

Clinical Updates From San Antonio

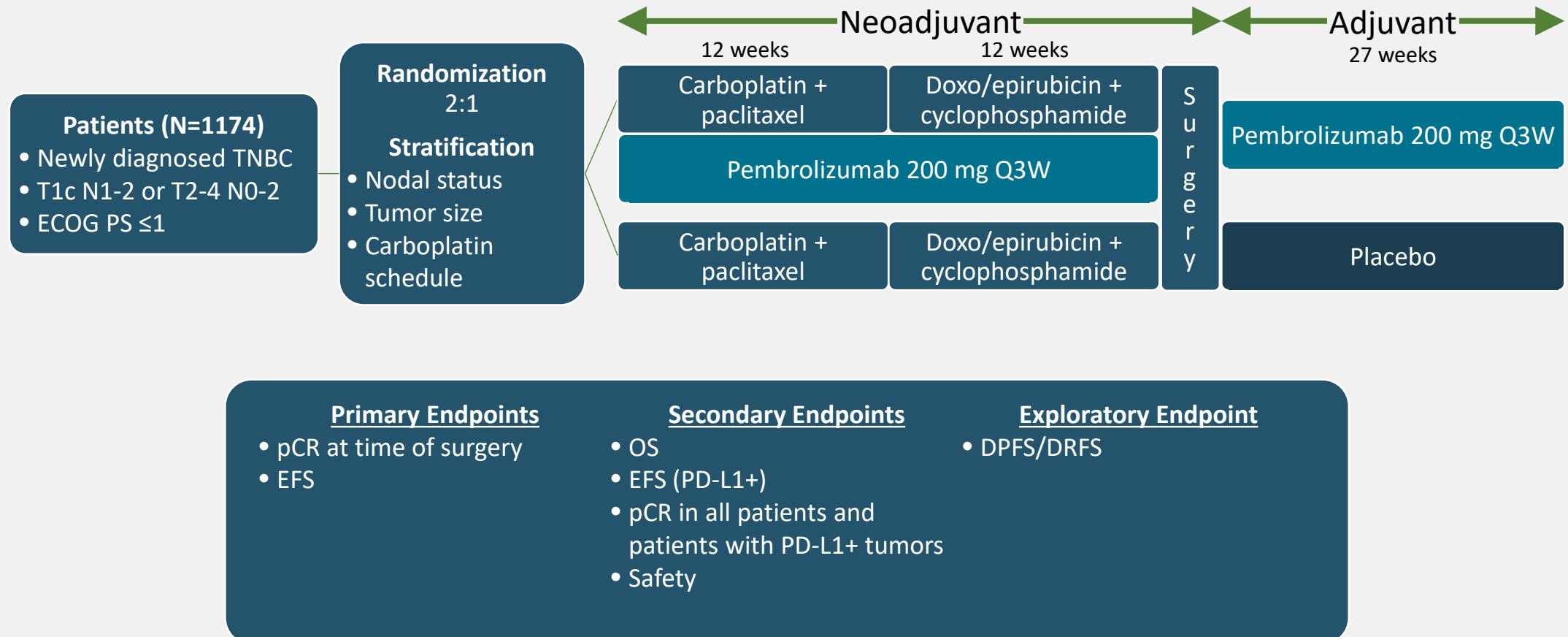
Triple-Negative Breast Cancer: Neoadjuvant/Adjuvant Immunotherapy

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KEYNOTE-522 Trial Design



DPRS, distant progression-free survival; DRFS, distant recurrence-free survival; Doxo, doxorubicin; ECOG, Eastern Cooperative Group; EFS, event-free survival; N0, no regional LN metastases; N1-2, metastases to axillary lymph nodes or ipsilateral internal mammary nodes in the absence of axillary lymph node metastases; OS, overall survival; pCR, pathologic complete response; PD-L1, programmed death-ligand 1; PS, performance status; Q3W, every 3 weeks; T1c, tumor >1 cm but ≤2 cm; T2-4, tumor >2 cm but ≤5 cm, >5 cm, or any size with extension to the chest wall and/or skin; TNBC, triple-negative breast cancer.

Schmid P, et al. *N Engl J Med.* 2020;382(9):810-821; Schmid P, et al. *N Engl J Med.* 2022;386(6):556-567.

Comparison of Real-World and Phase 3 Efficacy and Safety of Treatment per KEYNOTE-522* in Patients With eTNBC[†]

| Study | EHR Analysis (n=66) | | KEYNOTE-522 (n=781) | |
|-------------------------------|--------------------------|----------|------------------------|----------|
| Population characteristics | T3/4: 48.1% N+: 53.2% | | T3/4: 26% N+: 51.7% | |
| pCR rate, % | 45.7 (n=55) | | 64.8 (N=784) | |
| AEs, % | Any grade | Grade ≥3 | Any grade | Grade ≥3 |
| Nausea | 57.6 | 13.6 | 62.7 | 3.3 |
| Neutropenia | 78.7 | 62.1 | 46.7 | 34.6 |
| Fatigue | 71.2 | 27.3 | 41.1 | 3.5 |
| Diarrhea | 40.9 | 15.2 | 29.4 | 2.2 |
| Peripheral neuropathy | 39.4 | 9 | 19.7 | 1.9 |
| Rash | 12.1 | 4.5 | 21.8 | 0.9 |
| imAEs, % | Any grade | Grade ≥3 | Any grade | Grade ≥3 |
| Hypothyroidism | 18.2 | 4.5 | 13.7 | 0.4 |
| Hyperthyroidism | 1.5 | 1.5 | 4.6 | 0.3 |
| Type 1 diabetes | 3 | 3 | 0.3 | 0.3 |
| Primary adrenal insufficiency | 9 | 3 | 2.3 | 1.3 |
| Colitis | 9 | 7.6 | 1.7 | 0.9 |
| Pancreatitis | 3 | 1.5 | 0.5 | 0.5 |
| Hepatitis | 3 | 1.5 | 1.4 | 1.2 |
| Pneumonitis | 1.5 | 0 | 1.3 | 0.4 |
| Ocular toxicity | 3 | 1.5 | 0 | 0 |

*4 cycles of neoadjuvant pembrolizumab plus paclitaxel and carboplatin, followed by 4 cycles of pembrolizumab plus cyclophosphamide plus doxorubicin or epirubicin, and adjuvant pembrolizumab for up to 9 cycles; [†]Stage 2 or 3.

AE, adverse event; EHR, electronic health record; eTNBC, early triple-negative breast cancer; imAE; immune-mediated adverse event; N+, node positive; T3/4, tumor >5 cm/any size with extension to the chest wall and/or skin.

Hofherr M, et al. SABCS 2022. Abstract P3-06-06; Schmid P, et al. *N Engl J Med.* 2020;382(9):810-821; Schmid P, et al. *N Engl J Med.* 2022;386(6):556-567.

P3-06-06, Real-World Analysis of Adverse Events of Patients With Triple-Negative Breast Cancer Receiving Therapy per KEYNOTE-522

Patients Identified (N=79)

- Retrospective, single-center EHR analysis
- eTNBC with planned treatment per KN522*

Outcomes of Interest

- Treatment delays: number and length
- Treatment-related toxicities
- pCR rate

Hofherr M, Hedgecorth J, Ademuyiwa FO, Peterson LL, Bagegni NA, Suresh R, Frith A, Bose R, Weilbaecher K, Ma CX, Davis AA, Clifton KK

| Study | EHR Analysis (N=79) |
|--------------------------------|--------------------------|
| Population characteristics | T3/4: 48.1% N+: 53.7% |
| pCR rate, % | 45.7 (n=55) |
| Hospitalizations ≥ 1 | 38 |
| Emergency room visits ≥ 1 | 27 |
| Treatment delays | 40 |
| Dose reductions | 26 |

*4 cycles of neoadjuvant pembrolizumab plus paclitaxel and carboplatin, followed by 4 cycles of pembrolizumab plus cyclophosphamide plus doxorubicin or epirubicin, and adjuvant pembrolizumab for up to 9 cycles.

KN522, KEYNOTE-522.

Hofherr M, et al. SABCS 2022. Abstract P3-06-06.

P3-06-09, Real-World Toxicity of Pembrolizumab-Based Neoadjuvant Regimen in Patients With Early Triple-Negative Breast Cancer

Patients Identified (N=51)

- Ambispective, single-center analysis of clinical data
- eTNBC
- Treated per KN522*

Outcomes of Interest

- Efficacy
- Safety

Arnaud E, Alaoui K, Vaflard P, Korbi S, Meziani D, Thibault L, Desmaris R-P, Feron J-G, Pierga J-Y, Bidard F-C, Cottu P, Loirat D

| Study | Ambispective cohort (N=51) | |
|------------------------------------|----------------------------|--------------------|
| Population characteristics | T3/4: 25% N+: 72.5% | |
| Median follow-up, months | 5 | |
| pCR rate, % | 78.3 (KN522: 64.8) | |
| Postponement or discontinuation, % | 34 | |
| Dose reduction, % | 19 | |
| AEs ($\geq 5\%$), % | Any grade | Grade ≥ 3 |
| Any AE | 100 | 76.5 (KN522: 76.8) |
| Anemia | 100 | 37.3 |
| Thrombocytopenia | 37.3 | 11.8 |
| Neutropenia | 82.4 | 62.8 |
| Febrile neutropenia | 19.7 | 19.7 |
| Peripheral neuropathy | 47.1 | 2 |
| imAEs ($\geq 5\%$), % | 49 | 17.6 |
| Hypothyroidism | 9.8 | NA |
| Hyperthyroidism | 7.8 | 3.9 |
| Hypophysitis | 5.9 | NA |
| Troponine elevation | 7.8 | 2 |

*4 cycles of neoadjuvant pembrolizumab plus paclitaxel and carboplatin, followed by 4 cycles of pembrolizumab plus cyclophosphamide plus doxorubicin or epirubicin, and adjuvant pembrolizumab for up to 9 cycles.

P3-06-10, Immune-Related Adverse Events (irAEs) and Pathological Complete Response (pCR) Rates in Patients Receiving Neoadjuvant Chemotherapy (CHT) and Pembrolizumab (PEM) for Early Triple-Negative Breast Cancer (eTNBC)

Patients Identified (N=22)

- Prospective, multicenter analysis
- eTNBC
- Receiving treatment per KN522*

Outcomes of Interest

- imAE rate
- pCR rate

Marhold M, Udvica S, Wimmer K, Bago-Horvath Z, Robinson T, Fitzal F, Strasser-Weippl K, Bartsch R

| Study | Austrian (N=22) |
|-----------------------------------|---------------------------------|
| Population characteristics | Mean T: 29mm (10-75) N+: 41% |
| pCR rate, % | 50 |
| Discontinuation before week 18, % | 32 |
| imAEs grade ≤2, % | 50 |
| Hypothyroidism | 14 |
| Arthritis | 14 |
| Hepatitis | 5 |
| Pneumonitis | 5 |
| imAEs grade ≥3, % | 14 |
| Myocarditis | 9 |
| Nephritis | 5 |

*4 cycles of neoadjuvant pembrolizumab plus paclitaxel and carboplatin, followed by 4 cycles of pembrolizumab plus cyclophosphamide plus doxorubicin or epirubicin, and adjuvant pembrolizumab for up to 9 cycles.