Cancer is the disease in which the promise of drug and companion biomarker outputs from the human genome sequence has been best exemplified so far (1). Remarkable progress has been made in “drugging the cancer genome” and developing personalized or precision cancer medicine in which treatment is increasingly based on the genetic makeup of the individual tumor (2). Yet, we have arguably now reached a major inflection point. Thus, notwithstanding a veritable deluge of data from massively parallel sequencing and other large-scale omics initiatives, we remain far from achieving a complete systems biology understanding of human cancers (3). Furthermore, although many molecularly targeted cancer drugs have been approved, median survival is generally enhanced only by a matter of months and drug resistance is just as problematic as with traditional cytotoxic drugs—attributable to tumor heterogeneity, Darwinian selection of resistant clones, and biochemical feedback loops (4). New approaches are urgently needed to tackle drug resistance. A report in this issue of Cancer Discovery describes the first successful use of systematic, high-throughput, unbiased combinatorial screening of small-molecule agents to discover new and unexpected genotype-selective 2- and 3-drug cocktails with encouraging therapeutic activity in models of mutant BRAF and RAS melanoma resistant to the BRAF inhibitor vemurafenib and other treatments (5).

Despite initial hopes for single-agent therapies targeting addictions to unique driver oncogenes, best exemplified by the BCR-ABL inhibitor imatinib in chronic myeloid leukemia, it is increasingly clear that the challenges posed by genetic and phenotypic heterogeneity within individual tumors mean that the use of intelligent targeted drug combinations, longitudinally adapted to meet the changing biology of the cancer, will be the only way to circumvent drug resistance and hence achieve long-term remission and potential cure (4, 6). But how do we identify the best drug combination for a particular cancer genotype?

The classical pragmatic approach to clinical cytotoxic drug cocktails was to combine agents with different molecular mechanisms, aiming to enhance antitumor effects and reduce cross-resistance while avoiding overlapping toxicity (4). Commercial factors and regulatory approval strategies also have an impact. These considerations can lead to relatively ad hoc combinations that fail to exploit our growing molecular knowledge. However, the high number of potential drug combinations precludes their thorough evaluation in preclinical models, let alone clinically.

The common 40-digit combination lock generates 64,000 permutations, which is why deciphering the code is essentially impossible if it is forgotten. For the approximately 250 approved cancer drugs, 3,125 2-way combinations and 2,573,000 3-way combinations are possible, whereas for the approximately 1,200 developmental oncology drugs, the respective numbers are 719,000 and 287,280,400 (4). Viewed in another way, 7,626 2-way and 310,124 3-way potential combinations are possible for our estimate of 124 conventionally druggable targets among the 479 genes reported to have causal genetic alterations in cancer (4). How can we reduce these prohibitively large numbers to actionable clinical combinations? The main options currently are as follows: (i) hypothesis-driven approaches based on our increasing knowledge of the molecular basis of drug resistance; or (ii) large-scale strategies centered on in silico modeling, genetic technologies such as synthetic lethal screens, and unbiased combinatorial drug screens (4).

In the new work, Held and colleagues (5) used large-scale unbiased combinatorial drug screening to identify cocktails active against melanomas with activating BRAF or RAS mutations, including those with intrinsic or acquired resistance to the approved breakthrough BRAF inhibitor vemurafenib—a major problem that is now seen routinely in the clinic (7, 8). Their workflow is shown in Fig. 1A.

First, to establish an unbiased chemical library for the initial single-agent screen, Held and colleagues (5) curated...
**Figure 1.** Study workflow for unbiased combinatorial screening, leading to effective genotype-selective combinations discovered by Held and colleagues (5), and the emerging pathways and interaction network derived from their data. **A,** a mechanistically broad library of 150 small-molecule compounds was screened against 28 early-passage human melanoma cell lines, including mutant *BRAF* or *RAS* and wild type (WT). A representative set of 40 compounds that were broadly or selectively active against the melanoma panel were chosen for the combinatorial screen. The most synergistic combinations were carried through for further confirmatory analyses, including elucidation of target engagement and pathway modulation; confirmation of on-target activity through the use of alternative chemotype compounds against the same target; identification of a promising triple combination in *BRAF*-mutant cells; exploration of avoidance of combination-resistant clones arising; and finally, in vivo demonstration of the promising therapeutic effect of a selected 2-drug combination in a mutant NRAS melanoma xenograft in nude mice. **B,** signaling pathways involved in the effective triple combination active in vemurafenib-resistant *BRAF*-mutant cells. The combination of the *BRAF, EGFR/ERBB2,* and *AKT* inhibitors was found to be highly synergistic and most effective in overcoming resistance when compared with 2-agent combinations. Resistant clones could not be generated when melanoma cells were exposed to the triple-drug regimen. Biomarker data suggest that the triple combination is very effective at reducing signaling output in the phosphoinositide 3-kinase (PI3K)-*AKT* pathway and the mutant *BRAF*-MEK-ERK MAPK pathway. Dotted lines represent omitted pathway components. RAF homo- and heterodimers are not shown for simplicity. **C,** 2-drug combinations found to be effective in NRAS-mutant cells exploit both signaling and metabolic pathways. HMGCR is a key component of the mevalonate pathway, leading to the production of geranyl-geranyl pyrophosphate, required for the prenylation of NRAS, and hence for localizing NRAS proximally to the plasma membrane. Inhibition of prenylation through the inhibition of HMGCR alone did not have a major growth inhibition effect on *RAS*-mutant cells; however, combining it with either an HSP90 molecular chaperone inhibitor or a pan-cyclin-dependent kinase (CDK) inhibitor gave significant synergy. It is likely that the synergy seen within NRAS-mutant melanomas involves the cooperative effects of blocking the essential NRAS prenylation through HMGCR inhibition, thus preventing membrane localization and reducing NRAS-RAF-MEK MAPK signaling output, coupled with depletion of RAF isoforms, CDK4 and other relevant HSP90 clients in the case of 17-DMAG, or the inhibition of cell cycle and transcriptional kinases for flavopiridol. Dotted lines represent omitted pathway components; brown hexagons represent metabolites. (continued on following page)
150 small molecules selected to represent diverse biologic mechanisms. One-third comprised a wide range of kinase inhibitors, whereas 20% were nontargeted or broadly cytotoxic agents. The screening library contained approved drugs plus compounds under clinical investigation and chemical tools. High-throughput screening (HTS) was carried out with cell-growth inhibition as the end point, testing a range of concentrations against a panel of 28 patient-derived early-passage melanoma cell lines harboring either BRAF- or RAS-activating mutations or WT for both oncogenes (Fig. 1A, left).

To identify whether any tested agents exhibited selective effects in a genotype-specific pattern, Held and colleagues (5)
clustered the compounds based on their growth inhibition profiles and classified them into 3 major categories: those with little effect across the whole melanoma panel, those active across the majority of the panel, and those showing clear differential effects.

Of obvious interest, certain single-agent compounds were selective toward BRAF-mutant cells. As expected (9), these included the BRAF inhibitors vemurafenib, PLX4720, and GDC-0879. However, growth inhibition was often incomplete, and Held and colleagues (5) also confirmed previous observations that a significant fraction (25%) of BRAF-mutant cell lines exhibited innate resistance to vemurafenib, whereas growth stimulation was seen in some mutant RAS and WT lines (10). In addition, loss of PTEN or RB1, known vemurafenib resistance mechanisms, was not necessary for reduced sensitivity, consistent with a role for additional resistance factors such as MET/HGFR (6, 8). These various results emphasize the need for new therapies or combinations in melanomas of all genotypes.

Additional compounds showing selectivity toward mutant BRAF melanomas included the SRC/ABL inhibitor bosutinib, the FGFR inhibitor dovitinib, and the EGFR inhibitor gefitinib—all effectively inactive against BRAF, indicating that sensitivity does not derive from off-target BRAF inhibition. Two MEK inhibitors were active in both mutant BRAF and RAS melanomas, especially the former. Interestingly, melanoma lines resistant to BRAF inhibitors (including the 25% of mutant BRAF cultures with intrinsic resistance) were also resistant to MEK inhibitors. Most intriguingly, the approved 3-hydroxy-3-methylglutaryl-CoA reductase (HMGCR) inhibitor simvastatin, a cholesterol-lowering agent which indirectly blocks prenylation and essential membrane localization of RAS, was the only single agent showing a trend to greater growth inhibition in RAS-mutant melanomas, albeit at high concentrations.

Of note, no compounds selective for the WT cells were found, likely due to the genetic heterogeneity of this group.

The rather lackluster performance of compounds in the single-agent screen, and especially the relatively poor activity of compounds against the RAS-mutant and WT melanomas, further highlighted the need to identify effective combinations. In the second phase of their workflow, Held and colleagues (5) selected 40 representative compounds, chosen because of their broad growth-inhibitory effect or genotype-specific profiles, as well as representing the diverse molecular and biologic mechanisms of the full compound library, and carried these through to the combinatorial HTS (cHTS) campaign (Fig. 1A, middle). In total, they examined more than 7,000 pairwise combinations of the 40 agents, with 3 concentrations per drug, against 19 representative melanoma cell lines, comprising 8 BRAF-mutant, 5 NRAS-mutant, one HRAS-mutant, and 5 WT lines.

The cHTS identified many new, highly effective, and biologically interesting genotype-selective 2-drug combinations. Surprisingly, the selective combinations revealed for BRAF-mutant melanomas (1,021) massively outnumbered those selective for either RAS-mutant (75) or WT cells (22). Combinations found to be selective for BRAF-mutant cells involved a range of primary targets, albeit dominated by protein kinase inhibitors such as vemurafenib, the SRC/ABL inhibitor bosutinib, the FGFR inhibitor PD-173074, the MET inhibitor PHA-665752, the AKT inhibitor MK-2206, the ERBB2/EGFR inhibitor lapatinib, and the relatively unselective vascular endothelial growth factor receptor (VEGFR) inhibitor sunitinib, together with the proposed pan-BCL2 inhibitor obitoclax.

Remarkably, the HMGCR inhibitor simvastatin was involved in half of the 2-drug combinations selective against RAS-mutant melanomas. In contrast to the combination pairs in BRAF-mutant–selective combinations that were dominated by protein kinase inhibitors, simvastatin was effective against mutant RAS melanomas in combination with a far wider mechanistic range of compounds, including, particularly, inhibitors of the molecular chaperone HSP90 and the ubiquitin ligase XIAP, as well as those for MEK, paxilin-dependent kinases (CDK), and IGF1R. No combinations were found to be selective for WT melanomas, but some combinations were effective across all 3 melanoma genotypes, the strongest of which was the combination of a STAT inhibitor (static) with FAK inhibitor 14.

Moving on to the third phase of the study, Held and colleagues (5) characterized promising 2-way combinations in greater biologic detail, carried out mechanistic biomarker studies to show target engagement and pathway inhibition, tested additional compounds acting on highlighted targets, progressed promising members of effective 2-drug combinations into triple combination studies for mutant BRAF melanoma lines, and assessed ability to suppress development of resistant clones (Fig. 1A, right).

To identify the 2-way combinations with the most added value, Held and colleagues (5) looked for additive, synergistic, or antagonistic interactions according to the Bliss independence method. Intriguingly, most combinations selective toward BRAF-mutant cells showed additivity or synergy, whereas most combinations in RAS-mutant melanomas were formally antagonistic; that is, although effective, they were less active than predicted from the sum of single-agent sensitivities.

Among the many effective combinations in BRAF-mutant melanomas, the strongest 2-drug efficacies and synergies observed were between the allosteric AKT inhibitor MK-2206 combined with either the ERBB2/EGFR inhibitor lapatinib or the SRC/ABL inhibitor bosutinib. RAS-mutant (and WT) melanomas were generally less sensitive and the most efficacious and synergistic 2-way combinations for RAS-mutant melanomas involved the HMGCR inhibitor simvastatin plus either the HSP90 molecular chaperone inhibitor 17-DMAG or the pan-CDK inhibitor flavopiridol. It was reassuring that combinations using alternative compounds acting against the same targets gave similar synergistic effects, increasing confidence that the synergy derives from on-target, not off-target, modulation.

Importantly, Held and colleagues (5) identified a key triple combination, involving BRAF+EGFR+AKT inhibition, which was highly effective in BRAF-mutant melanomas and overcame vemurafenib resistance. Vemurafenib sensitivity was reestablished by blocking EGFR and AKT, with biomarker studies indicating that this was most likely because the combination suppressed both the phosphoinositide-3 kinase (PI3K) and RAS–RAF–MEK–ERK MAPK pathways, which makes obvious mechanistic sense (Fig. 1B). The results are consistent with the known effects of EGFR inhibition in blocking the feedback activation loop between MEK and EGFR (potentially via CDC25C) which activates CRAF and MEK following inhibition of mutant BRAF in colorectal cancer.
greater antitumor activity over the single agents. Further
nude mice. It was encouraging that the 2-drug combination
the effects of a triple combination of simvastatin, flavopiridol,
of the 3-drug combination of BRAF+EGFR+AKT inhibitors
cially interesting to pursue
combinations that merit progression toward clinical investigation
Thus, the synergistic effects of simvastatin in the combination
are likely mediated by reduced NRAS activity, which may also
apply to the other HMGC-R-containing combinations, for
example, with the HSP90 inhibitor 17-DMAG (Fig. 1C). The
beneficial interactions seen with HMGC-R inhibition in NRAS-
mutant melanomas may be hypothesized to involve the coop-
erative effects of blocking NRAS–RAF–MEK MAPK signaling
output coupled to the depletion of RAF isoforms, CDK4, and
other relevant HSP90 clients in the case of 17-DMAG or the
inhibition of cell cycle and transcriptional kinases for flavopri-
dol (Fig. 1C). The potentially interesting 3-drug combination
for NRAS-mutant melanomas was not investigated.

Finally, Held and colleagues (5) translated one exemplar key
finding into an in vivo NRAS-mutant melanoma xenograft in
nude mice. It was encouraging that the 2-drug combination
of simvastatin plus flavopiridol was well tolerated and showed
greater antitumor activity over the single agents. Further in vivo
animal model studies are now required, including evaluation of
the 3-drug combination of BRAF+EGFR+AKT inhibitors
identified in vitro for BRAF-mutant melanomas. Investigation of
the effects of a triple combination of simvastatin, flavopiridol,
and 17-DMAG in RAS-mutant melanoma would also be espe-
cially interesting to pursue in vivo (Fig. 1C), as would additional
detailed exploration of the mechanisms underlying the observed
selective combinations. Our own initial analysis of the network
of the targets of drugs for which the combinations showed the
highest efficacy and synergy in BRAF- or NRAS-mutant melano-
mas illustrates connected but distinct processes involved in the
combinatorial treatment of the 2 genotypes (Fig. 1D).

The new work illustrates the power of systematic large-scale
combinatorial screening of unbiased 2-way combinations for
discovering novel therapeutically actionable genotype-selective
combinations that merit progression toward clinical investigation
in mutant BRAF and RAS melanoma, including vemurafenib-
resistant disease. Significantly, the effective combinations
were not predicted by single-agent screening. Of special note
are the 2-drug combination of statins and pan-CDK inhibi-
tors (with additional potential of HSP90 inhibition) for RAS-
murant, and the 3-drug regimen of vemurafenib plus EGFR
and AKT inhibition for BRAF-mutant melanoma. In another
recent large-scale compound screen of pairwise combinations of
300 compounds in 9 established melanoma cell lines, Roller
and colleagues (12) did not find synergistic interactions
specific for BRAF, RAS, or WT genotypes but did identify
robust synergy across all genotypes between the multikinase
inhibitor sorafenib and the nonsteroidal anti-inflammatory
drug (NSAID) diclofenac, with drug substitution experiments
indicating combined effects on MAPK and cyclooxygenase
(COX). Differences seen may reflect the distinct properties of
early-passage versus established melanoma lines. The combi-
natorial small-molecule screening approach complements the
use of genome sequencing of vemurafenib-resistant melanoma,
examination of signaling crosstalk and feedback loops, and
unbiased genome- and kinase-wide loss-of-function RNAi
screening—which have also led to clinically actionable combi-
nations such as BRAF plus MEK or receptor tyrosine kinase
inhibitors (4, 6, 8, 11)– and certainly yielded some unexpected
combinations. Especially given the eye-wathering number of
possible drug–genotype combinations, rapid progress in the
search for the best genotype-selective combinations to over-
come resistance in multiple cancer types, which will likely
reveal many unpredicted results, as here, will require glo-
bal collaborations and data sharing (13). Furthermore, some
combinatorial targets revealed by unbiased screens will not
have clinical drugs available, emphasizing the importance
of systematic approaches to extending the druggable cancer
genome on the way to achieving the reality of combinatorial
precision cancer medicine (2, 14).

Disclosure of Potential Conflicts of Interest

P. Workman and B. Al-Lazikani are employees of The Institute of
Cancer Research which has a commercial interest in the discovery
and development of anticancer drugs, including kinase inhibitors,
and operates a Rewards to Inventors scheme. P. Workman is a former
employee of AstraZeneca and declares commercial interactions with
Yamanouchi (now Astellas), Piramed Pharma (acquired by Roche),
Genentech, Vernalis, Novartis, Chroma Therapeutics, Astex Phar-
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Unpicking the Combination Lock for Mutant BRAF and RAS Melanomas

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