

Duvelisib (follicular lymphoma, after \geq 2 prior therapies)

Resolution of:21 July 2022Valid until: unlimitedEntry into force on:21 July 2022Federal Gazette, BAnz AT 23 08 2022 B2

Therapeutic indication (according to the marketing authorisation of 19 May 2021):

Copiktra monotherapy is indicated for the treatment of adult patients with:

- relapsed or refractory chronic lymphocytic leukaemia (CLL) after at least two prior therapies.
- follicular lymphoma (FL) that is refractory to at least two prior systemic therapies.

Therapeutic indication of the resolution (resolution of 21.07.2022):

Copiktra monotherapy is indicated for the treatment of adult patients with follicular lymphoma (FL) that is refractory to at least two prior systemic therapies.

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

Adults with follicular lymphoma that is refractory to at least two prior systemic therapies

Appropriate comparator therapy:

Patient-individual therapy taking into account prior therapy, course of the disease and general condition

Extent and probability of the additional benefit of duvelisib compared to the appropriate comparator therapy:

An additional benefit is not proven.

Study results according to endpoints:1

Adults with follicular lymphoma that is refractory to at least two prior systemic therapies

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

¹ Data from the dossier assessment of the Institute for Quality and Efficiency in Health Care (IQWiG) (A22-14) unless otherwise indicated.

Summary of results for relevant clinical endpoints

| Endpoint category | Direction of effect/ | Summary | | | |
|--|------------------------------------|--------------------------------------|--|--|--|
| | risk of bias | | | | |
| Mortality | n.a. | There are no assessable data. | | | |
| Morbidity | n.a. | There are no assessable data. | | | |
| Health-related quality | n.a. | There are no assessable data. | | | |
| of life | | | | | |
| Side effects | n.a. There are no assessable data. | | | | |
| Explanations: | | | | | |
| 个: statistically significant a | nd relevant positive effect | with low/unclear reliability of data | | | |
| ↓: statistically significant and relevant negative effect with low/unclear reliability of data | | | | | |
| $\uparrow\uparrow$: statistically significant and relevant positive effect with high reliability of data | | | | | |
| $\downarrow \downarrow$: statistically significant and relevant negative effect with high reliability of data | | | | | |
| ↔: no statistically significant or relevant difference | | | | | |
| arnothing: There are no usable data for the benefit assessment. | | | | | |
| n.a.: not assessable | | | | | |

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

Adults with follicular lymphoma that is refractory to at least two prior systemic therapies

approx. 380 – 5,170 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 Copiktra (active ingredient: duvelisib) at the following publicly accessible link (last access: 7 April 2022):

https://www.ema.europa.eu/en/documents/product-information/copiktra-epar-productinformation_en.pdf

Treatment with duvelisib should only be initiated and monitored by specialists in internal medicine, haematology and oncology experienced in the treatment of adults with follicular lymphoma.

4. Treatment costs

Annual treatment costs:

The annual treatment costs shown refer to the first year of treatment.

Adults with follicular lymphoma that is refractory to at least two prior systemic therapies

| Designation of the therapy Annual treatment costs/ patient Medicinal product to be assessed: | | | | | |
|---|--|--|--|--|--|
| € 68,451.58 ² | | | | | |
| | | | | | |
| Appropriate comparator therapy: | | | | | |
| Patient-individual therapy ^a Bendamustine + rituximab | | | | | |
| | | | | | |
| € 6,143.00 | | | | | |
| € 24,768.89 | | | | | |
| € 30,911.89 | | | | | |
| € 67,05 | | | | | |
| | | | | | |
| € 6,143.00 | | | | | |
| € 38,365.91 | | | | | |
| € 44,508.91 | | | | | |
| € 11.40 | | | | | |
| vincristine, prednisone) + rituximab | | | | | |
| € 192.94 | | | | | |
| € 1,703.88 | | | | | |
| € 207.66 | | | | | |
| € 41.12 | | | | | |
| € 2,145.60 | | | | | |
| € 27,142.15 | | | | | |
| € 29,287.75 | | | | | |
| € 73.13 | | | | | |
| CVP (cyclophosphamide, vincristine, prednisolone) + rituximab | | | | | |
| € 288.96 | | | | | |
| | | | | | |

² Duvelisib is currently not sold in Germany. LAUER-TAXE® last revised: 15.April 2022.

³ Instead of prednisone, the comparable and less expensive prednisolone was presented due to the principle of economic efficiency.

| Designation of the therapy | Annual treatment costs/ patient | | |
|--|---------------------------------|--|--|
| Vincristine | € 276.88 | | |
| Prednisolone | € 56.08 | | |
| CVP total | € 621.92 | | |
| Rituximab | € 27,142.15 | | |
| CVP + rituximab | € 27,764.07 | | |
| Additionally required SHI costs | € 73.13 | | |
| FCM (fludarabine, cyclophosphamide, m | nitoxantrone) + rituximab | | |
| Fludarabine | € 1,558.92 - € 2,598.20 | | |
| Cyclophosphamide | € 147.00 - € 294.00 | | |
| Mitoxantrone | € 892.52 - € 1,785.04 | | |
| FCM total | € 2,589.44 - € 4,677.24 | | |
| Rituximab | € 10,856.86 - € 21,713.72 | | |
| FCM + rituximab | € 13,455.30 - € 26,390.96 | | |
| Additionally required SHI costs | € 36.67 - € 60.98 | | |
| Chlorambucil + rituximab | | | |
| Chlorambucil | € 166.85 | | |
| Rituximab | € 19,968.67 | | |
| Total | € 20,135.52 | | |
| Additionally required SHI costs | € 48.83 | | |
| Cyclophosphamide + rituximab | | | |
| Cyclophosphamide | € 288.96 | | |
| Rituximab | € 27,142.15 | | |
| Total | € 27,431.11 | | |
| Additionally required SHI costs | € 73.13 | | |
| MCP (mitoxantrone, chlorambucil, pred | nisone) + rituximab | | |
| Mitoxantrone | € 2,677.56 - € 3,570.08 | | |
| Chlorambucil | € 200.22 - € 266.96 | | |
| Prednisolone ³ | € 28.04 | | |
| Rituximab | € 16,285.29 - € 21,713.72 | | |
| MCP total | € 2,905.82 - € 3,865.08 | | |
| MCP + rituximab | € 19,191.11 - € 25,578.80 | | |
| Additionally required SHI costs€ 48.83 - € 60.98 | | | |
| Lenalidomide + rituximab | | | |

| Designation of the therapy | Annual treatment costs/ patient | | | | | |
|---|--|--|--|--|--|--|
| Lenalidomide | € 2,219.16 | | | | | |
| Rituximab | € 21,713.72 | | | | | |
| Total | € 23,932.88 | | | | | |
| Additionally required SHI costs | € 60.98 | | | | | |
| Rituximab monotherapy | | | | | | |
| Rituximab € 21,713.72 | | | | | | |
| Additionally required SHI costs | € 60.98 | | | | | |
| Yttrium-90 radiolabelled ibritumomab | tiuxetan pretreated with rituximab | | | | | |
| Ibritumomab tiuxetan | € 13,865.50 | | | | | |
| Yttrium-90 | incalculable | | | | | |
| Rituximab | € 3,382.70 | | | | | |
| Total | incalculable | | | | | |
| Additionally required SHI costs € 48,51 | | | | | | |
| Idelalisib monotherapy | | | | | | |
| Idelalisib € 52,043.65 | | | | | | |
| vincristine, prednisone) + obinutuzuma obinutuzumab, FM (fludarabine + mit carboplatin, etoposide) + rituximab/ mitoxantrone) + obinutuzumab, MCP (mit DHAP (dexamethasone, Ara-C/cytarabi comparators for the present benefit a | f active ingredients CHOP (cyclophosphamide, doxorubicin, b, CVP (cyclophosphamide, vincristine, prednisolone) + oxantrone) + rituximab/obinutuzumab, ICE (ifosfamide, obinutuzumab,—FCM (fludarabine, cyclophosphamide, coxantrone, chlorambucil, prednisone) + obinutuzumab and ne, cisplatin) + rituximab/obinutuzumab are suitable ssessment in the context of patient-individual therapy. | | | | | |

However, these active ingredients or combinations of active ingredients are not approved in the

present therapeutic indication, and therefore, no costs are presented for these active ingredients or combinations of active ingredients.

Costs after deduction of statutory rebates (LAUER-TAXE®) as last revised: 1 July 2022)

Other SHI services:

| Designation of the therapy | Type of service | Costs/ unit | Number/ cycle | Number/ patient/ year | Costs/ patient/ year | | |
|--|---|----------------|------------------------------|-----------------------------|----------------------------|--|--|
| Medicinal product to | Medicinal product to be assessed: Duvelisib | | | | | | |
| Incalculable. | | | | | | | |
| Appropriate compar | ator therapy: | | | | | | |
| Bendamustine | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 2 | 12 | €972 | | |
| Cyclophosphamide (in combination with rituximab) | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 1 | 8 | € 648 | | |
| Obinutuzumab (in combination with bendamustine) | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | Cycle 1: 3, cycle 2– 9: 1 | 11 | € 781 | | |
| Rituximab (in combination with chlorambucil) | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 1 | 6 | € 426 | | |
| Rituximab (in combination with cyclophosphamide) | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 1 | 10 | €710 | | |

| Rituximab (in combination with lenalidomide and rituximab as monotherapy) Rituximab (in | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies Surcharge for the | €71 | Cycle 1: 4 Cycle 2 - 5: 1 | 8 | € 568 € 639 |
|--|--|-----|------------------------------|----|----------------|
| combination with MCP and in combination with bendamustine) | preparation of a parenteral solution containing monoclonal antibodies | | 1 | | |
| СНОР | | | | | |
| Cyclophosphamide | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 1 | 6 | € 486 |
| Doxorubicin | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 1 | 6 | € 486 |
| Vincristine | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 1 | 6 | € 486 |
| Rituximab (in combination with CHOP) | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 1 | 10 | €710 |
| CVP | | | | | |

| Cyclophosphamide | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 1 | 8 | € 648 |
|---|---|-----|---|---------|--------------------|
| Vincristine | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 1 | 8 | € 648 |
| Rituximab (in combination with CVP) | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 1 | 10 | € 710 |
| FCM | | | | | |
| Fludarabine | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 3 | 4 - 8 | € 243 - € 648 |
| Cyclophosphamide | Surcharge for production of a parenteral preparation containing cytostatic agents | €81 | 3 | 12 - 24 | € 972 - € 1,944 |
| Rituximab (in combination with FCM) | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 1 | 4 - 8 | €284- €568 |
| Yttrium-90 radiolabelled ibritumomab tiuxetan pretreated with rituximab | | | | | |

| Yttrium-90 radiolabelled ibritumomab tiuxetan | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 1 | 1 | €71 |
|--|---|-----|---|---|-------|
| Rituximab | Surcharge for the preparation of a parenteral solution containing monoclonal antibodies | €71 | 2 | 2 | € 142 |