

ORIGINAL ARTICLE

Improved Survival with Vemurafenib in Melanoma with BRAF V600E Mutation

Paul B. Chapman, M.D., Axel Hauschild, M.D., Caroline Robert, M.D., Ph.D., John B. Haanen, M.D., Paolo Ascierto, M.D., James Larkin, M.D., Reinhard Dummer, M.D., Claus Garbe, M.D., Alessandro Testori, M.D., Michele Maio, M.D., David Hogg, M.D., Paul Lorigan, M.D., Celeste Lebbe, M.D., Thomas Jouary, M.D., Dirk Schadendorf, M.D., Antoni Ribas, M.D., Steven J. O'Day, M.D., Jeffrey A. Sosman, M.D., John M. Kirkwood, M.D., Alexander M.M. Eggermont, M.D., Ph.D., Brigitte Dreno, M.D., Ph.D., Keith Nolop, M.D., Jiang Li, Ph.D., Betty Nelson, M.A., Jeannie Hou, M.D., Richard J. Lee, M.D., Keith T. Flaherty, M.D., and Grant A. McArthur, M.B., B.S., Ph.D., for the BRIM-3 Study Group*

ABSTRACT

BACKGROUND

Phase 1 and 2 clinical trials of the BRAF kinase inhibitor vemurafenib (PLX4032) have shown response rates of more than 50% in patients with metastatic melanoma with the BRAF V600E mutation.

METHODS

We conducted a phase 3 randomized clinical trial comparing vemurafenib with dacarbazine in 675 patients with previously untreated, metastatic melanoma with the BRAF V600E mutation. Patients were randomly assigned to receive either vemurafenib (960 mg orally twice daily) or dacarbazine (1000 mg per square meter of body-surface area intravenously every 3 weeks). Coprimary end points were rates of overall and progression-free survival. Secondary end points included the response rate, response duration, and safety. A final analysis was planned after 196 deaths and an interim analysis after 98 deaths.

RESULTS

At 6 months, overall survival was 84% (95% confidence interval [CI], 78 to 89) in the vemurafenib group and 64% (95% CI, 56 to 73) in the dacarbazine group. In the interim analysis for overall survival and final analysis for progression-free survival, vemurafenib was associated with a relative reduction of 63% in the risk of death and of 74% in the risk of either death or disease progression, as compared with dacarbazine ($P < 0.001$ for both comparisons). After review of the interim analysis by an independent data and safety monitoring board, crossover from dacarbazine to vemurafenib was recommended. Response rates were 48% for vemurafenib and 5% for dacarbazine. Common adverse events associated with vemurafenib were arthralgia, rash, fatigue, alopecia, keratoacanthoma or squamous-cell carcinoma, photosensitivity, nausea, and diarrhea; 38% of patients required dose modification because of toxic effects.

CONCLUSIONS

Vemurafenib produced improved rates of overall and progression-free survival in patients with previously untreated melanoma with the BRAF V600E mutation. (Funded by Hoffmann–La Roche; BRIM-3 ClinicalTrials.gov number, NCT01006980.)

The authors' affiliations are listed in the Appendix. Address reprint requests to Dr. Chapman at the Department of Medicine, Memorial Sloan-Kettering Cancer Center, 1275 York Ave., New York, NY 10065, or at chapmanp@mskcc.org.

Drs. Flaherty and McArthur contributed equally to this article.

*Members of the BRAF Inhibitor in Melanoma 3 (BRIM-3) study group are listed in the Supplementary Appendix at NEJM.org.

This article (10.1056/NEJMoal103782) was published on June 5, 2011, and updated on March 13, 2012, at NEJM.org.

N Engl J Med 2011;364:2507-16.

Copyright © 2011 Massachusetts Medical Society.

METASTATIC MELANOMA HAS A POOR prognosis, with the median survival for patients with stage IV melanoma ranging from 8 to 18 months after diagnosis, depending on the substage.¹ In the United States last year, 8700 deaths from melanoma were projected, with an estimated rate of death of 2.6 in 100,000.² Rates of death from melanoma in Australia and New Zealand are slightly higher (3.5 in 100,000), whereas rates in Western Europe are slightly lower (1.8 in 100,000).³

In phase 3 studies, dacarbazine, the only chemotherapeutic agent approved by the Food and Drug Administration for the treatment of metastatic melanoma, was associated with a response rate of 7 to 12% and a median overall survival of 5.6 to 7.8 months after the initiation of treatment.⁴⁻⁷ Although higher response rates can be achieved with combination chemotherapy, these combinations have not resulted in improved rates of overall survival. Recently, the use of ipilimumab, a monoclonal antibody that blocks cytotoxic T-lymphocyte–associated antigen 4 (CTLA4) on lymphocytes, has been associated with improved overall survival, as compared with a peptide vaccine,⁸ and in combination with dacarbazine has been associated with better overall survival than dacarbazine alone.⁹

Approximately 40 to 60% of cutaneous melanomas carry mutations in *BRAF* that lead to constitutive activation of downstream signaling through the MAPK pathway.^{10,11} Approximately 90% of these mutations result in the substitution of glutamic acid for valine at codon 600 (*BRAF* V600E), although other activating mutations are known (e.g., *BRAF* V600K and *BRAF* V600R).

Vemurafenib (PLX4032) is a potent inhibitor of mutated *BRAF*.¹² It has marked antitumor effects against melanoma cell lines with the *BRAF* V600E mutation but not against cells with wild-type *BRAF*.¹²⁻¹⁴ A phase 1 trial established the maximum tolerated dose to be 960 mg twice daily and showed frequent tumor responses.¹⁵ A phase 2 trial involving patients who had received previous treatment for melanoma with the *BRAF* V600E mutation showed a confirmed response rate of 53%, with a median duration of response of 6.7 months.¹⁶ We conducted a randomized phase 3 trial to determine whether vemurafenib would prolong the rate of overall or progression-free survival, as compared with dacarbazine.

METHODS

PATIENTS

All patients in our study had unresectable, previously untreated stage IIIC or stage IV melanoma that tested positive for the *BRAF* V600E mutation on real-time polymerase-chain-reaction assay (Cobas 4800 *BRAF* V600 Mutation Test, Roche Molecular Systems). The test was performed at one of five central laboratories in the United States, Germany, and Australia. In approximately one third of the patients, *BRAF* was sequenced retrospectively by Sanger and 454 sequencing at a central laboratory. Other inclusion criteria were age of 18 years or older, a life expectancy of 3 months or longer, an Eastern Cooperative Oncology Group (ECOG) performance status of 0 (fully active and able to carry on all performance without restriction) or 1 (restricted in physically strenuous activity but ambulatory and able to carry out work of a light or sedentary nature), and adequate hematologic, hepatic, and renal function. Patients were excluded if they had a history of cancer within the past 5 years (except for basal- or squamous-cell carcinoma of the skin or carcinoma of the cervix) or metastases to the central nervous system, unless such metastases had been definitively treated more than 3 months previously with no progression and no requirement for continued glucocorticoid therapy. Concomitant treatment with any other anticancer therapy was not allowed.

The protocol was approved by the institutional review board at each participating institution and was conducted in accordance with the ethical principles of the Declaration of Helsinki and within the Good Clinical Practice guidelines, as defined by the International Conference on Harmonization. All patients provided written informed consent before enrollment.

STUDY DESIGN AND TREATMENT

From January 2010 through December 2010, a total of 2107 patients underwent screening at 104 centers in 12 countries worldwide. The most common reason for screening failure was a negative test for the *BRAF* V600 mutation. A total of 675 patients were randomly assigned in a 1:1 ratio to receive either vemurafenib (at a dose of 960 mg twice daily orally) or dacarbazine (at a dose of 1000 mg per square meter of body-surface area by intravenous infusion every 3 weeks) (Fig. A in the Supplementary Appendix, available with the full text of this article at NEJM.org). These pa-

tients included 20 with non-V600E mutations (19 with V600K and 1 with V600D), as identified on Sanger and 454 sequencing. Baseline characteristics of the patients were well balanced (Table 1).

Study patients were stratified according to American Joint Committee on Cancer stage (IIIC, M1a, M1b, or M1c), ECOG performance status (0 or 1), geographic region (North America, Western Europe, Australia or New Zealand, or other region), and serum lactate dehydrogenase level (normal or elevated).

Dose reductions for both vemurafenib and dacarbazine were prespecified for intolerable grade 2 toxic effects or worse. The development of cutaneous squamous-cell carcinoma did not require dose modification. The administration of vemurafenib was interrupted until the resolution of the toxic effect to at least grade 1 and restarted at 720 mg twice daily (480 mg twice daily for grade 4 events), with a dose reduction to 480 mg twice daily if the toxic effects recurred. If the toxic effect did not improve to grade 1 or lower or recurred at the 480-mg twice-daily dose, treatment was discontinued permanently. The administration of dacarbazine was interrupted for grade 3 or 4 toxic effects and could be restarted on recovery within 1 week to grade 1 (at full dose) or grade 2 (at 75% dose) or at 75% dose for grade 4 neutropenia or febrile neutropenia. A second dose reduction was allowed, if needed. Antiemetics and granulocyte colony-stimulating factor were administered according to standards at each study center. Treatment was discontinued on disease progression unless continued treatment was in the best interest of the patient in the judgment of the investigator and the sponsor.

ASSESSMENTS

At baseline, patients underwent computed tomography with contrast material or magnetic resonance imaging of the brain, chest, abdomen, pelvis, and other anatomical regions, as clinically indicated. Patients also underwent physical and dermatologic examinations and electrocardiography. Patients were examined every 3 weeks; tumor assessments were performed at baseline, at weeks 6 and 12, and every 9 weeks thereafter. Tumor responses were determined by the investigators according to the Response Evaluation Criteria in Solid Tumors (RECIST), version 1.1. Electrocardiograms were repeated every other cycle. Blood counts, biochemical analyses, and measurements of lactate dehydrogenase levels were performed at each visit.

Table 1. Baseline Demographic and Clinical Characteristics of Patients in the Intention-to-Treat Population.*

Characteristic	Vemurafenib (N = 337)	Dacarbazine (N = 338)
Median age (range) — yr	56 (21–86)	52 (17–86)
Male sex — no. (%)	200 (59)	181 (54)
White race — no. (%)†	333 (99)	338 (100)
Geographic region — no. (%)		
Australia or New Zealand	39 (12)	38 (11)
North America	86 (26)	86 (25)
Western Europe	205 (61)	203 (60)
Other	7 (2)	11 (3)
ECOG performance status — no. (%)‡		
0	229 (68)	230 (68)
1	108 (32)	108 (32)
Extent of metastatic melanoma — no. (%)§		
M1c	221 (66)	220 (65)
M1b	62 (18)	65 (19)
M1a	34 (10)	40 (12)
Unresectable IIIC	20 (6)	13 (4)
Lactate dehydrogenase — no. (%)¶		
≤Upper limit of the normal range	142 (42)	142 (42)
>Upper limit of the normal range	195 (58)	196 (58)

* Percentages may not total 100 because of rounding.

† Race was self-reported.

‡ An Eastern Cooperative Oncology Group (ECOG) performance status of 0 indicates that the patient is fully active and able to carry on all predisease activities without restriction; an ECOG performance status of 1 indicates that the patient is restricted in physically strenuous activity but is ambulatory and able to carry out work of a light or sedentary nature, such as light housework or office work.

§ The M1a stage is characterized by metastasis of the tumor to the skin, subcutaneous tissues, or distant lymph nodes, with a normal lactate dehydrogenase level; M1b by metastasis to the lung, with a normal lactate dehydrogenase level; and M1c by metastasis to any other visceral site or to any site, with an elevated lactate dehydrogenase level. In unresectable stage IIIC disease, melanoma has spread to at least three lymph nodes, which are enlarged because of the cancer.

¶ The upper limit of the normal range varied according to the reference values at each study center.

Adverse events were graded according to the National Cancer Institute's Common Terminology Criteria for Adverse Events, version 4.0. Monitoring of adverse events continued for up to 28 days after the last dose of a study drug had been administered or until any ongoing event resolved or stabilized. An independent data and safety monitoring board provided oversight and evaluated interim results on efficacy data.

STUDY OVERSIGHT

The trial was designed jointly by the senior academic authors and representatives of the sponsor, Hoffmann–La Roche. Data were collected by

the sponsor and analyzed in collaboration with the senior academic authors, who vouch for the completeness and accuracy of the data and analyses and for the conformance of this report to the protocol, as amended. The corresponding academic author prepared an initial draft of the manuscript in collaboration with the sponsor. All the authors contributed to subsequent drafts and made the decision to submit the manuscript for publication. The protocol and statistical analysis plan are available at NEJM.org.

STATISTICAL ANALYSIS

The original primary end point was the rate of overall survival. The statistical plan was revised in October 2010 on the basis of phase 1 and 2 efficacy and safety results and after consultation with global regulatory authorities. Under the revised plan, the rates of overall survival and progression-free survival were coprimary end points. The final analysis was planned after 196 deaths, and an interim analysis was planned after 50% of the projected deaths had occurred (Pocock boundary, $P \leq 0.028$ at the interim analysis and $P \leq 0.0247$ at the final analysis by the log-rank test). According to the revised plan, the final analysis of progression-free survival would be performed at the time of the interim analysis of overall survival. Secondary end points included the confirmed response rate, duration of response, and time to response.

The trial was designed for 680 patients to be randomly assigned to receive either vemurafenib or dacarbazine. The trial had a power of 80% to detect a hazard ratio of 0.65 for overall survival with an alpha level of 0.045 (an increase in median survival from 8 months for dacarbazine to 12.3 months for vemurafenib) and a power of 90% to detect a hazard ratio of 0.55 for progression-free survival with an alpha level of 0.005 (an increase in median survival from 2.5 months for dacarbazine to 4.5 months for vemurafenib). Survival was defined as the time from randomization to death from any cause. Progression-free survival was the time from randomization to documented disease progression or death. We used a two-sided unstratified log-rank test to compare survival rates in the two study groups. Hazard ratios for treatment with vemurafenib, as compared with dacarbazine, were estimated with the use of unstratified Cox regression. We estimated event-time distributions using the Kaplan–Meier method. All reported P values are two-sided, and

confidence intervals are at the 95% level. Descriptive statistics are used for adverse events.

This report is based on data as of December 30, 2010. Efficacy analyses were performed in the intention-to-treat population. In order to ensure adequate follow-up for each efficacy end point, patients could be evaluated for the analysis of overall survival, progression-free survival, and confirmed response if they had undergone randomization at least 2, 9, and 14 weeks, respectively, before the cutoff date. The safety analysis was performed in all patients who received a study drug and who had undergone at least one assessment during the study.

RESULTS

PATIENTS AND TREATMENTS

A total of 118 patients had died at the time of the interim analysis. The data and safety monitoring board determined that both the overall survival and progression-free survival end points had met the prespecified criteria for statistical significance in favor of vemurafenib. The board recommended that patients in the dacarbazine group be allowed to cross over to receive vemurafenib, and the protocol was amended accordingly on January 14, 2011. Median follow-up for the interim analysis was 3.8 months for patients in the vemurafenib group and 2.3 months for those in the dacarbazine group.

EFFICACY

A total of 672 patients were evaluated for overall survival. The hazard ratio for death in the vemurafenib group was 0.37 (95% confidence interval [CI], 0.26 to 0.55; $P < 0.001$) (Fig. 1A). The survival benefit in the vemurafenib group was observed in each prespecified subgroup, according to age, sex, ECOG performance status, tumor stage, lactate dehydrogenase level, and geographic region (Fig. 1B). At the time of the interim analysis, there were an inadequate number of patients in follow-up beyond 7 months in either study group to provide reliable Kaplan–Meier estimates of the survival curves.¹⁷ At 6 months, overall survival was 84% (95% CI, 78 to 89) in the vemurafenib group and 64% (95% CI, 56 to 73) in the dacarbazine group. Further follow-up is required.

Progression-free survival could be evaluated in 549 patients. The hazard ratio for tumor progression in the vemurafenib group was 0.26 (95% CI, 0.20 to 0.33; $P < 0.001$) (Fig. 2A). The esti-

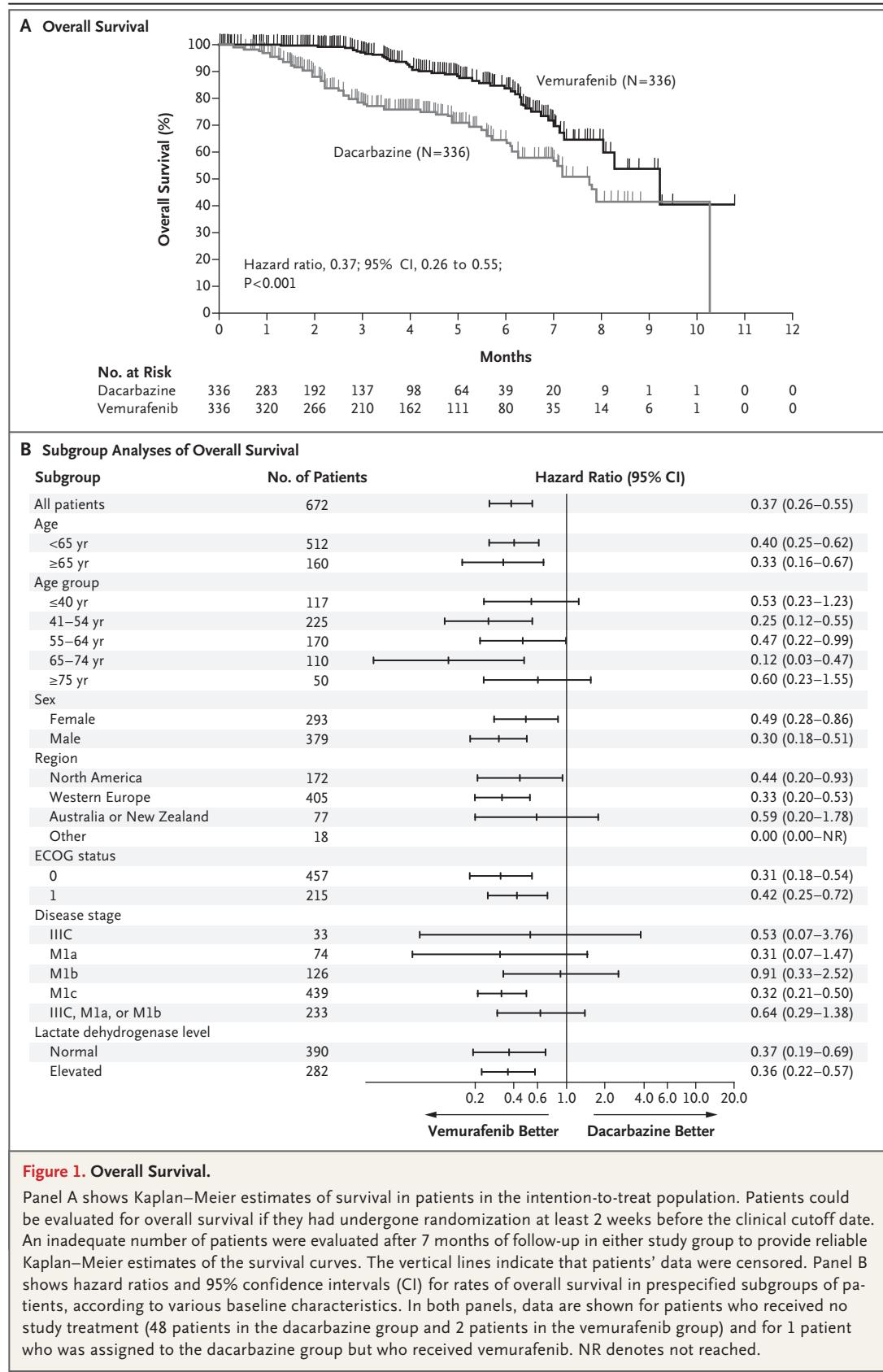
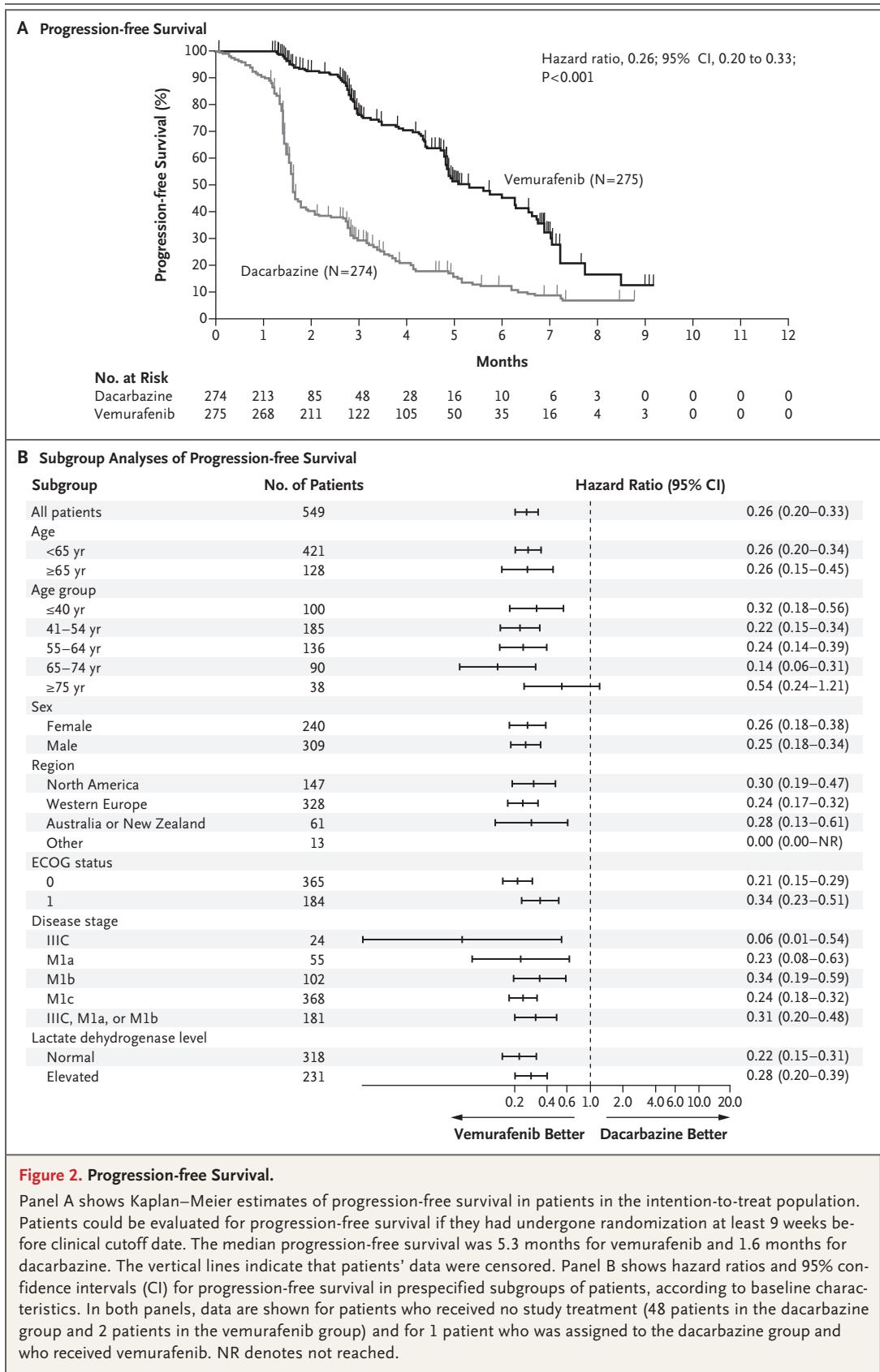
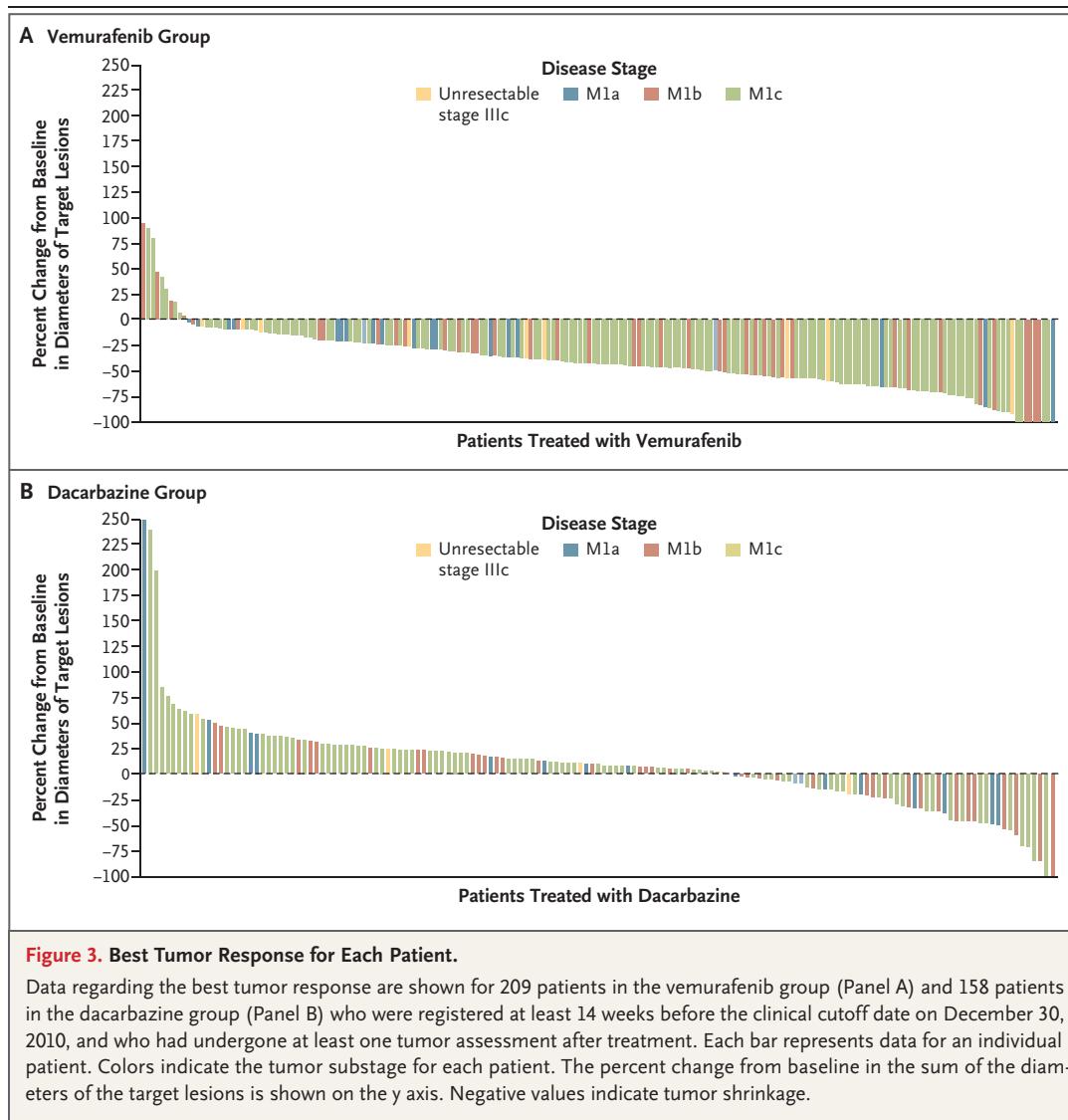


Figure 1. Overall Survival.

Panel A shows Kaplan–Meier estimates of survival in patients in the intention-to-treat population. Patients could be evaluated for overall survival if they had undergone randomization at least 2 weeks before the clinical cutoff date. An inadequate number of patients were evaluated after 7 months of follow-up in either study group to provide reliable Kaplan–Meier estimates of the survival curves. The vertical lines indicate that patients’ data were censored. Panel B shows hazard ratios and 95% confidence intervals (CI) for rates of overall survival in prespecified subgroups of patients, according to various baseline characteristics. In both panels, data are shown for patients who received no study treatment (48 patients in the dacarbazine group and 2 patients in the vemurafenib group) and for 1 patient who was assigned to the dacarbazine group but who received vemurafenib. NR denotes not reached.





mated median progression-free survival was 5.3 months in the vemurafenib group and 1.6 months in the dacarbazine group. Superior progression-free survival with vemurafenib over dacarbazine was observed in all subgroups that were analyzed (Fig. 2B).

A total of 439 patients (65%) could be evaluated for tumor response on the basis of having undergone randomization at least 14 weeks before the clinical cutoff date of December 30, 2010. In the vemurafenib group, most patients had a detectable decrease in tumor size (Fig. 3A), and 106 of 219 patients (48%; 95% CI, 42 to 55) had a confirmed objective response (including 2 patients with a complete response and 104 with a partial response), with a median time to response

of 1.45 months. Ten patients in the vemurafenib group were later found to have BRAF V600K mutations; of these patients, 4 had a partial response (40%). In the dacarbazine group, a minority of patients had a detectable decrease in tumor size (Fig. 3B), and only 12 of 220 patients (5%; 95% CI, 3 to 9) met the criteria for a confirmed response (all partial responses), with a median time to response of 2.7 months. The difference in confirmed response rates between the two study groups (48% vs. 5%) was highly significant ($P < 0.001$ by the chi-square test).

ADVERSE EVENTS

A total of 618 patients (92%) underwent at least one assessment as of the clinical cutoff date and

Table 2. Adverse Events in 618 Patients.*

Adverse Event	Vemurafenib (N = 336) [†] <i>no. of patients (%)</i>	Dacarbazine (N = 282)
Arthralgia		
Grade 2	60 (18)	1 (<1)
Grade 3	11 (3)	2 (<1)
Rash		
Grade 2	33 (10)	0
Grade 3	28 (8)	0
Fatigue		
Grade 2	38 (11)	33 (12)
Grade 3	6 (2)	5 (2)
Cutaneous squamous-cell carcinoma [‡]		
Grade 3	40 (12)	1 (<1)
Keratoacanthoma [§]		
Grade 2	7 (2)	0
Grade 3	20 (6)	0
Nausea		
Grade 2	25 (7)	32 (11)
Grade 3	4 (1)	5 (2)
Alopecia		
Grade 2	26 (8) [¶]	0
Pruritus		
Grade 2	19 (6)	0
Grade 3	5 (1)	0
Hyperkeratosis		
Grade 2	17 (5)	0
Grade 3	4 (1)	0
Diarrhea		
Grade 2	16 (5)	4 (1)
Grade 3	2 (<1)	1 (<1)
Headache		
Grade 2	15 (4)	5 (2)
Grade 3	2 (<1)	0
Vomiting		
Grade 2	9 (3)	14 (5)
Grade 3	4 (1)	3 (1)
Neutropenia		
Grade 2	1 (<1)	4 (1)
Grade 3	0	15 (5)
Grade 4	1 (<1)	8 (3)
Grade 5	0	1 (<1)

* Listed are all adverse events of grade 2 or higher that were reported in more than 5% of patients in either study group.

[†] One patient in the dacarbazine group who was treated with vemurafenib in error was included in the vemurafenib group for the assessment of adverse events.

[‡] The criteria for the diagnosis of cutaneous squamous-cell carcinoma were defined in the protocol and were reported as grade 3, according to the National Cancer Institute Common Terminology Criteria for Adverse Events. These events were evaluated by the investigators as grade 1 in one patient and as grade 2 in one patient.

[§] Three patients with keratoacanthomas that were assessed by the investigator as grade 1 are included among the grade 2 keratoacanthomas.

[¶] In one patient, alopecia that was scored as grade 3 by the investigator was re-scored as grade 2 since the Common Terminology Criteria for Adverse Events do not include grade 3 alopecia.

were evaluated for toxic effects. Adverse events of grade 2 or more that were reported in more than 5% of the patients in either study group are shown in Table 2; adverse events of any grade that were reported in more than 5% of the patients are shown in Table A in the Supplementary Appendix. The most common adverse events in the vemurafenib group were cutaneous events, arthralgia, and fatigue; photosensitivity skin reactions of grade 2 or 3 were seen in 12% of the patients, with grade 3 reactions characterized by blistering that often could be prevented with sunblock. As expected, the most common severe toxic effects in the dacarbazine group were fatigue, nausea, vomiting, and neutropenia. Adverse events led to dose modification or interruption in 129 of 336 patients (38%) in the vemurafenib group and in 44 of 282 patients (16%) in the dacarbazine group.

In the vemurafenib group, a cutaneous squamous-cell carcinoma, keratoacanthoma, or both developed in 61 patients (18%). All lesions were treated by simple excision. Pathological analyses of skin-biopsy specimens from these patients are currently being performed by an independent dermatology working group.

DISCUSSION

In our study, vemurafenib was associated with a relative reduction of 63% in the risk of death and of 74% in the risk of tumor progression in patients with previously untreated, unresectable stage IIIC or stage IV melanoma with the BRAF V600E mutation, as compared with dacarbazine. Benefit was seen in all subgroups of patients who were included in the analysis, including patients with stage M1c disease or an elevated lactate dehydrogenase level, both of which are associated with particularly poor prognoses.

Recently, ipilimumab, an anti-CTLA4 antibody, was shown to improve overall survival in patients with metastatic melanoma, as compared with a peptide vaccine, although there was only a modest effect on rates of response and progression-free survival.⁸ The use of ipilimumab combined with dacarbazine has also been associated with improved rates of survival over dacarbazine alone.⁹

Overall, 48% of the patients who were treated with vemurafenib met the criteria for a confirmed response, although most patients had some tumor shrinkage. This finding was consistent with the confirmed response rates seen in the phase 1 extension cohort¹⁵ and in a recent

phase 2 trial involving previously treated patients.¹⁶ Furthermore, 4 of 10 patients with the BRAF V600K mutation had a response to vemurafenib, indicating that melanomas with this variant are also sensitive to vemurafenib.

The confirmed response rate in the dacarbazine cohort was 5%, which is slightly lower than that in recent phase 3 trials.⁴⁻⁷ Our study was a randomized trial comparing vemurafenib with dacarbazine in which only patients with BRAF-mutated melanomas were treated. Recent studies have raised the possibility that melanomas with the BRAF V600E mutation are more aggressive^{18,19} and less sensitive to chemotherapy^{18,20} than BRAF wild-type melanomas. Also, 48 patients (14%) in the dacarbazine group (29 of whom were available for evaluation in this report) did not receive any treatment, most commonly because the patient withdrew consent (Fig. A in the Supplementary Appendix). Patients receiving vemurafenib reported relatively few grade 3 or worse adverse events. Other than cutaneous squamous-cell carcinoma and keratoacanthomas, the most common drug-related grade 3 (or worse) toxic effects were rash, arthralgias, photosensitivity, and fatigue. Overall, 38% of the patients receiving vemurafenib required dose modification because of adverse events.

Among patients treated with vemurafenib, 18% were reported to have at least one squamous-cell carcinoma of the skin or keratoacanthoma. These lesions were excised, and none required dose modification of vemurafenib. These rates are slightly lower than those in the phase 1 and 2 trials of vemurafenib,^{15,16} probably because of shorter follow-up in our study. Cutaneous squamous-cell cancer and keratoacanthomas have also been seen in patients treated with sorafenib,^{21,22} another compound with inhibitory activity against RAF kinases. No other secondary neoplasia was observed in our patients.

The mechanism of the induction of cutaneous neoplasia is under investigation, but it is speculated to involve the activating effect of vemurafenib on preneoplastic cells in which wild-type BRAF is further primed by upstream pathway activation. Several investigators have shown that vemurafenib and other inhibitors of RAF kinases can potentiate the activity of the MAPK pathway in cells with wild-type BRAF.²³⁻²⁵ This finding might explain the favorable therapeutic index of vemurafenib in patients who have melanoma with the BRAF V600E mutation but also suggests

that vemurafenib could accelerate the growth of some tumors with wild-type BRAF.

An important, related ongoing effort by many research groups is to clarify how melanomas become resistant to vemurafenib. Initial studies from several groups have indicated that the MAPK pathway is reactivated in resistant tumors.²⁶⁻²⁸ Although the precise mechanisms of reactivation are still being investigated, gatekeeper mutations in BRAF, which would prevent vemurafenib from binding BRAF, have not been observed.

Our results show that single-agent vemurafenib improved the rates of response and of both progression-free and overall survival, as compared with dacarbazine, in patients with metastatic melanoma with the BRAF V600E mutation. These findings provide a solid foundation for the development of future combination therapies.

Supported by Hoffmann–La Roche.

Dr. Chapman reports receiving consulting fees, payment for serving on scientific advisory boards, and grant support from Roche and consulting fees from GlaxoSmithKline; Dr. Hauschild, receiving consulting fees from Abraxis/Celgene, AstraZeneca, Bayer, Biogen, Bristol-Myers Squibb, Boehringer Ingelheim, Eisai, GlaxoSmithKline, Merck, Novartis, and Roche, grant support from Bayer and Merck, and lecture fees from Abraxis/Celgene, AstraZeneca, Biogen, Bristol-Myers Squibb, Merck, Novartis, and Roche; Dr. Robert, receiving consulting fees from Roche, Bristol-Myers Squibb, GlaxoSmithKline, and Merck and travel payments from Bristol-Myers Squibb; Dr. Ascierto, serving as a board member for Bristol-Myers Squibb, GlaxoSmithKline, Schering-Plough, Merck, and Roche and lecture fees from Schering-Plough and Merck; Dr. Garbe, receiving consulting fees from Bristol-Myers Squibb, Merck, GlaxoSmithKline, and Philogen, grant support from Bristol-Myers Squibb, Philogen, and Swedish Orphan, and travel payments from Bristol-Myers Squibb, Roche, and Merck; Dr. Maio, serving on advisory boards for Bristol-Myers Squibb and Pfizer and receiving lecture fees from Bristol-Myers Squibb; Dr. Hogg, receiving consulting fees from Roche, GlaxoSmithKline, and Bristol-Myers Squibb and lecture fees from Roche; Dr. Lorigan, receiving consulting fees from Roche and an educational grant from Roche; Dr. Lebbe, serving on paid advisory boards for Roche and Bristol-Myers Squibb and receiving payments for the development of educational presentations and travel payments from Bristol-Myers Squibb; Dr. Jouary, serving on an advisory board for Bristol-Myers Squibb and receiving consulting fees from Bristol-Myers Squibb and Schering-Plough; Dr. Schadendorf, receiving consulting fees from Roche, GlaxoSmithKline, Bristol-Myers Squibb, and Merck, serving on advisory boards for Roche, GlaxoSmithKline, Bristol-Myers Squibb, Merck, and Schering-Plough, receiving grant support from Schering-Plough and Plexikon, receiving lecture fees from Bristol-Myers Squibb, Merck, Schering-Plough, and Roche, and receiving payment for developing presentations for Bristol-Myers Squibb; Dr. Ribas, receiving consulting fees from Roche/Genentech; Dr. O'Day, receiving consulting fees from GlaxoSmithKline, receiving consulting fees, grant support, lecture fees, and payment for the development of educational presentations from Bristol-Myers Squibb, and serving as an expert witness in a private lawsuit; Dr. Sosman, receiving grant support from Roche, Millennium, and GlaxoSmithKline and consulting fees from Roche and Millennium; Dr. Kirkwood, serving as a board member for Morphotek and receiving consulting fees from GlaxoSmithKline and Vical, lecture fees from

Merck, and payment for the development of educational presentations from Imedex; Dr. Eggermont, receiving consulting fees from Bristol-Myers Squibb, Merck, and GlaxoSmithKline; Dr. Dreno, receiving consulting fees from Roche; Dr. Nolop, being an employee of and having an equity interest in Plexxikon; Mrs. Li and Dr. Lee, being employees of Roche; Miss Nelson and Dr.

Hou, being employees of Genentech and having an equity interest in Roche; Dr. Flaherty, receiving consulting fees from GlaxoSmithKline; and Dr. McArthur, receiving grant support from Pfizer. No other potential conflict of interest relevant to this article was reported. Disclosure forms provided by the authors are available with the full text of this article at NEJM.org.

APPENDIX

The authors' affiliations are as follows: the Department of Medicine, Memorial Sloan-Kettering Cancer Center, New York (P.B.C.); University of Kiel, Kiel (A.H.), University Hospital Essen, Essen (D.S.), and University of Tübingen, Tübingen (C.G.) — all in Germany; Institut Gustave Roussy (C.R., A.M.M.E.) and Service de Dermatologie, Hôpital Saint Louis (C.L.) — both in Paris; the Netherlands Cancer Institute, Amsterdam (J.B.H.); Istituto Nazionale Tumori Fondazione Pascale, Naples (P.A.), and Istituto Toscano Tumori, Florence (M.M.) — both in Italy; the Department of Medical Oncology, Royal Marsden Hospital, London (J. Larkin); the Department of Dermatology, University of Zurich, Zurich, Switzerland (R.D.); Istituto Europeo di Oncologia, Milan (A.T.); Princess Margaret Hospital and University Health Network, Toronto (D.H.); University of Manchester, Manchester, United Kingdom (P.L.); Saint Andre Hospital, Bordeaux (T.J.), and the Department of Dermatology, Nantes University Hospital, Nantes (B.D.) — both in France; the Department of Medicine, David Geffen School of Medicine at UCLA (A.R.), and the Angeles Clinic and Research Institute (S.J.O.) — both in Los Angeles; the Department of Medicine, Vanderbilt University School of Medicine, Nashville (J.A.S.); the Department of Medicine, University of Pittsburgh School of Medicine, Pittsburgh (J.M.K.); Plexxikon, Berkeley, CA (K.N.); Hoffmann-La Roche, Nutley, NJ (J. Li, R.J.L.); Genentech, San Francisco (B.N., J.H.); the Department of Medicine, Massachusetts General Hospital, Boston (K.T.F.); and the Peter MacCallum Cancer Center, Melbourne, VIC, Australia (G.A.M.).

REFERENCES

- Balch CM, Gershenwald JE, Soong S-J, et al. Final version of 2009 AJCC melanoma staging and classification. *J Clin Oncol* 2009;27:6199-206.
- Jemal A, Siegel R, Xu J, Ward E. Cancer statistics, 2010. *CA Cancer J Clin* 2010;60:277-300. [Erratum, *CA Cancer J Clin* 2011;61:133-4.]
- Ferlay J, Shin HR, Bray F, Forman D, Mathers C, Parkin DM. GLOBOCAN 2008, cancer incidence and mortality worldwide: IARC CancerBase no. 10. Lyon, France: International Agency for Research on Cancer, 2010. (<http://globocan.iarc.fr>)
- Chapman PB, Einhorn LH, Meyers ML, et al. Phase III multicenter randomized trial of the Dartmouth regimen versus dacarbazine in patients with metastatic melanoma. *J Clin Oncol* 1999;17:2745-51.
- Middleton MR, Grob JJ, Aaronson N, et al. Randomized phase III study of temozolomide versus dacarbazine in the treatment of patients with advanced metastatic malignant melanoma. *J Clin Oncol* 2000;18:158-66.
- Avril MF, Aamdal S, Grob JJ, et al. Fotemustine compared with dacarbazine in patients with disseminated malignant melanoma: a phase III study. *J Clin Oncol* 2004;22:1118-25.
- Bedikian AY, Millward M, Pehamberger H, et al. Bcl-2 antisense (oblimersen sodium) plus dacarbazine in patients with advanced melanoma: the Oblimersen Melanoma Study Group. *J Clin Oncol* 2006;24:4738-45.
- Hodi FS, O'Day SJ, McDermott DF, et al. Improved Survival with Ipilimumab in Patients with Metastatic Melanoma. *N Engl J Med* 2010;363:711-23. [Erratum, *N Engl J Med* 2010;363:1290.]
- Wolchok JD, Thomas L, Bondarenko IN, et al. A phase 3 randomized study of ipilimumab (IPI) plus dacarbazine (DTIC) versus DTIC alone as first-line treatment in patients with unresectable stage III or IV melanoma. *J Clin Oncol* 2011;29:Suppl: LBA5. abstract.
- Davies H, Bignell GR, Cox C, et al. Mutations of the BRAF gene in human cancer. *Nature* 2002;417:949-54.
- Curtin JA, Fridlyand J, Kageshita T, et al. Distinct sets of genetic alterations in melanoma. *N Engl J Med* 2005;353:2135-47.
- Bollag G, Hirth P, Tsai J, et al. Clinical efficacy of a RAF inhibitor needs broad target blockade in BRAF-mutant melanoma. *Nature* 2010;467:596-9.
- Tsai J, Lee JT, Wang W, et al. Discovery of a selective inhibitor of oncogenic B-Raf kinase with potent antimelanoma activity. *Proc Natl Acad Sci U S A* 2008;105:3041-6.
- Joseph EW, Pratilas CA, Poulidakos PI, et al. The RAF inhibitor PLX4032 inhibits ERK signaling and tumor cell proliferation in a V600E BRAF-selective manner. *Proc Natl Acad Sci U S A* 2010;107:14903-8.
- Flaherty KT, Puzanov I, Kim KB, et al. Inhibition of mutated, activated BRAF in metastatic melanoma. *N Engl J Med* 2010;363:809-19.
- Ribas A, Kim KB, Schuchter LM, et al. BRIM-2: an open-label, multicenter phase II study of vemurafenib in previously treated patients with BRAFV600E mutation-positive melanoma. *J Clin Oncol* 2011;29:Suppl:8509. abstract.
- Pocock SJ, Clayton TC, Altman DG. Survival plots of time-to-event outcomes in clinical trials: good practice and pitfalls. *Lancet* 2002;359:1686-9.
- Long GV, Menzies AM, Nagrial AM, et al. Prognostic and clinicopathologic associations of oncogenic BRAF in metastatic melanoma. *J Clin Oncol* 2011;29:1239-46.
- Arozarena I, Sanchez-Laorden B, Packer L, et al. Oncogenic BRAF induces melanoma cell invasion by downregulating the cGMP-specific phosphodiesterase PDE5A. *Cancer Cell* 2011;19:45-57.
- Kumar R, Angelini S, Czene K, et al. BRAF mutations in metastatic melanoma: a possible association with clinical outcome. *Clin Cancer Res* 2003;9:3362-8.
- Williams VL, Cohen PR, Stewart DJ. Sorafenib-induced premalignant and malignant skin lesions. *Int J Dermatol* 2011;50:396-402.
- Kwon EJ, Kish LS, Jaworsky C. The histologic spectrum of epithelial neoplasms induced by sorafenib. *J Am Acad Dermatol* 2009;61:522-7.
- Poulidakos PI, Zhang C, Bollag G, Shokat KM, Rosen N. RAF inhibitors transactivate RAF dimers and ERK signalling in cells with wild-type BRAF. *Nature* 2010;464:427-30.
- Hatzivassiliou G, Song K, Yen I, et al. RAF inhibitors prime wild-type RAF to activate the MAPK pathway and enhance growth. *Nature* 2010;464:431-5.
- Heidorn SJ, Milagre C, Whittaker S, et al. Kinase-dead BRAF and oncogenic RAS cooperate to drive tumor progression through CRAF. *Cell* 2010;140:209-21.
- Johannessen CM, Boehm JS, Kim SY, et al. COT drives resistance to RAF inhibition through MAP kinase pathway reactivation. *Nature* 2010;468:968-72.
- Nazarian R, Shi H, Wang Q, et al. Melanomas acquire resistance to B-RAF(V600E) inhibition by RTK or N-RAS upregulation. *Nature* 2010;468:973-7.
- Villanueva J, Vultur A, Lee JT, et al. Acquired resistance to BRAF inhibitors mediated by a RAF kinase switch in melanoma can be overcome by cotargeting MEK and IGF-1R/PI3K. *Cancer Cell* 2010;18:683-95.

Copyright © 2011 Massachusetts Medical Society.