

**Tremelimumab** (non-small cell lung cancer, EGFR/ALK negative, first-line, combination with durvalumab and platinum-based chemotherapy)

Resolution of: 5 October 2023  
Entry into force on: 5 October 2023  
Federal Gazette, BAnz AT 05 02 2024 B3

valid until: unlimited

**Therapeutic indication (according to the marketing authorisation of 20 February 2023):**

Tremelimumab AstraZeneca in combination with durvalumab and platinum-based chemotherapy is indicated for the first-line treatment of adults with metastatic non-small cell lung cancer (NSCLC) with no sensitising EGFR mutations or ALK positive mutations.

**Therapeutic indication of the resolution (resolution of 5 October 2023):**

See therapeutic indication according to marketing authorisation.

**1. Additional benefit of the medicinal product in relation to the appropriate comparator therapy**

a) Adults with metastatic NSCLC with PD-L1 expression  $\geq$  50% with no genomic EGFR or ALK tumour mutations, first-line therapy

**Appropriate comparator therapy:**

– pembrolizumab as monotherapy

*or*

– atezolizumab as monotherapy

*or*

– cemiplimab as monotherapy

*or*

– nivolumab in combination with ipilimumab and 2 cycles of platinum-based chemotherapy (only for patients with ECOG PS 0-1)

*or*

– pembrolizumab in combination with carboplatin and either paclitaxel or nab-paclitaxel (only for patients with ECOG PS 0-1 and a squamous NSCLC)

*or*

– pembrolizumab in combination with pemetrexed and platinum-containing chemotherapy (only for patients without ECOG PS 0-1 and a non-squamous NSCLC)

*or*

– atezolizumab in combination with bevacizumab, paclitaxel and carboplatin (only for patients with ECOG PS 0-1 and a non-squamous NSCLC)

*or*

- atezolizumab in combination with nab-paclitaxel and carboplatin (only for patients with ECOG PS 0-1 and a non-squamous NSCLC)

**Extent and probability of the additional benefit of tremelimumab in combination with durvalumab and platinum-based chemotherapy versus pembrolizumab:**

An additional benefit is not proven.

b) Adults with metastatic NSCLC with PD-L1 expression < 50% with no genomic EGFR or ALK tumour mutations, first-line therapy

**Appropriate comparator therapy:**

- pembrolizumab in combination with pemetrexed and platinum-containing chemotherapy (only for patients without ECOG PS 0-1 and a non-squamous NSCLC)

*or*

- pembrolizumab in combination with carboplatin and either paclitaxel or nab-paclitaxel (only for patients with ECOG PS 0-1 and a squamous NSCLC)

*or*

- atezolizumab as monotherapy (only for patients with PD-L1 expression  $\geq 10\%$  in tumour-infiltrating immune cells)

*or*

- atezolizumab in combination with bevacizumab, paclitaxel and carboplatin (only for patients with ECOG PS 0-1 and a non-squamous NSCLC)

*or*

- atezolizumab in combination with nab-paclitaxel and carboplatin (only for patients with ECOG PS 0-1 and a non-squamous NSCLC)

*or*

- nivolumab in combination with ipilimumab and 2 cycles of platinum-based chemotherapy (only for patients with ECOG PS 0-1)

*or*

- carboplatin in combination with a third-generation cytostatic (vinorelbine or gemcitabine or docetaxel or paclitaxel or pemetrexed) cf. Annex VI to Section K of the Pharmaceuticals Directive (only for patients with ECOG PS 2)

*or*

- carboplatin in combination with nab-paclitaxel (only for patients with ECOG PS 2)

**Extent and probability of the additional benefit of tremelimumab in combination with durvalumab and platinum-based chemotherapy compared to the appropriate comparator therapy:**

An additional benefit is not proven.

## Study results according to endpoints:<sup>1</sup>

- a) Adults with metastatic NSCLC with PD-L1 expression  $\geq$  50% with no genomic EGFR or ALK tumour mutations, first-line therapy

### Summary of results for relevant clinical endpoints

Endpoint category	Direction of effect/ risk of bias	Summary
Mortality	$\leftrightarrow$	No relevant difference for the benefit assessment.
Morbidity	n.a.	There are no assessable data.
Health-related quality of life	n.a.	There are no assessable data.
Side effects	$\leftrightarrow$	No relevant differences for the benefit assessment.

Explanations:  
↑: statistically significant and relevant positive effect with low/unclear reliability of data  
↓: statistically significant and relevant negative effect with low/unclear reliability of data  
↑↑: statistically significant and relevant positive effect with high reliability of data  
↓↓: statistically significant and relevant negative effect with high reliability of data  
 $\leftrightarrow$ : no statistically significant or relevant difference  
∅: No data available.  
n.a.: not assessable

### Adjusted indirect comparison

Durvalumab + tremelimumab + platinum-based chemotherapy vs pembrolizumab via the bridge comparator platinum-based chemotherapy:

POSEIDON study: Durvalumab + tremelimumab + platinum-based chemotherapy vs platinum-based chemotherapy; open-label RCT

KEYNOTE-024 study: Pembrolizumab vs platinum-based chemotherapy; open-label RCT

KEYNOTE-042 study: Pembrolizumab vs platinum-based chemotherapy; open-label RCT

<sup>1</sup> Data from the dossier assessment of the IQWiG (A23-29 | A23-31) and from the addenda (A23-84 and G23-20), unless otherwise indicated.

## Mortality

Endpoint	Durvalumab + tremelimumab + platinum-based chemotherapy or Pembrolizumab		Platinum-based chemotherapy (bridge comparator)		Intervention vs control
	N	Median survival time in months [95% CI] <i>Patients with event n (%)</i>	N	Median survival time in months [95% CI] <i>Patients with event n (%)</i>	HR [95% CI] p value
<b>Overall survival</b>					
Tremelimumab + durvalumab + platinum-based chemotherapy vs platinum-based chemotherapy					
POSEIDON (data cut-off 12.03.2021)	101	n.d. <i>69 (68.3)</i>	97	n.d. <i>80 (82.5)</i>	0.65 [0.47; 0.89]; n.d.
Pembrolizumab vs platinum-based chemotherapy					
KEYNOTE-024 (data cut-off 09.05.2016)	154	n.r. <i>44 (28.6)</i>	151	n.r. [9.4; n.c.] <i>64 (42.4)</i>	0.60 [0.41; 0.89]; 0.010
KEYNOTE-042 (data cut-off 26.02.2018)	299	20.0 [15.4; 24.9] n.d.	300	12.2 [10.4; 14.2] n.d.	0.69 [0.56; 0.85]; < 0.001
Total					0.67 [0.56; 0.80] < 0.001 <sup>a</sup>
Adjusted indirect comparison via bridge comparators <sup>b</sup> : Tremelimumab + durvalumab + platinum-based chemotherapy vs pembrolizumab					0.97 [0.67; 1.41]; 0.873 <sup>c</sup>

## Morbidity

Endpoint	
Symptomatology (EORTC QLQ-C30, EORTC QLQ-LC13)	No suitable data for indirect comparison <sup>d</sup>
Health status (EQ-5D VAS)	No suitable data for indirect comparison <sup>d</sup>
Health status (PGIC)	No suitable data for indirect comparison <sup>d</sup>

## Health-related quality of life

Endpoint	
Functional scales (EORTC QLQ-C30)	No suitable data for indirect comparison <sup>d</sup>

## Side effects

Endpoint	Durvalumab + tremelimumab + platinum-based chemotherapy or Pembrolizumab		Platinum-based chemotherapy (bridge comparator)		Intervention vs control
	N	Median time to event in months [95% CI] <i>Patients with event n (%)</i>	N	Median time to event in months [95% CI] <i>Patients with event n (%)</i>	HR [95% CI] p value Absolute difference (AD) <sup>k</sup>
<b>Total adverse events (presented additionally)</b>					
Tremelimumab + durvalumab + platinum-based chemotherapy vs platinum-based chemotherapy					
POSEIDON <sup>e</sup> (data cut-off 12.03.2021)	99	0.1 [0.1; 0.3] <i>98 (99.0)</i>	93	0.2 [0.1; 0.3] <i>88 (94.6)</i>	–
Pembrolizumab vs platinum-based chemotherapy					
KEYNOTE-024 (data cut-off 09.05.2016)	154	1.1 [0.7; 1.7] <i>148 (96.1)</i>	150	0.6 [0.4; 0.9] <i>145 (96.7)</i>	–
KEYNOTE-042 (data cut-off 26.02.2018) <sup>f</sup>	299	n.d.	300	n.d.	n.d.
<b>Serious adverse events (SAE)</b>					
Tremelimumab + durvalumab + platinum-based chemotherapy vs platinum-based chemotherapy					
POSEIDON <sup>e</sup> (data cut-off 12.03.2021)	99	15.4 [7.2; n.c.] <i>47 (47.5)</i>	93	18.3 [6.8; n.c.] <i>37 (39.8)</i>	0.92 [0.59; 1.43] 0.697
Pembrolizumab vs platinum-based chemotherapy					
KEYNOTE-024 (data cut-off 09.05.2016)	154	54.1 [27.1; n.c.] <i>68 [44.2]</i>	150	65.4 [23.1; n.c.] <i>66 (44.0)</i>	1.00 [0.71; 1.41] 0.994
KEYNOTE-042 (data cut-off 26.02.2018) <sup>f</sup>	299	n.d.	300	n.d.	n.d.

Endpoint	Durvalumab + tremelimumab + platinum-based chemotherapy or Pembrolizumab		Platinum-based chemotherapy (bridge comparator)		Intervention vs control
	N	Median time to event in months [95% CI] <i>Patients with event n (%)</i>	N	Median time to event in months [95% CI] <i>Patients with event n (%)</i>	HR [95% CI] p value Absolute difference (AD) <sup>k</sup>
Adjusted indirect comparison via bridge comparators <sup>b</sup> : Tremelimumab + durvalumab + platinum-based chemotherapy vs pembrolizumab					0.92 [0.53; 1.61] 0.770 <sup>i</sup>
<b>Severe adverse events (CTCAE grade ≥ 3)</b>					
Tremelimumab + durvalumab + platinum-based chemotherapy vs platinum-based chemotherapy					
POSEIDON <sup>h</sup> (data cut-off 12.03.2021)	No suitable data for indirect comparison				
Pembrolizumab vs platinum-based chemotherapy					
KEYNOTE-024 (data cut-off 09.05.2016)	154	27.1 [18.1; 44.4] 82 (53.2)	150	5.9 [4.4; 9.0] 109 (72.7)	0.49 [0.36; 0.66]; < 0.001 AD: 21.2 months
KEYNOTE-042 (data cut-off 26.02.2018) <sup>f</sup>	299	n.d.	300	n.d.	n.d.
Adjusted indirect comparison via bridge comparators <sup>b</sup> : Tremelimumab + durvalumab + platinum-based chemotherapy vs pembrolizumab					– <sup>j</sup>
<b>Therapy discontinuation due to adverse events</b>					
Tremelimumab + durvalumab + platinum-based chemotherapy vs platinum-based chemotherapy					
POSEIDON <sup>e</sup> (data cut-off 12.03.2021)	99	n.r. [16.4; n.c.] 31 (31.3)	93	n.r. 16 (17.2)	1.31 [0.72; 2.50]; 0.385
Pembrolizumab vs platinum-based chemotherapy					
KEYNOTE-024 (data cut-off 09.05.2016)	154	n.r. 14 (9.1)	150	n.r. 21 (14.0)	0.60 [0.31; 1.19]; 0.144
KEYNOTE-042 (data cut-off 26.02.2018) <sup>e</sup>	299	n.d.	300	n.d.	n.d.

Endpoint	Durvalumab + tremelimumab + platinum-based chemotherapy or Pembrolizumab		Platinum-based chemotherapy (bridge comparator)		Intervention vs control
	N	Median time to event in months [95% CI] <i>Patients with event n (%)</i>	N	Median time to event in months [95% CI] <i>Patients with event n (%)</i>	HR [95% CI] p value Absolute difference (AD) <sup>k</sup>
Adjusted indirect comparison via bridge comparators <sup>b</sup> : Tremelimumab + durvalumab + platinum-based chemotherapy vs pembrolizumab					- <sup>g</sup>
<b>Specific adverse events</b>					
PRO-CTCAE	No suitable data for indirect comparison <sup>j</sup>				
Immune-mediated AEs	No suitable data for indirect comparison <sup>j</sup>				
<p>a Meta-analysis calculated using a model with fixed effects</p> <p>b Indirect comparison according to Bucher</p> <p>c IQWiG calculation (effect, CI, p value)</p> <p>d The data of the Poseidon study are not assessable due to asynchronous data collection time points of the PRO data and the resulting different representation of the patients' burden; PCIG was only collected in the POSEIDON study</p> <p>e No data are available for the relevant sub-population for the predefined final data cut-off from 12.03.2021. The data from the dossier are used since the information on the total population between the final predefined data cut-off and the data cut-off submitted by the pharmaceutical company in the dossier (25.10.2021) is not relevantly different or identical.</p> <p>f These analyses only cover just under 50% of the relevant sub-population and are only available separately by histology.</p> <p>g The requirement for certainty of results for carrying out an adjusted indirect comparison is not met</p> <p>h No data on the relevant sub-population are available for the final data cut-off from 12.03.2021. In relation to the total population, the number of patients with an event at the data cut-off from 25.10.2021 deviates relevantly from the number of patients in the predefined final data cut-off, which is why the data from the dossier are not used.</p> <p>i IQWiG calculation</p> <p>j Infeasible as no suitable data are available or none at all.</p> <p>k Indication of absolute difference (AD) only in case of statistically significant difference; own calculation</p> <p>Abbreviations used: CTCAE = Common Terminology Criteria for Adverse Events; ECOG-PS = European Cooperative Oncology Group Performance Status; EORTC = European Organisation for Research and Treatment of Cancer; HR = hazard ratio; n.d.= no data available; CI = confidence interval; N = number of patients evaluated; n = number of patients with (at least 1) event; n.c. = not calculable; n.r. = not reached; PD L1 = Programmed Cell Death-Ligand-1; PGIC = Patient Global Impression of Change; PRO-CTCAE = Patient-reported Outcome - CTCAE; QLQ-C30 = Quality of Life Questionnaire - Cancer 30; QLQ-LC13 = Quality of Life Questionnaire - Lung Cancer 13; RCT = randomised controlled trial; SAE = serious adverse event; AE = adverse event; VAS = visual analogue scale; vs = versus</p>					

- b) Adults with metastatic NSCLC with PD-L1 expression < 50% with no genomic EGFR or ALK tumour mutations, first-line therapy

No adequate data are available to allow an assessment of the additional benefit.

### Summary of results for relevant clinical endpoints

Endpoint category	Direction of effect/ risk of bias	Summary
Mortality	n.a.	There are no assessable data.
Morbidity	n.a.	There are no assessable data.
Health-related quality of life	n.a.	There are no assessable data.
Side effects	n.a.	There are no assessable data.
Explanations: ↑: statistically significant and relevant positive effect with low/unclear reliability of data ↓: statistically significant and relevant negative effect with low/unclear reliability of data ↑↑: statistically significant and relevant positive effect with high reliability of data ↓↓: statistically significant and relevant negative effect with high reliability of data ↔: no statistically significant or relevant difference ∅: No data available. n.a.: not assessable		

## 2. Number of patients or demarcation of patient groups eligible for treatment

- a) Adults with metastatic NSCLC with PD-L1 expression ≥ 50% with no genomic EGFR or ALK tumour mutations, first-line therapy

approx. 3,860 – 6,850 patients

- b) Adults with metastatic NSCLC with PD-L1 expression < 50% with no genomic EGFR or ALK tumour mutations, first-line therapy

Approx. 10,610 to 17,810 patients

## 3. Requirements for a quality-assured application

The requirements in the product information are to be taken into account. The European Medicines Agency (EMA) provides the contents of the product information (summary of product characteristics, SmPC) for Tremelimumab AstraZeneca (active ingredient: tremelimumab) at the following publicly accessible link (last access: 26 May 2023):

[https://www.ema.europa.eu/en/documents/product-information/tremelimumab-astrazeneca-epar-product-information\\_en.pdf](https://www.ema.europa.eu/en/documents/product-information/tremelimumab-astrazeneca-epar-product-information_en.pdf)

Treatment with tremelimumab should only be initiated and monitored by specialists in internal medicine, haematology and oncology who are experienced in the treatment of patients with non-small cell lung cancer, as well as specialists in internal medicine and



pulmonology or specialists in pulmonary medicine and other doctors from specialist groups participating in the Oncology Agreement.

In accordance with the European Medicines Agency (EMA) requirements regarding additional risk minimisation measures, the pharmaceutical company must provide training material that contains information for medical professionals and patients (incl. patient card).

The training material contains, in particular, information and warnings about immune-mediated adverse reactions.

#### 4. Treatment costs

##### Annual treatment costs:

The costs for the first year of treatment are shown for the cost representation in the resolution.

a) Adults with metastatic NSCLC with PD-L1 expression  $\geq$  50% with no genomic EGFR or ALK tumour mutations, first-line therapy

Designation of the therapy	Annual treatment costs/ patient
Medicinal product to be assessed:	
Tremelimumab	€ 25,261.80
+ Durvalumab	€ 23,505.96
<i>Total</i>	€ 48,767.76
<i>+ 4 cycles of platinum-based chemotherapy (carboplatin or cisplatin + pemetrexed, carboplatin or cisplatin + gemcitabine, nab-paclitaxel + carboplatin)</i>	
<i>+ carboplatin + pemetrexed</i>	
Carboplatin	€ 2,004.20
Pemetrexed	€ 4,352.32
<i>Total</i>	€ 6,356.52
<i>Tremelimumab + durvalumab + carboplatin + pemetrexed</i>	€ 55,124.28
Additionally required SHI costs	€ 27.34 - € 36.02
<i>+ cisplatin + pemetrexed</i>	
Cisplatin	€ 456.12
Pemetrexed	€ 4,352.32
<i>Total</i>	€ 4,808.44
<i>Tremelimumab + durvalumab + cisplatin + pemetrexed</i>	€ 53,576.20
Additionally required SHI costs	€ 162.02 - € 192.28
<i>+ carboplatin + gemcitabine</i>	
Carboplatin	€ 2,004.20

Designation of the therapy	Annual treatment costs/ patient
Gemcitabine	€ 1,852.00
<i>Total</i>	€ 3,856.20
<i>Tremelimumab + durvalumab + carboplatin + gemcitabine</i>	€ 52,623.96
<i>+ cisplatin + gemcitabine</i>	
Cisplatin	€ 456.12
Gemcitabine	€ 1,852.00
<i>Total</i>	€ 2,308.12
<i>Tremelimumab + durvalumab + cisplatin + gemcitabine</i>	€ 51,075.88
Additionally required SHI costs	€ 134.68 - € 156.26
<i>+ carboplatin + nab-paclitaxel</i>	
Carboplatin	€ 2,004.20
nab-paclitaxel	€ 9,780.48
<i>Total</i>	€ 11,784.68
<i>Tremelimumab + durvalumab + carboplatin + nab-paclitaxel</i>	€ 60,552.44
<i>Antibody maintenance treatment (without histology-based maintenance treatment with pemetrexed)</i>	
Single dose of tremelimumab	€ 6,315.45
+ durvalumab	€ 58,764.90
<i>Total</i>	€ 65,080.35
<i>Antibody maintenance treatment and histology-based maintenance treatment with pemetrexed</i>	
Single dose of tremelimumab	€ 6,315.45
+ durvalumab	€ 58,764.90
Pemetrexed	€ 10,880.80
<i>Total</i>	€ 75,961.15
Additionally required SHI costs	€ 74.33 - € 103.37
<i>Overall presentation: Tremelimumab + durvalumab + 4 cycles of platinum-based chemotherapy + maintenance treatment</i>	
<i>Tremelimumab + durvalumab + carboplatin + pemetrexed + maintenance treatment</i>	
Tremelimumab + durvalumab + carboplatin + pemetrexed (4 cycles)	€ 55,124.28
Additionally required SHI costs	€ 27.34 - € 36.02
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy) + pemetrexed	€ 75,961.15
Additionally required SHI costs	€ 74.33 - € 103.37

Designation of the therapy	Annual treatment costs/ patient
<i>Total</i>	€ 131,085.43
<i>Total additionally required SHI costs</i>	€ 101.67 - € 139.39
<i>Tremelimumab + durvalumab + cisplatin + pemetrexed + maintenance treatment</i>	
Tremelimumab + durvalumab + cisplatin + pemetrexed (4 cycles)	€ 53,576.20
Additionally required SHI costs	€ 162.02 - € 192.28
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy) + pemetrexed	€ 75,961.15
Additionally required SHI costs	€ 74.33 - € 103.37
<i>Total</i>	€ 129,537.35
<i>Total additionally required SHI costs</i>	€ 236.35 - € 295.65
<i>Tremelimumab + durvalumab + carboplatin + gemcitabine + maintenance treatment</i>	
Tremelimumab + durvalumab + carboplatin + gemcitabine (4 cycles)	€ 52,623.96
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy)	€ 65,080.35
<i>Total</i>	€ 117,704.31
<i>Tremelimumab + durvalumab + cisplatin + gemcitabine + maintenance treatment</i>	
Tremelimumab + durvalumab + cisplatin + gemcitabine (4 cycles)	€ 51,075.88
Additionally required SHI costs	€ 134.68 - € 156.26
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy)	€ 65,080.35
<i>Total</i>	€ 116,156.23
<i>Total additionally required SHI costs</i>	€ 134.68 - € 156.26
<i>Tremelimumab + durvalumab + carboplatin + nab-paclitaxel + maintenance treatment</i>	
Tremelimumab + durvalumab + carboplatin + nab-paclitaxel (4 cycles)	€ 60,552.44
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy)	€ 65,080.35
<i>Total</i>	€ 125,632.79
Appropriate comparator therapy:	
<i>Monotherapies</i>	
Atezolizumab	€ 64,877.81 - € 68,557.39
Cemiplimab	€ 80,879.55
Pembrolizumab	€ 93,515.26

Designation of the therapy	Annual treatment costs/ patient
<i>Nivolumab + ipilimumab + 2 cycles of platinum-based chemotherapy (only for patients with ECOG-PS 0-1)</i>	
Nivolumab	€ 73,035.63
+ ipilimumab	€ 54,832.10
Total	€ 127,867.73
<i>Carboplatin + paclitaxel</i>	
Carboplatin	€ 1,002.10
Paclitaxel	€ 1,911.38
Total	€ 2,913.48
<i>Nivolumab + ipilimumab + carboplatin + paclitaxel</i>	€ 130,781.21
Additionally required SHI costs	€ 64.01
<i>Carboplatin + pemetrexed</i>	
Carboplatin	€ 1,002.10
Pemetrexed	€ 2,176.16
Total	€ 3,178.26
<i>Nivolumab + ipilimumab + carboplatin + pemetrexed</i>	€ 131,045.99
Additionally required SHI costs	€ 34.93 - € 41.13
<i>Cisplatin + pemetrexed</i>	
Cisplatin	€ 228.06
Pemetrexed	€ 2,176.16
Total	€ 2,404.22
<i>Nivolumab + ipilimumab + cisplatin + pemetrexed</i>	130271.95
Additionally required SHI costs	€ 147.82 - € 164.81
<i>Atezolizumab + bevacizumab + paclitaxel + carboplatin (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>	
<i>Induction therapy (4 -6 cycles)</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 10,506.88 - € 15,760.32 (840 mg; 4 -6 cycles) or € 14,914.44 - € 22,371.66 (1,200 mg; 4 -6 cycles) or € 21,013.76 - € 31,520.64 (1,680 mg; 4 -6 cycles)
+ bevacizumab (7.5 mg/kg or 15.5 mg/kg)	€ 8,486.88 - € 12,730.32 (7.5 mg/kg; 4 -6 cycles) or € 16,858.80 - € 25,288.20

Designation of the therapy	Annual treatment costs/ patient
	(15 mg/kg; 4 -6 cycles)
+ paclitaxel	€ 3,822.76 - € 5,734.14
+ carboplatin	€ 2,004.20 - € 3,006.30
<i>Maintenance treatment</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 52,797.07 - € 58,050.51 (840 mg; 20.1 -22.1 cycles) or € 42,506.15 - € 49,963.37 (1,200 mg; 11.4 -13.4 cycles) or € 36,774.08 - € 47,280.96 (1,680 mg; 7 -9 cycles)
+ bevacizumab (840 mg or 1,200 mg or 1,680 mg)	€ 24,187.61 – € 28,431.05 (7.5 mg/kg; 11.4 - 13.4 cycles) or € 48,047.58 - € 56,476.98 (15 mg/kg; 11.4 - 13.4 cycles)
<i>Total</i> <i>(Cost range taking into account the number of induction cycles and atezolizumab as well as bevacizumab dosing regimens)</i>	<u>Combination with 7.5 mg/kg bevacizumab:</u> € 111,302.28 - € 114,215.76 (4 - 6 induction cycles with 840 mg atezolizumab) or € 107,622.70 - € 110,536.18 (4 - 6 induction cycles with 1,200 mg atezolizumab) or € 111,039.61 - € 113,953.09 (4 - 6 induction cycles with 1,680 mg atezolizumab)  or <u>Combination with 15 mg/kg bevacizumab:</u> € 147,720.13 - € 150,633.61 (4 - 6 induction cycles with 840 mg atezolizumab) € 144,040.55 - € 146,954.03 (4 - 6 induction cycles with 1,200 mg atezolizumab) or € 147,457.46 - € 150,370.94 (4 - 6 induction cycles with 1,680 mg atezolizumab)
<i>Additionally required SHI costs</i>	80.20 - 135.46
<i>Atezolizumab + carboplatin + nab-paclitaxel</i> <i>(only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>	

Designation of the therapy	Annual treatment costs/ patient
<i>Induction therapy (4 -6 cycles)</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 10,506.88 - € 15,760.32 (840 mg; 4 -6 cycles) or € 14,914.44 - € 22,371.66 (1,200 mg; 4 -6 cycles) or € 21,013.76 - € 31,520.64 (1,680 mg; 4 -6 cycles)
+ carboplatin	€ 2,004.20 - € 3,006.3
+ nab-paclitaxel	€ 9,780.48 - € 14,670.72
<i>Maintenance treatment</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 52,797.07 - € 58,050.51 (840 mg; 20.1 -22.1 cycles) or € 42,506.15 - € 49,963.37 (1,200 mg; 11.4 -13.4 cycles) or € 36,774.08 - € 47,280.96 (1,680 mg; 7 -9 cycles)
<i>Total</i> <i>(Cost range taking into account the number of induction cycles and atezolizumab as well as bevacizumab dosing regimens)</i>	€ 80,342.07 - € 86,234.41 (4 - 6 induction cycles with 840 mg atezolizumab) or € 76,662.49 - € 82,554.83 (4 - 6 induction cycles with 1,200 mg atezolizumab) or € 80,079.40 - € 85,971.74 (4 - 6 induction cycles with 1,680 mg atezolizumab)
<i>Pembrolizumab + carboplatin + (nab)-paclitaxel</i> <i>(only for patients with ECOG-PS 0-1 and squamous NSCLC)</i>	
<i>Pembrolizumab + carboplatin + paclitaxel</i>	
Pembrolizumab	€ 93,515.26
Carboplatin	€ 8,718.27
Paclitaxel	€ 16,629.01
<i>Total</i>	€ 118,862.53
Additionally required SHI costs	€ 254.58
<i>Pembrolizumab + carboplatin + nab-paclitaxel</i>	
Pembrolizumab	€ 93,515.26
Carboplatin	€ 8,718.27
nab-paclitaxel	€ 42,545.09

Designation of the therapy	Annual treatment costs/ patient
<i>Total</i>	€ 144,778.61
<i>Pembrolizumab + pemetrexed + platinum-containing chemotherapy (only for patients with ECOG-PS 0-1 and non-squamous NSCLC)</i>	
<i>Pembrolizumab + pemetrexed + cisplatin</i>	
Pembrolizumab	€ 93,515.26
Pemetrexed	€ 18,932.59
Cisplatin	€ 1,984.12
<i>Total</i>	€ 114,431.97
Additionally required SHI costs	€ 445.50 - € 576.26
<i>Pembrolizumab + pemetrexed + carboplatin</i>	
Pembrolizumab	€ 93,515.26
Pemetrexed	€ 18,932.59
Carboplatin	€ 8,718.27
<i>Total</i>	€ 121,166.12
Additionally required SHI costs	€ 116.92 - € 154.64

Costs after deduction of statutory rebates (LAUER-TAXE®) as last revised: 15 September 2023

Costs for additionally required SHI services: not applicable

Other SHI services:

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
<b>Medicinal product to be assessed</b>					
<i>Induction</i>					
Tremelimumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0	€ 400
+ durvalumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0	€ 400
+ carboplatin	Surcharge for production of a parenteral	€ 100	1	4.0	€ 400

	preparation containing cytostatic agents				
+ cisplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0	€ 400
+ gemcitabine	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	2	4.0	€ 800
+ pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0	€ 400
+ nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	3.0	€ 300
<i>Antibody maintenance treatment and histology-based maintenance treatment with pemetrexed</i>					
Tremelimumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	10.0	€ 1,000
+ durvalumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	1.0	€ 100
+ pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	10.0	€ 1,000
<b>Appropriate comparator therapy</b>					
<i>Monotherapies</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	26.1	€ 2,610
				or	
				17.4	€ 1,740
				or	
				13.0	€ 1,300
Cemiplimab	Surcharge for the preparation of a	€ 100	1	17.4	€ 1,740



	parenteral solution containing monoclonal antibodies				
Pembrolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	8.7	€ 870
Pembrolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
<i>Nivolumab + ipilimumab + 2 cycles of platinum-based chemotherapy (only for patients with ECOG-PS 0-1)</i>					
Nivolumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
Ipilimumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	8.7	€ 870
Cisplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
Pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
<i>Atezolizumab + bevacizumab + paclitaxel + carboplatin (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>					
<i>Induction therapy</i>					

Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0 - 6.0	€ 400 - € 600
Bevacizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0 - 6.0	€ 400 - € 600
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0 - 6.0	€ 400 - € 600
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0 - 6.0	€ 400 - € 600
<i>Maintenance treatment</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	20.1 - 22.1	€ 2,010 - € 2,210
				or	
				11.4 - 13.4	€ 1,140 - € 1,340
				or	
				7.0 - 9.0	€ 700 - € 900
Bevacizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	11.4 - 13.4	€ 1,140 - € 1,340
<i>Atezolizumab + carboplatin + nab-paclitaxel (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>					
<i>Induction therapy</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0 - 6.0	€ 400 - € 600
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0 - 6.0	€ 400 - € 600

nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	3	4.0 - 6.0	€ 1,200 - € 1,800
<i>Maintenance treatment</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	20.1 - 22.1	€ 2,010 - € 2,210
				or	
				11.4 - 13.4	€ 1,140 - € 1,340
				or	
				7.0 - 9.0	€ 700 - € 900
<i>Pembrolizumab + carboplatin + (nab)-paclitaxel (only for patients with ECOG-PS 0-1 and squamous NSCLC)</i>					
Pembrolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
				or	
				8.7	€ 870
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	3	17.4	€ 5,220
<i>Pembrolizumab + pemetrexed + platinum-containing chemotherapy (only for patients with ECOG-PS 0-1 and non-squamous NSCLC)</i>					
Pembrolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
				or	
				8.7	€ 870
Pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Cisplatin	Surcharge for production of a parenteral	€ 100	1	17.4	€ 1,740

	preparation containing cytostatic agents				
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740

b) Adults with metastatic NSCLC with PD-L1 expression < 50% with no genomic EGFR or ALK tumour mutations, first-line therapy

Designation of the therapy	Annual treatment costs/ patient
Medicinal product to be assessed:	
Tremelimumab	€ 25,261.80
+ Durvalumab	€ 23,505.96
<i>Total</i>	€ 48,767.76
<i>+ 4 cycles of platinum-based chemotherapy (carboplatin or cisplatin + pemetrexed, carboplatin or cisplatin + gemcitabine, nab-paclitaxel + carboplatin)</i>	
<i>+ carboplatin + pemetrexed</i>	
Carboplatin	€ 2,004.20
Pemetrexed	€ 4,352.32
<i>Total</i>	€ 6,356.52
<i>Tremelimumab + durvalumab + carboplatin + pemetrexed</i>	€ 55,124.28
Additionally required SHI costs	€ 27.34 - € 36.02
<i>+ cisplatin + pemetrexed</i>	
Cisplatin	€ 456.12
Pemetrexed	€ 4,352.32
<i>Total</i>	€ 4,808.44
<i>Tremelimumab + durvalumab + cisplatin + pemetrexed</i>	€ 53,576.20
Additionally required SHI costs	€ 162.02 - € 192.28
<i>+ carboplatin + gemcitabine</i>	
Carboplatin	€ 2,004.20
Gemcitabine	€ 1,852.00
<i>Total</i>	€ 3,856.20
<i>Tremelimumab + durvalumab + carboplatin + gemcitabine</i>	€ 52,623.96
<i>+ cisplatin + gemcitabine</i>	
Cisplatin	€ 456.12

Designation of the therapy	Annual treatment costs/ patient
Gemcitabine	€ 1,852.00
<i>Total</i>	€ 2,308.12
<i>Tremelimumab + durvalumab + cisplatin + gemcitabine</i>	€ 51,075.88
Additionally required SHI costs	€ 134.68 - € 156.26
<i>+ carboplatin + nab-paclitaxel</i>	
Carboplatin	€ 2,004.20
nab-paclitaxel	€ 9,780.48
<i>Total</i>	€ 11,784.68
<i>Tremelimumab + durvalumab + carboplatin + nab-paclitaxel</i>	€ 60,552.44
<i>Antibody maintenance treatment (without histology-based maintenance treatment with pemetrexed)</i>	
Single dose of tremelimumab	€ 6,315.45
+ durvalumab	€ 58,764.90
<i>Total</i>	€ 65,080.35
<i>Antibody maintenance treatment and histology-based maintenance treatment with pemetrexed</i>	
Single dose of tremelimumab	€ 6,315.45
+ durvalumab	€ 58,764.90
Pemetrexed	€ 10,880.80
<i>Total</i>	€ 75,961.15
Additionally required SHI costs	€ 74.33 - € 103.37
<i>Overall presentation: Tremelimumab + durvalumab + 4 cycles of platinum-based chemotherapy + maintenance treatment</i>	
<i>Tremelimumab + durvalumab + carboplatin + pemetrexed + maintenance treatment</i>	
Tremelimumab + durvalumab + carboplatin + pemetrexed (4 cycles)	€ 55,124.28
Additionally required SHI costs	€ 27.34 - € 36.02
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy) + pemetrexed	€ 75,961.15
Additionally required SHI costs	€ 74.33 - € 103.37
<i>Total</i>	€ 131,085.43
<i>Total additionally required SHI costs</i>	€ 101.67 - € 139.39
<i>Tremelimumab + durvalumab + cisplatin + pemetrexed + maintenance treatment</i>	

Designation of the therapy	Annual treatment costs/ patient
Durvalumab + tremelimumab + cisplatin + pemetrexed (4 cycles)	€ 53,576.20
Additionally required SHI costs	€ 162.02 - € 192.28
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy) + pemetrexed	€ 75,961.15
Additionally required SHI costs	€ 74.33 - € 103.37
<i>Total</i>	€ 129,537.35
<i>Total additionally required SHI costs</i>	€ 236.35 - € 295.65
<i>Tremelimumab + durvalumab + carboplatin + gemcitabine + maintenance treatment</i>	
Tremelimumab + durvalumab + carboplatin + gemcitabine (4 cycles)	€ 52,623.96
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy)	€ 65,080.35
<i>Total</i>	€ 117,704.31
<i>Tremelimumab + durvalumab + cisplatin + gemcitabine + maintenance treatment</i>	
Tremelimumab + durvalumab + cisplatin + gemcitabine (4 cycles)	€ 51,075.88
Additionally required SHI costs	€ 134.68 - € 156.26
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy)	€ 65,080.35
<i>Total</i>	€ 116,156.23
<i>Total additionally required SHI costs</i>	€ 134.68 - € 156.26
<i>Tremelimumab + durvalumab + carboplatin + nab-paclitaxel + maintenance treatment</i>	
Tremelimumab + durvalumab + carboplatin + nab-paclitaxel (4 cycles)	€ 60,552.44
Maintenance treatment (Single dose of tremelimumab + durvalumab monotherapy)	€ 65,080.35
<i>Total</i>	€ 125,632.79
Appropriate comparator therapy:	
<i>Monotherapies</i>	
Atezolizumab	€ 64,877.81 - € 68,557.39
<i>Nivolumab + ipilimumab + 2 cycles of platinum-based chemotherapy (only for patients with ECOG-PS 0-1)</i>	
Nivolumab	€ 73035.63
+ ipilimumab	€ 54,832.10

Designation of the therapy	Annual treatment costs/ patient
Total	€ 127,867.73
<i>Carboplatin + paclitaxel</i>	
Carboplatin	€ 1,002.10
Paclitaxel	€ 1,911.38
Total	€ 2,913.48
<i>Nivolumab + ipilimumab + carboplatin + paclitaxel</i>	
	€ 130,781.21
Additionally required SHI costs	€ 64.01
<i>Carboplatin + pemetrexed</i>	
Carboplatin	€ 1,002.10
Pemetrexed	€ 2,176.16
Total	€ 3,178.26
<i>Nivolumab + ipilimumab + carboplatin + pemetrexed</i>	
	€ 131,045.99
Additionally required SHI costs	€ 34.93 - € 41.13
<i>Cisplatin + pemetrexed</i>	
Cisplatin	€ 228.06
Pemetrexed	€ 2,176.16
Total	€ 2,404.22
<i>Nivolumab + ipilimumab + cisplatin + pemetrexed</i>	
	€ 130,271.95
Additionally required SHI costs	€ 147.82 - € 164.81
<i>Atezolizumab + bevacizumab + paclitaxel + carboplatin (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>	
<i>Induction therapy (4 -6 cycles)</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 10,506.88 - € 15,760.32 (840 mg; 4 -6 cycles) or € 14,914.44 - € 22,371.66 (1,200 mg; 4 -6 cycles) or € 21,013.76 - € 31,520.64 (1,680 mg; 4 -6 cycles)
+ bevacizumab (7.5 mg/kg or 15.5 mg/kg)	€ 8,486.88 - € 12,730.32 (7.5 mg/kg; 4 -6 cycles) or € 16,858.80 - € 25,288.20 (15 mg/kg; 4 -6 cycles)
+ paclitaxel	€ 3,822.76 - € 5,734.14
+ carboplatin	€ 2,004.20 - € 3,006.30

Designation of the therapy	Annual treatment costs/ patient
<i>Maintenance treatment</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 52,797.07 - € 58,050.51 (840 mg; 20.1 -22.1 cycles) or € 42,506.15 - € 49,963.37 (1,200 mg; 11.4 -13.4 cycles) or € 36,774.08 - € 47,280.96 (1,680 mg; 7 -9 cycles)
+ bevacizumab (840 mg or 1,200 mg or 1,680 mg)	€ 24,187.61 – € 28,431.05 (7.5 mg/kg; 11.4 - 13.4 cycles) or € 48,047.58 - € 56,476.98 (15 mg/kg; 11.4 - 13.4 cycles)
<i>Total</i> <i>(Cost range taking into account the number of induction cycles and atezolizumab as well as bevacizumab dosing regimens)</i>	<u>Combination with 7.5 mg/kg bevacizumab:</u> € 111,302.28 - € 114,215.76 (4 - 6 induction cycles with 840 mg atezolizumab) or € 107,622.70 - € 110,536.18 (4 - 6 induction cycles with 1,200 mg atezolizumab) or € 111,039.61 - € 113,953.09 (4 - 6 induction cycles with 1,680 mg atezolizumab)  or <u>Combination with 15 mg/kg bevacizumab:</u> € 147,720.13 - € 150,633.61 (4 - 6 induction cycles with 840 mg atezolizumab) € 144,040.55 - € 146,954.03 (4 - 6 induction cycles with 1,200 mg atezolizumab) or € 147,457.46 - € 150,370.94 (4 - 6 induction cycles with 1,680 mg atezolizumab)
<i>Additionally required SHI costs</i>	80.20 - 135.46
<i>Atezolizumab + carboplatin + nab-paclitaxel (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>	



Designation of the therapy	Annual treatment costs/ patient
<i>Induction therapy (4 -6 cycles)</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 10,506.88 - € 15,760.32 (840 mg; 4 -6 cycles) or € 14,914.44 - € 22,371.66 (1,200 mg; 4 -6 cycles) or € 21,013.76 - € 31,520.64 (1,680 mg; 4 -6 cycles)
+ carboplatin	€ 2,004.20 - € 3,006.3
+ nab-paclitaxel	€ 9,780.48 - € 14,670.72
<i>Maintenance treatment</i>	
Atezolizumab (840 mg or 1,200 mg or 1,680 mg)	€ 52,797.07 - € 58,050.51 (840 mg; 20.1 -22.1 cycles) or € 42,506.15 - € 49,963.37 (1,200 mg; 11.4 -13.4 cycles) or € 36,774.08 - € 47,280.96 (1,680 mg; 7 -9 cycles)
<i>Total</i> (Cost range taking into account the number of induction cycles and atezolizumab as well as bevacizumab dosing regimens)	€ 80,342.07 - € 86,234.41 (4 - 6 induction cycles with 840 mg atezolizumab) or € 76,662.49 - € 82,554.83 (4 - 6 induction cycles with 1,200 mg atezolizumab) or € 80,079.40 - € 85,971.74 (4 - 6 induction cycles with 1,680 mg atezolizumab)
<i>Pembrolizumab + carboplatin + (nab)-paclitaxel</i> (only for patients with ECOG-PS 0-1 and squamous NSCLC)	
<i>Pembrolizumab + carboplatin + paclitaxel</i>	
Pembrolizumab	€ 93,515.26
Carboplatin	€ 8,718.27
Paclitaxel	€ 16,629.01
<i>Total</i>	€ 118,862.53
Additionally required SHI costs	€ 254.58
<i>Pembrolizumab + carboplatin + nab-paclitaxel</i>	
Pembrolizumab	€ 93,515.26

Designation of the therapy	Annual treatment costs/ patient
Carboplatin	€ 8,718.27
nab-paclitaxel	€ 42,545.09
<i>Total</i>	€ 144,778.61
<i>Pembrolizumab + pemetrexed + platinum-containing chemotherapy (only for patients with ECOG-PS 0-1 and non-squamous NSCLC)</i>	
<i>Pembrolizumab + pemetrexed + cisplatin</i>	
Pembrolizumab	€ 93,515.26
Pemetrexed	€ 18,932.59
Cisplatin	€ 1,984.12
<i>Total</i>	€ 114,431.97
Additionally required SHI costs	€ 445.50 - € 576.26
<i>Pembrolizumab + pemetrexed + carboplatin</i>	
Pembrolizumab	€ 93,515.26
Pemetrexed	€ 18,932.59
Carboplatin	€ 8,718.27
<i>Total</i>	€ 121,166.12
Additionally required SHI costs	€ 116.92 - € 154.64
<i>Carboplatin + nab-paclitaxel (only for patients with ECOG-PS 2)</i>	
Carboplatin	€ 8,718.27
nab-paclitaxel	€ 42,545.09
<i>Total</i>	€ 51,263.36
<i>Carboplatin + third-generation cytostatic (vinorelbine or gemcitabine or docetaxel or paclitaxel or pemetrexed) cf. Annex VI to Section K of the Pharmaceuticals Directive (only for patients with ECOG-PS 2)</i>	
<i>Carboplatin + vinorelbine</i>	
Carboplatin	€ 8,718.27
Vinorelbine	€ 4,717.11 - € 5,686.60
<i>Total</i>	€ 13,435.38 - € 14,404.87
<i>Carboplatin + gemcitabine</i>	
Carboplatin	€ 8,718.27
Gemcitabine	€ 8,056.20
<i>Total</i>	€ 16,774.47
<i>Carboplatin + docetaxel</i>	

Designation of the therapy	Annual treatment costs/ patient
Carboplatin	€ 8,718.27
Docetaxel	€ 8,523.22
<i>Total</i>	€ 17,241.49
<i>Carboplatin + paclitaxel</i>	
Carboplatin	€ 8,718.27
Paclitaxel	€ 16,629.01
<i>Total</i>	€ 25,347.28
Additionally required SHI costs	€ 254.58
<i>Carboplatin + pemetrexed</i>	
Carboplatin	€ 8,718.27
Pemetrexed	€ 18,932.59
<i>Total</i>	€ 27,650.86
Additionally required SHI costs	€ 116.92 - € 154.64

Costs after deduction of statutory rebates (LAUER-TAXE®) as last revised: 15 September 2023

Costs for additionally required SHI services: not applicable

Other SHI services:

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
<b>Medicinal product to be assessed</b>					
<i>Induction</i>					
Tremelimumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0	€ 400
+ durvalumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0	€ 400
+ carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0	€ 400

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
+ cisplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0	€ 400
+ gemcitabine	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	2	4.0	€ 800
+ pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0	€ 400
+ nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	3.0	€ 300
<i>Antibody maintenance treatment and histology-based maintenance treatment with pemetrexed</i>					
Tremelimumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	10.0	€ 1,000
+ durvalumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	1.0	€ 100
+ pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	10.0	€ 1,000
<b>Appropriate comparator therapy</b>					
<i>Monotherapy</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	26.1	€ 2,610
				or	
				17.4	€ 1,740
				or	
				13.0	€ 1,300
<i>Nivolumab + ipilimumab + 2 cycles of platinum-based chemotherapy</i>					

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
<i>(only for patients with ECOG-PS 0-1)</i>					
Nivolumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
Ipilimumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	8.7	€ 870
Cisplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
Pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	2.0	€ 200
<i>Atezolizumab + bevacizumab + paclitaxel + carboplatin (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>					
<i>Induction therapy</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0 - 6.0	€ 400 - € 600
Bevacizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0 - 6.0	€ 400 - € 600

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0 - 6.0	€ 400 - € 600
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0 - 6.0	€ 400 - € 600
<i>Maintenance treatment</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	20.1 - 22.1	€ 2,010 - € 2,210
				or	
				11.4 - 13.4	€ 1,140 - € 1,340
				or	
				7.0 - 9.0	€ 700 - € 900
Bevacizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	11.4 - 13.4	€ 1,140 - € 1,340
<i>Atezolizumab + carboplatin + nab-paclitaxel (only for patients with ECOG PS 0-1 and non-squamous NSCLC)</i>					
<i>Induction therapy</i>					
Atezolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	4.0 - 6.0	€ 400 - € 600
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	4.0 - 6.0	€ 400 - € 600
nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	3	4.0 - 6.0	€ 1,200 - € 1,800
<i>Maintenance treatment</i>					
Atezolizumab	Surcharge for the preparation of a	€ 100	1	20.1 - 22.1	€ 2,010 - € 2,210
				or	

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
	parenteral solution containing monoclonal antibodies			11.4 - 13.4	€ 1,140 - € 1,340
				or	
				7.0 - 9.0	€ 700 - € 900
<i>Pembrolizumab + carboplatin + (nab)-paclitaxel (only for patients with ECOG-PS 0-1 and squamous NSCLC)</i>					
Pembrolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
				or	
				8.7	€ 870
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	3	17.4	€ 5,220
<i>Pembrolizumab + pemetrexed + platinum-containing chemotherapy (only for patients with ECOG-PS 0-1 and non-squamous NSCLC)</i>					
Pembrolizumab	Surcharge for the preparation of a parenteral solution containing monoclonal antibodies	€ 100	1	17.4	€ 1,740
				or	
				8.7	€ 870
Pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Cisplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740

Designation of the therapy	Type of service	Costs/ unit	Number / cycle	Number/ patient/ year	Costs/ patient/ year
<i>Carboplatin + nab-paclitaxel (only for patients with ECOG-PS 2)</i>					
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
nab-paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	3	17.4	€ 5,220
<i>Carboplatin + third-generation cytostatic (vinorelbine or gemcitabine or docetaxel or paclitaxel or pemetrexed) cf. Annex VI to Section K of the Pharmaceuticals Directive (only for patients with ECOG-PS 2)</i>					
Carboplatin	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Vinorelbine	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	2	17.4	€ 3,480
Gemcitabine	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	2	17.4	€ 3,480
Docetaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Paclitaxel	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740
Pemetrexed	Surcharge for production of a parenteral preparation containing cytostatic agents	€ 100	1	17.4	€ 1,740



**5. Designation of medicinal products with new active ingredients according to Section 35a, paragraph 3, sentence 4 SGB V that can be used in a combination therapy with the assessed medicinal product**

In the context of the designation of medicinal products with new active ingredients pursuant to Section 35a, paragraph 3, sentence 4 SGB V, the following findings are made:

a) Adults with metastatic NSCLC with PD-L1 expression  $\geq$  50% with no genomic EGFR or ALK tumour mutations, first-line therapy

The following medicinal products with new active ingredients that can be used in a combination therapy with tremelimumab in the therapeutic indication of the resolution on the basis of the marketing authorisation under Medicinal Products Act are named (active ingredient and invented names) in accordance with Section 35a, paragraph 3, sentence 4 SGB V:

Durvalumab (Imfinzi)

b) Adults with metastatic NSCLC with PD-L1 expression  $<$  50% with no genomic EGFR or ALK tumour mutations, first-line therapy

The following medicinal products with new active ingredients that can be used in a combination therapy with tremelimumab in the therapeutic indication of the resolution on the basis of the marketing authorisation under Medicinal Products Act are named (active ingredient and invented names) in accordance with Section 35a, paragraph 3, sentence 4 SGB V:

Durvalumab (Imfinzi)

The designation of combinations exclusively serves the implementation of the combination discount according to Section 130e SGB V between health insurance funds and pharmaceutical companies. The findings made neither restrict the scope of treatment required to fulfil the medical treatment mandate, nor do they make statements about expediency or economic feasibility.