-throughput small-molecule analytic platform (Metabolon, Inc.; see Supplementary Materials and Methods), a total of 457 biochemicals were detected in lung and 324 were detected in blood, including 238 compounds common to both lung and blood, 219 compounds unique to lung, and 86 compounds unique to blood (Supplementary Tables S1 and S2). Statistical comparisons were used to identify biochemicals whose relative levels differed between experimental groups (Supplementary Table S3 and Supplementary Materials and Methods). In lung tissue, there were extensive metabolomic alterations consistent with tumor development within 2 weeks of induction of the EGFR transgene, which were typically stronger at the terminal time point (data summarized in Fig. 3A). These changes were characterized by a shift to high anabolic activity, including diversion of glucose oxidation to the pentose phosphate pathway, elevated amino acid and nucleotide pools, higher levels of polyamines, glycerolipids, and long-chain fatty acids.

Lipid pathway metabolites were upregulated as a result of tumor induction, including fatty acids (used in cell membrane construction, energy metabolism, and as substrates for signaling molecules), and phospholipids (essential for membrane biosynthesis, Fig. 3A). For example, CDP-choline and intermediates to CDP-choline synthesis (choline and choline-phosphate) were increased in tumor tissue compared with control lung and decreased upon rapamycin or combination drug treatment. MTORC1 was recently shown to directly induce lipid biosynthesis (15) and therefore, decrease in these metabolites may indicate effective inhibition of mTOR by rapamycin.

Treatment with either BIBW 2992 or rapamycin tended to reverse the metabolic shifts observed during tumor induction in lung tissue; however, rapamycin (or the combination) caused a more dramatic reversal for the vast majority of metabolites than BIBW 2992 alone (Fig. 3A). The majority of
Figure 2. Supervised clustering shows correlation between tumor status, treatment, and pathology score and pathways. A, gene expression profiles of normal lung, lung tumors (2 and 4 weeks postinduction) from untreated EGFR-TL mice, and vehicle, BIBW 2992, rapamycin, or BIBW 2992 plus rapamycin–treated mice at indicated time points. At the bottom of the heat map are the corresponding pathology scores for each lung sample. B, single sample gene set enrichment analysis scores of samples from A. A positive enrichment score (red) indicates a positive correlation between genes in the gene set and the tumor sample expression profile; a negative enrichment score (blue) indicates the reverse (see also Materials and Methods).
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### A

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<th>Pathway</th>
<th>Metabolite</th>
<th>E-R+ 2w</th>
<th>T</th>
<th>B</th>
<th>R</th>
<th>B+R</th>
<th>E+R+ 2w</th>
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<th>V</th>
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### B

- N-acetyl aspartate (NAA)
- Glutathione, reduced (GSH)
- Proline
- Alanine
- Threonine
- N-acetylglutamine
- S-glutathione
- Asparagine
- Aspartate
- Asymmetric dimethylarginine (ADMA)
- Phenylalanine
- Proline
- Glutathione, oxidized (GSSG)
- Putrescine
- Arginine
- S-adenosylmethionine (SAM)
- Aspartate
- Tyrosine
- Leucine
- Methionine
- Isoleucine
- Glutamine
- Lysine
- Histidine
- Serine
- Leucine
- Cysteine-glutathione disulfide
- N-acetylornithine
- Cystine
- C-glutamylcysteine
- Tyrosine
- Homocysteine
- Citrulline
- Lactate
- Taurine
- N-acetylserotonin
- GABA

- Control lung
- untreated tumor (E+R+)
- B6W2593+Rapamycin-treated lung (E+R+)
metabolomic alterations were in pathways affected downstream of mTOR, for example amino acid synthesis. Individual amino acid pathway metabolites significantly altered in lung tumors as compared with control lung were plotted as Z-scores (Fig. 3B) to examine specific effects of the individual versus the combination treatments. The overall pattern of response is similar in all 3 treatment regimens, not unexpected as both drugs target nodes in the EGFR signaling pathway. Notably, citrate, an intermediate in the tricarboxylic acid (TCA) cycle, is significantly increased in BIBW 2992 and BIBW 2992 + rapamycin combination–treated lung tumors but not by rapamycin alone. As citrate is also an intermediate in fatty acid and phospholipid synthesis, it is possible that the general decrease in lipids upon drug treatment correlates to less diversion of citrate from the TCA cycle to membrane biosynthetic processes.

Thus, this lung cancer model reflects the abnormal metabolism expected in cancer cells, including aberrant nutrient uptake and increased macromolecular synthesis. Moreover, treatment with drugs targeting the EGFR signaling pathway results in reversal of these effects, highlighting the importance of identifying specific metabolites involved in the response.

### Table 1. Metabolite changes detectable in blood and lung tissue 2 weeks postinduction

<table>
<thead>
<tr>
<th>Subpathway</th>
<th>Metabolite</th>
<th>E + R+/E - R+ Blood (exp1)†</th>
<th>E + R+/E - R+ Blood (exp2)§</th>
<th>E + R+/E - R+ Lung (exp1)</th>
<th>E + R+/E - R+ Lung (exp2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alanine and aspartate</td>
<td>Aspartate</td>
<td>0.75 (P = 0.0036)</td>
<td>0.71 (P = 0.0054)</td>
<td>2.13 (P = 0.0551)</td>
<td>0.59 (P = 0.0086)</td>
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<td>Glutamate metabolism</td>
<td>Glutamine</td>
<td>1.14 (P = 0.0012)</td>
<td>1.32 (P = 0.0709)</td>
<td>1.52 (P = 0.0147)</td>
<td>1.58 (P = 0.001)</td>
</tr>
<tr>
<td>Glycine, serine, and</td>
<td>Threonine</td>
<td>1.55 (P &lt; 0.001)</td>
<td>1.57 (P = 0.0148)</td>
<td>2.50 (P = 0.0252)</td>
<td>1.7 (P = 0.0368)</td>
</tr>
<tr>
<td>Threonine metabolism</td>
<td>Serine</td>
<td>1.35 (P = 0.0437)</td>
<td>1.36 (P = 0.0006)</td>
<td>2.60 (P = 0.0359)</td>
<td>n.d.</td>
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<tr>
<td>Tryptophan metabolism</td>
<td>Tryptophan</td>
<td>1.40 (P &lt; 0.001)</td>
<td>1.45 (P = 0.0086)</td>
<td>3.31 (P = 0.0099)</td>
<td>1.8 (P = 0.0002)</td>
</tr>
<tr>
<td>Tryptophan metabolism</td>
<td>3-Indoxyl sulfate</td>
<td>1.51 (P = 0.0254)</td>
<td>1.47 (P = 0.0309)</td>
<td>1.62 (P = 0.0078)</td>
<td>n.d.</td>
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<tr>
<td>Urea cycle: arginine-,</td>
<td>Arginine</td>
<td>1.29 (P = 0.0062)</td>
<td>1.40 (P = 0.0254)</td>
<td>1.44 (P = 0.0225)</td>
<td>1.45 (P = 0.0014)</td>
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<td>proline-, metabolism</td>
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<tr>
<td>Carnitine metabolism</td>
<td>Deoxycarnitine</td>
<td>1.28 (P &lt; 0.001)</td>
<td>1.28 (P = 0.0057)</td>
<td>1.68 (P = 0.0857)</td>
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<td>Carnitine metabolism</td>
<td>Propionylcarnitine</td>
<td>1.16 (P &lt; 0.001)</td>
<td>1.19 (P = 0.0032)</td>
<td>2.39 (P = 0.0148)</td>
<td>n.d.</td>
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<tr>
<td>Lysolipid</td>
<td>1-Arachidoylglycerophosphocholine</td>
<td>0.28 (P = 0.0037)</td>
<td>0.34 (P = 0.0406)</td>
<td>n.d.</td>
<td>n.d.</td>
</tr>
<tr>
<td>Sterol/steroid</td>
<td>Corticosterone</td>
<td>2.41 (P &lt; 0.001)</td>
<td>2.24 (P = 0.2535)</td>
<td>2.67 (P = 0.0108)</td>
<td>n.d.</td>
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Abbreviation: n.d., not done.

†exp1: experiment 1, n = 8 E – R+, 8 E+R+ mice
§exp2: experiment 2, n = 15 E – R+, 15 E+R+ mice

Key metabolite signature observed in blood and tissue samples during lung tumorigenesis and treatment

In addition to lung tissue, we sampled blood from tumor-bearing animals to determine whether there were metabolic changes that could be used as potential biomarkers indicating the presence of a tumor. Blood was sampled from E – R+ and E+R+ mice at 2 weeks and at terminal stages postinduction for metabolic profiling. The unbiased screen recovered 26 metabolites with statistical significance of P ≤ 0.05. The changes after 2 weeks were more profound, possibly due to wasting at the terminal stages. Relative fold-changes were much lower in blood than tissue, most likely due to the dilution of organ-specific effects by the entire blood volume as well as the homeostatic mechanism, which changes or removes many biochemicals that were identified in blood samples from the first study exhibited similar trends to levels measured in samples collected for the validation study. We identified 11 metabolites from both studies that were significantly changed in blood as well as in the induced lung tissue (Table 1).

Figure 3. Widespread metabolomic alterations consistent with tumor development are found within 2 weeks of induction of the EGFR transgene and are reversed upon treatment. A, metabolite group means in lung tissue from control [E – R: 2 weeks (2w) and terminal (T)], tumored untreated [E – R: 2 weeks (2w) and terminal (T)] as well as tumored treated mice [E – R: vehicle-treated (V) and BIBW 2992 (B), rapamycin (R), or BIBW 2992 + rapamycin–treated (B + R)]. The heat map colors display each group mean level relative to the median for that compound, such that white represents the median, saturated red represents a 2-fold increase relative to the median, and saturated blue is one-half the median value. B, Z-score plot of amino acid pathway only from A. Each point represents 1 metabolite in 1 sample, expressed as the number of SDs from the mean of the control group for that comparison and colored by type (left, control E–R+ lung; yellow, tumored E+R+ lung; red, right, vehicle-treated lung tumors, yellow; BIBW 2992+ rapamycin–treated tumors, pink). For clarity, the plot was truncated at 20 SDs above the mean of the control and vehicle-treated samples, respectively.
Evidence of elevated amino acid pools observed in lung were reflected in blood. The branched chain amino acid conjugates propionyl-carnitine and deoxycarnitine were significantly increased in E+R+/ blood and lung tissue compared with control E–R++. These intermediates of valine, leucine, and isoleucine catabolism are indicative of oxidative stress or a shortage of coenzyme A and can build up when the pathway proceeds at rates in excess of downstream capacity to use metabolites in the TCA cycle. The amino acids glutamine, threonine, serine, arginine, and tryptophan were also increased (Table 1), suggestive of increased amino acid synthesis or enhanced protein turnover due to muscle wasting during cancer development. Unlike the other amino acids, aspartate was upregulated in lung and downregulated in blood. Using liquid chromatography-tandem mass spectrometry (LC-MS/MS) quantification, we verified that glutamine, tryptophan, and arginine were elevated in E+R+ blood and tumors compared with E–R++ control samples induced for 2 weeks, whereas aspartate was upregulated in lung only (Table 1; MS-targeted assay). In addition, we detected the levels of these amino acids in induced, tumor-bearing lungs after 2 weeks of BBW 2992, rapamycin, or combination drug treatment (Supplementary Fig. S5). As expected, drug treatment reversed the higher levels of the metabolites observed in the vehicle-treated tumors, though in some cases (aspartate, glutamate) the combination treatment did not decrease the levels beyond the individual treatments. This could be due to the natural variation in a smaller number of samples, or a difference in the 2 week treatment compared with the original analysis after 4 weeks of treatment. The amino acids identified are members of key pathways affected in the treatment response (see below) underscoring the potential significance of small changes in their relative quantities.

**Integration of metabolite and gene expression data highlights pathways essential to drug response**

While the examination of gene expression or metabolic signatures was informative, we speculated that combining the datasets could reveal the most significant changes. To test this, we conducted pathway enrichment analysis on metabolic and transcriptomic signatures. Gene expression and metabolite data from control E–R++ mice, induced E+R+ mice and mice treated with the drug combination were mapped together onto Kyoto Encyclopedia of Genes and Genomes (KEGG) pathways (Supplementary Fig. S6). Each dataset alone did not show significant enrichment in most pathways, but merging the 2 datasets points to key pathways in cancer cell energy production, such as the pentose phosphate pathway, glutathione metabolism, and glycolysis and gluconeogenesis that are differentially regulated in untreated versus treated tumors. There were 46 significantly enriched pathways that are upregulated in E+R+ tumors compared with E–R++ and of these, 29 were reversed in direction by drug treatment. Two of the pathways, "alanine, aspartate, and glutamate metabolism" and "glycolysis and gluconeogenesis" are also of note as, in a study of gene expression profiles in tumor samples from patients with lung adenocarcinoma, gene sets representing those pathways were enriched in patients with poor outcome (11).

Individual pathway maps are informative to highlight the concordance between the metabolites altered in key energy pathways and changes in expression of the enzymes that control production of individual metabolites. For example, both oxidized (GSSG) and reduced glutathione (GSH), as well as amino acids used to synthesize them, were elevated in tumor development (Fig. 4A). Antioxidant enzymes related to glutathione synthesis and coordinately regulated through antioxidant response elements (AREs) and NRF2 binding, such as gamma-glutamylcysteine synthetase (γ-GCS), glutathione reductase, and glutaredoxin were correspondingly upregulated in tumors (Fig. 4A). The glutathione pathway is involved in protection from oxidative stress, a characteristic of rapidly growing cells, by providing a substrate for neutralization of reactive oxygen species (ROS; ref. 16). ROS production and subsequent oxidative stress have been implicated specifically in carcinogenesis driven by oncogenes such as ARAS (17) and EGFR (18). Alternatively, ROS production can occur due to intratumoral hypoxic conditions, leading to oxidative stress, which causes the cells to oxidize GSH to GSSG and results in the activation of glutathione synthesis (19). A hypoxia-related gene signature was overexpressed in E+R+ versus E–R++ lung (Supplementary Fig. S7). For example, GLUT1, a target of hypoxia-inducible factor-1 and mediator of glucose uptake (20), was upregulated in tumors reflecting hypoxic conditions and correlated to an increase in glycolytic metabolites and other glycolytic enzymes (Fig. 4B). After treatment with the drug combination, oxidized, and reduced glutathione as well as the hypoxia-inducible factor-1 target gene signature were downregulated (Fig. 4B, Supplementary Fig. S7), indicating a decrease in energy needs in the tissue.

Glutamine levels were significantly lowered by BBW 2992 treatment or BBW 2992 + rapamycin, but not by rapamycin alone (Fig. 4A). Glutamine is the major nutrient consumed by cancer cells besides glucose and is a precursor of glutathione as well as a source of TCA cycle intermediates. Glutamine metabolism is increased by C-MYC, and downstream gene targets of C-MYC like the glycolytic enzyme enolase, are upregulated in
Figure 5. Oncogenic transcription factor activity is linked to downstream alteration of key metabolic pathways. A, gene set enrichment analysis of TFBS in expression profiles from normal lung, lung tumors from untreated EGFR-TL mice, and vehicle, BIBW 2992, rapamycin or BIBW 2992 plus rapamycin–treated EGFR-TL mice. Heat map shows significant enrichments of TFBS across indicated comparisons. Color scale depicts the logarithm of $P$ value (red, overrepresentation; blue, underrepresentation). B, leading edge analysis of metabolic genes that have TFBS in their promoters, in normal lung, lung tumors from untreated, and vehicle, BIBW 2992, rapamycin or BIBW 2992 plus rapamycin–treated EGFR-TL mice. Heat map shows relative expression levels of genes annotated to KEGG metabolic pathways across sample set. Black boxes (middle) represent the presence of an indicated TFBS in the promoters.
tumor versus control samples and reversed in drug-treated samples (Fig. 4B and see below).

Several markers of high glucose mobilization, including glucose-6-phosphate, glucose-1-phosphate, and UDP-N-acetyl-glucosamine, which is an important precursor for protein glycosylation and polysaccharide synthesis, were present in higher levels in E+R+ tumor tissue versus E+R− control lungs (Fig. 4B). While glucose itself is lower in tumor tissue, it is not a good indicator of tumor glucose levels as it may simply indicate binding of glucose to dead cells. The phosphorylated products fructose-6-phosphate (F-6-P) and fructose-1,6-bisphosphate (F-1,6-BP) are higher in tumors and phosphofructokinase (PFK), which catalyzes the conversion of F-6-P to F-1,6-BP, is correspondingly upregulated. There was also strong evidence for activation of the pentose phosphate pathway, which provides precursors for growth as well as the NADPH needed for various anabolic reactions (Fig. 4B). For example, expression of transketolase (TKT), which catalyzes the formation of sedoheptulose as well as the conversion of xylose-5-phosphate back to F-6-P in the pentose phosphate pathway, was oppositely regulated in tumors versus combination-treated lungs.

Of note, pyruvate and lactate, the oxidation product of pyruvate, were higher in tumor tissue, as was lactate dehydrogenase (LDH) gene expression. This is consistent with previous observations made in tumor tissue, in which pyruvate is converted to lactate (Warburg effect) avoiding energy-producing oxidative phosphorylation. Pyruvate was lowered only by the combination drug treatment, indicating that full suppression of the pathway may be necessary to inhibit glycolysis.

**Oncogenic transcription factor activity may be responsible for downstream alteration of key metabolic pathways**

Recent studies have suggested that oncoproteins can directly reprogram tumor cell metabolism, and several oncogenes and tumor suppressors have been linked to the regulation of downstream metabolic processes through transcription factor activity (21). To identify candidate transcription factors downstream of oncogenic EGFR activation in our model, we used GSEA and a library of gene sets containing genes with a cis-regulatory element and conserved DNA motifs, representing known or predicted regulatory elements in promoters and 3′-UTRs (13). This bioinformatic methodology was used to identify overrepresented transcription factor binding sites (TFBS) that were present in the promoters of the genes included in gene expression profiles of tumors from untreated E+R+ mice, and vehicle, BIBW 2992, rapamycin or BIBW 2992 plus rapamycin-treated E+R+ mice. Our analysis revealed a cluster of 12 transcription factors that were significantly enriched (FDR value < 0.05) in EGFR tumors compared with normal lung, for example MYC and SREBP1 (Fig. 5A). In contrast, after the first week of drug treatment, the targets of some of these transcription factors were underrepresented, for example NRF1 and NRF2.

To identify metabolic genes that drive the enrichment for each transcription factor gene set, we ran a leading edge analysis of GSEA results described above. By grouping those leading edge genes (i.e., genes with the highest enrichment scores in EGFR tumors) included in TFBS gene sets that were significantly underrepresented after drug treatments (GABP, NRF1, NRF2, SREBP, ELK1 and C-MYC), we were able to categorize them into metabolic processes. As shown in Fig. 5B, the strongest downregulation of metabolic genes activated in EGFR-TL tumors seemed to be after only 1 week of treatment, particularly in response to the combination of BIBW and rapamycin. Key genes, such as hexokinase 2 (HK2) and lactate dehydrogenase A (LDHA; glycolysis), and GPX1 and GSS (glutathione metabolism) were affected. After 4 weeks of treatment, the single-drug regimens were unable to repress most of the metabolic genes included in the analysis. Only the drug combination reduced gene expression levels more consistently, suggesting that transcriptional control of metabolic pathways through C-MYC and SREBP, as well as ELK1, NRF1 and NRF2, depends on both EGFR and mTORC1 signaling.

**Discussion**

Presently, human clinical trials using BIBW 2992 (afatinib) and rapamycin combination are ongoing (22, 23). The unbiased profiling approach described here highlights the utility of GEMs in rapidly identifying clinically testable hypotheses that might immediately impact the analysis of these ongoing clinical trials, and the design of subsequent human studies. To that point, we observed that several of the prominent pathways modulated during EGFR-driven tumorigenesis were reversed upon combination drug treatment, revealing possible reasons for increased efficacy.

Because of their controlled genetics and environment, mouse models represent ideal tools for the identification of candidate biomarkers. Recent work comparing the plasma proteome of mouse ovarian cancer models with human ovarian cancer cell lines and tissue samples from patients resulted in promising biomarker candidates (24); another study in a mouse pancreatic cancer model revealed proteins that could discriminate human pancreatic cancer sera from controls (25); and protein signatures have been found that distinguish between different types of lung cancer in the mouse (26). We previously identified metabolomic changes in a mouse model of serous epithelial ovarian cancer that overlapped with findings obtained from tissue of human patients with metastatic ovarian cancer (27, 28). As energy metabolism has long been known to be aberrant in cancer, in this study we conducted metabolomic profiling in the EGFR-TL mouse model to identify biochemical processes that might be perturbed both in tumor development and by drug treatment. By comparing tissue and blood metabolomic changes during tumor progression we identified disease-associated changes. Several of the same metabolites have recently been shown to be altered in human lung cancer tissue and sera from patients with squamous cell carcinoma and small cell lung cancer as well as adenocarcinoma (29), indicating fidelity of the mouse models in mimicking the human condition. Furthermore, they show that the molecular interactions between the tumor and host may be similar in these 2 species. Because the dilution factor represented by the movement of a compound from the tumor...
to blood is large, and at the same time other organs may be acting to clear or modify the compound, the ability to validate the blood markers by correlation to lung tumor tissue metabolites in the mouse model is crucial.

As seen in patients (29), murine lung cancer tissue contained higher levels of most amino acids compared with normal lung tissue, including higher levels of glutamine and tryptophan. It is remarkable that high levels of glutamine are maintained although large amounts of glutamine are consumed in cancer cells through glutamine metabolism. Notably, treatment with BIBW 2992 and the combination of BIBW 2992 and rapamycin downregulated the levels of glutamine in the lung tissue, indicating the potential importance of this key source of energy in maintaining cancer cell growth. This could also offer a clue to the synergistic effects of the drug treatment with rapamycin, as glutamine depletion can cause apoptosis in lung cells by affecting pathways that do not require mTOR activity; instead they are MYC-regulated (30). Glutamine can also maintain the TCA cycle when glucose levels are limiting. Growth factor signaling through the PI3K-Akt-mTOR pathway in EGFR-over-expressing mice may be further activated by higher glutamine levels, resulting in increased protein translation (31).

mTORC1, the target of rapamycin and a key regulator of cell growth and proliferation downstream of oncogenic signaling pathways, including EGFR, has been linked previously to cellular metabolism regulation through the transcriptional control of metabolic genes (32, 33), and other recent studies have also linked metabolism to specific oncogenic signaling pathways (34, 35). In our analysis, relating transcriptional control elements from the gene expression data to metabolic processes revealed functional connections between known oncogenes and metabolite changes in the lung tumors. After the first week of drug treatment, binding sites for several transcription factors enriched in tumor samples were underrepresented, for example NRF1 and NRF2. Both BIBW 2992 and rapamycin treatments result in a decrease in the target genes for NRF2 after 1 week, but after 4 weeks only, the dual treatment suppresses expression of those genes. NRF2 is a transcriptional activator of cytoprotective enzymes including those involved in glutathione synthesis, and has increased activation in lung cancer tissue and cell lines (36). Therefore, NRF2 activation of glutathione metabolism may be a previously unappreciated mediator of oncogenesis which is inhibited by the combination drug regimen used here.

It was recently observed that mTORC1 signaling increases flux through the pentose phosphate pathway in some measure through activation of SREBP1 (15). SREBP1 is a transcriptional regulator of genes involved in de novo lipid and sterol biosynthesis and is itself regulated by mTOR (32). Interestingly, although rapamycin alone can inhibit mTOR, the SREBP1 gene target set is underrepresented in rapamycin-treated lungs only after 4 weeks of treatment, and even more significantly in the combination-treated samples. Targets of C-MYC, a key transcription factor driver in metabolic processes in cancer, are enriched in E-/R+ tumors and underrepresented in rapamycin and combination-treated lungs after 1 or 4 weeks of drug treatment.

In summary, this study combines transcriptional and metabolic analyses of a complex in vivo model system to examine temporal responses after drug treatment. Examination of lung tumor progression in EGFR-overexpressing TKI-resistant mice as well as a drug regimen known to result in tumor regression revealed marked changes in gene expression and biochemical pathways during tumor growth that were in many cases reversed when treatment was effective, and were different depending on the specific treatment and duration of treatment. These studies can be used to guide future analyses of drug combinations for human disease with targeted agents aimed at resistance by simultaneous blockade of relevant pathways.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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References


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