Curative-Intent Treatment with Durvalumab in Early-Stage Cancers

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ABSTRACT

The introduction of immunotherapy has fundamentally transformed the treatment landscape in cancer, providing long-term survival benefit for patients with advanced disease across multiple tumor types, including non-small cell lung cancer (NSCLC). In the placebo-controlled phase 3 PACIFIC trial, the PD-L1 inhibitor durvalumab demonstrated significant improvements in progression-free survival and overall survival in patients with unresectable, stage III NSCLC who had not progressed after platinum-based chemoradiotherapy (CRT). These findings have led to the widespread acceptance of the ‘PACIFIC regimen’ (durvalumab after CRT) as the standard of care in this setting. Moreover, the PACIFIC trial is the first study to demonstrate a proven survival advantage with an immunotherapy in a curative-intent setting, thereby providing a strong rationale for further investigation of durvalumab in early-stage cancers. Herein, we describe the extensive clinical development program for durvalumab across multiple tumor types in curative-intent settings, outlining the scientific rationale(s) for its use and highlighting the innovative research (e.g., personalized cancer monitoring) advanced by these trials.

Keywords: Bladder cancer; Cervical cancer; Curative intent; Durvalumab; Early-stage cancer; Esophageal cancer; Gastric cancer; Hepatocellular carcinoma; Lung cancer; PACIFIC
Immunotherapy has fundamentally transformed the treatment landscape in cancer.

In the placebo-controlled phase 3 PACIFIC trial, the programmed cell death-ligand 1 inhibitor durvalumab demonstrated significant improvements in survival in patients with unresectable, stage III non-small lung cancer.

This has led to the widespread acceptance of the ‘PACIFIC regimen’ (durvalumab after chemoradiotherapy) as the standard of care in this setting.

As durvalumab is the first immunotherapy with a proven survival advantage in a curative-intent setting, there is a strong rationale for its further investigation in early-stage cancers.

An extensive clinical development program, as described herein, is investigating durvalumab across multiple tumor types in curative-intent settings.

DIGITAL FEATURES

This article is published with digital features, including a summary slide, to facilitate understanding of the article. To view digital features for this article, go to https://doi.org/10.6084/m9.figshare.14035628.

INTRODUCTION

The introduction of immunotherapy has fundamentally transformed the treatment landscape in cancer, providing long-term survival benefit for patients with metastatic disease across a range of tumor types. In particular, immune checkpoint blockade (ICB) of the programmed cell death-ligand 1 (PD-L1) and PD-1 pathway, and of cytotoxic T-lymphocyte-associated protein 4 (CTLA-4), has shown survival benefit in many types of advanced cancer, including non-small cell lung cancer (NSCLC) [1, 2]. The success of these agents has led to several regulatory approvals and, in advanced NSCLC, has recently led to a shift in the use of ICB therapy from the second- to first-line setting as the preferred treatment option (either alone or in combination with chemotherapy [CT]) for patients without genomic-driven tumors [3].

Durvalumab is a selective, high-affinity, human IgG1 monoclonal antibody that targets PD-L1 and occludes its binding to PD-1 and CD80, allowing T cells to recognize and kill tumor cells [4–6]. In an early phase 1/2 study of patients with a range of advanced solid tumors, including NSCLC [7, 8], durvalumab demonstrated encouraging antitumor activity and, based on this study, was granted accelerated approval in the US for patients with locally advanced or metastatic urothelial carcinoma after failure of platinum-based CT [9, 10]. In the subsequent phase 3 PACIFIC trial, durvalumab versus placebo demonstrated significant improvements in the primary end points of progression-free survival (PFS) and overall survival (OS) in patients with unresectable, stage III NSCLC who had not progressed after platinum-based chemoradiotherapy (CRT) [11–14].

These findings have led to global regulatory approvals and the widespread acceptance of the ‘PACIFIC regimen’ (durvalumab after CRT) as the standard of care in this setting. Moreover, the PACIFIC trial is the first study to demonstrate a proven survival advantage with an immunotherapy in a curative-intent setting, thereby providing a strong rationale for further investigation of durvalumab in early-stage cancers. Two other checkpoint inhibitors have been approved by global regulatory authorities for use in the curative-intent setting: nivolumab and pembrolizumab as adjuvant therapy in patients with resected melanoma [15, 16] and pembrolizumab as treatment of patients with bacillus Calmette-Guerin (BCG)-unresponsive, high-risk non-muscle invasive bladder cancer (HR-NMIBC) with carcinoma in situ (CIS), who are ineligible for or elect not to undergo
However, the focus of this review is further evaluation of durvalumab in early-stage cancers.

Herein, we describe the extensive clinical development program for durvalumab across multiple tumor types in curative-intent settings, including several trials in early-stage lung cancer, as well as in other cancers (e.g., bladder cancer, hepatocellular carcinoma [HCC], and gastric and esophageal cancer). In addition, we will outline the scientific rationale(s) for use of durvalumab in these settings and highlight the innovative research in personalized cancer monitoring that will be advanced by these studies. This program will not only shed light on the potential use of durvalumab to reduce recurrence rates and improve survival outcomes, but also increase our understanding of patient selection and optimal use of immunotherapy in this setting where we have the best chance to cure patients.

This article is based on previously conducted studies and does not contain any new studies with human participants or animals performed by any of the authors.

**EARLY-STAGE CANCER: DIAGNOSIS, PROGNOSIS AND RATIONALE FOR CURATIVE-INTENT IMMUNOTHERAPY**

A patient’s treatment goals and prognosis can vary significantly, depending on his/her disease stage at diagnosis. Localized, early-stage disease can be treated with curative intent, whereas the goal for metastatic disease is to extend survival and, in some cases, is palliative only [17–19]. For patients diagnosed with early-stage cancer, surgery remains the mainstay of curative treatment for most common tumor types. However, recurrence after curative surgery remains a major concern, with or without the use of neoadjuvant or adjuvant treatment [20–26]. It is hypothesized that, based on the more amenable biology of early-stage cancer, there may be an increased likelihood of cure with immunotherapy.

**Diagnosis and Treatment Outcomes in Early-Stage Cancer**

Across all tumor types, a common goal is screening and early detection in order to cure patients by revealing the malignancy, or precursor lesion, at an early prior stage, when treatment is most effective. This is reflected in survival rates, which can vary significantly depending on the stage at diagnosis. For example, among patients with bladder cancer, the 5-year survival rates range from as high as 96% for patients with in situ disease to 36% for patients with regional disease and as low as 5% for patients with metastatic disease [27]. Similarly, among NSCLC patients, 5-year survival rates can reach up to 92% for patients with stage IA1 diagnosis to as low as 10% for patients with stage IV diagnosis (i.e., metastatic disease), with significant variation by, and within, the stages in between [28].

However, the proportion of patients diagnosed with early-stage cancer varies by tumor type and depends on several factors, including ease of detection, the stage at which patients become symptomatic (e.g., early-stage disease may remain asymptomatic and undetected) and whether reliable screening programs are in place [20, 27, 29, 30]. For example, only 30% of patients with NSCLC present with resectable disease at diagnosis [20, 29, 31, 32]; most patients are diagnosed at a later stage, associated with an unfavorable prognosis, as reflected in the overall 5-year survival rate of 18% [33]. In contrast, for example, the overall 5-year survival rate for patients with bladder cancer is 77%, since >90% of patients are diagnosed with resectable disease [19, 27].

Screening has been shown to reduce mortality for several types of cancer. For example, the National Lung Screening Trial in the US and the Dutch-Belgian NELSON lung cancer screening trial have shown a reduction in lung cancer-related deaths of 20–25% using low-dose computed tomography compared with control groups [30]. Further efforts to increase lung screening and earlier diagnosis are being championed by the Lung Ambition Alliance [34], of which AstraZeneca is a founding partner.
Finally, a major area of research in improving earlier diagnosis (i.e., ‘stage shifting’) is utilization of blood-based circulating tumor cell-free DNA (cfDNA) to simultaneously detect and localize multiple cancer types. In a recently published analysis from the Circulating Cell-free Genome Atlas study, based on 6689 participants (2482 with previously untreated cancer and 4207 without cancer), target methylation analysis of plasma cfDNA was used to simultaneously detect > 50 cancer types, across all stages, and localize the tissue of origin with > 90% accuracy [35].

Rationale for Curative-Intent Immunotherapy

First, there are several theoretical advantages for use of immunotherapy, early in the course of tumor development, including (1) lower tumor volume, (2) reduced tumor heterogeneity [36–40] and (3) largely intact host fitness, not yet impaired by multiple rounds of CT.

There is also a strong scientific rationale for the use of immunotherapy in the curative-intent setting, based on potential synergistic effects when used as part of a multi-model treatment approach. For example, it has been suspected for some time that tumor resection may somewhat paradoxically promote tumor growth. Recent findings suggest that surgery may suppress antitumor immunity and induce the development of new postoperative metastases as well as promote the growth of micrometastases and residual disease [41–43]. However, in a preclinical model for surgical stress, treatment with a PD-1 inhibitor boosted CD8+ T cell numbers and function [44]. In addition, preclinical evidence suggests that CRT causes upregulation of PD-L1 expression and induction of immunogenic cell death as well as activation of a post-CRT inflammatory response [45–50].

Finally, based on several studies of postoperative adjuvant CT in NSCLC [51–53], which have demonstrated absolute risk reductions in 5-year survival rates of 5–15%, we can extrapolate that immunotherapy in the early-stage setting may confer similar benefit, given the improvement already demonstrated by immunotherapy in the treatment of some metastatic cancers. However, because such benefit is reliant upon immune cell infiltration of the tumor, it cannot be guaranteed that this will be replicated in all early-stage cancers (e.g., if infiltration is deficient in so-called ‘cold tumors’) [54].

That said, all of these factors could serve to improve the likelihood of a cure, using a treatment capable of harnessing the body’s immune mechanisms.

THE PACIFIC STUDY: EVIDENCE OF DURVALUMAB’S POTENTIAL AS CURATIVE-INTENT TREATMENT

Historically, among stage III NSCLC patients who are considered unresectable, platinum-based doublet CT, administered with definitive-dose RT (concurrent CRT), was the standard of care. The treatment goal was with curative intent; however, outcomes were poor because most patients had disease progression after CRT, with approximately 15–32% of patients alive at 5 years [55, 56]. Prior to the PACIFIC trial, several studies had investigated systemic therapies with curative intent after patients had disease control with CRT. However, none had proven effective [57–61].

The phase 3 PACIFIC trial was designed to exploit the potential synergy between immunotherapy and CRT (Fig. 1). In PACIFIC, patients with unresectable, stage III NSCLC (and any tumor PD-L1 status) who had not progressed after platinum-based CRT were randomized in a 2:1 ratio to receive durvalumab 10 mg/kg or matching placebo given intravenously every 2 weeks for up to 12 months [11–14]. Durvalumab versus placebo significantly improved the primary end points of PFS (stratified HR 0.52, 95% CI 0.42–0.65; \( P < 0.001 \); median 16.8 vs. 5.6 months; data cutoff [DCO], 13 February 2017) and OS (stratified HR 0.68, 95% CI 0.53–0.87; \( P = 0.0025 \); median not reached vs. 28.7 months; DCO, 22 March 2018) [11, 12] and was observed across most patient subgroups. In addition, durvalumab exhibited a manageable safety profile and
had no detrimental effect on patient-reported outcomes (PROs) [63], which was notable since the previous standard of care in this setting was observation alone with no presumed detriment to PROs.

In addition, updated survival analyses at approximately 4 years after the last patient was randomized (median follow-up, 34.2 months; DCO, 20 March 2020) demonstrated consistent and durable benefit with durvalumab [14]. Updated OS (stratified HR, 0.71; 95% CI 0.57–0.88) and PFS (stratified HR, 0.55; 95% CI 0.44–0.67) remained consistent with the primary analyses, and median OS for durvalumab was reached, 47.5 months (versus 29.1 months for placebo). The estimated 4-year OS rates were 49.6% for durvalumab versus 36.3% for placebo, and the 4-year PFS rates were 35.3% versus 19.5%, respectively. With approximately 50% of patients alive at 4 years and approximately 35% progression-free, despite completing their last treatment, 3+ years earlier, these findings have further established the PACIFIC regimen as the standard of care in this setting.

The PACIFIC trial, and subsequent worldwide approvals of the PACIFIC regimen for patients in this setting, has resulted in a paradigm shift in the treatment landscape, setting the stage for the investigation and potential use of durvalumab as curative-intent treatment in other early-stage cancer contexts.

**DURVALUMAB CLINICAL DEVELOPMENT PROGRAM IN EARLY-STAGE LUNG CANCER**

The role of durvalumab treatment in early-stage lung cancer is undergoing extensive investigation within several ongoing and planned phase 2 and 3 trials as (1) neoadjuvant or adjuvant treatment of resectable NSCLC, (2) concurrent treatment with or following CRT in unresectable NSCLC and (3) treatment after CRT in limited stage SCLC (Table 1).

**Resectable Stage I–III NSCLC**

The ongoing phase 3 BR.31 study [68] (Table 1), led by the Canadian Cancer Trials Group, will be a key trial in the adjuvant setting, enrolling approximately 1360 patients with NSCLC, classified as stage IB (≥ 4 cm in the longest diameter), II or IIIA, to be randomized to either durvalumab or placebo; prior, postoperative platinum-based doublet CT, per standard of care, is allowed but not mandatory. The BR.31 study will investigate the importance of PD-L1 testing at time of diagnosis, assessing disease-
**Table 1** Summary of ongoing and planned key durvalumab trials in early-stage lung cancer

| Diagnosis and staging | Setting | Trial | Patient population | Estimated (or actual) enrollment | Design | Primary end points |
|-----------------------|---------|-------|--------------------|----------------------------------|--------|-------------------|
| Resectable Stage I–III NSCLC | Neoadjuvant | NeoCOAST [64, 65] (NCT03794544) | Stage I (≥ 2 cm) to IIIA | 80 | Phase 2, open-label, randomized, multicenter, multidrug platform trial |
| | | | | | Arm A: Durvalumab |
| | | | | | Arm B: Durvalumab + olaparib |
| | | | | | Arm C: Durvalumab + monalizumab |
| | | | | | Arm D: Durvalumab + danvatirsen |
| Neoadjuvant/adjuvant | AEGEAN [66, 67 Data on file] (NCT03800134) | Stage IIA to select (i.e., N2) Stage IIIb | 800 | Phase 3, double-blind, randomized, international trial |
| | | | | | Arm 1: Neoadjuvant platinum doublet CT + durvalumab, with adjuvant durvalumab |
| | | | | | Arm 2: Neoadjuvant platinum doublet CT + placebo, with adjuvant placebo |
| Adjuvant | BR.31 [68] (NCT02273375) Partnered (CCTG) | Stage IB (≥ 4 cm in the longest diameter) to IIA | 1360 | Phase 3, double-blind, randomized, international trial |
| | | | | | Arm 1: Durvalumab (after optional SOC adjuvant platinum doublet CT) |
| | | | | | Arm 2: Placebo (after optional SOC adjuvant platinum doublet CT) |
| | MERMAID-1 [69] (NCT04385368) | Stage II–III | 332 | Phase 3, double-blind, randomized, international trial |
| | | | | | Arm 1: Adjuvant SOC platinum-based CT + durvalumab, followed by durvalumab |
| | | | | | Arm 2: Adjuvant SOC platinum-based CT + placebo, followed by placebo |
| | MERMAID-2 [70] (NCT04642469) | Stage II–III | 284 | Phase 3, double-blind, randomized, international trial |
| | | | | | Arm 1: Adjuvant durvalumab after local SOC curative intent therapy |
| | | | | | Arm 2: Adjuvant placebo after local SOC curative intent therapy |
| Inoperable Stage I/II NSCLC | Concurrent with SBRT | PACIFIC-4 [71, 72] (NCT03833154) | Lymph-node negative | 706 | Phase 3, double-blind, randomized, international trial |
| | | | | | Arm 1: Definitive SOC SBRT + durvalumab, followed by durvalumab |
| | | | | | Arm 2: Definitive SOC SBRT + placebo, followed by placebo |
| Diagnosis and staging | Setting | Trial | Patient population | Estimated (or actual) enrollment | Design | Primary end points |
|------------------------|---------|-------|--------------------|----------------------------------|--------|-------------------|
| Unresectable Stage III NSCLC | Following CRT | PACIFIC [11–14] (NCT02125461) | No progression after cCRT | 713 | Phase 3, randomized, double-blind, placebo-controlled trial<br>Arm 1: Durvalumab<br>Arm 2: Placebo | PFS<br>OS |
| PACIFIC-5 [73, 74] (NCT03706690) | No progression after cCRT or sCRT | 360 | Phase 3, double-blind, randomized, international trial<br>Arm 1: Durvalumab<br>Arm 2: Placebo | PFS |
| PACIFIC-6 [75, 76] (NCT03693300) | No progression after sCRT | 120 | Phase 2, open-label, international study<br>Cohort 1 (WHO/ECOG PS 0–1; n ~ 100–120): durvalumab<br>Cohort 2 (WHO/ECOG PS 2; n ≤ 30): durvalumab | Grade 3/4 TRAE rate within 6 months |
| COAST [77, 78] (NCT03822351) | No progression after cCRT | 189 | Phase 2, open-label, randomized, multicenter, multikrug platform trial<br>Control arm: Durvalumab<br>Arm A: Durvalumab + oleclumab<br>Arm B: Durvalumab + monalizumab | ORR |
| Concurrent with CRT | PACIFIC-2 [79, 80] (NCT03519971) | No prior or current cancer Tx | 328 | Phase 3, double-blind, randomized, international trial<br>Arm 1: Durvalumab + SOC platinum-based cCRT, followed by durvalumab<sup>b</sup><br>Arm 2: Placebo + SOC platinum-based cCRT, followed by placebo<sup>c</sup> | PFS |
| Following RT | DUART [81] (NCT04249362) | CT-ineligible with no progression after RT | 150 | Phase 2 open-label, single-arm, multicenter, international study<br>Cohort A: Durvalumab (following standard RT)<br>Cohort B: Durvalumab (following palliative RT) | Grade 3/4 PRAE rate |
| Limited Stage SCLC | Following cCRT<sup>a</sup> | ADRIATIC [82, 83] (NCT03703297) | No progression after cCRT | 724 | Phase 3, double-blind, randomized, international trial<br>Arm 1: Durvalumab<br>Arm 2: Durvalumab + tremelimumab<br>Arm 3: Placebo | PFS<br>OS |

The current status of each trial (i.e., recruitment status and estimated primary completion date) can be obtained by accessing its study record on the ClinicalTrials.gov registry using its unique NCT number.

**CCTG Canadian Cancer Trials Group, cCRT (concurrent) chemoradiotherapy, CR complete response, CT chemotherapy, DFS disease-free survival, ECOG Eastern Cooperative Oncology Group, EFS event-free survival, G3/4 grade 3 or 4, IASLC International Association for the Study of Lung Cancer, mPR major pathologic response, MRD minimal residual disease, NSCLC non-small cell lung cancer, ORR objective response rate, OS overall survival, pCR pathologic complete response, PFS progression-free survival, PR partial response, PRAE possibly related adverse event, PS performance status, RT radiation therapy, SBRT stereotactic body radiation therapy, SCLC small cell lung cancer, SD stable disease, SOC standard of care, TAE treatment-related adverse event, Tx treatment, WHO World Health Organization**

<sup>a</sup> Although T3N2M0 tumors have been reclassified to Stage IIIB in the 8th edition of the IASLC staging system, these patients remain eligible (as Stage IIIA under the 7th edition criteria)

<sup>b</sup> Complete resection ± neoadjuvant and/or adjuvant therapy

<sup>c</sup> Consolidation treatment in patients with CR, PR or SD only
free survival (DFS) in all patients and in patients with either PD-L1 tumor cell (TC) expression ≥ 25% or ≥ 1%.

In addition to the effects of surgery, recent evidence indicates that, like advanced tumors, early-stage tumors may have an immunosuppressive microenvironment [84]. Emerging clinical data show that PD-1/PD-L1 inhibition (alone or combined with CT) in the neoadjuvant setting in patients with resectable NSCLC is associated with expansion of CD8+ T cell clones in both tumor and peripheral blood expansion and with encouraging rates of major pathologic response (mPR) [85–87].

The phase 3 AEGEAN trial [66, 67, Data on file] (Table 1) will assess the clinical benefit of combining durvalumab with CT in the neoadjuvant setting in patients with resectable stage IIA to select IIIB NSCLC; patients will continue their randomized treatment (durvalumab or placebo) after surgery. The primary end points are pathologic complete response (pCR) and event-free survival (EFS).

Minimal residual disease (MRD), as indicated by the presence of circulating tumor DNA (ctDNA), has shown promise as a prognostic marker for disease recurrence [88]. Its detection, in the absence of radiologic evidence of disease, may improve clinical outcomes via earlier detection and selective intervention with adjuvant therapy, minimizing overtreatment of MRD-negative (MRD−) patients, most of whom are expected to have been cured by their surgery. In addition, MRD detection may help identify patients at high risk of clinical relapse who may benefit from more intense or earlier therapeutic intervention. Such a novel proposed management strategy would replace the current paradigms in which treatment is often reactive, unable to catch-up with the progressing disease, and allow identification of patients at risk for relapse who may need additional therapies (e.g., to maximize curative intent) in a setting where cure is still potentially achievable (Fig. 2).

The phase 3 MERMAID-1 and -2 studies [69, 70] (Table 1) will aim to build a better understanding of the value of MRD in identifying the risk of relapse and the need for therapeutic intervention in patients with completely resected stage II–III NSCLC. In both studies, each patient’s post-surgery MRD status will be assessed using an innovative, highly sensitive, personalized assay developed by ArcherDX that uses whole exome sequencing (WES) of the patient’s resected tumor tissue to detect ctDNA in collected plasma samples. In MERMAID-1, a ‘landmark’ approach with respect to MRD detection immediately after surgery will be utilized, whereas, in MERMAID-2, longitudinal monitoring for ‘emergent’ relapse will be used.

In MERMAID-1 [69], patients will be randomized following successful surgery to
standard-of-care adjuvant CT plus either durvalumab or placebo. The primary end point is DFS in patients who have detectable MRD (MRD+) following surgery, who are at high risk of relapse. In MERMAID-2 [70], patients who have completed curative-intent therapy (complete resection + optional neoadjuvant and/or adjuvant therapy) will be enrolled in a 96-week surveillance period during which they will be monitored for the emergence of MRD; patients who subsequently become MRD+ will then be rescreened and, if eligible, randomized to treatment. The primary end point in MERMAID-2 is DFS in patients with PD-L1 TC ≥ 1%.

Although combinations of neoadjuvant PD-1/PD-L1 inhibitors with CT seem to improve efficacy (as measured by mPR rate) compared with immunotherapy alone [89], improved efficacy may come at risk of combined and increased toxicity; as such, novel combinations of immunotherapy may reduce the level of preoperative toxicity.

In this context, the phase 2 multidrug platform NeoCOAST study [64, 65] (Table 1) will explore the efficacy of durvalumab alone or combined with novel agents (known to be additive to or synergistic with durvalumab) in the neoadjuvant setting. These agents include oleclumab, a monoclonal antibody that binds to CD73, inhibiting production of immunosuppressive adenosine; monalizumab, an immune checkpoint inhibitor targeting NKG2A receptors; or danvatisen, an antisense oligonucleotide that inhibits the STAT3 transcription factor.

**Inoperable Stage I/II NSCLC**

Most patients with unresectable NSCLC have stage III disease, although some patients with localized, stage I or II NSCLC will be medically inoperable because of comorbidities such as heart disease or emphysema. Stereotactic body RT (SBRT) is a recommended standard-of-care treatment for these patients [20, 31], with local tumor control of > 90% at 5 years, and a 5-year OS rate of 40% in stage I medically inoperable patients [90]. The efficacy of durvalumab, compared with placebo, given concurrently with definitive SBRT to patients who have refused surgery or with medically inoperable stage I/II, lymph node-negative NSCLC, will be assessed in the phase 3 PACIFIC-4 trial [71, 72] (Table 1).

**Unresectable Stage III NSCLC**

Stage III NSCLC is heterogeneous in presentation, and the majority of cases are unsuitable for curative surgery as the primary treatment modality [91]. Historically, definitive treatment for unresectable stage III NSCLC had been concurrent CRT; however, based on the PACIFIC trial, durvalumab after CRT is now the standard of care for eligible patients who have not progressed following platinum-based concurrent CRT.

Several hypothesized mechanisms may have contributed to the unprecedented efficacy observed with durvalumab in the PACIFIC trial. These include upregulation of tumor PD-L1 expression by CT and/or RT [45–48] and/or increased availability of tumor neo-antigens (as a result of DNA damage and cell death from RT), promoting both the priming and effector phases of the antitumor immune response mediated by T cells [49, 50]. Moreover, in vivo experiments have demonstrated the synergistic effects of RT and, specifically, PD-1/PD-L1 blockade [46, 47]. However, the optimum sequencing of durvalumab and CRT was not explored in PACIFIC and has not yet been fully investigated in a clinical setting; in addition, preclinical studies suggest that efficacy may be increased when immune checkpoint inhibitors are administered concurrently with RT [47, 50, 92].

In this context, the next key study anticipated to read out in this setting is the PACIFIC-2 trial [79, 80] (Table 1), which will assess whether durvalumab given concurrently with CRT (followed by durvalumab consolidation treatment) provides additional benefit versus concurrent CRT alone.

Although concurrent CRT is superior to sequential CRT for treatment of unresectable stage III NSCLC with respect to survival, it is associated with greater acute
esophageal toxicity [93]; sequential CRT may therefore be preferable for some older patients, those with poorer performance status or those with specific comorbidities unable to tolerate the increased toxicity [31, 94, 95]. The ongoing phase 3 PACIFIC-5 trial [73, 74] (Table 1) was therefore designed to assess the efficacy and safety of durvalumab in patients with no disease progression following either sequential or concurrent CRT. However, unlike the PACIFIC trial, the window for randomization in PACIFIC-5 is up to 28 days post-CRT, rather than up to 42 days, and consolidation treatment with durvalumab or placebo is continued until disease progression (because of the poorer 5-year OS and DFS rates seen for this broader population, which includes patients who progress during the induction CRT therapy [11–14, 57, 93, 96, 97]), unacceptable toxicity or other discontinuation criteria.

Finally, additional studies of durvalumab in patients with unresectable, stage III NSCLC include the three phase 2 trials, PACIFIC-6 [75, 76], COAST [77, 78] and DUART [81] (Table 1).

**Limited-Stage SCLC**

Approximately one-third of patients diagnosed with SCLC present with limited-stage disease, for which the curative-intent standard of care is generally platinum-based CT with concurrent thoracic RT, followed by prophylactic cranial irradiation (PCI) if indicated [98, 99]. However, recurrence of disease is common with a 5-year survival rate of 31–34% [100].

The phase 3 CASPIAN study demonstrated a significant improvement in OS for patients with extensive-stage SCLC who received first-line durvalumab, in combination with a choice of platinum-etoposide (EP) chemotherapy (followed by durvalumab maintenance) versus EP alone (HR, 0.73, 95% CI 0.59–0.91; P = 0.0047) [101], leading to worldwide regulatory approvals for durvalumab as a new standard of care in this setting.

The positive OS outcomes demonstrated in both the CASPIAN and PACIFIC studies suggest the potential for durvalumab to provide benefit for the treatment of limited-stage SCLC. The ADRIATIC study [82, 83] (Table 1), therefore, builds on these studies and will enroll patients with limited-stage SCLC (stages I–III) and no disease progression after concurrent CRT; patients will be randomized to receive durvalumab, durvalumab plus tremelimumab or placebo up to 24 months or until disease progression or intolerable toxicity.

**DURVALUMAB CLINICAL DEVELOPMENT PROGRAM IN OTHER EARLY-STAGE CANCERS**

**Bladder Cancer**

As noted above, there is significant variation in survival rates for patients with bladder cancer, depending on the stage at diagnosis. Management of low-risk disease focuses on preventing recurrence or progression to HR-NMIBC or muscle-invasive bladder cancer (MIBC) for which the 5-year mortality rate is 50–70% (even after radical cystectomy) [102, 103]. The most common treatment for early-stage or superficial (non-muscle invasive) bladder cancers is transurethral resection of the bladder tumor (TURBT), which is commonly followed by intravesical therapy with CT or immunotherapy (as induction and/or maintenance therapy) [19].

It has long been known that bladder cancer is immuno-responsive, when it was demonstrated that intravesical instillation of BCG, a live-attenuated tuberculosis-related bacteria, could be used to stimulate an immunologic reaction in NMIBC, thereby, inducing a pro-inflammatory cytokine and direct cell-to-cell cytotoxicity [104]. BCG is the most common intravesical immunotherapy for treating early-stage bladder cancer. In addition, among all cancer types, bladder has one of the highest mutational burdens and, as such, is likely to elicit a relatively large T-cell mediated antitumor immune response and benefit from checkpoint inhibition [105]. Higher mutation rates have been shown to be associated with higher responses to ICB therapy in several cancer types.
including melanoma, lung cancer and urothelial carcinoma [106–108]. Notably, recent analyses of patients with urothelial carcinoma from a phase 1 trial of durvalumab plus tremelimumab and a phase 1/2 trial of durvalumab monotherapy demonstrated that the tumor mutation burden (TMB) was associated with survival benefit [109]. Finally, upregulation of the PD-1 pathway has also been observed in BCG-resistant NMIBC [110].

Based on the preceding data, combined with the incentive to treat patients with bladder cancer early, there is a strong rationale to study durvalumab in this setting. As such, durvalumab is being assessed in two early-stage bladder cancer trials (Table 2). The ongoing phase 3 NIAGARA trial [111, 112] in patients with resectable MIBC is investigating the effect of adding durvalumab to standard-of-care cisplatin-based neoadjuvant treatment [113, 114] followed by radical cystectomy with extended lymphadenopathy and then adjuvant durvalumab monotherapy. The phase 3 POTOMAC trial [115, 116] will assess whether the addition of durvalumab to standard-of-care BCG, after complete resection of papillary tumors, improves DFS compared with BCG alone.

Liver Cancer

The development of HCC is facilitated by the intrinsic intra-hepatic immunosuppressive microenvironment and chronic inflammation, making HCC an appropriate candidate for immune checkpoint inhibitors [117]. A phase 2 study of patients with advanced HCC (Study 22) recently demonstrated promising clinical activity for durvalumab in combination with the anti-CTLA4 tremelimumab [118]. Durvalumab plus tremelimumab was recently granted orphan drug designation by the US FDA for patients with advanced HCC [119].

Transarterial chemoembolization (TACE) is a standard locoregional treatment for patients with intermediate-stage HCC, but many patients relapse within a year (e.g., based on a systematic review of efficacy across multiple studies of TACE using lipiodal-based regimens, the 1-year PFS rate was 40.6% and estimated 2- and 5-year OS rates were 51.8% and 32.4%, respectively [120]). Combination treatment with a PD-L1 inhibitor, atezolizumab and the VEGF inhibitor, bevacizumab, improved OS in advanced HCC compared with sorafenib [121]; these data provide a rationale for combining durvalumab with VEGF inhibition in locoregional HCC.

Two phase 3 trials of durvalumab in early-stage HCC are currently recruiting patients, EMERALD-1 and EMERALD-2 (Table 2). EMERALD-1 [122, 123] is a placebo-controlled, phase 3 trial of TACE in combination with durvalumab with or without bevacizumab, compared with TACE alone, in patients with locoregional HCC. EMERALD-2 [124, 125] is a placebo-controlled, phase 3 trial evaluating durvalumab with or without bevacizumab as adjuvant therapy in patients with HCC after curative resection or ablation who are at high risk of recurrence.

Other Cancers

Additional phase 3 trials are investigating durvalumab in patients with other early stage cancers, including unresectable esophageal squamous cell carcinoma (e.g., the placebo-controlled KUNLUN study [126], which is assessing durvalumab in combination with definitive CRT), resectable gastric and gastroesophageal junction cancer (e.g., a placebo-controlled trial of perioperative durvalumab in combination with FLOT [fluorouracil + leucovorin + oxaliplatin + docetaxel] CT [127]), and locally advanced cervical cancer (e.g., the placebo-controlled CALLA study [128, 129], which is assessing durvalumab with concurrent CRT, followed by durvalumab, versus concurrent CRT alone (Table 2)).

CONCLUSION

Patients diagnosed with early-stage cancers frequently relapse with locoregional or distant disease. The practice-changing survival benefit shown with durvalumab following CRT in the phase 3 PACIFIC trial of patients with stage III, unresectable NSCLC supports the strategy of
| Diagnosis and staging | Setting         | Trial                  | Patient population                                                                 | Estimated (or actual) enrollment | Design                                                                 | Primary end points |
|-----------------------|-----------------|------------------------|------------------------------------------------------------------------------------|----------------------------------|-----------------------------------------------------------------------|--------------------|
| Resectable muscle-invasive bladder cancer | Neoadjuvant/adjuvant | NIAAGARA [111, 112] (NCT03732677) | Stage T2-4aN0/1M0 with transitional and mixed transitional cell histology          | 1050                              | Phase 3, open-label, randomized, global trial                         | pCR                |
|                        |                 |                        |                                                                                      |                                  | Arm 1: Neoadjuvant durvalumab + cisplatin-gemcitabine, with adjuvant durvalumab | EFS                |
|                        |                 |                        |                                                                                      |                                  | Arm 2: Neoadjuvant cisplatin-gemcitabine, with observation after surgery |                    |
| Non-muscle invasive bladder cancer | BCG-naive      | POTOMAC [115, 116] (NCT03528694) | High risk<sup>a</sup>                                                                | 973                               | Phase 3, open-label, randomized, global trial                         | DFS                |
| Locoregional hepatocellular carcinoma | TACE-naive     | EMERALD-1 [122, 123] (NCT03778957) | Not amenable to curative therapy                                                    | 600                               | Phase 3, double-blind, randomized, global trial                         | PFS (Arm B vs. Arm C) |
| Hepatocellular carcinoma | Adjuvant       | EMERALD-2 [124, 125] (NCT03847428) | Prior curative resection or ablation and high risk of recurrence                     | 888                               | Phase 3, double-blind, randomized, global trial                         | RFS (Arm B vs. Arm C) |
| Locally advanced cervical cancer | CRT-naive     | CALLA [128, 129] (NCT03830866) | Cervical adenocarcinoma or squamous carcinoma FIGO (2009) Stages IB2–IIA node positive or FIGO (2009) Stages IIIA–IVA any node | 714                               | Phase 3, double-blind, randomized, global trial                         | PFS                |
|                        |                 |                        |                                                                                      |                                  | Arm 1: Durvalumab + SOC cCRT, followed by durvalumab                     |                    |
|                        |                 |                        |                                                                                      |                                  | Arm 2: Placebo                                                         |                    |

The current status of each trial (i.e., recruitment status and estimated primary completion date) can be obtained by accessing its study record on the ClinicalTrials.gov registry using its unique NCT number.

*BCG bacillus Calmette-Guerin, CIS carcinoma in situ, (c)CRT concurrent chemoradiotherapy, DFS disease-free survival, EFS event-free survival, FIGO International Federation of Gynecology and Obstetrics (staging system), G3 histologic grade 3, HCC hepatocellular carcinoma, MIBC muscle-invasive bladder cancer, NMIBC non-muscle-invasive bladder cancer, pCR pathologic complete response, PFS progression-free survival, RFS recurrence-free survival, RT radiation therapy, SOC standard of care, TACE transarterial chemoembolization*

<sup>a</sup> Defined as any of the following: T1 tumor; high grade/G3 tumor; CIS; multiple, recurrent and large (i.e., diameter of largest evaluable node ≥ 3 cm; all three criteria must be met here) tumors.
using immunotherapy in a curative-intent paradigm. There may be inherent challenges associated with conducting a clinical development program in early-stage cancers, e.g., ensuring harmonization of locoregional treatments and coordinating among members of the multidisciplinary teams, which are integral to the patient journey. However, it is anticipated that the program for durvalumab will add considerably to our knowledge regarding its potential use (and immunotherapies, in general) in both resectable and unresectable early-stage disease. For example, it will address questions related to optimal sequencing or combination of durvalumab with other standard-of-care treatments (e.g., as adjuvant or neoadjuvant therapy), the optimal length of treatment with durvalumab, and the efficacy and safety of durvalumab in novel combinations with other agents.

In addition, the trials in the durvalumab development program are at the forefront of innovative, personalized cancer monitoring (e.g., via experimental use of ctDNA to monitor patients’ risk of relapse). These efforts, combined with other investigative trends, such as ‘stage shifting’ via population-scale early cancer detection, hold the promise to, once again, shift the treatment landscape in the curative-intent setting.

Finally, the extensive clinical development program for durvalumab is complemented by trials of other compounds in early-stage cancers, such as the recently reported phase 3 ADAURA trial, which assessed the EGFR inhibitor osimertinib as adjuvant treatment, showing a significantly prolonged DFS for patients with early-stage EGFR-mutant NSCLC [130]. The overall breadth and depth of AstraZeneca’s cancer trial portfolio have the potential for significant advances in the treatment of these early-stage tumors over the years to come.

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