













Phase II Trial of Atezolizumab Combined With Carboplatin and Pemetrexed for Patients With Advanced Nonsquamous Non–Small-Cell Lung Cancer With Untreated Brain Metastases (Atezo-Brain, GECP17/05)

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ABSTRACT

PURPOSE The Atezo-Brain study evaluated atezolizumab combined with chemotherapy in patients with advanced non–small-cell lung cancer (NSCLC) with untreated brain metastases, a population traditionally excluded from trials.

METHODS This single-arm phase II clinical trial enrolled patients with advanced non-squamous NSCLC with untreated brain metastases without neurologic symptoms or asymptomatic with medical treatment. Dexamethasone was allowed up to 4 mg once daily. Atezolizumab plus carboplatin and pemetrexed was given for four to six cycles followed by atezolizumab plus pemetrexed until progression for a maximum of 2 years. The primary end points were to determine the progression-free survival (PFS) rate at 12 weeks and the incidence of grade ≥ 3 adverse events during the first 9 weeks. Intracranial outcomes were assessed using response assessment in neuro-oncology brain metastases criteria.

RESULTS Forty patients were enrolled and 22 (55%) were receiving corticosteroids at baseline. The overall 12-week PFS rate was 62.2% (95% credibility interval [CrI], 47.1 to 76.2). The rate of grade 3/4 adverse events during the first 9 weeks was 27.5%. Most neurologic events were grade 1 and 2 but five patients (12.5%) experienced grade 3–4 neurologic events. With a median follow-up of 31 months, intracranial median PFS was 6.9 months and response rate was 42.7% (95% CrI, 28.1 to 57.9). Systemic median PFS was 8.9 months and response rate was 45% (95% CrI, 28.1 to 57.9). The median overall survival (OS) was 11.8 months (95% CI, 7.6 to 16.9) and the 2-year OS rate was 27.5% (95% CI, 16.6 to 45.5).

CONCLUSION Atezolizumab plus carboplatin and pemetrexed demonstrates activity in patients with advanced nonsquamous NSCLC with untreated brain metastases with an acceptable safety profile.

ACCOMPANYING CONTENT

 Editorial, p. 4462

 Data Sharing Statement

 Data Supplement

 Protocol

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INTRODUCTION

Lung cancer is the most common solid tumor to metastasize to the CNS and about one quarter of patients diagnosed with stage IV non–small-cell lung cancer (NSCLC) have brain metastases at diagnosis.¹ Historically, whole-brain radiotherapy (WBRT) had been the cornerstone of treatment for brain metastases in patients with advanced NSCLC, especially when stereotactic radiosurgery (SRS) and surgery were not feasible or indicated. However, the use of WBRT has diminished because of modest survival benefit and its negative impact on cognitive function and quality of life

(QOL), and because of emerging evidence that SRS may benefit patients with higher disease burden in the brain metastases.^{2–5} In addition, the incorporation of highly CNS-penetrant targeted therapies in oncogene-addicted NSCLC modified the clinical management of patients with asymptomatic brain metastases harboring *EGFR* and *ALK* positive tumors such that local therapy could be deferred.⁶

Although most clinical trials exclude or under-represent patients with NSCLC who have previously untreated brain metastases,⁷ immunotherapy has shown encouraging intracranial efficacy in patients with oncogene driver–negative

CONTEXT

Key Objective

There is a high need for effective and safe treatment options for patients with lung cancer with brain metastases. In this Bayesian clinical trial, we evaluated the combination of atezolizumab plus carboplatin and pemetrexed in patients with advanced nonsquamous non–small-cell lung cancer, who had brain metastases that were neurologically asymptomatic or controlled with corticosteroids and had not previously received any local brain therapies.

Knowledge Generated

Initiating systemic therapy with chemoimmunotherapy in patients with untreated brain metastases was safe and yielded relevant intracranial and systemic responses that were concordant in most patients. When disease progressed in the brain, most patients were rescued with brain radiotherapy. We observed that chemoimmunotherapy made it possible to delay whole-brain radiotherapy without a decline in quality of life and neurologic status.

Relevance (T.E. Stinchcombe)

This study provides prospective data about the activity and safety of chemotherapy and immunotherapy in patients with untreated brain metastases. Chemotherapy and immunotherapy as the initial therapy with imaging surveillance of the brain metastases is an option in select patients.*

*Relevance section written by JCO Associate Editor Thomas E. Stinchcombe, MD.

PD-L1–positive NSCLC who have brain metastases that are untreated or progressing after previous radiotherapy.⁸ Several post hoc exploratory analyses of phase III clinical trials showed that chemotherapy combined with immunotherapy improved overall survival (OS) compared with chemotherapy alone regardless of the presence of brain metastases.^{9,10} However, these trials were not specifically designed to evaluate the intracranial efficacy of chemoimmunotherapy and excluded patients receiving corticosteroids at doses higher than 10 mg daily of prednisone or equivalent.

We designed this phase II clinical trial to evaluate the safety and the efficacy of atezolizumab combined with carboplatin and pemetrexed in patients with advanced nonsquamous NSCLC with untreated brain metastases. We present the final analysis in this report.

METHODS

Study Design and Eligibility

This was an open-label, single-arm, phase II clinical trial conducted at 15 hospitals in Spain. Patients were age 18 years and older, had Eastern Cooperative Oncology Group performance status (ECOG PS) 0–1, and had stage IV nonsquamous NSCLC with untreated brain metastases that did not exhibit neurologic symptoms or that were controlled with anticonvulsants or dexamethasone at a maximum dose of 4 mg daily, with measurable disease in the body by computed tomography (CT) per RECIST version 1.1 criteria in the body and in the brain by magnetic resonance imaging (MRI) per response assessment in neuro-oncology brain metastases (RANO-BM), with

at least one measurable lesion with a minimum size of 10 mm.¹¹ There was no limitation on the maximum number of brain metastases or the maximum size per lesion. Patients had adequate bone marrow, liver, and renal function. Exclusion criteria included known *EGFR* mutations or *ALK* rearrangements; presence of leptomeningeal carcinomatosis or metastases in the brainstem, medulla, or lesions causing obstructive hydrocephalus; contraindication for immunotherapy (history of active autoimmune, infectious disease, or interstitial lung disease); previous malignancies within 3 years of study entry; previous treatment with immune checkpoint inhibitors; and hepatitis B or C infection or HIV positivity.

The study complied with the Declaration of Helsinki, International Council on Harmonisation Good Clinical Practice, local laws, and regulatory requirements. Independent ethics committees or institutional review boards approved the Protocol (online only). Patients provided written informed consent.

Study Procedures and Treatment

Patients were treated with atezolizumab 1,200 mg intravenously once every 3 weeks in combination with carboplatin (AUC 5) plus pemetrexed (500 mg per square meter). All the patients received premedication with folic acid, vitamin B12, and dexamethasone according to guidelines for pemetrexed use. After completing four to six cycles, patients continued with pemetrexed plus atezolizumab maintenance until unacceptable toxicity, disease progression, patient decision, completion of 2 years of therapy, or patient withdrawal of consent (Data Supplement [Fig S1], online only).

A body CT scan and a brain MRI were performed at baseline, every 6 weeks until the 12th week, and thereafter every 9 weeks until disease progression. Patients with progression exclusively in the brain could receive brain radiotherapy while staying within the study if they continued to derive clinical benefit and had ECOG PS ≤ 2 . In case of systemic progression without brain progression, a novel line of systemic treatment was considered.

Adverse events and abnormal laboratory findings were graded according to the National Cancer Institute Common Terminology Criteria for Adverse Events (NCI-CTCAE) version 4.0. Investigators determined whether adverse events were treatment-related according to the study protocol and standard regulatory requirements. Dose reductions were not permitted for atezolizumab, which could be interrupted, delayed, or discontinued depending on tolerability. Reductions were permitted for pemetrexed and carboplatin in accordance with two levels of dosage specified in the trial protocol. PD-L1 tumor proportion score expression was conducted at each center by immunohistochemical analysis after clinical practice using the Dako 22C3 and Ventana SP263 antibodies. PD-L1 expression was considered positive when $\geq 1\%$ of tumor cells had PD-L1 membranous staining.

Study Outcomes

The two primary end points were safety assessed by NCI-CTCAE version 4.0 defined as the incidence of grade 3 or higher adverse events during the first 9 weeks and progression-free survival (PFS) according to RANO-BM and RECIST v1.1 criteria. Overall PFS was defined as the time from time from inclusion until death from any cause or objective tumor progression evaluated by the investigators using RANO-BM in the brain and RECIST v1.1 in the body, whichever occurred first.¹¹

Secondary end points were the intracranial and systemic response rate (overall response rate [ORR]), the duration of response ([DOR] time from first documented complete or partial response [PR] to disease progression or death), and OS. OS was defined as the time from inclusion to death from any cause. The exploratory end points consisted of determining the time to brain radiotherapy and the event-free survival (EFS) defined as time from inclusion to brain radiotherapy or death, and to evaluate the correlation between PD-L1 expression in tumor tissue and efficacy end points. Additional exploratory end points were the assessment of QOL using the European Organisation for Research and Treatment of Cancer QLQ-C30 questionnaire that was administered at baseline and every 3 months. Global health status and overall QOL were analyzed in patients who had a baseline measurement.

Statistics

Safety and efficacy were assessed in the intention-to-treat population cohort of 40 patients using the Bayesian Multicohort design (Data Supplement [Methods]).^{12,13} Details about

previous and posterior distributions are detailed in the protocol. A sample size of 40 patients ensured that, if the trial is not terminated early, a posterior 90% credibility interval (CrI) would have width of 0.257 at most, under the assumption of a 50% PFS rate at 12 weeks. Futility and excess of toxicity stopping rules on the basis of posterior probabilities were evaluated sequentially every five patients (Data Supplement [Fig S2]). Posterior distributions with CrI at 95% were calculated for response or toxicity rates. Kaplan-Meier method was used to estimate PFS and OS. Data for patients who were alive or lost to follow-up were censored for OS at the time they were last known to be alive. Data for patients who were alive and did not have disease progression or who were lost to follow-up were censored for the analysis of PFS at the time of the last imaging assessment. When patients had tumor progression in one compartment (brain or systemic), they were censored for the PFS analysis in the other compartment. The stratified log-rank test was used to assess differences in PFS and OS among different subgroups. Hazard ratios and associated 95% CI were calculated with the use of a stratified Cox proportional-hazards model.

Data collection and data quality was ensured by the Spanish Lung Cancer Group. All analyses were based on the study database lock on March 31, 2022. All statistical analysis was performed using R Statistical Software (v 4.1.3; San Francisco, CA).

RESULTS

Patients and Disease Characteristics

Study enrollment was completed between November 2018 and December 2019. Forty-three patients were registered, but 40 patients met eligibility criteria and were enrolled at 13 sites (Fig 1; Data Supplement [Table S1]). Baseline patient characteristics are shown in Table 1. The median age was 62.5 years. Twenty-nine (72.5%) patients were male and most had a smoking history (85.0%). Brain metastases were diagnosed concurrently with lung cancer in most patients (92.5%). PD-L1 expression was $\geq 1\%$ in 20 patients (50.0%), negative in 18 patients (45.0%), and unknown in two patients (5.0%). The median total number of brain metastases per patient was 5, ranging from 1 to 20. The median size of the sum of all target lesions per patient was 13 mm, ranging from 10 to 42 mm. Twenty-two (55.0%) patients were receiving dexamethasone during the week before inclusion in the study to control neurologic symptoms with continued administration throughout the study, regardless of pemetrexed premedication. Sixteen patients were receiving 4 mg of dexamethasone once daily while six patients received < 4 mg once daily.

Efficacy

The study was completed since the boundaries for futility or unacceptable toxicity were not reached (NR) at any moment. The overall PFS rate at 12 weeks was 62.2% (95% CrI, 47.1 to

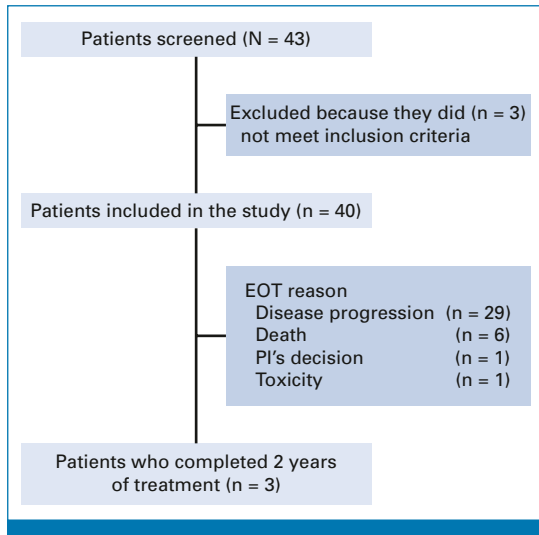


FIG 1. Study flowchart. EOT, end of treatment; PI, principal investigator.

TABLE 1. Baseline Patient Characteristics

Characteristic	Overall (N = 40)
Age, years, median (IQR)	62.6 (11.5)
Sex, No. (%)	
Female	11 (27.5)
Male	29 (72.5)
Race, No. (%)	
Caucasian	40 (100)
Smoking status, No. (%)	
Never	6 (15.0)
Former	11 (27.5)
Current	23 (57.5)
ECOG PS, No. (%)	
0	14 (35.0)
1	26 (65.0)
Histology, No. (%)	
Lung adenocarcinoma	39 (97.5)
NOS	1 (2.5)
PD-L1 expression, No. (%)	
≥50%	10 (25.0)
1%-49%	10 (25.0)
0%	18 (45.0)
Unknown	2 (5.0)
Baseline corticosteroids, No. (%)	
Yes	22 (55.0)
No	18 (45.0)
Diagnosis of brain metastases, No. (%)	
Synchronous	37 (92.5)
Metachronous	3 (7.5)
Total No. of brain lesions per patient, median (range) ^a	5 (1-20)
Total No. of target brain lesions per patient, median (range)	1 (1-4)

Abbreviations: ECOG, Eastern Cooperative Oncology Group; NOS, not otherwise specified; PS, performance status.

^aIncluding target and nontarget lesions.

Treatment at Intracranial Progression

During the study, 24 patients (60%) received brain radiotherapy because of PD. Sixteen patients received WBRT and eight patients received SRS. Median time to brain radiotherapy was 10.9 months (95% CI, 7.8 to 15.9; Data Supplement [Fig S7]). Thirteen patients died before receiving brain radiotherapy mainly because of complications related with extracranial progression. Median EFS, defined as the time from inclusion to brain radiotherapy or death, was 7.6 months (95% CI, 5.5 to 10.9).

Toxicity

The median number of cycles of carboplatin was four and of atezolizumab and pemetrexed was 8; three patients completed the planned 2 years of treatment. The rate of grade 3-4

76.2; Data Supplement [Fig S3]), above the expected 50%. Seventeen patients (42.7%; 95% CrI, 28.1 to 57.9; Table 2) had a confirmed intracranial response (12 [70.6%] PRs and 5 [29.4%] complete responses [CRs]), while 17 patients had stable disease (SD) in the brain and five had progressive disease (PD). The median time to intracranial response was 82 days and the median DOR in the brain among patients with CNS response was 14 months (95% CI, 10 to NR; Data Supplement [Fig S4]). Eighteen patients (45%; 95% CrI, 28.1 to 57.9; Table 2) achieved a systemic response (17 [94%] PRs and 1 [6%] CR), while 16 patients achieved SD and four patients had PD. Most responses were concordant in the brain and in the body, except for 6 (15%) patients (Fig 2). Three patients had systemic progression with SD in the brain and three patients had intracranial progression with PR or SD in the body. In an exploratory analysis, the intracranial ORR was similar among patients receiving corticosteroids (50%) compared with those who did not receive them (38.9%). The systemic ORR was also consistent between patients who were being treated (52.6%) or not with corticosteroids (44.4%). No differences were observed in the intracranial or systemic ORR according to PD-L1 expression (Data Supplement [Fig S5]).

With a minimum follow-up of 27 months, the median intracranial PFS was 6.9 months (95% CI, 4.7 to 11.9; Fig 3A) and the median systemic PFS was 8.9 months (95% CI, 6.7 to 13.8; Fig 3B). The median OS was 11.8 months (95% CI, 7.6 to 16.9; Fig 3C). Estimated 1- and 2-year OS rates were 50% (95% CI, 36.7 to 68.2) and 27.5% (95% CI, 16.6 to 45.5), respectively. No significant differences were observed in OS according to PD-L1 expression and baseline corticosteroids (Data Supplement [Fig S6]). Nevertheless, the 2-year OS rate for patients with PD-L1 ≥1% was 40.0% (95% CI, 23.4 to 68.4), while in patients with PD-L1 <1%, it was 16.7% (95% CI, 5.9 to 46.8).

TABLE 2. Efficacy Results

Result (N = 40)	Intracranial	Systemic
Best overall response, No. (%)		
CR	5 (12.5)	1 (2.5)
PR	12 (30.0)	17 (42.5)
SD	17 (42.5)	16 (40.0)
PD	5 (13.0)	4 (10.0)
NE	1 (2.5)	2 (5.0)
ORR, % (95% CrI)	42.7 (28.1 to 57.9)	45.0 (28.1 to 57.9)
Median DOR, months (95% CI)	14 (10 to NR)	11.9 (8.9 to NR)
12-week PFS rate, % (95% CrI)	62.2 (47.1 to 76.2)	
Median PFS, months (95% CI)	6.9 (4.7 to 11.9)	8.9 (6.7 to 13.8)

NOTE. Intracranial outcomes were evaluated using RANO-BM, while systemic outcomes were assessed by RECIST v1.1. Posterior CrIs were computed.

Abbreviations: CR, complete response; CrI, credibility interval; DOR, duration of response; NE, not evaluable; NR, not reached; ORR, overall response rate; PD, progressive disease; PFS, progression-free survival; PR, partial response; RANO-BM, response assessment in neuro-oncology brain metastases; SD, stable disease.

toxicity during the first 9 weeks was 27.5%, below the pre-defined boundary of 35%. No grade 5 adverse events were reported during the first 9 weeks. All neurologic adverse events, irrespective of causality to systemic therapy and time of onset, are shown in Table 3. Most neurologic adverse events were grade 1 and 2, and only five patients (12.5%) experienced grade 3-4 neurologic toxicity consisting of grade 4 hallucinations in one patient, grade 3 seizure in two patients, and grade 3 sciatica and spinal cord compression in one patient

each. The median time to the appearance of treatment-related neurologic toxicity was NR and the median time to the appearance of any neurologic toxicity was 15.7 months (Data Supplement [Figs S8A and S8B]). Global health status and overall QOL scores were maintained throughout the study (Data Supplement [Figs S9A and S9B]).

Twenty-eight patients (70%) had grade 3-4 treatment-related adverse events, but only 7 (17.5%) were deemed serious: acute kidney injury and pneumonitis in two patients each, while nephritis, pulmonary embolism, and febrile neutropenia were seen in one patient each (Data Supplement [Table S2]). One patient experienced grade 5 toxicity, consisting of febrile neutropenia and sepsis, that was considered related to chemotherapy.

DISCUSSION

The results of this phase II clinical trial show that patients with advanced NSCLC with untreated brain metastases can benefit from initiating systemic treatment with atezolizumab combined with chemotherapy. Previous exploratory analysis from pivotal phase III clinical trials have demonstrated that patients with brain metastases benefited from the addition of immunotherapy to chemotherapy versus chemotherapy alone. However, these reports were based on subgroup analyses with significant attrition since most studies excluded patients with untreated brain lesions or receiving corticosteroids or were not designed to evaluate the efficacy outcomes in the CNS.

To our knowledge, this is the first study centered on this special population that evaluates the activity and safety of a PD-L1

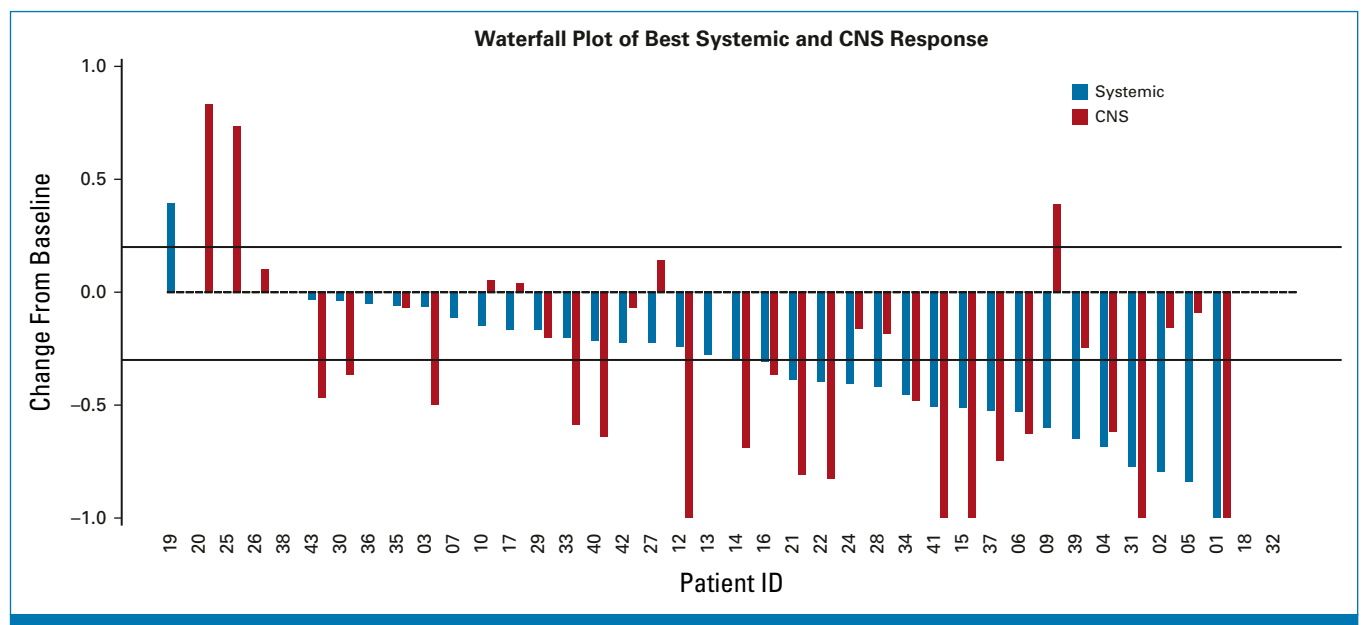


FIG 2. Waterfall plot of best ORR according to RECIST version 1.1 (depicted by the red bars) and RANO-BM (depicted by blue bars). Two patients are not shown because of on-treatment assessments not being available. ORR, overall response rate; RANO-BM, response assessment in neuro-oncology brain metastases.

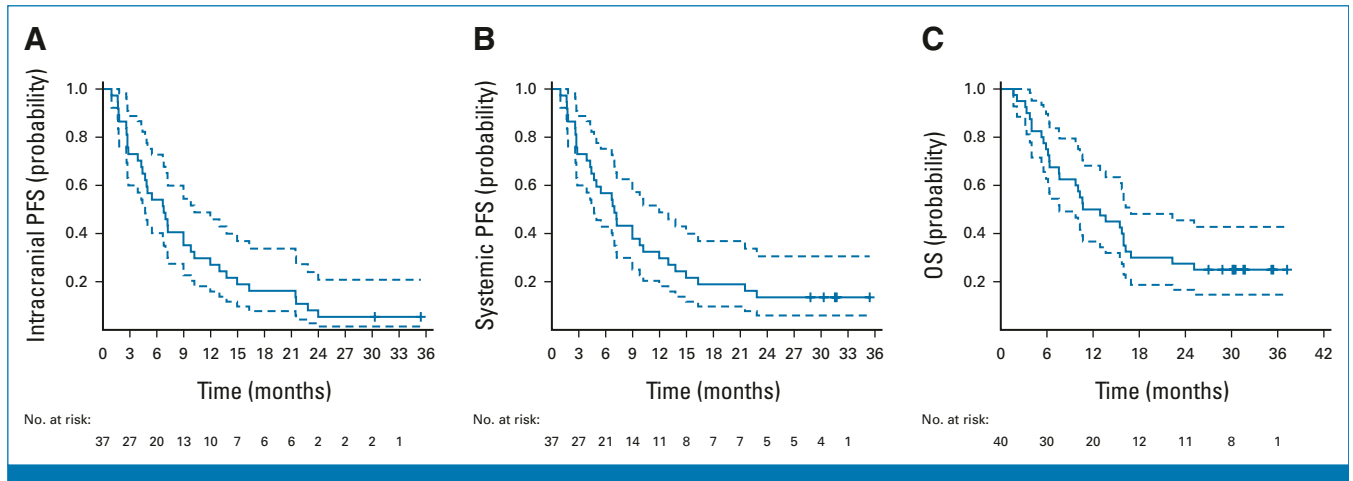


FIG 3. Kaplan-Meier plots of (A) intracranial PFS according to RANO-BM and (B) systemic PFS according to RECIST v1.1. (C) Kaplan-Meier plot of OS. Dotted lines indicate 95% CI. OS, overall survival; PFS, progression-free survival; RANO-BM, response assessment in neuro-oncology brain metastases.

inhibitor combined with platinum-based chemotherapy in patients with NSCLC with untreated brain metastases. This phase II trial has a Bayesian sequential design to minimize the risk of harm or futility on this vulnerable population. Forty patients were included, and the study was successfully completed. Some patients had significant disease burden on the basis of the total number of brain metastases and half of the patients were receiving corticosteroids at baseline. The patients received atezolizumab combined with carboplatin and pemetrexed, which was previously evaluated in the IMpower132 trial in advanced nonsquamous NSCLC; however, this study excluded patients with untreated brain metastases and those requiring corticosteroids.¹⁴

We assessed efficacy outcomes in the brain using RANO-BM to allow future cross-trial comparisons.¹¹ In our study, chemoimmunotherapy yielded similar ORR in the brain and in the body, and responses were highly concordant in both

compartments. Chemoimmunotherapy led to a complete intracranial response in five patients. Brain pseudoprogression is uncommon but has been reported with single immunotherapy in a large NSCLC cohort.¹⁵ In our study, we did not observe any case of pseudoprogression. An exploratory analysis did not show significant correlation between intracranial response and PD-L1 expression or corticosteroids treatment at baseline.

In our study, the median intracranial PFS by RANO-BM was 6.9 months, consistent with previous PFS reported in the pooled analysis of Keynote-021G, -189, and -407 trials evaluating pembrolizumab plus chemotherapy.⁹ This exploratory analysis included a subset of patients with untreated brain metastases but did not report the intracranial efficacy, and PFS was exclusively measured by RECIST v1.1. In an exploratory analysis of the CheckMate-9LA trial, the combination of two cycles of chemotherapy plus nivolumab and ipilimumab yielded a promising median intracranial PFS of 13.5 months in

TABLE 3. Frequency of Neurologic Events in All Patients Regardless of Their Relationship and Their Frequency at Any Time

NCI-CTCAE Version 4.0 (N = 40)	Grade 1, No. (%)	Grade 2, No. (%)	Grade 3, No. (%)	Grade 4, No. (%)	Grade 5, No. (%)
Hallucinations	0	0	0	1 (2.5)	0
CNS disorders—other ^a	3 (7.5)	0	2 (5.0)	0	0
Seizure	1 (2.5)	0	2 (5.0)	0	0
Headache	8 (20.0)	1 (2.5)	0	0	0
Insomnia	2 (5.0)	1 (2.5)	0	0	0
Anxiety	0	1 (2.5)	0	0	0
Edema cerebral	0	1 (2.5)	0	0	0
Dizziness	4 (10.0)	0	0	0	0
Peripheral sensory neuropathy	1 (2.5)	0	0	0	0
Confusion	1 (2.5)	0	0	0	0
Depression	1 (2.5)	0	0	0	0
Vertigo	1 (2.5)	0	0	0	0

Abbreviation: NCI-CTCAE, National Cancer Institute Common Terminology Criteria for Adverse Events.

^aSciatica and spinal cord compression.

patients with previously treated and stable brain metastases, without corticosteroids and asymptomatic.¹⁰ However, this study population was highly selected and cannot be directly compared with our study, which is enriched with more vulnerable patients. Nevertheless, in our study, the 2-year OS was 27.5% in the overall population and 40% in patients with PD-L1 expression $\geq 1\%$, similar to that reported in the seminal study of pembrolizumab enrolling patients with PD-L1-positive NSCLC with untreated brain metastases.⁸ Because of the limited statistical power of our study, it is not possible to determine whether the baseline corticosteroids have an effect on ORR or OS.

Most patients who progressed in the brain were considered candidates for salvage brain radiotherapy, predominantly consisting of WBRT, although some were treated with SRS. The greater use of WBRT was attributed to regional practice trends and to a high number of brain metastases in certain patients. Median time to brain radiotherapy was 10.9 months. In our study, we observed that systemic therapy allowed for the deferral of WBRT without a decline in QOL.¹⁶ However, when considering the initiation of initiating systemic therapy and the deferral of brain-directed local therapy in patients with lung cancer, it is important to discuss within a multidisciplinary tumor board that includes radiation oncologists and neurosurgeons.⁶

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Most neurologic adverse events were mild or moderate, except four patients who had grade 3 and 1 with grade 4 neurologic events that fully recovered. Treatment-related serious adverse events were observed only in 17.5% and there was only one patient with grade 5 toxicity related to chemotherapy. This safety profile is consistent with the previously published data using this treatment combination.¹⁴

The major limitation of the study is the single-arm design, which precludes us from establishing the optimal treatment strategy for patients with NSCLC and synchronous brain metastases. Randomized clinical trials would help to understand how to integrate brain local therapy with immunotherapy or chemoimmunotherapy in this patient population. Another limitation is the relatively small sample size, which reduces the statistical power to conduct subgroup analyses.

In summary, atezolizumab combined with carboplatin and pemetrexed has activity in patients with treatment-naïve advanced nonsquamous NSCLC with untreated brain metastases. Systemic and intracranial efficacy was similar, thus highlighting that atezolizumab plus chemotherapy could be a therapeutic option in this highly vulnerable population of patients. Future studies targeting additional immune checkpoints or integrating systemic therapies with stereotactic radiotherapy to improve the control rate and minimize the risk of brain toxicity are warranted.

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DISCLAIMER

The content is solely the responsibility of the authors. Roche had no role in data collection, data analysis, data interpretation, or writing of the report. All authors had full access to all the data in the study and provided approval to submit the manuscript for publication. The corresponding author had final responsibility for the decision to submit for publication.

PRIOR PRESENTATION

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CLINICAL TRIAL INFORMATION

[NCT03526900](https://clinicaltrials.gov/ct2/show/study/NCT03526900)

AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

Disclosures provided by the authors are available with this article at DOI <https://doi.org/10.1200/JCO.22.02561>.

DATA SHARING STATEMENT

A data sharing statement provided by the authors is available with this article at DOI <https://doi.org/10.1200/JCO.22.02561>.

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AUTHORS' DISCLOSURES OF POTENTIAL CONFLICTS OF INTEREST

Phase II Trial of Atezolizumab Combined With Carboplatin and Pemetrexed for Patients With Advanced Nonsquamous Non–Small-Cell Lung Cancer With Untreated Brain Metastases (Atezo-Brain, GECP17/05)

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