At the time of acquired resistance to the first generation epidermal growth factor receptor (EGFR) tyrosine kinase inhibitors (TKIs), gefitinib, erlotinib or afatinib (1), 50–60% of EGFR mutant non-small cell lung cancer (NSCLC) patients have developed the secondary gatekeeper T790M point mutation on exon 20 of the EGFR gene (2,3).

Osimertinib (AZD9291) is a third-generation EGFR TKI that targets EGFR mutant (including T790M positive) tumors (4). It was initially evaluated in EGFR mutant NSCLC patients who had disease progression after previous treatment with an EGFR TKI, in the phase I part of the phase I/II AURA study (5). A response rate of 51% was achieved for all patients treated across all dose levels. Among the 222 patients of the expansion cohorts, the response rate to osimertinib was 61% for EGFR T790M-positive patients while the EGFR T790M-negative patients were not able to achieve a response rate greater than 21% (Table 1). No dose limiting toxicities were observed at any dose level and, based on tumor growth inhibition, the dose of 80 mg once daily was selected for being further evaluated for the treatment of EGFR T790M-positive NSCLC patients (5). On November 13th 2015, and on February 3rd 2016, FDA and EMA approved osimertinib 80 mg once daily for the treatment of EGFR T790M-positive NSCLC patients, respectively, based on the data from two phase II studies (AURA extension and AURA2) and the AURA phase I expansion study. Sixty-three EGFR T790M-positive, dose expansion cohort patients receiving 80 mg of osimertinib once daily in the phase I part of the AURA study demonstrated an objective response rate (ORR) of 71% [95% confidence interval (CI), 57–82] and a median progression-free survival (PFS) of 9.7 months (95% CI, 8.3–13.6). In a pre-planned pooled analysis of the AURA extension phase II study and the AURA2 phase II study with a total of 411 EGFR T790M-positive patients, ORR was 66% and median PFS was 11.0 months (95% CI, 9.6–12.4) (6).

Goss et al., published in *Lancet Oncology* the final results of the phase II, open-label, single-arm AURA2 study, which assessed the efficacy and safety of osimertinib in patients with EGFR T790M-positive NSCLC, who had progressed after previous therapy with EGFR TKIs. In less than 6 months, more than 472 patients were screened, 210 EGFR T790M-positive patients were treated with osimertinib and 199 patients were evaluable for response analysis in the AURA II study (7). The FDA approved Cobas EGFR mutation test v2 was used for the central confirmation of the EGFR T790M mutation. An ORR of 70% and a disease control rate of 92% were achieved, as evaluated by blinded independent central review. Six...
(3%) patients achieved complete response with osimertinib treatment and 134 (67%) achieved partial response. The median duration of response was 11.4 months (95% CI, 9.0–not calculable). There was a high concordance between the responses obtained by investigator assessment and by the blinded independent central review. The median PFS was 9.9 months (95% CI, 8.5–12.3) (7). The treatment was well tolerated with the most common treatment related grade 3–4 adverse events being prolonged electrocardiogram QT, decreased neutrophil count, and thrombocytopenia. Interstitial lung disease occurred in 2% of the patients.

Only 2 months after the publication of the AURA2 study, the results of the phase III AURA3 clinical trial became available (8). In this study, 419 EGFR T790M-positive NSCLC patients who had progressed during first-line were assigned in a 2:1 ratio to receive osimertinib or platinum-pemetrexed chemotherapy. Median PFS was significantly longer with osimertinib compared with chemotherapy [10.1 vs. 4.4 months; hazard ratio (HR) 0.30; 95% CI, 0.23–0.41; P<0.001] (8). Osimertinib-treated patients achieved an ORR of 71% (95% CI, 65–76) in comparison to an ORR of 31% (95% CI, 24–40) for those who received chemotherapy (odds ratio 5.39; 95% CI, 3.47–8.48; P<0.001) (8).

It was previously demonstrated that patients with the T790M mutation detected by plasma ctDNA respond equally to osimertinib as those whose mutation is detected in a tumor tissue biopsy (9-11). Similar findings were reported in the AURA3 trial (8). On September 29th, 2016, the FDA approved an expansion of the Cobas EGFR blood mutation test v2 to include testing of the T790M mutation in order to confirm the presence of the EGFR T790M mutation and qualify patients for treatment with osimertinib. Due to the relatively high false negative rates with plasma T790M testing, it is highly recommended that patients with a negative liquid biopsy for the presence of the T790M to be reevaluated for the feasibility of a tissue biopsy (10).

More than 30% of EGFR mutant NSCLC patients, whose disease progressed during or after first line EGFR TKI, have brain metastases (12). In comparison with other EGFR TKIs, including gefitinib, rociletinib or afatinib, osimertinib has demonstrated greater blood-brain barrier penetration in preclinical models (13). In the AURA2 trial, patients with brain metastases obtained an ORR of 69% (95% CI, 58–79) and a median PFS of 9.2 months (95% CI, 7.7–11.1) (7). In the AURA3 study, among the 144 patients who had central nervous system (CNS) metastases, median PFS was 8.5 months (95% CI, 6.8–12.3) for those treated with osimertinib and 4.2 months (95% CI, 4.1–5.6) for those who received chemotherapy (HR 0.32; 95% CI, 0.21–0.49) (8). Interestingly, due to the high risk of development CNS metastases in EGFR mutant NSCLC, an agent with high blood-brain barrier penetration, AZD3759, has been recently developed and is currently being evaluated in a phase I clinical trial (14). Unavoidably, as happens with the first generation EGFR TKIs, osimertinib treated patients develop resistance to treatment after less than one year (8). An additional EGFR mutation, the C797S, can cause resistance to third generation EGFR TKIs and its allelic

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NR, not reached; DoR, duration of response; Ref., reference; vs. versus; HR, hazard ratio; OS, overall survival.
context can define sensitivity to subsequent treatments (15,16) (Figure 1).

Osimertinib is now being evaluated in the first line setting of EGFR mutant NSCLC patients. According to the results of two (80 and 160 mg) phase I expansion cohorts of the AURA study including 6 EGFR mutant patients, an overall median PFS of 19.3 months (95% CI, 13.7–not reached) and an ORR of 77% (95% CI, 64–87) were obtained (17). The phase III FLAURA study (NCT02296125) compares osimertinib with gefitinib or erlotinib in treatment-naïve EGFR mutant NSCLC patients. PFS in patients with tumors harboring T790M is a key secondary objective of the study (18). The coexistence of the pretreatment T790M mutation has been under appreciated, in spite of accumulative evidence that it is present in a frequency of 35–60%, using different detection methods (19-21). In the Spanish Lung Cancer Group (SLCG) and the European Thoracic Oncology Platform (ETOP) BELIEF trial, pretreatment T790M mutations were centrally identified in 34% of the patients who reached a median PFS of 16 months (95% CI, 13.1–not reached) with the combination of erlotinib plus bevacizumab (21). We have recently started the AZENT study (NCT02841579), an investigator initiated study that explores the safety and efficacy of osimertinib as first line therapy for patients with metastatic EGFR mutant NSCLC and concomitant pretreatment T790M mutation (Figure 1).

Without any doubt, osimertinib has made a breakthrough

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**Figure 1** Therapeutic approach for advanced NSCLC with EGFR mutations. NSCLC, non-small cell lung cancer; EGFR, epidermal growth factor receptor; TKIs, tyrosine kinase inhibitors.
in lung cancer therapy. Other third generation EGFR TKIs are in clinical development, including olmutinib (Hanmi Pharmaceutical Company), EGF816 (Novartis Pharmaceuticals), naquotinib (Astellas Pharma Global Development), PF06747775 (Pfizer) and avitinib (Hangzhou ACEA Pharmaceutical Research) (22,23). Still, there are several issues to be resolved in the treatment of EGFR mutant patients. One important issue is the discovery of the best therapeutic approach, besides chemotherapy, for those patients who are not EGFR T790M-positive at the time of progression to first and second generation EGFR TKIs. A second issue is that, even if osimertinib is an efficient treatment for T790M driven acquired resistance to initial EGFR inhibition, still the number of complete responses reported in the phase II and III clinical trials is very small, indicating that we are far from the cure of this disease. We have shown that co-targeting STAT3 and Src with EGFR can more efficiently abrogate tumor growth than EGFR inhibition alone (24). Resistance to first, second and third EGFR TKIs are heterogeneous and complex, evolving dynamically under the pressure of each generation’s inhibitor and is a challenge for the development of novel targeted combinations.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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