

Atezolizumab 1200 / Carboplatin 6 / Nab-Paclitaxel 100, Non-Small Cell Lung Cancer

Protocol-ID: 1238 V1.2 (Complete), ATEZ1200/CRBP6/NPAC100, NSCLC

Indication(s)

• Lung Carcinoma, Non-Small Cell (non-squamous); ICD-10 C34.-

Protocol classification

Classification: alternative
Intensity: Standard dose
Therapy mode: First line
Therapy intention: palliative

Cycles

Cycle length 21 days, recommended cycles: 6

Protocol sequences

 IMpower130: ATEZ1200/CRBP6/NPAC100, NSCLC (PID1238) -|- ATEZ1200/CRBP6/NPAC100 - ATEZ1200 maint. (PID1272)

Risks

- Emetogenicity (MASCC/ESMO): high (>90%) Carboplatinkombination
- Emetogenicity (MASCC/ESMO): low (10-30%) Nab-paclitaxel
- Neutropenia: very high (>41%) °3-4: 44%
- Thrombocytopenia below 50 000/μl: very high (>41%) °3-4: 45%
- Anemia Hb below 8g/dl: high (16-30%) °3-4: 29%
- Diarrhea: CTC AE °3-4: 5%
 Fatigue: CTC AE °3-4: 6%
 Nausea: CTC AE °3-4: 3%

Therapy

Hydra	tion: Balanced Crystalloid Solution						HYD
Hydration: Balanced Crystalloid Solution Access: peripheral venous Hydration before, during, or after antitumor therapy Day Substance Dosage Solution Appl. Inf. time Procedure Balanced Crystalloid Solution 500 ml i.v. 60 min 60 min before Atezolizumab (d1) 8,15 Balanced Crystalloid Solution 500 ml i.v. 60 min 60 min before Nab-paclitaxel (d8,15)							
Hydra	tion before, during, or after antitumor therapy						
Day	Substance	Dosage	Solution	Appl.	Inf. time	Procedure	
1	Balanced Crystalloid Solution	500 ml		i.v.	60 min	60 min before Atezolizumab (d1)	
8,15	Balanced Crystalloid Solution	500 ml		i.v.	60 min	60 min before Nab-paclitaxel (d8,15)	

Antiemesis: Emetogenicity high (CRBP), FOSAP, GRAN i.v., DEXA i.v.

Access: peripheral venous

DGHO 2016, DKG 2016, MASCC/ESMO 2016, carboplatin-containing combination therapies

Day	Substance	Dosage	Solution	Appl.	Inf. time	Procedure
1	Fosaprepitant	150 mg	NaCl 0.9% 150 ml	i.v.	20 min	30 min before Atezolizumab (d1)
1	Dexamethasone	12 mg	NaCl 0.9% 50 ml	i.v.	5 min	30 min before Atezolizumab (d1)
1	Granisetron	1 mg	NaCl 0.9% 50 ml	i.v.	5 min	15 min before Atezolizumab (d1)
8,15	Granisetron	1 mg	NaCl 0.9% 50 ml	i.v.	5 min	15 min before Nab-paclitaxel (d8,15)

Medical tumor therapy: ATEZ1200/CRBP6/NPAC100

CTX

ΑE

Access: peripheral venous

Atezolizumab, carboplatin, and nab-paclitaxel in non-small cell, non-plate epithelial lung cancer

Day	Substance	Dosage	Solution	Appl.	Inf. time	Procedure
1	Atezolizumab	1200 mg	NaCl 0.9% 250 ml	i.v.	60 min	Sequence
If the first i	nfusion was well tolerated	, the second infusion can be ç	given over 30 minutes.			
1	Carboplatin	6 AUC	Dextrose 5% 250 ml	i.v.	30 min	Sequence
1,8,15	Nab-paclitaxel	100 mg/m² BSA	none	i.v.	30 min	Sequence

Concomitant therapy supplements

Granisetron in place of Dexamethasone for antiemesis on days 8 and 15 to avoid immunosuppression and the risk of infection from Dexamethasone exposure.

Notes

4 or 6 induction cycles were administered, after which patients received atezolizumab as maintenance therapy. Therapy is continued until reduction of clinical benefit or the occurrence of undesirable side effects.

Cycle diagram

Hydration: Balanced Crystalloid Solution

			We	ek 1	/ d			Week 2 / d								Week 3 / d						
Substance	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	
Balanced Crystalloid Solution (i.v.)																						
Balanced Crystalloid Solution (i.v.)																						

Antiemesis: Emetogenicity high (CRBP), FOSAP, GRAN i.v., DEXA i.v.

			We	ek 1	/ d			Week 2 / d								Week 3 / d							
Substance	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21		
Fosaprepitant (i.v.)																							
Dexamethasone (i.v.)																							
Granisetron (i.v.)																							
Granisetron (i.v.)																							

Medical tumor therapy: ATEZ1200/CRBP6/NPAC100

			We	ek 1	/ d			Week 2 / d								Week 3 / d							
Substance	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21		
Atezolizumab (i.v.)																							
Carboplatin (i.v.)																							
Nab-paclitaxel (i.v.)																							

Cycles

Cycle length 21 days, recommended cycles: 6

Controls:

- · Blood count: on day 1 and subsequently weekly
- Echocardiography, ECG Nab-paclitaxel: monitoring for cardiac events, cases of left ventricular dysfunction and congestive heart failure occurred.
- Day 1: TSH, fT3, fT4 Monitor for changes in thyroid function and signs of thyroid disease. Monitor for immune-mediated endocrinopathies at baseline and during therapy.
- Day 1: GOT, GPT, GGT, Bilirubin, AP, Cholinesterase Nab-paclitaxel: Liver monitoring before and during therapy, dose adjustment if necessary. Impairment of liver function possible with carboplatin therapy.
- Day 1: Glomerular Filtration Rate (GFR) monitor immune-mediated nephritis, Carboplatin dose calculation according to AUC and Calvert's formula; in normal renal function, expect a maximum GFR of 125 ml/min to avoid overdoses.

Original indication

non-small cell lung cancer, adeno-, stage IV, first line, ECOG 0-1

Original author

West H (2019)

Origin

Thoracic Oncology Program, Swedish Cancer Institute, Seattle, USA, IMpower130

References

- West H, Atezolizumab in combination with carboplatin plus nab-paclitaxel chemotherapy compared with chemotherapy alone as first-line treatment for metastatic non-squamous non-small-cell lung cancer (IMpower130): a multicentre, randomised, open-label, phase 3 trial., Lancet Oncol 2019 May 20; [PMID]
- Arbour KC, Impact of Baseline Steroids on Efficacy of Programmed Cell Death-1 and Programmed Death-Ligand 1
 Blockade in Patients With Non-Small-Cell Lung Cancer., J Clin Oncol 2018 10 01;36(28):2872-2878 [PMID]

Recommendations

- 01/2023: European Society for Medical Oncology
- 02/2024: National Comprehensive Cancer Network

Status

Valid since 2024-02-15, Version 1.2, last updated 2024-02-15

Last modification: V1.2: Addition of the corticosteroid under immunotherapy V1.1: Cato test done. Removal of corticosteroid according to Della Corte 2019 / Arbour 2018, replacement of Granisetron with Palonosetron during immunotherapy. V1.0: Cato test done. V0.1: Runtimes according to summary of product characteristics.

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