

**Pfizer Forms Global Alliance with
Merck KGaA, Darmstadt, Germany
to Accelerate Presence in Immuno-Oncology**

November 17, 2014



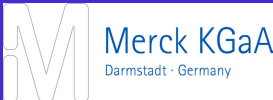
Forward-looking statements

- Our discussions during this conference call will include forward-looking statements. Actual results could differ materially from those projected in the forward-looking statements. The factors that could cause actual results to differ are discussed in Pfizer's 2013 Annual Report on Form 10-K and in our subsequent reports on Form 10-Q and Form 8-K, and in Pfizer's Press Release issued on November 17, 2014.
- Also, the discussions during this conference call may include certain financial measures that were not prepared in accordance with U.S. generally accepted accounting principles (GAAP). Reconciliations of those non-U.S. GAAP financial measures to the most directly comparable U.S. GAAP financial measures can be found in Pfizer's Current Report on Form 8-K dated October 28, 2014 and Pfizer's Quarterly Report on Form 10-Q for the fiscal quarter ended September 28, 2014.
- These reports are available on our website at www.pfizer.com in the "Investors—SEC Filings" section.

Pfizer and Merck KGaA, Darmstadt, Germany: A Great Match



- Strong global commitment to Oncology development and commercialization
- Investing heavily in immuno-oncology R&D



- Strong legacy and commitment to oncology with cetuximab ex-US
- Major player in immuno-oncology with their anti-PD-L1

**Focused on
improving the
lives
of patients
with cancer**

**Together, one team
committed to driving rapid
and broad development
and commercialization
of our anti-PD-L1 and
anti-PD-1 antibodies
as a key priority**

Key Deal Terms

Co-Development and Co-Commercialization of Anti-PD-L1/ Anti-PD-1

- Upfront payment of \$850 million to Merck KGaA, Darmstadt, Germany
- Merck KGaA, Darmstadt, Germany is eligible to receive regulatory and commercial milestone payments up to approximately \$2 billion
- Pfizer and Merck KGaA, Darmstadt, Germany will jointly fund all development and commercialization costs
- Collaboration revenues from any PD-L1 or PD-1 products will be shared equally

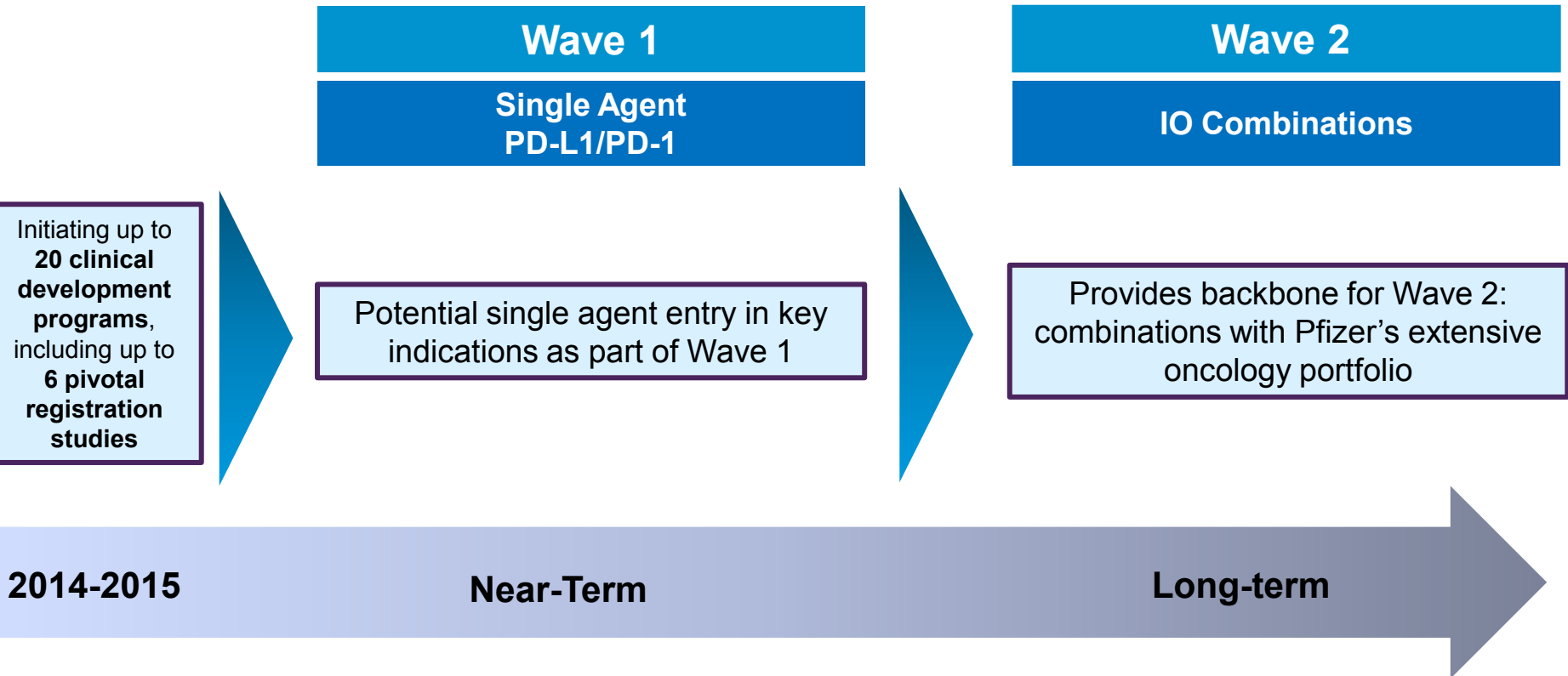
Xalkori Co-Promotion

- Merck KGaA, Darmstadt, Germany and Pfizer will co-promote Xalkori in the United States and several other key markets

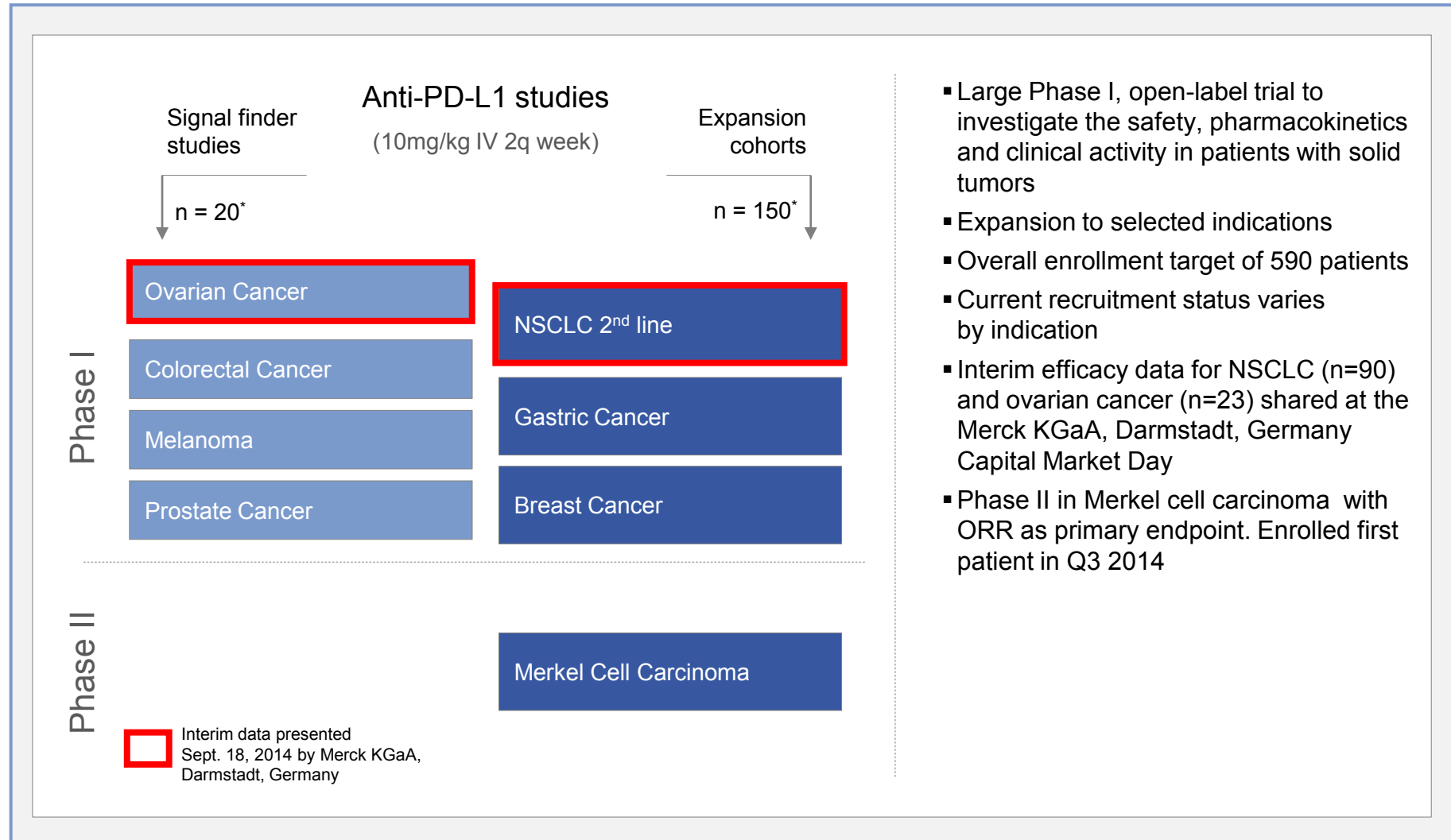
Governance

- Collaboration team and governance structure will operate with joint decision-making

Collaboration Positions both Companies into Potential Wave 1 of Immunotherapy Launches



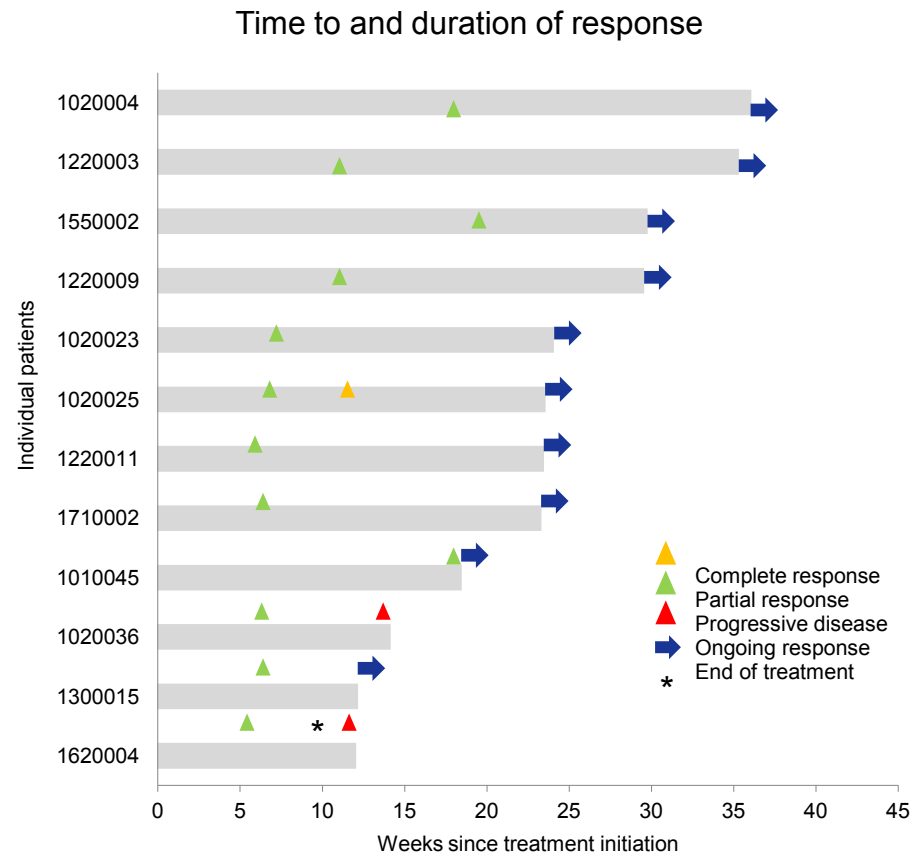
Current clinical program of Anti-PD-L1



- Large Phase I, open-label trial to investigate the safety, pharmacokinetics and clinical activity in patients with solid tumors
- Expansion to selected indications
- Overall enrollment target of 590 patients
- Current recruitment status varies by indication
- Interim efficacy data for NSCLC (n=90) and ovarian cancer (n=23) shared at the Merck KGaA, Darmstadt, Germany Capital Market Day
- Phase II in Merkel cell carcinoma with ORR as primary endpoint. Enrolled first patient in Q3 2014

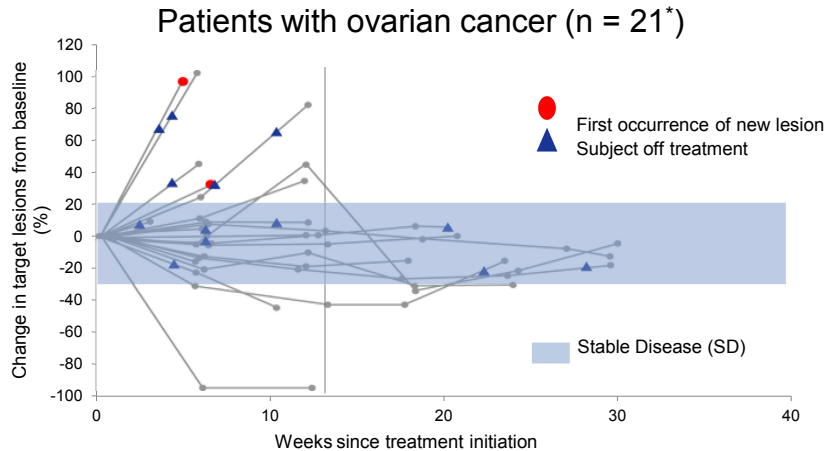
Phase I efficacy result: Response rates in NSCLC

Best Overall Response by RECIST 1.1 unconfirmed	NSCLC Intent-to-treat, n = 90 (%)
Complete Response (CR)	1 (1.1%)
Partial Response (PR)	11 (12.2%)
Stable Disease (SD)	30 (33.3%)
Progressive Disease (PD)	35 (38.9%)
Non-evaluable (NE)	13 (14.4%)
Objective response rate* (ORR) [95% CI**]	13.3% [7.1%, 22.1%]



With minimum follow-up time of 3 months, the ORR is similar to other anti-PD-1/PD-L1 agents

Phase I results in ovarian cancer: Tumor shrinkage and duration of response



Best Overall Response by RECIST 1.1
unconfirmed

Ovarian cancer
n = 23; n (%)

Complete Response (CR)

0

Partial Response (PR)

4 (17.4%)

Stable Disease (SD)

11 (47.8%)

Progressive Disease (PD)

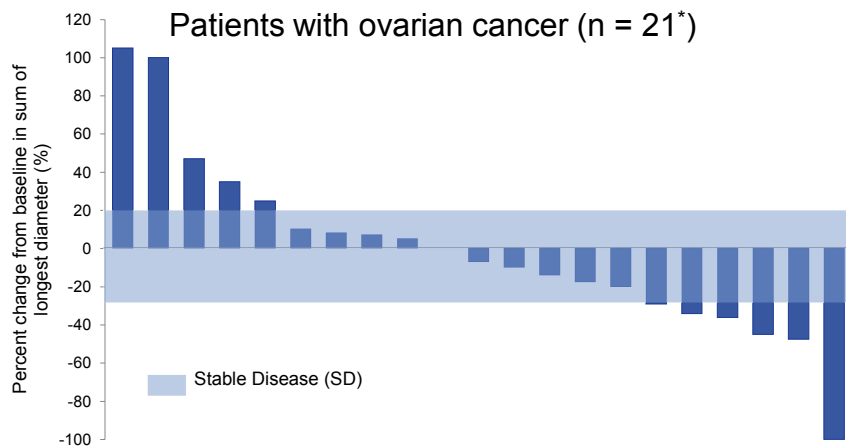
7 (30.4%)

Non-evaluable (NE)

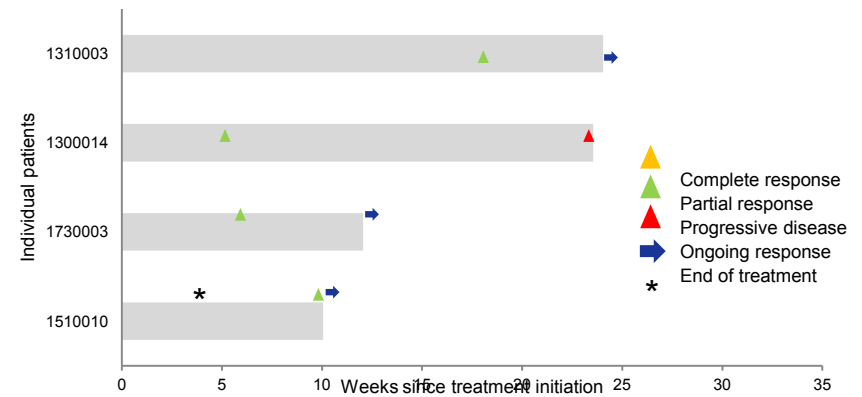
1 (4.3%)

Objective Response Rate (ORR)**
[95% CI***]

17.4%
[5.0%, 38.8%]



Time to and duration of response



Data presented at Merck KGaA, Darmstadt, Germany Capital Markets Day, September 18, 2014, and based on an interim analysis

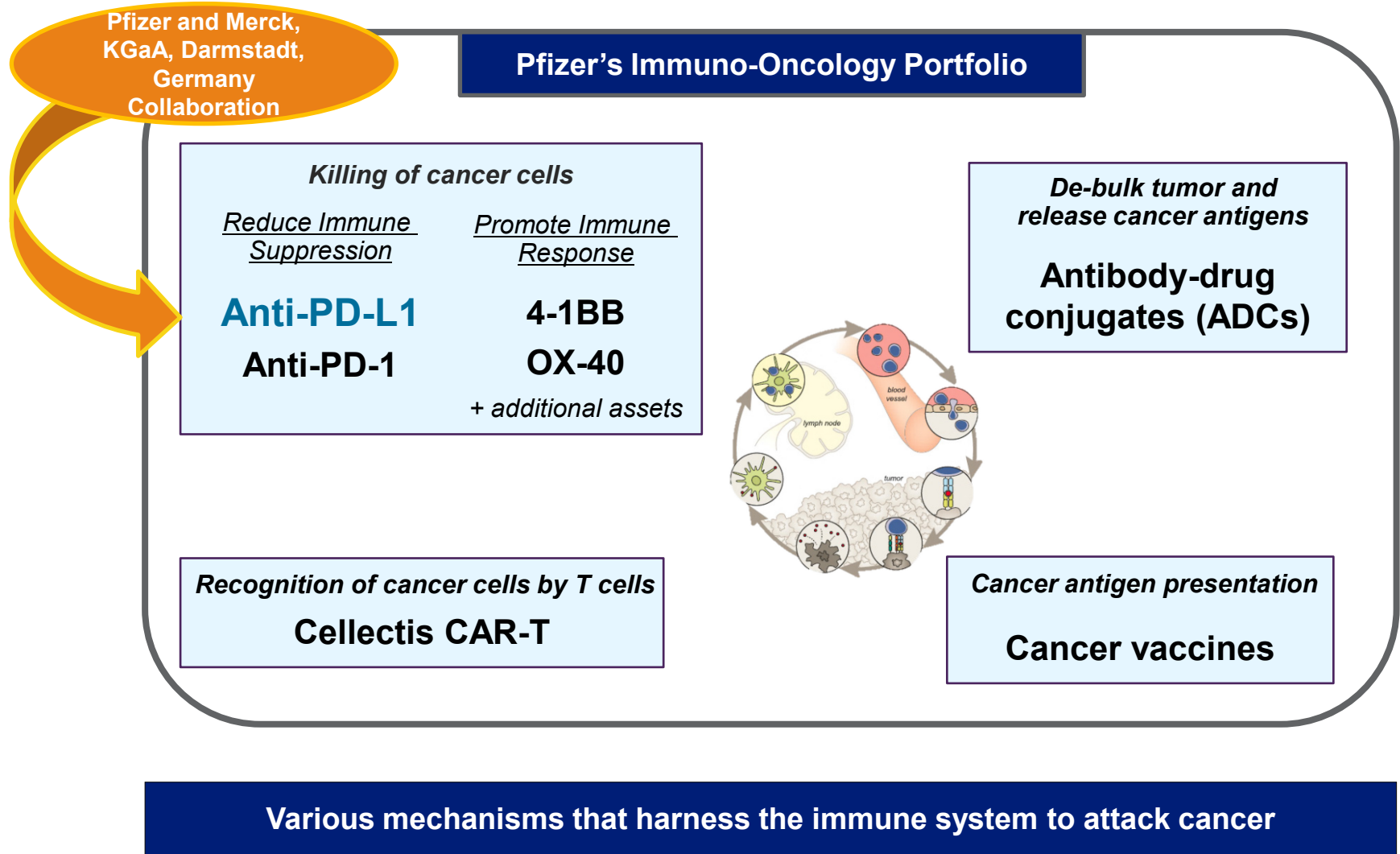
*Based on evaluable patients ; **Response rate per RECIST v1.1 is based on all treated patients. ORR includes both confirmed and unconfirmed responses (CR and PR); ***Confidence interval

Phase I safety results: Adverse Events

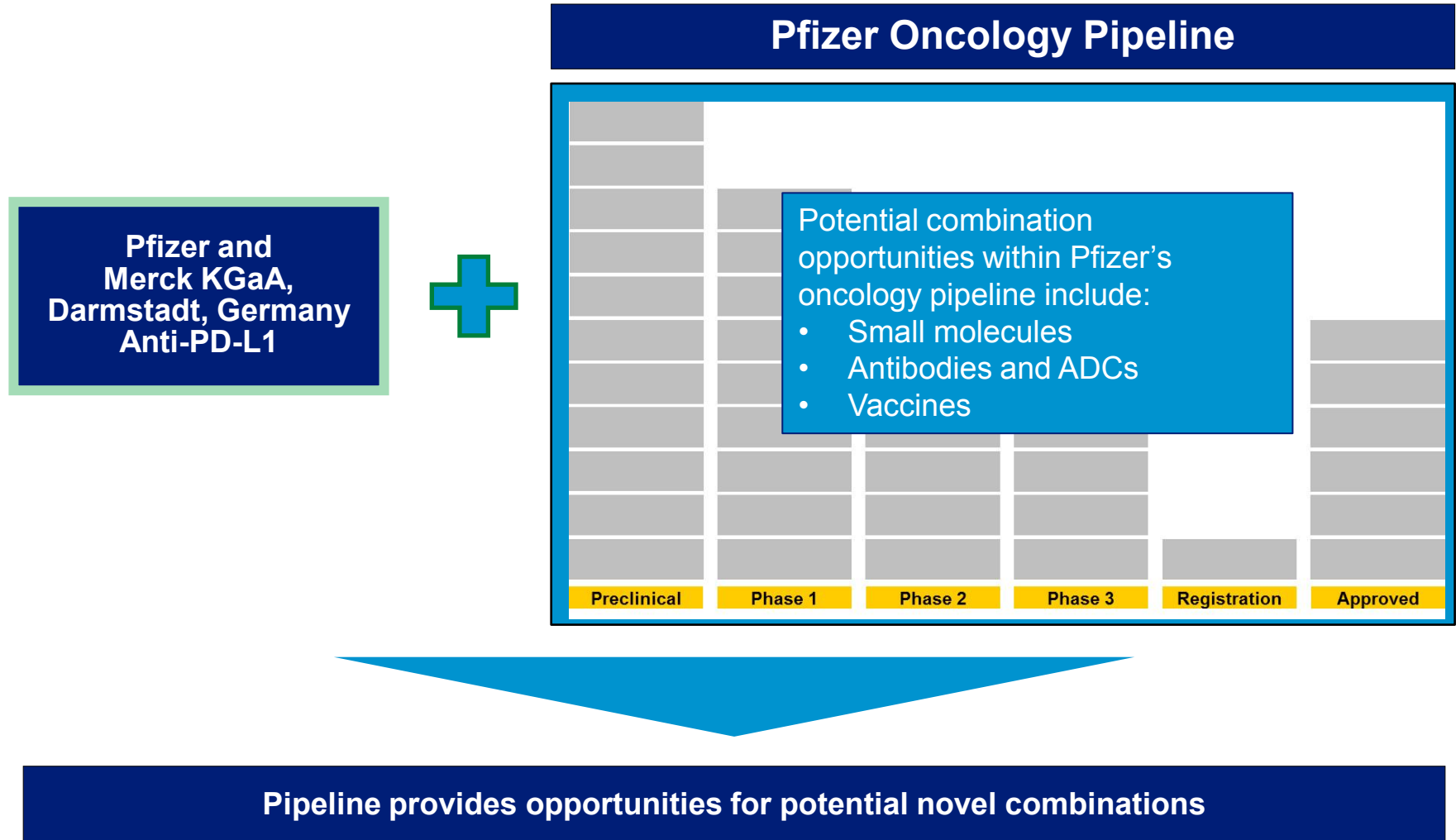
	Pooled expansion cohorts (n = 290) n (%)	NSCLC (n = 127) n (%)	Ovarian cancer (n = 23) n (%)
AEs	262 (90.3)	114 (89.8)	23 (100.0)
Related AEs	198 (68.3)	87 (68.5)	18 (78.3)
AEs, Grade ≥3	124 (42.8)	55 (43.3)	9 (39.1)
Related AEs, Grade ≥3	38 (13.1)	17 (13.4)	2 (8.7)

- Current safety information based on an analysis of 290 subjects (expansion part of study -001)
- Cut-off date: July 16, 2014
- Minimum follow-up time: 4 weeks

Pfizer's Existing Immuno-oncology Portfolio is Complemented by Merck KGaA's, Darmstadt, Germany Anti-PD-L1



Collaboration Will Leverage Pfizer's Broad Oncology Pipeline



Q&A