

# KEYNOTE-022 part 3: a randomized, double-blind, phase 2 study of pembrolizumab, dabrafenib, and trametinib in *BRAF*-mutant melanoma

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## ABSTRACT

**Background** In the KEYNOTE-022 study, pembrolizumab with dabrafenib and trametinib (triplet) improved progression-free survival (PFS) versus placebo with dabrafenib and trametinib (doublet) without reaching statistical significance. Mature results on PFS, duration of response (DOR), and overall survival (OS) are reported.

**Methods** The double-blind, phase 2 part of KEYNOTE-022 enrolled patients with previously untreated *BRAF*<sup>V600E/K</sup>-mutated advanced melanoma from 22 sites in seven countries. Patients were randomly assigned 1:1 to intravenous pembrolizumab (200 mg every 3 weeks) or placebo plus dabrafenib (150 mg orally two times per day) and trametinib (2 mg orally one time a day). Primary endpoint was PFS. Secondary endpoints were objective response rate, DOR, and OS. Efficacy was assessed in the intention-to-treat population, and safety was assessed in all patients who received at least one dose of study drug. This analysis was not specified in the protocol.

**Results** Between November 30, 2015 and April 24, 2017, 120 patients were randomly assigned to triplet (n=60) or doublet (n=60) therapy. With 36.6 months of follow-up, median PFS was 16.9 months (95% CI 11.3 to 27.9) with triplet and 10.7 months (95% CI 7.2 to 16.8) with doublet (HR 0.53; 95% CI 0.34 to 0.83). With triplet and doublet, respectively, PFS at 24 months was 41.0% (95% CI 27.4% to 54.2%) and 16.3% (95% CI 8.1% to 27.1%); median DOR was 25.1 months (95% CI 14.1 to not reached) and 12.1 months (95% CI 6.0 to 15.7), respectively. Median OS was not reached with triplet and was 26.3 months with doublet (HR 0.64; 95% CI 0.38 to 1.06). With triplet and doublet, respectively, OS at 24 months was 63.0% (95% CI 49.4% to 73.9%) and 51.7% (95% CI 38.4% to 63.4%). Grade 3–5 treatment-related adverse events (TRAEs) occurred in 35 patients (58%, including one death) receiving triplet and 15 patients (25%) receiving doublet. **Conclusion** In *BRAF*<sup>V600E/K</sup>-mutant advanced melanoma, pembrolizumab plus dabrafenib and trametinib substantially improved PFS, DOR, and OS with a higher incidence of TRAEs. Interpretation of these results is limited by the post hoc nature of the analysis.

## INTRODUCTION

Immunotherapies with antiprogrammed death 1 (PD-1) monoclonal antibodies, either alone or in combination with anticytotoxic T-lymphocyte-associated antigen 4 monoclonal antibodies, and targeted therapies with *BRAF* plus MEK inhibitors have significantly improved the outcomes of patients with advanced melanoma.<sup>1</sup> In particular, use of *BRAF* and MEK inhibitors generally results in higher objective response rates for patients with *BRAF*<sup>V600</sup>-mutant disease, whereas immune checkpoint inhibitors typically result in more durable responses. Importantly, these treatments have distinct mechanisms of action; thus, it has long been hypothesized that combining immune checkpoint inhibitors with *BRAF* and MEK inhibitors may result in a higher frequency of durable responses.<sup>2,3</sup> Evidence from previous preclinical modeling and analysis of patient biopsy specimens suggest that inhibition of the *BRAF*<sup>V600</sup> driver oncogene facilitates an antitumor immune response.<sup>4–7</sup>

Results of early clinical trials in patients with previously untreated metastatic *BRAF*<sup>V600</sup>-mutant melanoma highlight the promise of this strategy. For example, 15 patients were treated with the anti-PD-1 antibody pembrolizumab plus the *BRAF* inhibitor dabrafenib and the MEK inhibitor trametinib in the phase 1 portion of the KEYNOTE-022 study,<sup>8</sup> and 39 patients received the *BRAF* inhibitor vemurafenib and the MEK inhibitor cobimetinib in combination with the anti-PD ligand 1 (PD-L1) antibody atezolizumab in another phase 1 trial.<sup>9</sup> Efficacy data obtained from these two studies were remarkably consistent,

with the combination of BRAF, MEK, and immune checkpoint inhibitors producing objective response rates of 73% and 72%, respectively; median progression-free survival was 15.4 months and 12.9 months, respectively. Interestingly, tumor biopsy specimens showed increased tumor infiltration of CD8<sup>+</sup> T cells and expression of PD-L1 and a T cell-inflamed gene signature after treatment with the combination of pembrolizumab, dabrafenib, and trametinib.<sup>8</sup>

Other studies investigating combinations used the anti-PD1-L1 antibody durvalumab<sup>10</sup> or the anti-PD-1 antibody spartalizumab<sup>11</sup> combined with the BRAF inhibitor dabrafenib and the MEK inhibitor trametinib. In the latter study, the objective response rate was 75% (n=27), the complete response rate was 33% (n=12), and the 12 month progression-free survival rate was 65.3%.<sup>11</sup>

In addition, the randomized, double-blind, phase 2 part of KEYNOTE-022 compared the efficacy of the triplet combination of pembrolizumab, dabrafenib, and trametinib with the efficacy of the doublet combination of placebo with dabrafenib and trametinib.<sup>12</sup> At a median follow-up of 9.6 months, although the primary endpoint of progression-free survival did not show statistically significant improvement in the triplet arm compared with the doublet arm, numerically higher values were observed (16.0 months vs 10.3 months; HR, 0.66; p=0.043). After additional follow-up of a median 36.6 months, we report mature results showing clinically substantial improvement in duration of response and survival.

## METHODS

### Study design

KEYNOTE-022 was a five-part, multicenter, phase 1/2 dose-finding and preliminary efficacy study of pembrolizumab plus dabrafenib and trametinib in patients with unresectable or metastatic melanoma or solid tumors (parts 4 and 5 only) (online supplemental table S1). Part 3 was the randomized, double-blind, phase 2 portion in which patients with previously untreated advanced melanoma were randomly assigned 1:1 to receive pembrolizumab or placebo combined with dabrafenib and trametinib. Patients were enrolled from 22 medical centers in seven countries.<sup>12</sup>

### Patients

Eligible patients were aged  $\geq 18$  years and had unresectable stage III or stage IV cutaneous melanoma, with at least one measurable lesion as defined by Response Evaluation Criteria in Solid Tumours version 1.1 (RECIST V.1.1)<sup>13</sup> on imaging (CT or MRI). Patients were required to have *BRAF*-mutation positive (V600E or V600K) disease should not have previously received systemic therapy for advanced melanoma, and should not present with active central nervous system metastases and/or carcinomatous meningitis.

### Procedures

Patients were randomly assigned to receive pembrolizumab plus dabrafenib and trametinib or placebo plus

dabrafenib and trametinib using an interactive voice response system/integrated web response system. Randomization was stratified by Eastern Cooperative Oncology Group performance status (ECOG PS; 0 vs 1) and lactate dehydrogenase level ( $>1.1\times$  upper limit of normal (ULN) vs  $\leq 1.1\times$  ULN). Because of the low number of patients, the ECOG PS 1 and lactate dehydrogenase level  $>1.1\times$  ULN strata were combined.

Patients received pembrolizumab 2 mg/kg or placebo intravenously every 3 weeks combined with dabrafenib 150 mg orally two times per day and trametinib 2 mg orally one time a day. Treatment was continued for up to 24 months or until disease progression, unacceptable toxicity, or withdrawal from the study. Trametinib and/or dabrafenib treatment could be continued beyond 24 months per standard of care until disease progression. Dose modification or interruption of pembrolizumab, placebo, trametinib, or dabrafenib was allowed to manage toxicity per protocol-specified management guidelines.<sup>12</sup>

Response and disease progression were assessed radiologically per RECIST V.1.1 by investigator review. Tumor imaging was performed at baseline, at 12 weeks after the first study treatment, every 6 weeks following this up to 18 months, and every 12 weeks thereafter while the patient remained on study therapy. Patients were monitored throughout the study for adverse events, which were graded according to the National Cancer Institute Common Terminology Criteria for Adverse Events (V.4.0). Immune-mediated adverse events are defined in the online supplemental. PD-L1 expression in tumor samples was determined using the PD-L1 IHC 22C3 pharmDx assay (Agilent Technologies).

### Outcomes

The primary endpoint was progression-free survival, defined as the time from randomization to the first documented disease progression or death from any cause, whichever occurred first. Secondary endpoints were objective response rate, duration of response, and overall survival.

### Statistical analysis

Progression-free survival, objective response rate, and overall survival were assessed in the intention-to-treat population.<sup>12</sup> Safety was assessed in patients who received at least one dose of study drug. Clinical data were collected using InForm V.4.6.5 software (Oracle Health Sciences, Redwood City, California, USA), and SAS V.9.4 software (SAS Institute, Cary, North Carolina, USA) was used for all statistical analyses. Progression-free survival was estimated using the Kaplan-Meier method with patients censored at the last disease assessment date or at the last disease assessment before initiation of a new anticancer treatment. A stratified log-rank test was used to assess between-group differences in progression-free survival. A Cox proportional hazards model with Efron's method of tie-handling was used to estimate HRs and 95% CIs. The Kaplan-Meier

method was used for summary statistics for duration of response and overall survival (see protocol for censoring rules). The treatment comparison for objective response rate was based on Miettinen and Nurminen's method.<sup>14</sup> Assuming an HR of 0.5, the study needed approximately 74 progression-free survival events to have 80% power to reject the null hypothesis at a one-sided 0.025 type I error. The observed HR had to be approximately 0.62 or lower for the treatment effect to be statistically significant. The primary analysis was conducted using all available alphas; therefore, this subsequent post hoc analysis cannot be considered to carry any statistical significance. The study was overseen by a data monitoring committee and is registered at ClinicalTrials.gov (NCT02130466).

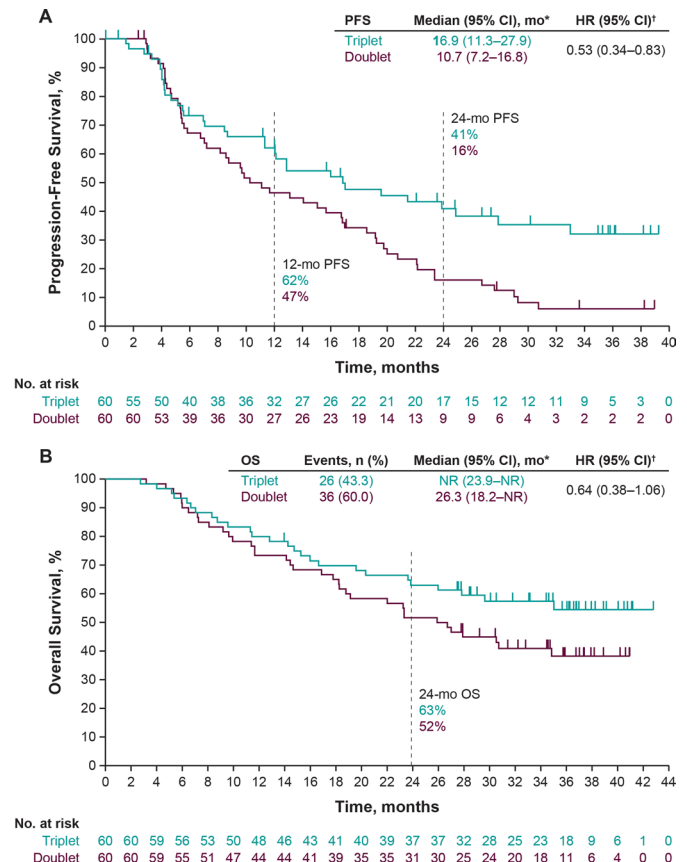
## RESULTS

Between November 30, 2015 and April 24, 2017, 180 patients were screened for eligibility and 120 were randomly assigned to receive pembrolizumab plus dabrafenib and trametinib (triplet therapy; n=60 (50%)) or placebo plus dabrafenib and trametinib (doublet therapy; n=60 (50%))<sup>12</sup> (online supplemental figure S1). All patients received at least one dose of study treatment. The median age was 56 years (range, 18–83), and 58% (n=69) of patients were men. Baseline characteristics were generally well balanced between arms,<sup>12</sup> apart from a higher proportion of patients with M1c disease in the triplet arm (82% (n=49) triplet arm, 63% (n=38) doublet arm) (online supplemental table S2). The median exposure duration for each study drug is provided in online supplemental table S3.

At data cut-off (June 26, 2019), the median time from randomization to data cut-off was 36.6 months (range, 26.1–42.9) in the triplet arm and 36.4 months (range, 27.2–41.6) in the doublet arm. Fifteen patients (13%) were receiving treatment: 11 (18%) in the triplet arm and 4 (7%) in the doublet arm. Forty-three patients (72%) had discontinued treatment in the triplet arm and 56 (93%) in the doublet arm, mainly because of adverse events (18 (30%) triplet arm, 10 (17%) doublet arm) and progressive disease (17 (28%) triplet arm, 44 (73%) doublet arm).

Thirty-four patients (57%) in the triplet arm and 53 (88%) in the doublet arm had a progression event. Median progression-free survival was 16.9 months (95% CI 11.3 to 27.9) in the triplet arm and 10.7 months (95% CI 7.2 to 16.8) in the doublet arm (HR 0.53; 95% CI 0.34 to 0.83) (figure 1A). The 12-month progression-free survival rates were 62% (95% CI 48.1% to 73.5%) in the triplet arm and 47% (95% CI 33.4% to 58.7%) in the doublet arm, and the 24-month progression-free survival rates were 41% (95% CI 27.4% to 54.2%) and 16% (95% CI 8.1% to 27.1%), respectively.

Exploratory subgroup analysis of progression-free survival showed that the HRs for the subgroups favored the triplet over the doublet arm in patients who were



**Figure 1** Kaplan-Meier estimates. (A) Progression-free survival, defined as time from randomization to disease progression or death, whichever came first. Median progression-free survival was based on Kaplan-Meier estimate per investigator assessment. (B) Overall survival, defined as time from randomization to death. \*Based on Kaplan-Meier estimates of progression-free survival (PFS) (investigator assessment by Response Evaluation Criteria in Solid Tumors, V.1.1) or overall survival (OS). †The HRs and 95% CIs were calculated using a Cox regression model with treatment as a covariate stratified by Eastern Cooperative Oncology Group performance status (ECOG PS) (0 vs 1) and lactate dehydrogenase (LDH) (>1.1× upper limit of normal (ULN) vs ≤1.1× ULN); because of the low number of patients enrolled in the ECOG PS 1 and LDH ≤1.1× ULN strata, these strata were combined. Doublet, placebo plus dabrafenib plus trametinib; mo, month; NR, not reached; triplet, pembrolizumab plus dabrafenib plus trametinib.

aged ≤65 years, men, had ECOG PS 0, or had elevated lactate dehydrogenase levels (online supplemental figure S2).

Twenty-six patients (43%) in the triplet arm and 36 (60%) in the doublet arm died. Median overall survival was not reached in the triplet arm and was 26.3 months in the doublet arm (HR, 0.64; 95% CI 0.38 to 1.06) (figure 1B). The 12-month overall survival rates were 80% (95% CI 67.5% to 88.1%) in the triplet arm and 73% (95% CI 60.2% to 82.7%) in the doublet arm, and the 24-month overall survival rates were 63% (95% CI 49.4% to 73.9%) and 52% (95% CI 38.4% to 63.4%), respectively. Exploratory subgroup analysis of overall survival



**Table 1** Best overall response (intention-to-treat population). Responses are based on investigator best assessment across time points per Response Evaluation Criteria in Solid Tumors v1.1 with confirmation

	Pembrolizumab + dabrafenib + trametinib n=60 n (%)	Placebo + dabrafenib + trametinib n=60 n (%)	Difference in rate,* % (95% CI)
Objective response <sup>†</sup>	38 (63.3)	43 (71.7)	-8.5 (-24.8 to 8.3)
Complete response	12 (20.0)	9 (15.0)	5.2 (-8.8 to 18.9)
Partial response	26 (43.3)	34 (56.7)	-13.8 (-30.9 to 4.3)
Disease control <sup>‡</sup>	51 (85.0)	56 (93.3)	-8.1 (-20.1 to 3.6)
Stable disease	13 (21.7)	13 (21.7)	0.5 (-14.6 to 15.6)
Progressive disease	5 (8.3)	3 (5.0)	3.2 (-7.0 to 13.8)
Not evaluable	2 (3.3)	0 (0.0)	3.4 (-2.8 to 11.6)
No assessment	2 (3.3)	1 (1.7)	1.4 (-6.1 to 9.6)

\*Difference in rate was based on Miettinen and Nurminen method stratified by Eastern Cooperative Oncology Group (ECOG) performance status (0 vs 1) and lactate dehydrogenase (LDH) (>1.1× upper limit of normal (ULN) vs ≤1.1× ULN); because of the small number of patients enrolled in the ECOG performance status 1 and LDH ≤1.1× ULN strata, these strata were combined.

<sup>†</sup>Complete response + partial response.

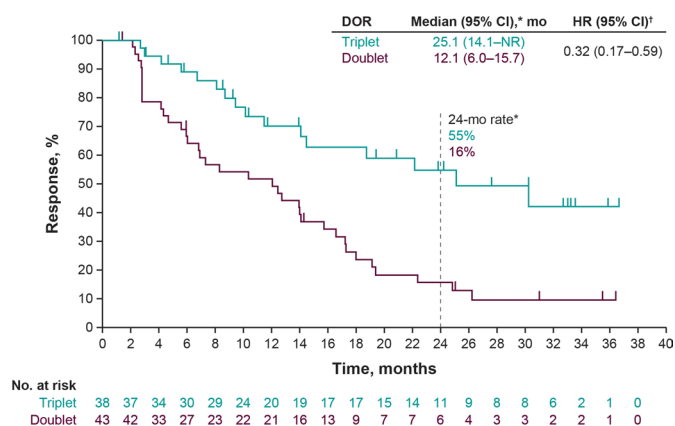
<sup>‡</sup>Complete response + partial response + stable disease.

showed no difference between the triplet and doublet arms in any subgroup (online supplemental figure S3).

The objective response rate was 63% in the triplet arm and 72% in the doublet arm (difference in rate -8.5%; 95% CI -24.8 to 8.3). A complete response was reported in 12 patients (20%) in the triplet arm and 9 patients (15%) in the doublet arm (table 1). The median duration of response was 25.1 months in the triplet arm and 12.1 months in the doublet arm (HR, 0.32; 95% CI 0.17 to 0.59) (figure 2). At

the data cut-off, responses were ongoing in 11 of 38 patients (29%) in the triplet arm and 4 of 43 patients (9%) in the doublet arm (online supplemental figure S4). A reduction in target lesion size of at least 50% was observed in 44 patients (73%) in the triplet arm and 38 patients (63%) in the doublet arm (online supplemental figure S5).

Grade 3–5 adverse events occurred in 42 patients (70%) in the triplet arm and 27 patients (45%) in the doublet arm; grade 3–5 treatment-related adverse events were reported in 35 patients (58%) and 15 patients (25%), respectively. Adverse events led to dose reductions in 16 patients (27%) and 9 patients (15%). Adverse events leading to dose interruption occurred in 50 patients (83%) and 41 patients (68%), respectively; the most common were fever (55%; n=33) and diarrhea (15%; n=9) in the triplet arm and fever (45%; n=27) and neutropenia (10%; n=6) in the doublet arm. Adverse events led to treatment discontinuation in 28 patients (47%) in the triplet arm and 12 patients (20%) in the doublet arm (online supplemental table S4). Treatment-related adverse events occurred in 57 patients (95%) in the triplet arm and 56 patients (93%) in the doublet arm (table 2); the most common were fever (72%; n=43), rash (37%; n=22), and chills (35%; n=21) in the triplet arm and fever (68%; n=41), chills (38%; n=23), and fatigue (38%; n=23) in the doublet arm. Immune-mediated adverse events occurred in 31 patients (52%) in the triplet arm and 9 patients (15%) in the doublet arm (table 3); the most frequently occurring in the triplet arm were pneumonitis (17%; n=10) and hypothyroidism (8%; n=5). Serious treatment-related adverse events occurred in 24 patients (40%) and 14 patients (23%), respectively. One patient (1.7%) in the triplet arm died of treatment-related pneumonitis.



**Figure 2** Kaplan-Meier estimates of duration of response. Duration of response (DOR) was defined as the time from the first response to disease progression or death, whichever occurred first. \*Median DOR was based on Kaplan-Meier estimate per investigator assessment. <sup>†</sup>The HRs and 95% CIs were calculated using a Cox regression model with treatment as a covariate stratified by Eastern Cooperative Oncology Group performance status (ECOG PS) (0 vs 1) and lactate dehydrogenase (LDH) (LDH>1.1× upper limit of normal (ULN) vs ≤1.1× ULN); because of the low number of patients enrolled in the ECOG PS 1 and LDH ≤1.1× ULN strata, these strata were combined. Doublet, placebo plus dabrafenib plus trametinib; mo, month; NR, not reached; Triplet, pembrolizumab plus dabrafenib plus trametinib.

## DISCUSSION

In our previous report on part 3 of the KEYNOTE-022 trial,<sup>12</sup> despite a median progression-free survival of

**Table 2** Treatment-related adverse events occurring in  $\geq 15\%$  of patients in either treatment arm (all-subjects-as-treated population)

Event, n (%)	Pembrolizumab + dabrafenib + trametinib n=60		Placebo + dabrafenib + trametinib n=60	
	Any grade	Grade 3–5*	Any grade	Grade 3–5
Any event	57 (95.0)	35 (58.3)	56 (93.3)	15 (25.0)
Gastrointestinal disorders				
Diarrhea	17 (28.3)	2 (3.3)	7 (11.7)	0 (0.0)
Nausea	16 (26.7)	0 (0.0)	18 (30.0)	0 (0.0)
Vomiting	12 (20.0)	0 (0.0)	11 (18.3)	0 (0.0)
General disorders and administration site conditions				
Asthenia	15 (25.0)	0 (0.0)	7 (11.7)	2 (3.3)
Chills	21 (35.0)	0 (0.0)	23 (38.3)	1 (1.7)
Fatigue	16 (26.7)	1 (1.7)	23 (38.3)	0 (0.0)
Fever	43 (71.7)	6 (10.0)	41 (68.3)	2 (3.3)
Investigations				
ALT increased	12 (20.0)	2 (3.3)	10 (16.7)	2 (3.3)
AST increased	12 (20.0)	4 (6.7)	12 (20.0)	2 (3.3)
Blood ALP	8 (13.3)	1 (1.7)	11 (18.3)	1 (1.7)
GGT increased	5 (8.3)	4 (6.7)	9 (15.0)	3 (5.0)
Musculoskeletal and connective tissue disorders				
Arthralgia	18 (30.0)	2 (3.3)	12 (20.0)	0 (0.0)
Myalgia	10 (16.7)	0 (0.0)	9 (15.0)	1 (1.7)
Skin and subcutaneous tissue disorders				
Dermatitis acneiform	9 (15.0)	0 (0.0)	4 (6.7)	0 (0.0)
Pruritus	9 (15.0)	0 (0.0)	6 (10.0)	0 (0.0)
Rash	22 (36.7)	3 (5.0)	16 (26.7)	0 (0.0)
Vitiligo	12 (20.0)	0 (0.0)	0 (0.0)	0 (0.0)

\*One patient in the triplet arm had grade 5 treatment-related pneumonitis.

ALP, alkaline phosphatase; ALT, alanine aminotransferase; AST, aspartate aminotransferase; GGT, gamma-glutamyltransferase.

16.0 months in the triplet arm versus 10.3 months in the doublet arm and an HR of 0.66, the study did not meet the prespecified primary endpoint of progression-free survival. Analysis was performed after 9.5 months of follow-up at the occurrence of 72 progression-free survival events; the study plan required 74 events to have 80% power to reject the null hypothesis. The short follow-up time could have influenced the results. Although the current analysis was not protocol-specified, as data matured with a longer median follow-up of 36.6 months, improvements in progression-free survival (16.9 months and 10.7 months, respectively), differences in duration of response (25.1 months and 12.1 months, respectively), and overall survival (not reached and 26.3 months, respectively) became more pronounced between the two arms.

Patients in the doublet arm had a higher initial objective response rate than patients in the triplet arm at the time of the primary analysis, which was considered to have been likely due to an imbalance in the baseline prognostic factors between the treatment arms (more patients

in the triplet arm had M1c disease).<sup>12</sup> This result persisted in the mature analysis, even though the survival benefit became more pronounced in the triplet arm. Although it is unclear whether other factors may have contributed to this dichotomy, the additional follow-up in the patients in the triplet arm showed deeper and more durable responses than in the doublet arm and a higher rate of complete responses, a lower rate of relapses, and progressive separation in the curves of duration of response. Improved depth of response was associated with improved survival in pooled analyses of trials in melanoma; therefore, it is likely that these response characteristics resulted in the further separation of the progression-free survival and overall survival curves.<sup>15 16</sup> These data confirm the rationale of the study and are also consistent with the notion that adding anti-PD-1 blocking antibodies to BRAF plus MEK inhibitor therapy may not improve the initial objective response rates but instead prevent or delay the onset of acquired resistance, as demonstrated by the progressive separation of the time-to-event curves.<sup>17 18</sup> Exploratory analysis of progression-free survival showed a substantial

**Table 3** Immune-mediated adverse events\* occurring in at least one patient in either treatment arm (all-subjects-as-treated population)

Event, n (%)	Pembrolizumab + dabrafenib + trametinib n=60	Placebo +dabrafenib + trametinib n=60
At least one immune-mediated adverse event	31 (51.7)	9 (15.0)
Pneumonitis	10 (16.7)	2 (3.3)
Hypothyroidism	5 (8.3)	1 (1.7)
Severe skin reactions	5 (8.3)	1 (1.7)
Hepatitis <sup>†</sup>	4 (6.7)	2 (3.3)
Hyperthyroidism	3 (5.0)	0 (0.0)
Uveitis	3 (5.0)	2 (3.3)
Colitis	2 (3.3)	1 (1.7)
Hypophysitis	2 (3.3)	0 (0.0)
Nephritis	2 (3.3)	0 (0.0)
Infusion reactions	1 (1.7)	1 (1.7)
Myositis	0 (0.0)	1 (1.7)

\*Immune-mediated adverse events were selected from a prespecified list and defined as events of unknown cause associated with drug exposure and consistent with an immune event.

<sup>†</sup>Includes autoimmune hepatitis, drug-induced liver injury, and hepatitis.

advantage of triplet versus doublet therapy in all patient subgroups, particularly in patients younger than 65 years of age, male patients, and patients with elevated lactate dehydrogenase levels at baseline. Patients with elevated lactate dehydrogenase levels have adverse features for durable response with dabrafenib and trametinib or anti-PD-1 therapy alone, have worse prognosis, and are at higher risk for early progression with immune or targeted therapies.<sup>19–21</sup> Our findings support the idea that these patients can be offered triplet combination therapy to circumvent the development of resistance.

Although the use of BRAF and MEK inhibitors is recommended for patients with BRAF-mutant disease,<sup>22</sup> immune checkpoint inhibitor combinations have also been shown to be effective in this population. In the CheckMate 067 trial, nivolumab plus ipilimumab (5-year overall survival rate, 60%) or nivolumab alone (46%) improved long-term survival compared with ipilimumab (30%) in patients with previously untreated BRAF-mutant melanoma.<sup>23</sup> The 5-year progression-free survival rate in CheckMate 067 was 38% with nivolumab plus ipilimumab, 22% with nivolumab alone, and 11% with ipilimumab in patients with previously untreated BRAF-mutant melanoma. Notably, 24-month overall survival and progression-free survival rates with nivolumab plus ipilimumab are similar to those with triplet therapy in the current analysis (63% and 41%, respectively). Robust antitumor

activity has also been observed with pembrolizumab plus reduced-dose ipilimumab in the phase 1b KEYNOTE-029 study of patients with BRAF-mutant melanoma in which the majority (87%) of patients had not received prior systemic therapy. Patients with BRAF-mutant melanoma who had not previously received BRAF or MEK inhibitor therapy had a response rate of 60%, whereas those who had previously received BRAF or MEK inhibitor therapy had a response rate of 38%.<sup>24</sup> While the results of these studies suggest that patients with BRAF-mutant melanoma may benefit from immune checkpoint inhibitor combinations, it remains unknown how such regimens compare with targeted therapies, or indeed with the triplet therapy investigated in the current analysis, as cross-trial comparisons are hindered by differences in study design and patient population. Trials directly comparing these therapeutic combinations are required to definitively ascertain which regimen is more effective in patients with BRAF-mutant melanoma. Meanwhile, several clinical trials currently underway are investigating the optimal sequencing of targeted and immune checkpoint inhibitor therapy, including COWBOY (NCT02968303), ImmunoCobiVem (NCT02902029), and EBIN (NCT03235245); the results of these should provide insight into which regimen is more effective as front-line therapy.

In the current analysis, the triplet combination was associated with greater toxicity compared with the doublet regimen. Patients who received pembrolizumab plus dabrafenib and trametinib experienced a higher rate of grades 3–5 treatment-related adverse events than patients who received placebo plus dabrafenib and trametinib (58% vs 25%), leading to a higher discontinuation rate because of toxicity. Both immune-mediated adverse events (52% vs 15%) and serious treatment-related adverse events (40% vs 23%) were also more common in the triplet arm compared with the doublet arm. Interestingly, discontinuation resulted primarily from disease progression (44 patients (73%)) in the doublet arm but from adverse events (18 patients (30%)) in the triplet arm. Studies are ongoing to address whether different intermittent or dose-sequencing regimens, combining immunotherapies with targeted therapies for melanoma, may be able to maintain the benefit of the concomitant triple therapy while reducing toxicity (NCT02858921, NCT02224781, and NCT02631447 (Sequential COMBo Immuno and Targeted therapy)).

Recently, positive data were reported from the phase 3 IMspire 150 trial comparing the frontline use of atezolizumab plus vemurafenib and cobimetinib with that of vemurafenib and cobimetinib alone in patients with BRAF<sup>V600</sup>-mutation-positive-advanced melanoma.<sup>25</sup> This study reported a significant improvement in progression-free survival with the combination of atezolizumab, vemurafenib, and cobimetinib versus vemurafenib and cobimetinib alone (15.1 vs 10.6 months) and a 2-year overall survival rate of 60% vs 53%, respectively; progression-free survival and overall survival rates were similar to those observed in the current analysis. While



the lead-in of targeted therapies before initiating atezolizumab in the IMspire150 trial could have affected the early efficacy results, it seems not to have influenced toxicity. In fact, the incidence of grade 3/4 treatment-related adverse events in IMspire 150 was substantially higher than in the current analysis (atezolizumab, vemurafenib, and cobimetinib arm, 79%; vemurafenib and cobimetinib arm, 73%). In contrast, a recent analysis of part 3 of the COMBI-I trial reported that spartalizumab plus dabrafenib and trametinib did not significantly improve progression-free survival compared with placebo plus dabrafenib and trametinib among patients with previously untreated *BRAF*<sup>V600</sup>-mutant melanoma.<sup>26</sup> In the COMBI-I study, the median progression-free survival was 16.2 months for spartalizumab plus dabrafenib and trametinib versus 12.0 months for placebo plus dabrafenib and trametinib (HR, 0.82; 95% CI 0.655 to 1.027).<sup>26</sup> Median overall survival was not reached in either arm, with analyses ongoing. Grade 3 or higher treatment-related adverse events were reported in 55% of patients receiving spartalizumab plus dabrafenib and trametinib compared with 33% of patients receiving placebo plus dabrafenib and trametinib. Although the results of these studies provide insight into the role of the combination of PD-1/PD-L1 inhibitors with BRAF and MEK inhibitors in patients with advanced, treatment-naive *BRAF*<sup>V600</sup>-mutant melanoma, differences in agents investigated, study design, and patient population preclude direct comparison. In conclusion, pembrolizumab plus dabrafenib and trametinib as first-line therapy in patients with advanced *BRAF*<sup>V600</sup>-mutant melanoma provided a clinically substantial improvement in progression-free survival, longer duration of response, and higher overall survival rate compared with placebo plus dabrafenib and trametinib. An important lesson from this analysis is the further improvement in outcomes observed with longer follow-up. However, the results should be interpreted with caution given the post hoc nature of the analysis. This finding should be considered in the design of future studies evaluating immunotherapy combinations, as duration of response may be a critical factor. Indeed, durable responses can allow the differences between treatments to become more pronounced over time, and an early data cut-off may not provide the full picture of the true clinical benefit.<sup>27</sup>

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## Correction: KEYNOTE-022 part 3: a randomized, double-blind, phase 2 study of pembrolizumab, dabrafenib, and trametinib in BRAF-mutant melanoma

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