Depatuxizumab-mafodotin in *EGFR*-amplified newly diagnosed glioblastoma: a phase III randomized clinical trial

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ABSTRACT

Background: Approximately 50% of newly diagnosed glioblastomas (GBMs) harbor *EGFR* gene amplification (*EGFR*-amp). Preclinical and early phase clinical data suggested efficacy of depatuxizumab mafodotin (depatux-m), an antibody drug conjugate (ADC) comprised of a monoclonal antibody that binds activated EGFR (overexpressed wild-type and EGFRvIII-mutant) linked to a microtubule-inhibitor toxin in *EGFR*-amp GBMs.

Methods: In this phase III trial, adults with centrally confirmed, *EGFR*-amp, newly diagnosed GBM were randomized 1:1 to radiotherapy, temozolomide, and depatux-m/placebo. Corneal epitheliopathy (CE) was treated with a combination of protocol-specified prophylactic and supportive measures. There was 85% power to detect a Hazard Ratio (HR) ≤ 0.75 for survival (OS) at a 2.5% one-sided significance level (i.e., traditional two-sided p ≤ 0.05) by log-rank testing.

Results: There were 639 randomized patients (median age 60, range 22-84; 62% men). Pre-specified interim analysis found no improvement in OS for depatux-m over placebo (median 18.9 vs. 18.7 months, HR 1.02, 95% CI 0.82-1.26, one-sided p= 0.63). Progression-free survival was longer for depatux-m than placebo (median 8.0 vs. 6.3 months; HR 0.84, 95% CI 0.70-1.01, p=0.029), particularly among those with *EGFRvIII* mutant (median 8.3 vs. 5.9 months, HR 0.72, 95% CI 0.56-0.93, p=0.002 one sided) or *MGMT* unmethylated (HR 0.77, 95% CI 0.61-0.97; p=0.012 one-sided) tumors but without an OS improvement. CE occurred in 94% of depatux-m treated patients (61% grade 3-4), causing 12% to discontinue.

Conclusions: Interim analysis demonstrated no OS benefit for depatux-m in treating *EGFR*-amp newly diagnosed GBM. No new important safety risks were identified.

Keywords: Glioblastoma, EGFR, depatuxizumab-mafodotin, antibody drug conjugate, phase III

Key Points:

Approximately 50% of newly diagnosed glioblastomas harbor *EGFR*-amplification (EGFR-amp).

The antibody-drug conjugate depatuxizumab mafodotin binds activated EGFR.

Depatuxizumab mafodotin did not improve overall survival in *EGFR*-amp newly diagnosed glioblastoma.

Importance of the Study:

In this phase III clinical trial, there was no improvement in survival from treatment with the EGFR-directed antibody-drug conjugate depatuxizumab-mafodotin (depatuix-m) over placebo in addition to standard chemoradiotherapy. Progression-free survival was longer among patients randomized to depatux-m, particularly in *EGFRvIII* mutant cases. Corneal epitheliopathy occurred in most depatux-m treated patients caused a small minority to discontinue.

INTRODUCTION

Glioblastoma (GBM) is the most common primary brain tumor in adults. Prognosis is poor; new approaches are needed. Focal epidermal growth factor receptor (*EGFR*) gene amplification on chromosome 7 (*EGFR*-amp) has long been observed[1] in approximately 50% of GBMs (although geographic differences exist).[2] EGFR variant 3 (EGFRvIII) mutation, a tumor-specific deletion of exons 2-7, is constitutively active and observed in approximately 50% of *EGFR*-amp GBMs (~25% overall).[3]. Several EGFR/EGFRvIII-directed therapeutic approaches have been used, including receptor tyrosine kinase inhibitors (RTKIs),[4] antibodies, [5-7] and vaccines.[8] Despite TKI success in molecularly selected non-small lung cancers[9] and with antibodies in other solid tumors,[10] these approaches have been disappointing for GBM.[4]

Depatuxizumab (depatux, formerly ABT-806) is a humanized recombinant monoclonal antibody originally generated against EGFRvIII in mice,[11] although it also binds to wild-type EGFR when present at high levels.[12] The epitope becomes accessible to the antibody when EGFR is activated, either by ligand for wild-type receptor or constitutive mutation (e.g., EGFRvIII).[12,13] The antibody-drug conjugate (ADC) depatuxizumab mafodotin (depatux-m, formerly ABT-414, **Figure S1**) links depatux to a microtubule cytotoxic payload, monomethyl auristatin F (MMAF, mafodotin).[14,15] Following binding to activated EGFR, the antibody and linked payload are endocytosed and degraded in acidic endocytic compartments, releasing the toxin causing cell death.[16] This direct cytotoxic effect of the ADC, therefore, does not rely on inhibition of EGFR signaling and does not cause rash, diarrhea or other toxicities typical of receptor tyrosine kinase inhibitors or monoclonal antibodies that bind to unamplified wild-type receptor in normal organs.[5] Although GBMs do not respond to the unconjugated antibody (depatux),[5] depatux-m is effective against

EGFR-amp and EGFRvIII harboring GBM cell lines and animal models, both alone and combined with radiotherapy (RT) and temozolomide.[15] In addition, ADCs have superior efficacy to unconjugated monoclonal antibodies in other solid tumors, with several under investigation in many cancers and conditions[17] including GBM.[18]

Therefore, we previously conducted a phase 1 trial of depatux-m and identified a recommended dose for use alone or in combination with RT and/or temozolomide. Radiographic responses were observed, mainly *EGFR*-amp disease.[19-22] Corneal epitheliopathy (CE, previously termed ocular side effects or keratopathy)[23] was very common but typically reversible. Of note, another mafodotin-containing biologic was US FDA approved for myeloma despite a similarly high frequency of CE.[24]

The encouraging pre-clinical[15] and early phase clinical data formed the basis of two large international randomized trials. The open label phase II European Organisation for the Research and Treatment of Cancer (EORTC) trial 1410 (AbbVie M14-483, INTELLANCE 2, NCT02343406) accrued patients with *EGFR*-amp recurrent GBM. In the primary analysis (median follow-up 15.0 months), results trended toward longer overall survival (OS) following treatment with depatux-m in combination with temozolomide compared to control of lomustine or temozolomide (Hazard Ratio, HR 0.71; 95% Confidence Interval, CI, 0.50-1.02; log rank p=.06). In the exploratory follow-up analysis (median follow-up 28.7 months), the HR was 0.66 for the comparison of the combination arm vs. control (95% CI 0.47-0.93, log rank p=.017).[25] Concurrently, we conducted a randomized, double-blind placebo-controlled phase III trial of depatux-m for newly diagnosed *EGFR*-amp GBM as an academic-industry collaboration between the Radiation Therapy Oncology Group Foundation

(RTOG-F 3508) and AbbVie (M13-813, INTELLANCE 1, NCT02573324) and report results here.

METHODS

Eligibility

Patients were ≥18 years old and had Karnofsky Performance Status ≥70, an RT and chemotherapy naïve unifocal GBM harboring *EGFR*-amp, and end-organ function. Laser-assisted in situ keratomileusis (LASIK) within the prior year, cataract surgery within the prior 3 months, and other contraindication to ocular corticosteroids required as supportive care for CE (below) were exclusionary. All subjects provided written informed consent prior to any study-specific procedures, and the study was approved by the Institutional Review Board of each participating institution. Detailed criteria are available in the protocol (**Supplemental Material**).

Biomarkers

Biomarkers (**Table S1**) and histology (GBM by World Health Organization 2016 criteria,[26] KA) were confirmed centrally before randomization as described previously: *EGFR*-amp by Fluorescence in Situ Hybridization (FISH),[2] *MGMT* by methylation-specific polymerase chain reaction (PCR)[19], and *EGFRvIII* mRNA by reverse-transcription-PCR.[19] *Isocitrate dehydrogenase* (*IDH*) mutation is typically mutually exclusive with *EGFR*-amp and was not assessed.[26]

Treatment

Up to 7 weeks following diagnostic surgery, eligible subjects were randomly assigned 1:1 to RT, temozolomide, and either depatux-m or placebo in a stratified (below) double-blind manner. RT was planned using a post-operative contrast-enhanced baseline brain MRI to a

total dose of 60 Gy in 30 fractions (or 59.4 Gy in 33 fractions) over approximately 6 weeks. A planning MRI (repeated if necessary) was obtained ≤ 4 weeks post-operatively and ≤ 3 weeks before RT. Either a sequential boost to the contrast-enhanced region of the target as per standard RTOG approach or single-phase technique as per the EORTC approach were permitted.

Temozolomide was dosed at 75mg/m2 daily during RT followed by 6 adjuvant cycles of 150-200 mg/m2 on days 1-5/28[27] with up to 12 adjuvant cycles allowed. Depatux-m was dosed at 2.0 mg/kg during RT then 1.25 mg/kg thereafter on days 1 and 15/28[19,21] and allowed to continue until disease progression. Post-progression treatment was at the discretion of the treating investigator except cross-over from placebo to depatux-m was disallowed.

Supportive care

Prophylactic ocular corticosteroids were mandatory with each dose of depatux-m/placebo to reduce the potential for CE as described previously.[28] Additional ocular supportive care measures (e.g., lubricating eye drops, therapeutic bandage contact lenses, punctal plugs, and/or antibiotic drops, etc.) were recommended for both symptomatic relief of CE (e.g., photophobia, blurry vision, and/or other eye discomfort) and to reduce side effect driven interruptions or reductions of depatux-m dosing.

Pneumocystis jirovecii pneumonia (PJP, previously PCP) prophylaxis during chemoradiotherapy[27] and antiemetic prophylaxis before temozolomide were recommended. Growth factor and transfusion support were permitted for cytopenias other than to induce eligibility or affect temozolomide cycle length or dose. Systemic corticosteroids and anticonvulsants were allowed without restriction.

Follow-up

In addition to serial ophthalmologic examinations, patients underwent routine physical, neurologic, bone marrow, serum chemistry, and hepatic function evaluations at baseline, before every cycle, and more frequently as clinically indicated. Dose interruptions and reductions of depatux-m/placebo were permitted for treatment related CTCAE grade 2-3 and required for grade 4 ocular adverse events (such as corneal perforation or acuity ≤ 20/200). Up to 3 consecutive depatux-m/placebo dose reductions during chemoradiotherapy (by -0.5 mg/kg each) and up to 4 during adjuvant treatment (by -0.25 mg/kg each) were permitted for treatment-related toxicities. Re-escalations were permitted only for improved CE and serum chemistry abnormalities but not for other adverse events. Temozolomide adjustments were allowed per local prescribing regulations.

Baseline contrast-enhanced brain MRI scans, neurocognitive function (NCF) tests, and patient reported outcomes (PROs) were required before chemoradiotherapy and serially before odd-numbered adjuvant cycles (1, 3, 5, etc.) of temozolomide and depatux-m/placebo, and then every 8 weeks thereafter. Progression as a study endpoint was assessed centrally and retrospectively using Response-Assessment in Neuro-Oncology (RANO) criteria,[29] but treatment decisions were made using local interpretation in real time with continuation encouraged in equivocal scenarios.

Results of Neuro-Cognitive Function (NCF) testing and Patient Reported Outcomes (PROs) were also performed at the time of locally determined progression, although scoring and results were not used in treatment decision making; rather, NCF results and PROs were verified and associations evaluated centrally. The M. D. Anderson Symptom Inventory – Brain Tumor (MDASI-BT) questionnaire is a validated PRO instrument used to assess the

severity of brain tumor related symptoms and its impact on daily function. It consists of 22 symptom items and 6 interference items, each rated from 0 (best) to 10 (worse).[30,31] The symptom severity score and symptom interference score are the average of the symptom and interference items, respectively.[32-34] The Hopkins Verbal Learning Test – Revised (HVLT-R)[35] is a sensitive, highly standardized, validated neurocognitive test to assess change in verbal episodic learning and memory over time. There are six alternate forms to limit practice effects. The Total Recall score was chosen a priori as a secondary endpoint and is the sum of the total number of words recalled across 3 trials.

Study design

In order to balance known and potential prognostic factors between arms, randomization (using permuted block[36] sizes of 4 that was generated by the AbbVie Data and Statistical Sciences department) was stratified by Region of world, Radiation Therapy Oncology Group - Recursive Partitioning Analysis (RTOG-RPA) class (which incorporates age, performance status, extent of resection, and neurological function, **Table 1** legend),[37] O^6-methylguanine-DNA-methyltransferase (*MGMT*) promoter methylation status, and *EGFRvIII* mutation (as a mechanistically predictive biomarker for enhanced depatux binding to tumor cells). All randomized subjects were included in the intent-to-treat analysis.

Originally, a phase II/III trial was planned but accrual to phase II rapidly outpaced both the planned (progression-free survival) PFS analysis and phase III accrual goals, despite the stringent requirements for central pathology review and biomarker testing (**Figure S2**). In addition, early results from the concurrently conducted INTELLANCE-2 trial in recurrent GBM suggested depatux-m in combination with temozolomide improved OS relative to control.[25] Therefore, the trial design was amended as a phase III with OS as the primary

endpoint, but pre-specified an interim analysis for futility (or overwhelming superiority, below).

Median OS with placebo was estimated as 16 months and hypothesized to improve to 21.3 months with depatux-m. With 441 deaths among 640 randomized patients, we had 85% power to detect a ≥25% reduction in risk of death (HR ≤0.75) at a 2.5% one-sided level of significance (i.e., traditional two-sided p≤0.05). Anticipating delayed treatment effect, a Fleming Harrington version of weighted log-rank test with parameters $\rho=0$ and $\gamma=0.2$ was used. Thus, at least 66% of information, due to increased weighting for later events, would be accumulated at the interim analysis and resulted in testing the futility bound at HR>0.9 or the efficacy bound (for superiority) at a one-sided significance level of 0.0058. [38,39] Secondary and exploratory endpoints included PFS, molecular subgroup analyses, NCF and PROs.

PFS was defined as the interval from randomization to first of either progressive disease (by blinded independent central review per RANO criteria) or death from any cause, and OS to death from any cause. Subjects not experiencing progression or death were censored. NCF and PRO were analyzed using deterioration free survival (DFS), with deterioration defined using the reliable change index criterion for the HVLT-R Total Recall (i.e., as a reduction of 5 points as compared to baseline)[40] and a decrease of 1 point as compared to baseline for the MDASI-BT symptom severity and symptom interference scores. DFS was defined as the interval from randomization to first occurrence of deterioration or death from any cause (**Figures S6-S7 and Table S15**). Subjects not experiencing an event were censored.

Time to event (PFS, OS, and DFS) analyses were performed using the Kaplan-Meier method[54] with HRs and 95% CIs estimated using Cox proportional hazards regression models

adjusting for stratification factors. An Independent Data Monitoring Committee (IDMC), managed by RTOG-F, reviewed unblinded data and interim results.

Importantly, hierarchical testing was used for all secondary and exploratory analyses (**Table S2**) to reduce the potential for falsely identifying a significant difference when conducting multiple comparisons.[41] In this manner, subsequent differences in outcome between arms could only be considered statistically significant (regardless of the HR or p), if the prior analysis in the hierarchy were significant (two-sided $p \leq 0.05$). However, we report the pre-planned secondary and exploratory analyses descriptively to understand the trial outcomes thoroughly. Details of the statistical collaboration between AbbVie and RTOG Foundation can be found in the **Supplementary Material**.

RESULTS

Accrual

Accrual occurred at 190 sites in 26 countries from September 11, 2015 to March 31, 2018 (**Figures 2, S1; Table S3**). Among 2229 screened, lack of *EGFR*-amp by central analysis was the most common reason for ineligibility (61% of excluded cases). Central histology review nearly always (98%) confirmed a GBM diagnosis. The pace of accrual exceeded projections (**Figure S1**). As a consequence, the phase II/III design was converted to a phase III trial as outlined above.

Median age was 60 years (range 22-84), 62% (394/639) of randomized subjects were men, and 13% (81) were ≥ 70 years old. Baseline characteristics of the study population were similar (53% *EGFRvIII*-mutant, 37% methylated *MGMT*) to those of other newly diagnosed GBM trials and reports.[3,42,43] Arms were well balanced (**Table 1**).

Safety

The most common adverse events were ocular (grouped under the general term of CE, **Table S4**) consistent with prior reports. [19-22,25] For example, CE of any grade occurred in 94% of subjects randomized to depatux-m, although was surprisingly reported in 36% on the placebo arm. Grade 3 CE (vision decline to worse than 20/40 but better than 20/200, or limiting self-care activities of daily living) was reported in 55%, and grade 4 perforation or blindness with acuity 20/200 or worse in 5% of patients randomized to depatux-m (**Table 2**). CE of all grades was managed by a combination of both prophylactic and supportive measures and by dose interruptions or delays (44%), although complete discontinuation of protocol therapy was required infrequently (12% in the depatux-m arm, 0% in the placebo arm). Thrombocytopenia was also more commonly observed among patients randomized to depatux-m than placebo (61% any grade with 14% each grade 3 and 4 vs. 36% any grade with 6% each grade 3 and 4).

Survival

The pre-planned interim analysis was conducted in May 2019 after 346 deaths among all randomized patients (> 332 required). At that time, slightly more than 50% of patients in each arm died (169/316 placebo, 177/323 depatux-m), and nearly 70% in each arm had progressed by central review (219/316 placebo, 221/323 depatux-m). After median follow-up of 18.1 months among 293 surviving patients, there was no OS improvement for depatux-m over placebo (median 18.9 months for depatux-m vs. 18.7 months for placebo, HR 1.02, 95% CI 0.82-1.26, one-sided log rank p= 0.63; **Figure 3**). As the primary analysis for OS failed to demonstrate a significant difference between arms, subsequent endpoint analyses were exploratory (**Table S2**). PFS (centrally determined) was longer following depatux-m

than placebo (median 8.0 months vs. 6.3 months; HR 0.84, 95% CI 0.70-1.01; p = 0.029 one-sided; **Figure 3**), driven at least in part by the *EGFRvIII* mutant subgroup (median 8.3 vs. 5.9 months, HR 0.72, 95% CI 0.56-0.93, p=0.002 one-sided; **Figure 4**). By contrast, among those without *EGFRvIII* mutant disease, there was no difference in PFS between arms (median 6.9 months for depatux-m vs. 7.9 months for placebo, HR 1.01, 95% CI 0.76-1.33, p=0.61 one-sided, **Figure 4**).

There was no improvement in OS by treatment for any subgroup, although, as above, the study was not powered to detect a statistically significant difference (**Figures S3-S5, Tables S5-S14)**.

Finally, to explore *EGFRvIII* for prognostic importance regardless of treatment, we analyzed survival by mutational status among patients randomized to placebo to eliminate potential confounding by treatment with depatux-m (**Figure S5**). PFS was longer among cases without (n=148) than with (n=168) documented *EGFRvIII* (median PFS 7.9 months vs. 5.9 months, HR 0.74, 95% CI 0.57-0.97, p=0.03) but without a difference in OS (HR 0.95, 95% CI 0.70-1.29, p=0.76) in this post-hoc, underpowered, univariate analysis.

NCF and PROs

The compliance for the HVLT-R and MDASI-BT was similar: ≥93% at baseline, >80% at adjuvant week 1, ≥70% at adjuvant week 9, ≥58% at adjuvant week 17, ≥51% at adjuvant week 25, and ≥47% at adjuvant week 33 (**Table S15**). There were no differences between treatment arms with respect to baseline HVLT-R Total Recall and MDASI-BT scores. There was no between arm difference in DFS for HVLT-R Total Recall, symptom severity, or

symptom interference (HR 1.14, 95% CI 0.92-1.40, p=0.81; HR 1.33, 95% CI 1.09-1.63, p=0.99; HR 1.19, 95% CI 0.97-1.45, p=0.94, respectively; **Figures S6-S7**).

DISCUSSION

In this phase III trial, survival was not improved by depatux-m for people with newly diagnosed *EGFR*-amp GBM; the study was stopped early and unblinded for futility. PFS (centrally determined) was longer with depatux-m than placebo, particularly in the *EGFRvIII*-mutant subgroup. No DFS differences between arms in verbal learning, symptoms or symptom interference were observed. No new important safety risks from depatux-m were identified with reversible CE (which were also reported in the placebo arm) and thrombocytopenia observed most commonly. Patients on active treatment were permitted to continue after unblinding and re-consent.

There are several potential explanations for the negative result. Most importantly, despite encouraging pre-clinical and early phase clinical data, it is possible that depatux-m is simply ineffective for treating glioblastoma, notwithstanding any potential enrichment strategy. Other potential biologic explanations include the possibility that depatux-m effectively killed off *EGFR*-amp (and particularly *EGFRvIII*-mutant) tumor cells, lengthening PFS, but resistant clones emerged and voided any OS benefit, a hypothesis supported by results from patient derived xenografts[44]; we also previously demonstrated that *EGFR*-amp was preferentially lost in GBMs following treatment with depatux-m among longitudinally sampled tumor tissue on an intra-patient basis.[45] Pre-clinical work from others also supports emergence clones as a mechanism of acquired resistance. Also, our focus on *EGFR*-amp for eligibility may have inadequately enriched the study population for benefit, particularly as *EGFR* gene amplification correlated only imperfectly with response in our prior studies. A

better strategy may have been to power the study for, or restrict eligibility to, the *EGFR-vIII* mutant subgroup or set a lower bound on the minimum number of patients with other potentially depatux-sensitizing *EGFR* mutations[46,47] Our observation that PFS was shorter among *EGFRvIII*-mutant cases randomized to placebo further supports our impression that improved PFS with depatux-m in this subgroup was not spurious. We also previously described payload sensitizing mutations,[48] and penetration of depatux-m into large tumors may be limited,[49] although neither of these biomarkers were screening criteria. Finally, limited penetration of the blood brain barrier by depatux may also impede efficacy against intracranial tumors, particularly in the non-enhancing part of the tumor; this is a critically important lesson for future studies of large molecules.[44]

Finally, other ADCs are being investigated for GBM and other solid tumors.[50] Higher affinity antibodies conjugated to cell-permeant payloads (permitting bystander killing of adjacent tumor cells) with different safety profiles may result in different outcomes.

In retrospect, it may have been prudent to complete the originally planned phase II study, suspending accrual and deferring phase III until analyses were complete. This is an important consideration for future studies with a phase II/III design.

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Figure legends

Figure 1: Consolidated Standards of Reporting Trials (CONSORT) diagram of RTOG-F 3508/AbbVie M13-813 (INTELLANCE-1) at the time of pre-planned interim analysis

Figure 2: Overall and Progression-Free Survival

Overall (A) and Progression-Free Survival (B, by central review) curves by treatment arm among all randomized patients (intent-to treat). HR, Hazard Ratio with 95% Confidence Intervals

Figure 3: Overall and Progression-Free Survival by *EGFRvIII* mutation

Overall (A, C) and Progression-Free Survival (B, D; by central review) by treatment arm for patients with (A, B) or without (C, D) *EGFRvIII* mutation on an intent-to treat basis. HR, HR, Hazard Ratio with 95% Confidence Intervals

Table 1: Patient characteristics among randomized patients, n (%)

Baseline Characteristics: randomized patients (n=639)	**Placebo (n=316)**	**Depatux-M (n=323)**
Age, years		
Median	60	59
Range	29-82	22-84
Gender, n (%)		
Male	188 (59)	206 (64)
Female	128 (41)	117 (36)
Histology, (central review) n (%)		
Glioblastoma	311 (98)	319 (99)
Gliosarcoma	1 (<1)	3 (1)
Other	1 (<1)	1 (<1)
Missing	3 (1)	0 (0)
Karnofsky Performance Status (KPS), n (%)		
70	38 (12)	44 (14)
80	80 (25)	76 (23)
90-100	198 (63)	203 (63)
Extent of Resection (EOR), n (%)		
Gross total Resection	181 (57)	185 (57)
Partial/Subtotal Resection	122 (39)	128 (40)
Biopsy	10 (3)	10 (3)
Missing	3 (1)	0 (0)
Impairment of Neurologic Function (INF), n (%)		
> minor	25 (8)	27 (8)
≤ minor	288 (91)	296 (92)
Missing	3 (1)	0 (0)
***Radiation Therapy Oncology Group – Recursive Partitioning Analysis (RTOG-RPA) Prognostic Class[37], n (%)**		
III	46 (14)	51 (16)
IV	233 (74)	236 (73)
V	37 (12)	36 (11)
HVLT-R		
Total Recall, mean (SD)	**-1.5 (2.2)**	**-1.4 (1.9)**
***Region of World n (%)**		
Other	214 (68)	216 (67)
US/Canada	102 (32)	107 (33)
****MGMT*, n (%)**		
Methylated	117 (37)	118 (37)
Unmethylated	199 (63)	205 (63)
****EGFRvIII*, n (%)**		
Mutated	168 (53)	164 (51)
Other	148 (47)	159 (49)

*Stratification Factor

RTOG-RPA Class definitions

- III: Age < 50, KPS ≥ 90
- IV: Age < 50, KPS < 90; OR Age ≥ 50, KPS ≥ 70, EOR > biopsy, INF ≤ minor
- V: Age ≥ 50, KPS ≥ 70, EOR > biopsy, INF > minor; OR Age ≥ 50, KPS ≥ 70, EOR = Biopsy

Table 2: Grade 3 and 4 adverse events reported in at least 5% of patients

	Placebo (n=313) n (%)		Depatux-M (n=323) n (%)	
Adverse Event	**Grade 3**	**Grade 4**	**Grade 3**	**Grade 4**
Any	135 (43.1)	47 (15.0)	191(59.1)	69 (21.4)
Corneal epitheliopathy (CE)*	2 (0.6)	0	179 (55.4)	16 (5.0)
Thrombocytopenia	20 (6.4)	18 (5.8)	44 (13.6)	46 (14.2)
Gamma-glutamyltransferase increased	2 (0.6)	1 (0.3)	33 (10.2)	2 (0.6)
Lymphopenia	37 (11.8)	4 (1.3)	23 (7.1)	4 (1.2)
Seizure	16 (5.4)	4 (1.3)	16 (5.0)	2 (0.6)
Alanine aminotransferase increased	5 (1.6)	0	17 (5.3)	0
Neutropenia	15 (4.8)	10 (3.2)	9 (2.8)	6 (1.9)

*Includes keratopathy, vision blurred, photophobia, dry eye, eye pain, keratitis, and punctate keratitis

Figure 1

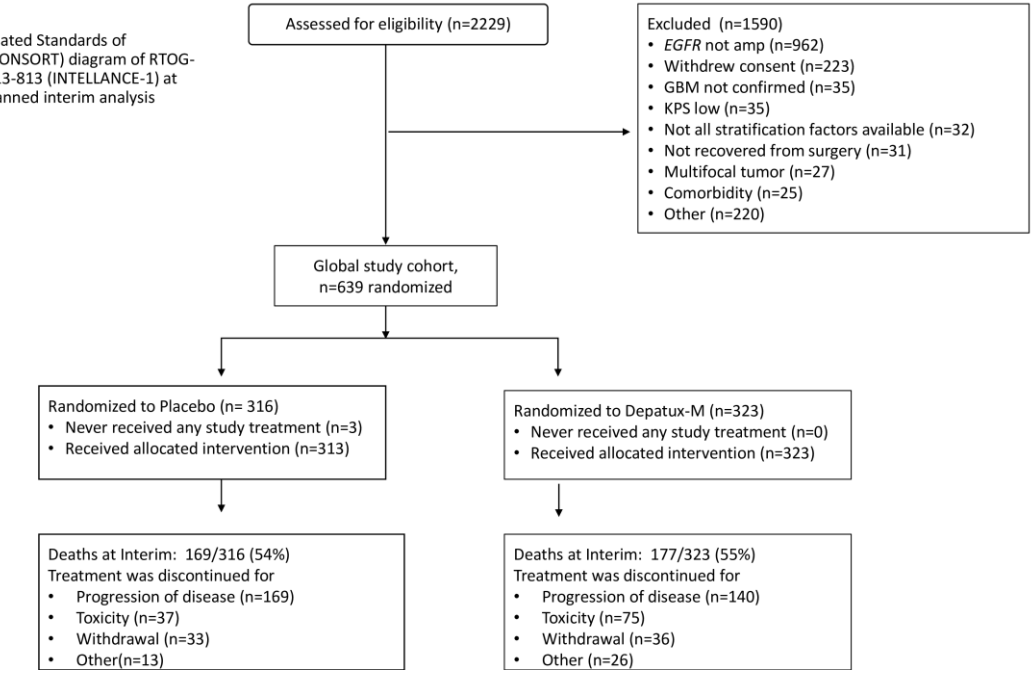

Figure 2: Consolidated Standards of Reporting Trials (CONSORT) diagram of RTOG-F 3508/AbbVie M13-813 (INTELLANCE-1) at the time of pre-planned interim analysis

Figure 2

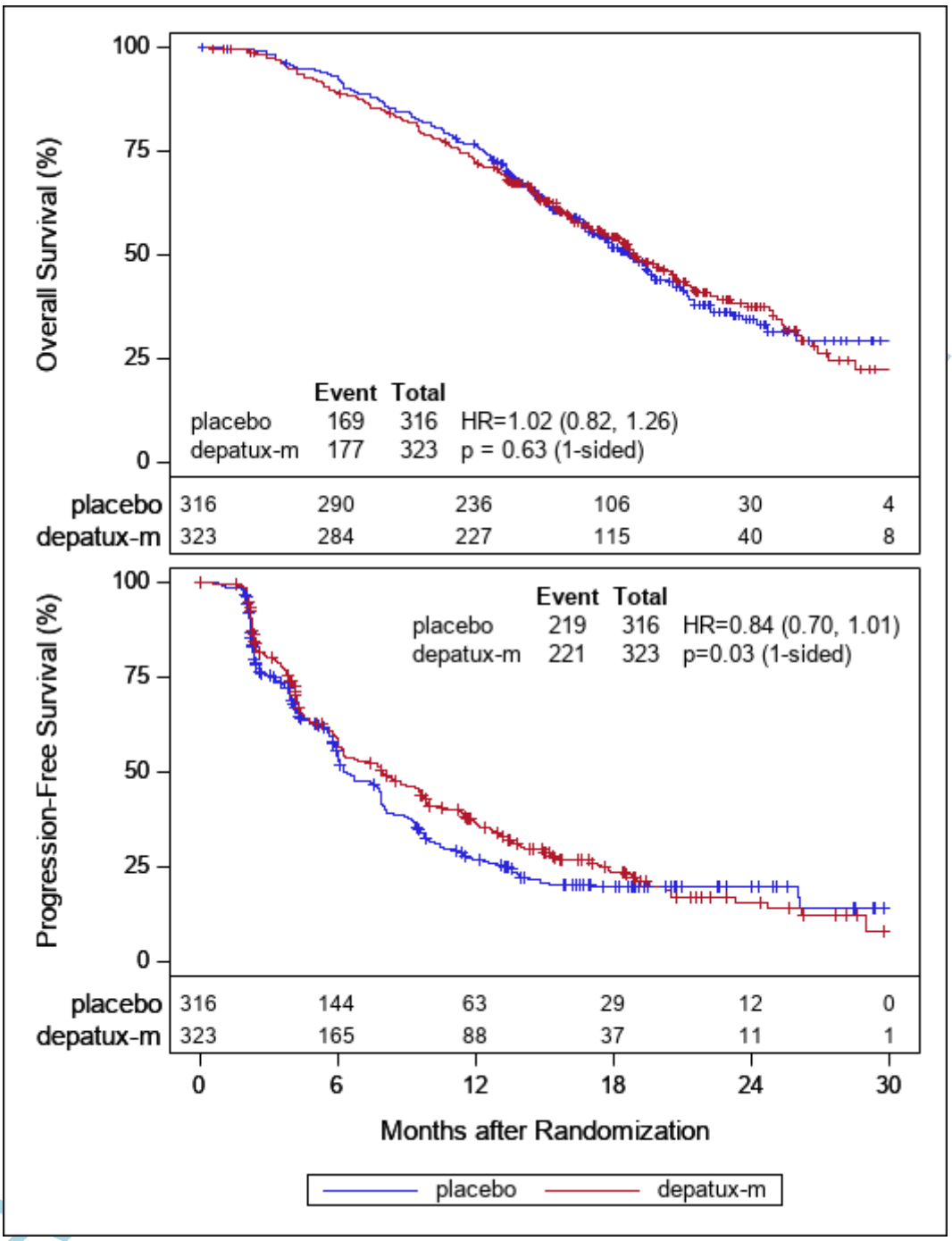

Figure 3