

# Efficacy of osimertinib in advanced T790M-positive NSCLC after progression to prior EGFR-TKI: real world data from a Brazilian cohort.

Eficácia de osimertinibe para CPNPC T790M-positivo avançado em progressão após EGFR-TKI prévia: dados do mundo real de um coorte brasileiro.

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## ABSTRACT

**Objectives:** Osimertinib is a third-generation EGFR inhibitor with activity against both sensitizing and resistance mutations. Following results of the phase III study, AURA3, which led to the approval of osimertinib worldwide, we have conducted ASTRIS with the aim of confirming the efficacy and safety of osimertinib. **Methods:** This is a phase IV, international, multicentric, open trial assessing the efficacy and safety of osimertinib at a dose of 80mg daily, orally. Eligible patients were those with diagnosis of T790M-positive NSCLC on progression after prior EGFR-TKI. Herein, we present the Brazilian experience at ASTRIS. **Results:** Eighty-eight patients were enrolled in Brazil between August, 2015 and March, 2017. The median age was 34-89 (years), and most were females (66%). Fifty-four patients (61%) had received prior therapy with erlotinib, forty-two (48%) with gefitinib, and 3 (3%) with afatinib. Exon 19 deletions were the most common primary mutation in EGFR, present in 55 cases (62.5%), followed by *L858R* in 24 cases (27%). The response rate was 58.2% (95%CI = 46.6-69.2), and median progression-free survival was 9.4 months (95%CI = 8.2-not reached). The most common AE was pneumonia (5 cases). Only 1 patient (1.1%) had a pneumonitis-like event and 2 patients (2.3%) had a prolongation of the QTc interval. **Conclusion:** In a real-world setting, osimertinib constitutes a safe and effective therapeutic option for Brazilian patients with advanced T790M-positive NSCLC after progression on a prior EGFR-TKI, including those patients with central nervous system metastasis. Our findings support previous observations and add valuable information regarding osimertinib effectiveness in Brazilian patients.

**Keywords:** Lung neoplasms; Genes, erbB-1; Molecular targeted therapy.

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## RESUMO

**Objetivos:** Osimertinibe é um inibidor de EGFR de terceira geração com atividade contra mutações sensibilizantes e resistentes. Após os resultados do estudo de AURA3, fase III, que levaram à aprovação do osimertinibe em todo o mundo, conduzimos o ASTRIS com o objetivo de confirmar a eficácia e a segurança do osimertinibe. **Métodos:** Trata-se de um estudo aberto, de fase IV, internacional, multicêntrico, que avalia a eficácia e a segurança do osimertinibe na dose de 80mg por dia, por via oral. Os pacientes elegíveis foram aqueles com diagnóstico de CPNPC T790M-positivo, em progressão após EGFR-TKI prévia. Aqui, apresentamos a experiência brasileira no ASTRIS. **Resultados:** Oitenta e oito pacientes foram matriculados no Brasil, entre agosto de 2015 e março de 2017. A idade média foi de 34-89 anos e a maioria era do sexo feminino (66%). Cinquenta e quatro pacientes (61%) haviam recebido terapia prévia com erlotinibe, quarenta e dois (48%) com gefinibe e 3 (3%) com afatinibe. As deleções do exão 19 foram as mutações primárias mais comuns no EGFR, presente em 55 casos (62,5%), seguida por L858R em 24 casos (27%). A taxa de resposta foi de 58,2% (IC95% = 46,6-69,2) e a sobrevida média livre de progressão foi de 9,4 meses (IC95% = 8,2 - não atingido). O EA mais comum foi pneumonia (5 casos). Apenas 1 paciente (1,1%) teve um evento semelhante à pneumonite e 2 pacientes (2,3%) prolongaram o intervalo QTc. **Conclusão:** Em um cenário do mundo real, o osimertinibe constitui uma opção terapêutica segura e eficaz para pacientes brasileiros com CPNPC T790M-positivo avançado em progressão após EGFR-TKI prévia, incluindo aqueles com metástase no sistema nervoso central. Nossos resultados apoiam observações anteriores e acrescentam informações valiosas sobre a eficácia de osimertinibe em pacientes brasileiros.

**Descritores:** Neoplasias pulmonares; Genes erbB-1; Terapia direcionada molecular.

## INTRODUCTION

Lung cancer is the most common cancer in the world, with projected 2 million incident cases and 1.7 million deaths in 2016, according to the 2016 Global Burden of Disease Analysis (Tracheal, Bronchus, and Lung Cancer Group).<sup>[1]</sup> Non-small cell lung cancer (NSCLC) represents 80-90% of all cases in Brazilian populations, with two thirds of patients presenting locally advanced (stage III) or metastatic (stage IV) disease at diagnosis.<sup>[2]</sup> Patients presenting advanced NSCLC have a median overall survival (OS) of 10 to 12 months.<sup>[3]</sup>

The development of epidermal growth factor receptor (EGFR)-targeted tyrosine kinase inhibitors (TKI) resulted in significant clinical improvements for patients with EGFR mutations (EGFRm) – 10-15% of NSCLC patients in Western high-income countries and 30-40% in Asia.<sup>[4,5]</sup> The National Comprehensive Cancer Network (NCCN) and the European Society of Medical Oncology (ESMO) currently recommend first-line treatment with an EGFR-TKI for patients with documented EGFRm.<sup>[6,7]</sup>

Even with high response rates among those patients (50-80% versus less than 30% with conventional chemotherapy), they eventually present acquired resistance with disease progression after 9-13 months.<sup>[8,9]</sup> The major explanation for the development of resistance among those patients is the emergence of T790M.<sup>[10,11]</sup> The standard of care for resistant patients has not been well established in clinical guidelines so far, thus therapeutic options for this selected subgroup are generally limited to chemotherapy or clinical trials.<sup>[12]</sup> Switching between EGFR-TKI confers response rates as low as 10% with median progression-free survival (PFS) of 1.7-6.2 months.<sup>[13,14]</sup>

Osimertinib is an oral, potent, selective, irreversible inhibitor of both EGFR-TKI sensitizing and resistance mutations in NSCLC with a significant selectivity margin over wild-type EGFR. The efficacy and safety of osimertinib in advanced NSCLC patients previously treated with first-generation EGFR-TKI and with acquired EGFR T790M mutation was assessed in several clinical trials. In the phase I, dose-escalation

AURA trial,<sup>[15]</sup> an open-label, multicenter study, 127 out of 253 enrolled patients had confirmed T790M mutation and presented an objective response rate (ORR) of 61% and progression-free survival (PFS) of 9.6 months. In the subsequent phase II

trial (AURA2),<sup>[16]</sup> 210 NSCLC patients with confirmed T790M after progression with EGFR-TKIs received osimertinib 80mg, once daily, and an ORR of 71% was observed. The phase III clinical trial AURA3<sup>[17]</sup> confirmed the efficacy of osimertinib in this subgroup

of patients, after randomly assigned 419 patients to either osimertinib 80mg once daily or intravenous pemetrexed plus carboplatin or cisplatin for up to six cycles. The median PFS was significantly better with osimertinib versus chemotherapy (10.1 vs. 4.4 months, hazard ratio (HR)=0.30;  $p<0.001$ ], as well as the ORR (71% versus 31%). Additionally, in patients with central nervous system disease (n=144), the PFS was also improved (8.5 vs. 4.2 months, HR=0.32).

Thus, the primary objective of this study was to describe the efficacy and safety of single agent osimertinib in a real world setting in adult Brazilian patients with advanced or metastatic EGFR T790M mutation-positive NSCLC, who have progressed after prior EGFR-tyrosine kinase inhibitor (TKI) therapy.

## METHODS

### Study design

ASTRIS(NCT02474355)<sup>[18,19]</sup> is a phase IV, multinational, open-label, real world trial of osimertinib for advanced T790M mutation NSCLC who received prior EGFR-TKI. Patients received osimertinib administered orally as one 80mg tablet once a day, as long as they continued to show clinical benefit, as judged by the investigator, and in the absence of discontinuation criteria. The projected maximum follow-up period was 18 months after the last patient was enrolled in the country. Patients withdrawing from the study treatment prior to the end of the study were followed up as part of this study until death or loss to follow-up, if they agreed with it.

### Population

Adult patients (aged 18 years or older) with locally advanced (stage IIIB) or metastatic (stage IV) NSCLC with confirmed T790M mutation on tissue or plasma samples, who have received prior EGFR-TKI therapy were enrolled. Other inclusion criteria included:

i) Provision of signed and dated written informed consent; ii) World Health Organization (WHO) performance status 0-2; iii) Adequate bone marrow reserve and organ function as demonstrated by complete blood count, biochemistry in blood and urine at baseline; iv) ECG recording at baseline showing absence of any cardiac abnormality; v) Female patients of childbearing potential must be using adequate contraceptive measures, must not be breast feeding, and must have a negative pregnancy test prior to start of dosing; or they must have evidence of non-childbearing potential; vi) Male patients must be willing to use barrier contraception.

Exclusion criteria were:

i) Previous (within 6 months) or current treatment with osimertinib; ii) Patients currently receiving any treatment known to be potent inhibitors or inducers of cytochrome P450; iii) Any evidence of severe or uncontrolled systemic diseases, including uncontrolled hypertension, active bleeding diatheses, active infection including hepatitis B, hepatitis C and

human immunodeficiency virus, or significantly impaired bone marrow reserve or organ function, including hepatic and renal impairment, which in the investigator's opinion would significantly alter the risk/benefit balance; iv) Patient with symptomatic central nervous system (CNS) metastases who is neurologically unstable or has required increasing doses of steroids to manage CNS symptoms within the 2 weeks prior to start study drug administration; v) Past medical history of Interstitial Lung Disease (ILD), drug-induced ILD, radiation pneumonitis requiring steroid treatment, or any evidence of clinically active ILD; vi) Any of the following cardiac criteria: mean resting corrected QT interval (QTcF) > 470ms using Fredericia's formula, any clinically important abnormalities in rhythm, conduction or morphology of resting ECG, any factors that increase the risk of QTc prolongation or risk of arrhythmic events; vii) Any unresolved toxicity from prior therapy CTCAE > grade 3 at the time of starting treatment; viii) History of hypersensitivity to excipients of osimertinib or to drugs with a similar chemical structure or class.

All patients signed informed consent form before any trial procedure and the trial was approved by local Ethics Committee of each participating site.

### Variables and outcomes

Data collected included demographics, variables related to eligibility criteria (medical history, past and current disease characteristics, and tumor EGFR mutational status), osimertinib exposure (starting dose, dose adjustments and discontinuations), investigator-reported efficacy (including tumor response and disease progression), overall survival (OS), and safety (including serious adverse events [SAEs] and adverse events leading to dose modification).

### Statistical analysis

Data presented here refers to the subset of Brazilian patients (n=88) enrolled to participate in the ASTRIS trial. The cutoff was October 20<sup>th</sup>, 2017. Data was examined for the full Brazilian analysis set. Descriptive statistics was used for all variables, as appropriate. Continuous variables were summarized by the number of observations, mean, standard deviation, median, minimum and maximum. Categorical variables were summarized by frequency counts and percentages for each category. Progression-free survival (PFS) was summarized using Kaplan-Meier estimates of the median time progression and quartiles together with their 95% confidence intervals.

## RESULTS

### Patient population

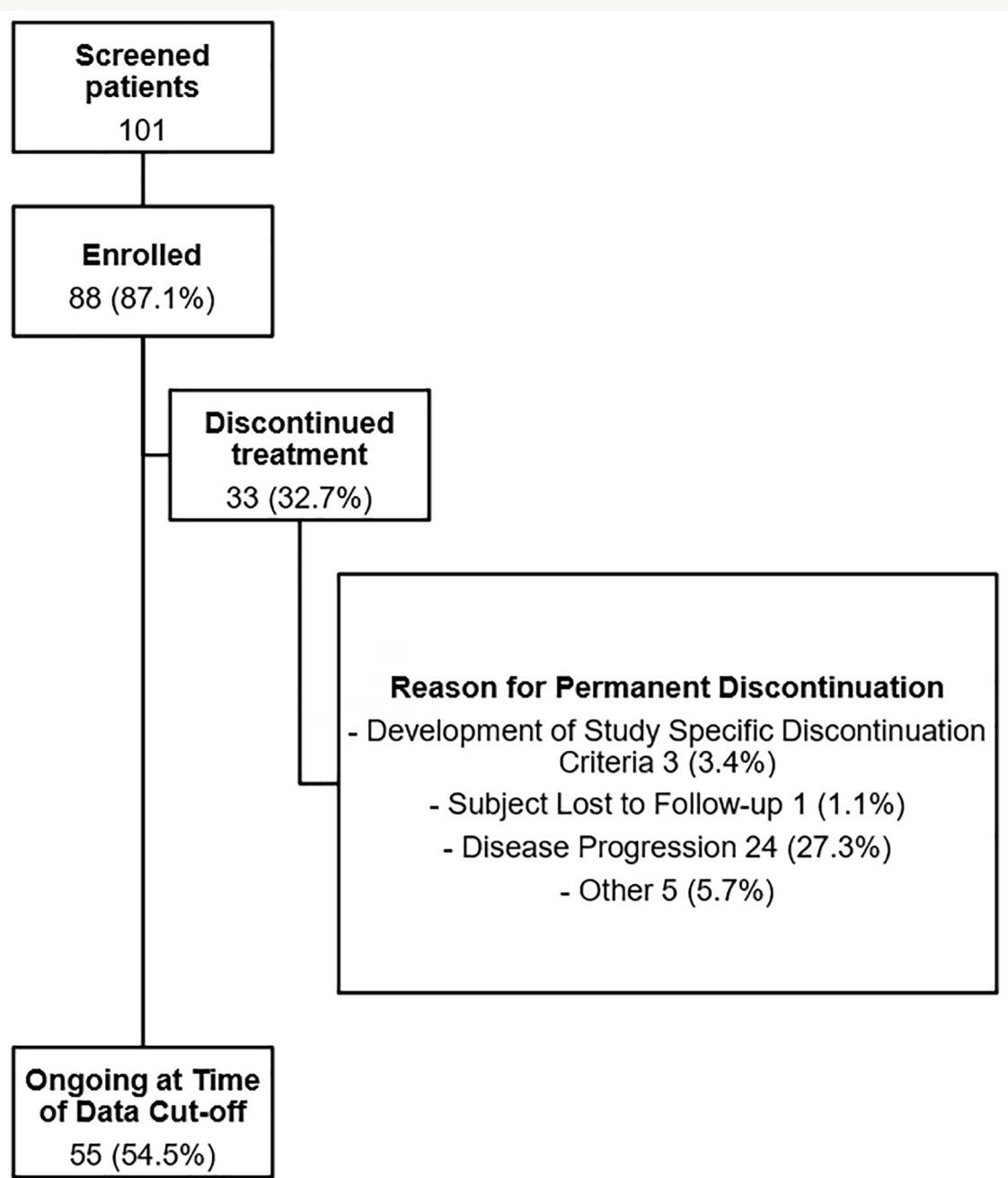
A total of 88 patients from 14 different Brazilian sites were included in this analysis. Patients' disposition is depicted in Figure 1 and their main clinical and demographic characteristics are described in Table 1. Mean age was 63.6 years (SD 11.6), most patients

were female (65.9%) with a higher frequency of white self-reported ethnicity (78.4%). According to WHO *performance status*, most patients (87.5%) presented normal or restricted activity (PS of 0-1). The most frequent EGFR mutation was exon 19 deletion (62.5%) followed by L858R (27.3%). Known brain or leptomeningeal metastases were observed in 28.4% of the sample. The most frequent EGFR-TKI patients had received was erlotinib (61.4%), and 56% had received a previous chemotherapy. The median time between initial diagnostic and enrollment was 23.7 months (range from 8 to 145 months).

### Efficacy endpoints

The median duration of total exposure to osimertinib in the sample was 9.3 months (range 0-16). **Figure 2** illustrates the percentage of patients still on treatment in each time point.

Overall response rate (ORR) was assessed for the full Brazilian analysis set (n=88) and also for the subgroup of patients with brain and/or leptomeningeal metastases (n=25). As shown in **Table 2**, ORR was 58.2% and 59.1%, respectively. Most patients had a best overall response, as assessed by the investigator, recorded as "Responding" (52.3% and 52.0%).



**Figure 1.** Patient disposition flowchart.

**Table 1.** Baseline characteristics of patients.

Characteristic	N (%)
Sex – female	58 (65.9)
Age, years (mean [standard deviation])	63.6 [11.6]
Self-reported ethnicity	
White	69 (78.4)
Black	7 (8.0)
Asian	7 (8.0)
Other	5 (5.7)
WHO Performance Status	
0 - Normal Activity	22 (25.0)
1 - Restricted Activity	55 (62.5)
2 - In Bed Less Than or Equal to 50% of the Time	11 (12.5)
3 - In Bed More Than 50% of the Time	0 (0.0)
4 - 100% Bedridden	0 (0.0)
ECG results	
Normal	56 (63.6)
Abnormal	32 (36.4)
Clinically relevant (n=32)	1 (3.1)
Receptor status (positivity) <sup>†</sup>	
T790M	88 (100.0)
G719X	2 (2.3)
EXON 19 Deletion	55 (62.5)
S768I	3 (3.4)
EXON 20 Insertion	2 (2.3)
L858R	24 (27.3)
Known Brain and Leptomeningeal Metastasis	
Any lesion present	25 (31.8)
Brain Metastases Only (n=25)	22 (25.0)
Leptomeningeal Disease Only (n=25)	1 (1.1)
Both Brain and Leptomeningeal (n=25)	2 (2.3)
Previous EGFR-TKI	
Erlotinib	54 (61.4)
Gefitinib	42 (47.7)
Afatinib	3 (3.4)

<sup>†</sup>Type of sample tested: Tissue (35.2%), Blood (63.6%), and FNA/Cytology (1.1%); WHO: World Health Organization.

Patients had a median PFS of 9.4 months (95% Confidence Interval [CI] 8.2-not reached) (Figure 3). Progression events occurred in 26 patients (29.5%), while 6 (6.8%) patients died due to progression and 4 (4.5%) due to other reasons.

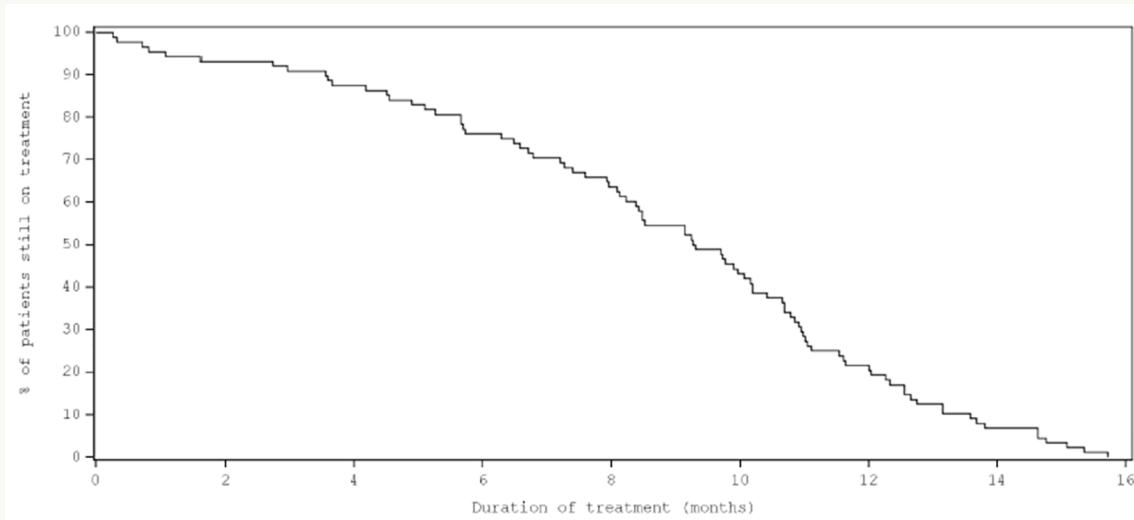
### Safety endpoints

Osimertinib was generally well tolerated and only 5.7% of patients discontinued treatment due to adverse events (AEs) (Table 3). The most common AE was pneumonia, and the five cases in the sample were classified as serious adverse events by the

investigators. In terms of AEs of special interest, only 1 patient (1.1%) had a pneumonitis-like event and 2 patients (2.3%) had a prolongation of the QTc interval. Three patients died due to AEs in the sample, for the following causes: pneumonia, pulmonary sepsis and respiratory failure (one each).

### DISCUSSION

The PFS, OR and tolerability profile results confirm the efficacy of osimertinib in patients with advanced T790M-positive NSCLC who progressed after receiving an EGFR-TKI. These findings are particularly relevant



**Figure 2.** Exposure plot (percentage of patients still on treatment against time).

**Table 2.** Response endpoints.

Endpoint	N (%)
<b>Total Sample (n=88)</b>	
Overall response rate (n=79) (% [95% CI])	58.2 [46.6; 69.2]
Best overall response (n=88)	
Responding	46 (52.3)
Stable Disease	25 (28.4)
Progressing	8 (9.1)
Unknown †	9 (10.2)
<b>Brain/Leptomeningeal Metastases Present (n=25)</b>	
Overall response rate (n=22) (% [95% CI])	59.1 [36.4; 79.3]
Best overall response (n=25)	
Responding	13 (52.0)
Stable Disease	5 (20.0)
Progressing	4 (16.0)
Unknown †	3 (12.0)

†Patients with no assessment were recorded as ‘Unknown’.

considering the real-world approach, indicating that even outside the context of clinical trial patients achieve clinically sound benefits. Estimated median PFS of 9.4 months are in line with those observed in osimertinib AURA clinical trials program<sup>[15-17]</sup> and superior to the one observed for patients switching to another EGFR-targeting therapy after an initial failure (1.7-6.2 months).<sup>[13,14]</sup>

ORR was also similar to the ones reported in clinical trials, for all patients and particularly for those with CNS metastasis. CNS metastasis are life-threatening complications of NSCLC present in approximately 20-40% of patients.<sup>[20]</sup> This incidence can reach 50-60% in patients with tumors expressing driver mutations, especially EGFR mutation or anaplastic lymphoma kinase (ALK) rearrangement.<sup>[21]</sup> The third-generation

EGFR inhibitor osimertinib had substantial clinical activity against brain metastases as shown in phase III trials<sup>[22]</sup> and corroborated here.

Osimertinib demonstrated an adequate toxicity profile with 5.7% of discontinuations due to AEs. A total of 21.6% of all patients had dose modifications (interruption or reduction) due to AE, similar to the rates reported by AURA2<sup>[16]</sup> and AURA3<sup>[17]</sup> trials (21.0% and 27.0%, respectively).

Limitations of our study can be related to the reduced sample size and the single-arm design. Despite these limitations, this is the first Brazilian study assessing efficacy and safety of osimertinib in this particular subgroup of lung cancer patients with acquired resistance to a previous line of EGFR-TKIs. These patients faced unmet needs with the previous

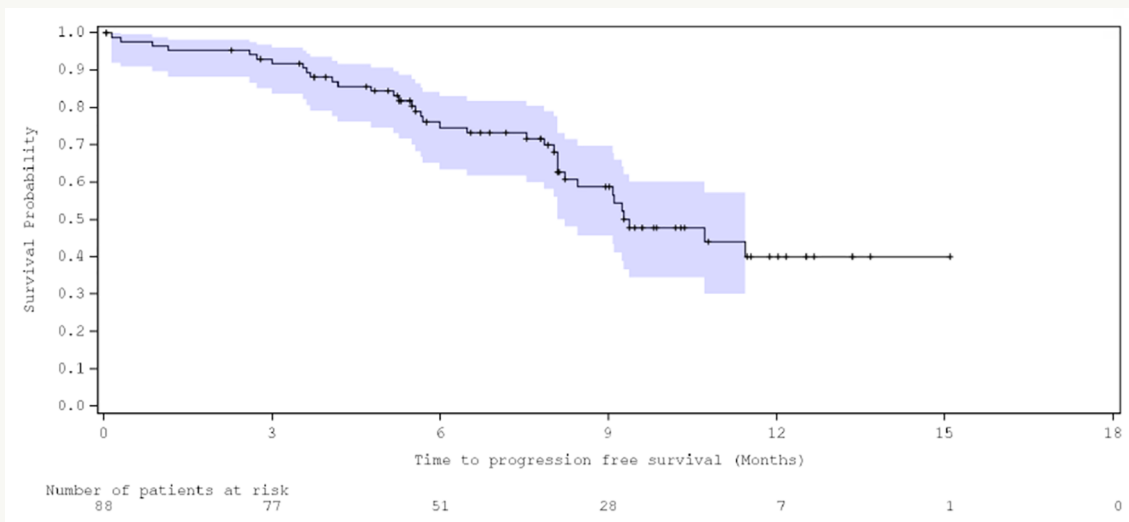


Figure 3. Kaplan Meier plot of time to progression free survival.

Table 3. Summary of adverse events results.

Endpoint	N (%)
<b>Number of patients with at least one:</b>	
Adverse Event	30 (34.1)
Adverse Event Leading to Dose Modification	14(15.9)
Adverse Event Leading to Discontinuation	5 (5.7)
Adverse Event of Special Interest	3 (3.4)
Serious Adverse Event	19(21.6)
<b>Adverse events in &gt; 2% of patients</b>	
Pneumonia	5 (5.7)
Diarrhea	3 (3.4)
Neutropenia	2 (2.3)
Thrombocytopenia	2 (2.3)
Blepharitis	2 (2.3)
Vomiting	2 (2.3)
Electrocardiogram QT prolonged	2 (2.3)
Deep vein thrombosis	2 (2.3)
<b>Death</b>	<b>3(3.4)</b>

available therapeutic options in the Brazilian healthcare system and local data may help guide prescribers and policy decision-makers to better address these needs. Phase IV real-world study also provides a more pragmatic overview of the treatment effects without the controlled environment and co-interventions typical of clinical trials, thus allowing care providers to understand how their patients can respond to therapy in real life.

More recently, osimertinib was tested as front-line therapy for advanced T790M- positive NSCLC in the FLAURA trial<sup>[22]</sup> with promising results. The median PFS was significantly longer as compared to standard EGFR-TKIs (18.9 months vs. 10.2 months; hazard

ratio [HR]=0.46; 95%CI = 0.37-0.57;  $p<0.001$ ), with similar ORR and longer duration of response (17.2 vs. 8.5 months). Thus, further research could elucidate the effectiveness of osimertinib as first-line option in the real-world local context. Also, aspects markedly relevant to lung cancer patients and the healthcare system, such as quality of life and cost-effectiveness, need to be addressed in future studies in the Brazilian setting.

## CONCLUSION

In a real-world setting, osimertinib constitutes a safe and effective therapeutic option for Brazilian patients with advanced T790M-positive NSCLC after

progression on a prior EGFR-TKI, including those patients with central nervous system metastasis. Our findings support previous observations and add valuable information regarding osimertinib effectiveness in Brazilian patients.

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