6.05 PEMBROLIZUMAB,   
Solution concentrate for I.V. infusion 100 mg in 4 mL,   
Keytruda®,

**Merck Sharp & Dohme (Australia) Pty Ltd**

1. Purpose of application
   1. The Category 2 submission requested an Authority Required (Streamlined) listing for pembrolizumab in combination with platinum-based chemotherapy for the treatment of locally advanced (Stage III) or metastatic (Stage IV) oesophageal adenocarcinoma (OAC) or oesophageal squamous cell carcinoma (OSCC), or human epidermal growth factor receptor 2 (HER2)-negative adenocarcinoma (AC) of the gastro-oesophageal junction (GOJ).
   2. The basis for the requested listing was a cost-utility analysis against the comparator, chemotherapy alone.
   3. Table 1 presents the key components of the clinical issue addressed by the submission.

Table 1: Key components of the clinical issue addressed by the submission

|  |  |
| --- | --- |
| **Component** | **Description** |
| Population | Patients with Stage III unresectable (advanced)a or Stage IV (metastatic) OAC or OSCC, or HER2-negativea AC of the GOJ. |
| Intervention | Pembrolizumab 200 mg IV Q3W for up to 35 cycles + 5-FU 800 mg/m2 for days 1-5 for up to 35 cycles + cisplatin 80 mg/m2 Q3W for up to 6 cycles |
| Comparator | Standard of Care. Due to the complexities of the various regimens, the following have been identified and included as representative of all comparators:  Cisplatin + 5-FU (represents other chemotherapy regimens including XELOX/CAPOX and FOLFOX) |
| Outcomes | Objective response rate, progression free survival, overall survival |
| Clinical claim | In patients with Stage III unresectable (advanced) or Stage IV (metastatic) oesophageal cancer or HER2 negative gastro-oesophageal junction cancer, pembrolizumab in combination with chemotherapy is more effective than standard chemotherapy at improving survival, progression free survival, and quality of life. The submission described pembrolizumab plus chemotherapy as inferior in terms of safety compared to chemotherapy alone. |

Source: Table 1.1.1, pp3-4 of the submission.

AC = adenocarcinoma; 5-FU = 5 fluorouracil; FOLFOX = oxaliplatin plus leucovorin plus 5-FU; GOJ = gastro-oesophageal junction; HER2 = human epidermal growth factor receptor 2; IV = intravenous; mg = milligram; OAC = oesophageal adenocarcinoma; OSCC = oesophageal squamous cell carcinoma; Q3W = every three weeks; XELOX/CAPOX = capecitabine plus oxaliplatin

a The submission was inconsistent in describing the stage of disease in the target population and specifying HER2-negative status for GOJ. The restriction included advanced and metastatic, whereas the PICO table only included metastatic. This has been amended in the table.

1. Background

***Registration status***

* 1. Pembrolizumab was registered for OAC, OSCC or GOJ on the 28 September 2021. The TGA application for pembrolizumab for this indication was granted priority review under Project ORBIS and received a designation of Orphan Drug status. The approved TGA indication is:

In combination with platinum- and 5-FU-based chemotherapy for the first-line treatment of patients with locally advanced or metastatic carcinoma of the oesophagus or HER2-negative gastroesophageal junction adenocarcinoma (tumour centre 1 to 5 centimetres above the gastroesophageal junction) that is not amenable to surgical resection or definitive chemoradiation.

1. Requested listing
   1. The requested listing for pembrolizumab is provided below. Secretariat suggestions and additions proposed are shown in italics and deletions are in strikethrough.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Name, restriction, manner of administration, form** | **Maximum amount (units)** | **No. of repeats** | **Dispensed price for maximum amount** | **Proprietary name and manufacturer** |
| Pembrolizumab  100mg injection, 1 vial | 200mg | 6 | $7,881.87 published price (private)  $7,733.78 published price (public) | Keytruda, Merck Sharpe & Dohme (Australia) Pty Ltd |
| **Category / Program:** Section 100 – Efficient Funding of Chemotherapy Public/Private hospitals | | | | |
| **Prescriber type:** Medical Practitioners | | | | |
| **Restriction Type:** Authority Required (STREAMLINED) [new code] | | | | |
| ***Administrative advice:*** *Applications for authorisation under this restriction may be made in real time using the Online PBS Authorities system (see www.servicesaustralia.gov.au/HPOS) or by telephone by contacting Services Australia on 1800 888 333.* | | | | |
| ***Administrative advice:*** *No increase in the maximum number of repeats may be authorised.* | | | | |
| ***Administrative advice:*** *Special Pricing Arrangements apply.* | | | | |
| ***Episodicity:*** *[blank]* | | | | |
| ***Severity:*** *Advanced (Stage III) or metastatic**(Stage IV)* | | | | |
| **Condition:** ~~Oesophageal/GOJ cancer~~ *Carcinoma of the following types: (i) Siewert type I adenocarcinoma of oesophagogastric junction, (ii) adenocarcinoma of oesophagus, (iii) squamous cell carcinoma of oesophagus* | | | | |
| **Indication:** ~~Unresectable locally advanced (III) or metastatic (IV) carcinoma of the oesophagus or HER2 negative adenocarcinoma of the gastroesophageal junction (GOJ)~~ *advanced (Stage III) or metastatic (Stage IV)* *carcinoma of the following types: (i) Siewert type I adenocarcinoma of oesophagogastric junction, (ii) adenocarcinoma of oesophagus, (iii) squamous cell carcinoma of oesophagus* | | | | |
| **Treatment Phase:** Initial *~~T~~treatment* | | | | |
| **Clinical criteria:** | | | | |
| ~~The condition must be locally advanced (Stage III) or metastatic (Stage IV) carcinoma of the oesophagus or adenocarcinoma of the gastroesophageal junction (GOJ) that is not amenable to surgical resection or definitive chemoradiation~~ *The condition must be unsuitable for each of: (i) surgical resection, (ii) chemoradiation* | | | | |
| **AND** | | | | |
| ~~Patient~~ *The condition* must have evidence of human epidermal growth factor receptor 2 (HER2) negativity as demonstrated by immunohistochemistry in tumour material*, in any diagnosis of Siewert type I adenocarcinoma of oesophagogastric junction – retain this evidence on the patient’s medical records; do not submit a copy of this evidence in this authority application* | | | | |
| **AND** | | | | |
| ~~The patients should not have had prior systemic therapy administered in the metastatic setting~~ | | | | |
| ***Treatment criteria:*** | | | | |
| *Patient must be undergoing treatment with this drug for the first time* | | | | |
| ***AND*** | | | | |
| *Patient must be undergoing treatment with this drug for metastatic disease (Stage IV disease) that is untreated with drug therapy; or* | | | | |
| **~~AND~~** | | | | |
| ~~Patients must not have had a recurrence within 6 months of platinum-based chemotherapy in the locally advanced setting~~  *Patient must be undergoing treatment with this drug for locally advanced disease (Stage III disease) that is untreated with drug therapy* | | | | |
| **AND** | | | | |
| ~~The treatment must be commenced in combination with platinum- and fluoropyrimidine-based chemotherapy~~ | | | | |
| ***Treatment criteria:*** | | | | |
| *Patient must be undergoing concomitant treatment with chemotherapy, at leas at treatment initiation with this drug, containing a minimum of: (i) a platinum agent, plus (ii) a fluoropyrimidine agent* | | | | |
| **AND** | | | | |
| ~~Patient must have a WHO performance status of 0 or 1~~  *Patient must have WHO performance status no higher than 1 at treatment initiation with this drug* | | | | |
| **~~Treatment criteria:~~**  ~~The treatment must not exceed a total of 7 doses under this restriction~~**~~.~~** | | | | |
| **Administrative Advice:**  In the first few months after start of immunotherapy, some patients can have a transient tumour flare with subsequent disease response. When progression is suspected, this should be confirmed through a confirmatory scan, taken at least 4 weeks later  ~~Following completion of the initial PBS subsidised course, further applications for treatment will be assessed under the continuing treatment restriction.~~ | | | | |
|  | | | | |
| **Restriction Type:** Authority Required (STREAMLINED) [new code] | | | | |
| 1. **Indication:** ~~Unresectable locally advanced (III) or metastatic (IV) carcinoma of the oesophagus or HER2 negative adenocarcinoma of the gastroesophageal junction (GOJ)~~ 2. *Advanced (Stage III) or metastatic (Stage IV) carcinoma of the following types: (i) Siewert type I adenocarcinoma of oesophagogastric junction, (ii) adenocarcinoma of oesophagus, (iii) squamous cell carcinoma of oesophagus* | | | | |
| 1. **Treatment Phase:** Continuing *~~T~~treatment* | | | | |
| 1. **Clinical criteria:** | | | | |
| 1. ~~Patient must have previously received PBS-subsidised treatment with this drug for this condition~~ | | | | |
| 1. **~~AND~~** | | | | |
| 1. ~~Patient must not have progressive disease while receiving PBS-subsidised treatment with this drug for this condition~~ | | | | |
| 1. *The condition must not have progressed* | | | | |
| 1. ***AND*** | | | | |
| 1. ***Treatment criteria:*** | | | | |
| 1. *Patient must be undergoing continuing treatment with this drug, with PBS-subsidised treatment having commenced through one of: (i) the ‘Initial treatment’ phase listing, (ii) ‘Grandfather arrangements’ listing; do not commence PBS-subsidised treatment through this treatment phase* | | | | |
| 1. ***AND*** | | | | |
| 1. **Treatmentcriteria:** | | | | |
| 1. ~~The treatment must not exceed a total of 35 cycles of treatment in a lifetime~~ ~~under this restriction~~. | | | | |
| 1. *Patient must not be undergoing continuing treatment through the PBS such that the total duration of treatment (as measured from the first dose of this drug, regardless if it was PBS-subsidised/non-PBS subsidised) goes beyond whichever comes first out of the following: (i) 24 months, (ii) 35 doses (based on a 3-weekly dose regimen), (iii) disease progression* | | | | |
|  | | | | |
| **Restriction Type:**  Authority Required (Streamlined) [new code] | | | | |
| **Indication:** ~~Unresectable locally advanced (III) or metastatic (IV) carcinoma of the oesophagus or HER2 negative adenocarcinoma of the gastroesophageal junction (GOJ)~~  *Advanced (Stage III) or metastatic (Stage IV) carcinoma of the following types: (i) Siewert type I adenocarcinoma of oesophagogastric junction, (ii) adenocarcinoma of oesophagus, (iii) squamous cell carcinoma of oesophagus* | | | | |
| **Treatment Phase:** ~~Grandfather Treatment~~ *Transitioning from non-PBS to PBS-subsidised supply – ‘Grandfather’ arrangements* | | | | |
| **Clinical criteria:** | | | | |
| Patient must ~~have previously received non-PBS subsidised treatment with this drug for this condition prior to~~ *be currently receiving treatment with this drug for this PBS indication, with treatment having commenced prior to* [PBS listing date] | | | | |
| **AND** | | | | |
| ~~Patient must not have had been treated for this condition in the metastatic setting prior to initiating non-PBS subsidised treatment with this drug for this condition~~ | | | | |
| **~~AND~~** | | | | |
| ~~Patient must not have experienced recurrence within 6 months of completion of systemic therapy if treated in the locally advanced setting prior to non-PBS subsidised treatment with this drug for this condition~~ | | | | |
| **~~AND~~** | | | | |
| ~~The treatment must be commenced in combination with platinum-based chemotherapy~~ | | | | |
| **~~AND~~** | | | | |
| ~~Patient must have stable or responding disease~~ | | | | |
| **~~AND~~** | | | | |
| ~~Patient must have a WHO performance status of 0 or 1~~ | | | | |
| ***Clinical criteria:*** | | | | |
| *Patient must have met all other PBS eligibility criteria that a non-‘Grandfather’ patient would ordinarily be required to meet, meaning that at the time non-PBS supply was commenced, the patient: (i) had a WHO performance status no greater than 1, (ii) was unsuitable for both surgical resection plus chemoradiation, (iii) for a diagnosis of Siewert type I adenocarcinoma of oesophagogastric junction, there was evidence confirming HER2 negativity via immunohistochemistry in tumour material, (iv) was untreated with this drug, (v) for metastatic disease (Stage IV disease), the metastatic disease had yet to be treated with drug therapy, (vi) for locally advanced disease (Stage III disease), the locally advanced disease had yet to be treated with drug therapy, (vii) was treated concomitantly with chemotherapy containing at least each of a platinum agent plus a fluoropyrimidine agent* | | | | |
| ***AND*** | | | | |
| ***Clinical criteria:*** | | | | |
| *The condition must not have progressed* | | | | |
| **~~AND~~** | | | | |
| **~~Treatment criteria:~~** | | | | |
| ~~The treatment must not exceed a total of 35 cycles of treatment in a lifetime under this restriction.~~ | | | | |
| ***Treatment criteria:*** | | | | |
| *Patient must not be undergoing continuing treatment through the PBS such that the total duration of treatment (as measured from the first dose of this drug, regardless if it was PBS-subsidised/non-PBS subsidised) goes beyond whichever comes first out of the following: (i) 24 months, (ii) 35 doses (based on a 3-weekly dose regimen), (iii) disease progression* | | | | |
| **~~Administrative Advice:~~**  ~~In the first few months after start of immunotherapy, some patients can have a transient tumour flare with subsequent disease response. When progression is suspected, this should be confirmed through a confirmatory scan, taken at least 4 weeks later.~~  ~~A patient may only qualify for PBS-subsidised treatment under this restriction once.~~  ~~Following completion of the initial PBS subsidised course, further applications for treatment will be assessed under the continuing treatment restriction.~~ | | | | |
| ***Administrative advice:***  *Patients may qualify for PBS-subsidised treatment under this restriction once only. For continuing PBS-subsidised treatment, a 'Grandfathered' patient must qualify under the 'Continuing treatment' criteria.* | | | | |
| ***Administrative advice:***  *This grandfather restriction will cease to operate from 12 months after the date specified in the clinical criteria.* | | | | |

* 1. The submission stated that a Special Pricing Arrangement (SPA) will be required. The submission included an ex-manufacturer price per 100 mg vial of $''''''''''''''''' and the Pre-Sub-Committee Response (PSCR) proposed a revised price of $'''''''''''''''''
  2. The requested restriction is generally consistent with the approved TGA indication and the inclusion criteria of the KN590 trial.
  3. HER2-negative status is only relevant to patients with AC of the GOJ. Patients with AC of the GOJ would receive a HER2 test to determine eligibility for treatment with trastuzumab (which is PBS listed for HER2-positive GOJ), but patients with OAC and OSCC may not routinely receive a HER2 test, and thus have unknown HER2 status.
  4. The requested restriction stated: “The treatment must be commenced in combination with platinum- and fluoropyrimidine-based chemotherapy.” The Secretariat suggested rewording to “Patient must be undergoing concomitant treatment with chemotherapy, at least at treatment initiation with this drug, containing a minimum of: (i) a platinum agent, plus (ii) a fluoropyrimidine agent”. The PBAC considered this could be simplified to “Patient must be undergoing concomitant chemotherapy with a platinum agent plus a fluoropyrimidine agent at treatment initiation with this drug”. The criterion permits use of pembrolizumab with cisplatin + 5-FU, FOLFOX, or XELOX/CAPOX (paragraph 5.2).
  5. The PBAC considered that the restriction should reflect that a requirement for HER2 testing should only apply to patients with AC of the GOJ, and to fully align with the TGA outcome, the restriction should specify the location of the tumour centre above the junction (1 to 5 cm) by stating “Siewert type I adenocarcinoma of oesophagogastric junction” as part of the restriction condition. The PBAC considered the restriction should specify Siewert type I GOJ cancer. The PBAC noted that even with inclusion of this wording there was a risk pembrolizumab would be used in Siewert type II and III GOJ and gastric cancer.

*For more detail on PBAC’s view, see section 7 PBAC outcome.*

1. Population and disease
   1. Oesophageal cancer is the eighth most common cancer and the sixth most common cause of cancer death worldwide. Squamous cell carcinoma and adenocarcinoma account for over 95 per cent of oesophageal malignant tumours (Saltzman 2021[[1]](#footnote-1)). Smoking and alcohol are major risk factors for OSCC, while Barrett's oesophagus with intestinal metaplasia (a complication of gastroesophageal reflux disease), obesity, and smoking are risk factors for OAC.
   2. In a review of all oesophageal cancer patients presented to the South Australia state-wide upper gastrointestinal cancer multi-disciplinary team from 2012 to 2014 (Nguyen 2019[[2]](#footnote-2)), the diagnosis was OAC in 69.6%, OSCC in 24.8% and high grade dysplasia (HGD) in 5.6%. The majority of patients presented with Stage II and III disease (56.8%). A total of 51.5% were treated with curative intent, with 28.8% undergoing surgery and/or neoadjuvant therapy. Treatment was palliative in 48.5%, with chemoradiotherapy utilised in 20.8%.

*For more detail on PBAC’s view, see section 7 PBAC outcome.*

1. Comparator
   1. The submission noted that there is no single main comparator, since the choice of agent in first line is determined by the patients’ performance status, organ function and the ability to swallow. There are currently three main chemotherapy regimens used in Australia and recommended by Australian Guidelines:

* Cisplatin and fluorouracil (5-FU);
* Oxaliplatin and capecitabine (XELOX/CAPOX); and
* Leucovorin calcium (folinic acid), 5-FU and oxaliplatin (FOLFOX).
  1. The submission stated that cisplatin + 5-FU has been selected as representative of all platinum- and 5-FU-based regimens and is therefore representative of all standard of care comparators. The submission noted that all three regimens are used in practice, with clinicians considering them to be interchangeable with respect to efficacy and safety. The PSCR stated that while FOLFOX and XELOX/CAPOX are the most commonly used regimens, the PBAC has previously deemed all platinum doublets to be comparable and clinically non-inferior (Section 6, gefitinib Public Summary Document (PSD), July 2013) and fluoropyrimidines to be interchangeable (Table 1, trastuzumab PSD, July 2015). The ESC agreed with the PSCR that most patients would receive FOLFOX or XELOX/CAPOX regimens, as they have comparable efficacy and reduced toxicity compared with cisplatin + 5-FU. The ESC agreed with the submission that cisplatin + 5-FU can be considered representative of all platinum- and 5-FU-based regimens.
  2. The submission also noted that nivolumab plus chemotherapy may be a near term comparator. The PBAC noted that nivolumab was being considered at the same PBAC meeting for advanced or metastatic gastric cancer, GOJ cancer and OAC. The PBAC noted the indications for pembrolizumaband nivolumab (which were based on the clinical trial populations) do not fully align; however, the PBAC considered there was likely to be significant overlap in clinical practice due to overlap in anatomical distribution of these cancers.

*For more detail on PBAC’s view, see section 7 PBAC outcome.*

1. Consideration of the evidence

## Sponsor hearing

* 1. There was no hearing for this item.

## Consumer comments

* 1. The PBAC noted and welcomed the input from an individual, health care professionals (HCPs) (2) and organisations (2) via the Consumer Comments facility on the PBS website.
  2. The comments from two HCPs emphasised the aggressive nature of the disease and the poor prognosis for patients. One HCP stated that there is an urgent need for newer agents to be available, and commented that unlike some other cancers, there are few survivors to advocate for this patient population, as the majority of patients die within 12 months of diagnosis. Another HCP described the course of the disease where patients have simultaneous issues with metastatic cancer spread and difficulty swallowing from the primary cancer, leading to rapid weight loss, decreased quality of life and side effects from chemotherapy. The HCP emphasised that the current options for treatment are very limited, and commented that the KEYNOTE 590 trial (along with the CheckMate 648 nivolumab trial) was one of the only positive clinical trials in oesophageal cancer for a long time.
  3. The PBAC noted the advice received from the Pancare Foundation clarifying the likely use of pembrolizumab in clinical practice. In addition to similar issues listed in paragraph 6.3, Pancare described oesophageal carcinoma and HER-2 negative gastro-oesophageal carcinoma as having one of the worst survival rates of all cancers in Australia (5-year survival 22.5%). Pancare stated that a PBS listing for pembrolizumab would enable access to a superior first-line treatment and offer much-needed hope for Australians with oesophageal cancer. The PBAC noted that this advice was supportive of the evidence provided in the submission.
  4. The Medical Oncology Group of Australia (MOGA) also expressed its strong support for the use of pembrolizumab in first line (1L) oesophageal carcinoma (OC) and HER2-negative AC of the GOJ, categorising it as one of the therapies of “highest priority for PBS listing” for patients with PD-L1 CPS ≥ 10% and “high priority for PBS listing” for all patients (regardless of PD-L1 CPS expression) on the basis of the KEYNOTE 590 trial. The PBAC noted that the MOGA presented a European Society for Medical Oncology Magnitude of Clinical Benefit Scale (ESMO-MCBS) for pembrolizumab (+ chemotherapy) in 1L OC and HER2-negative AC of the GOJ, which was limited to 4 (out of a maximum of 5, where 4 and 5 represent the grades with substantial improvement)[[3]](#footnote-3), based on a comparison with chemotherapy alone.
  5. One individual who has used pembrolizumab provided insight into their experience with oesophageal cancer and treatment with this medicine. The individual reported their positive experience with treatment including being able to work, a reduction in pain and weight gain. The patient advocated that access to pembrolizumab should be available through the PBS.

## Clinical trials

* 1. The submission was based on KEYNOTE 590 (KN590) (n=749), a randomised, double blind trial comparing pembrolizumab + chemotherapy (n=373) to cisplatin + 5-FU (n=376) in patients with locally advanced unresectable or metastatic OAC or OSCC or advanced/metastatic Siewert type I AC of the GOJ. The ESC noted that 73% of patients enrolled in KN590 were diagnosed with OSCC, 15% had OAC and 12% had AC of the GOJ.
  2. Details of the KN590 trial presented in the submission are provided in Table 2.

Table 2: Trials and associated reports presented in the submission

| **Trial ID** | **Protocol title/ Publication title** | **Publication citation** |
| --- | --- | --- |
|  | A Phase III randomised study of chemotherapy + pembrolizumab vs chemotherapy + placebo as first-line therapy for patients with advanced oesophageal or esophagogastric junction (E/EGJ) cancer. | CSR. September 2020 |
| Keynote 590 (KN590) | Kato K, Shah MA, Enzinger P, Bennouna J, Shen L, Adenis A, Sun JM, Cho BC, Özgüroğlu M, Kojima T, Kostorov V, Hierro C, Zhu Y, McLean LA, Shah S, Doi T. KEYNOTE-590: Phase III study of first-line chemotherapy with or without pembrolizumab for advanced esophageal cancer. | Future Oncol. 2019 Apr;15(10):1057-1066. Epub 2019 Feb 8. |
|  | Kato K, Sun J, Shah MA, Enzinger PC, Adenis A, Doi T, Kojima T, Metges J, Li Z, Kim S, Chul Cho BC, Mansoor W, Li S, Sunpaweravong P, Maqueda MA, Goekkurt E, Liu Q, Shah S, Bhagia P, Shen L, | Annals of Oncology (2020) 31 (suppl\_4): S1142-S1215. [abstract] |

Source: Table 2.2-1, p10 of the submission.

* 1. The key features of the KN590 trial are summarised in Table 3.

Table 3: Key features of the included evidence

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **Trial** | **N** | **Design/duration** | **Risk of bias** | **Patient population** | **Outcome(s)** | **Use in modelled evaluation** |
| **Pembrolizumab plus chemo vs. chemo alone** | | | | | | |
| KN590 | 749 | R, DB, MC  ~12 mthsa | Low | First line | OS, PFS | OS, PFS, ToT |

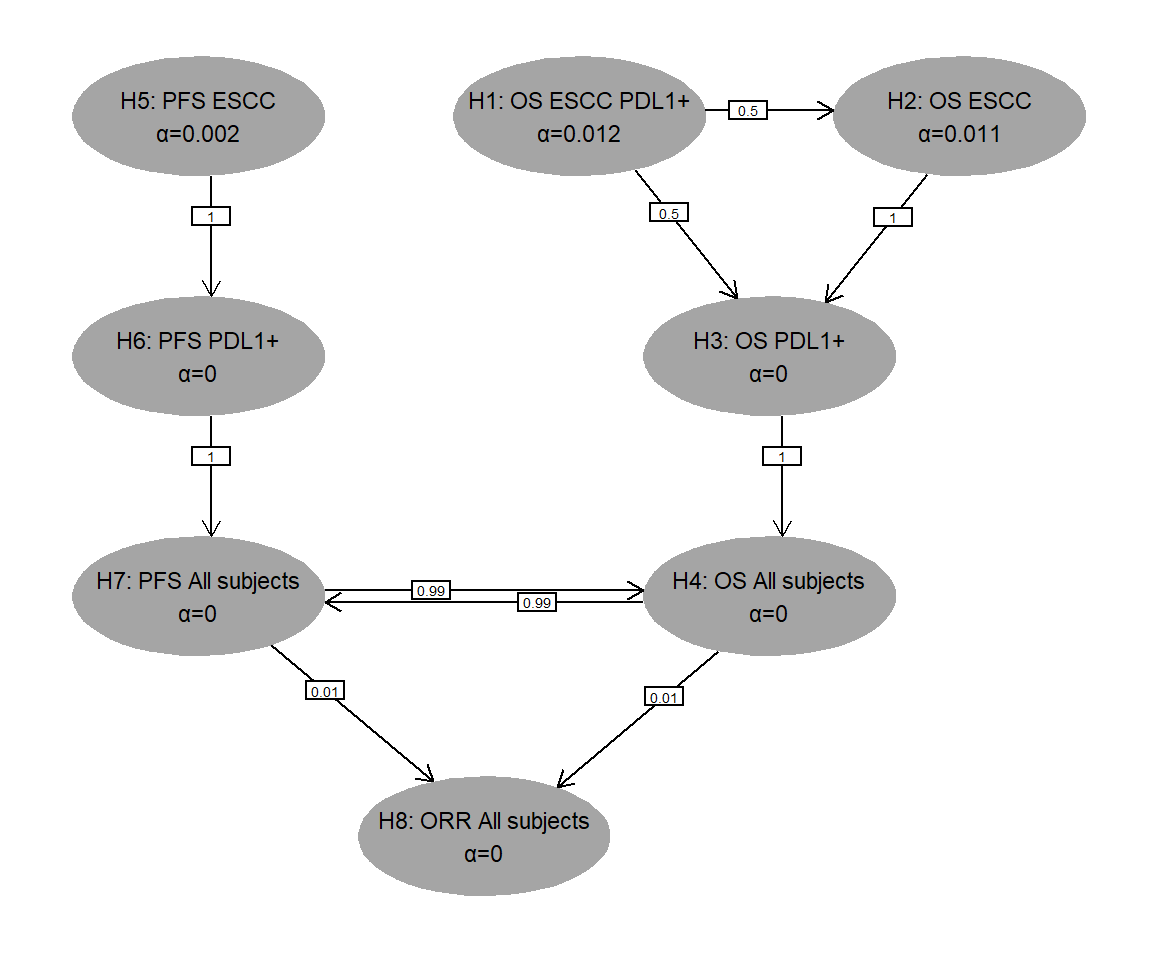
Source: pp10-20 of the submission.

DB = double blind; MC = multi-centre; OL = open label; OS = overall survival; PFS = progression-free survival; R = randomised; ToT = time on treatment

a Median follow-up in treatment arm

* 1. The dual primary efficacy outcomes for KN590 were overall survival (OS) and progression free survival (PFS). Overall Response Rate (ORR) was the key secondary efficacy endpoint.
  2. The KN590 trial was evaluated in a stepwise fashion, with sequential hypothesis testing. This included key primary and secondary outcomes in the following pre-specified subgroup populations:
* Participants with OSCC in the intention-to-treat (ITT) population.
* Participants with OSCC with programmed death-ligand 1 (PD-L1) combined positive score (CPS) ≥10 in the ITT population (OSCC CPS ≥10).
* Participants with PD-L1 CPS ≥10 in the ITT population.
  1. Protocol Amendment 8 included the addition of three primary objectives and their corresponding hypotheses.
* OS in OSCC population.
* OS in OSCC CPS ≥10 population.
* PFS in OSCC population.
  1. Figure 1 illustrates the sequential hypothesis testing in the KN590 trial.

**Figure 1**: Sequential hypothesis testing and alpha spending in the KN590 trial



Source: Figure 2.4-2, p23 of the submission.

ESCC = oesophageal squamous cell carcinoma; H = hypothesis; ORR = objective response rate; OS = Overall survival; PDL+ = programmed death-ligand 1 positive; PFS = progression free survival.

## Comparative effectiveness

* 1. Key survival results of the KN590 trial are summarised in Table 4.

Table 4: Summary of ITT survival outcomes in KN590

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | **Pembrolizumab plus chemo**  **N=376** | **Chemo alone**  **N=376** | **Absolute difference** | **HRb (95% CI)**  **p-value** |
| **Progression-free survival** | | | | |
| Patients with event n (%) | 297 (79.6) | 333 (88.6) | *-* | - |
| Mediana PFS months (95% CI) | 6.3 (6.2, 6.9) | 5.8 (5.0, 6.0) | 0.5 months | **0.65 (0.55, 0.76)**  p-valuec <0.0001 |
| Proportion progression free at 12 months (95% CI) | 62.4 (57.1, 67.3) | 48.7 (43.4, 53.7) | 13.7% | **-** |
| **Overall survival** | | | | |
| Patients with event n (%) | 262 (70.2) | 309 (82.2) | - | - |
| Mediana OS months (95% CI) | 12.4 (10.5, 14.0) | 9.8 (8.8, 10.8) | 2.6 months | **0.73 (0.62, 0.86)**  p-valuec <0.0001 |
| Proportion alive at 12 months (95% CI) | 50.6 (45.4, 55.6) | 39.4 (34.4, 44.3) | 11.2% | **-** |

Source: Table 2.5.2, p30 and Table 2.5.3, p32 of the submission.

Chemo = chemotherapy; CI = confidence interval; HR = hazard ratio; OS = overall survival; PFS = progression free survival.

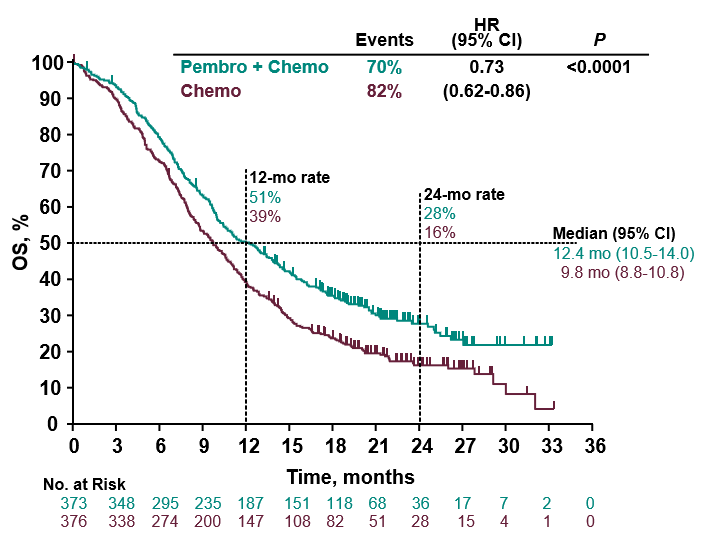
a Median survival was estimated through Kaplan Meier product-limit method of censored data.

b Hazard ratio was based on Cox regression model with Efron’s method of tie handling with treatment as a covariate stratified by geographic region (Asia versus Rest of the World) and ECOG performance status (0 versus 1).

c The one-sided p-value based on log-rank test stratified by geographic region (Asia versus Rest of the World) and ECOG performance status (0 versus 1).

* 1. The KN590 results showed statistically significant improvements in both PFS and OS in the ITT population.
  2. Figure 2 and Figure 3 present the KN590 Kaplan Meier survival curves for OS and PFS, respectively.

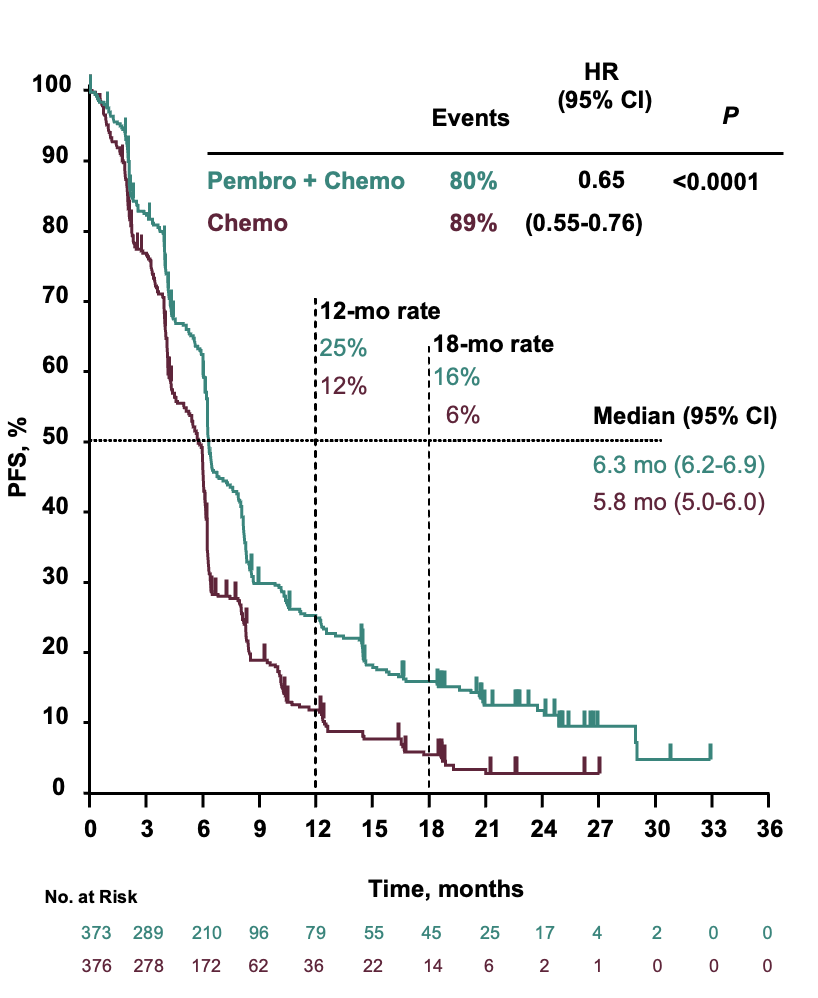
Figure 2: Kaplan-Meier Estimates of Overall Survival (ITT Population)



Source: Figure 2.5-1, p31 of the submission.

CI = confidence interval; HR = hazard ratio; mo = month; No = number; OS = overall survival

Figure 3: Kaplan-Meier Estimates of Progression-Free Survival Based on Investigator Assessment per RECIST 1.1 (ITT Population)



Source: Slide 9 of Kato 2020 ESMO presentation.

CI = confidence interval; HR = hazard ratio; mo = month; No = number

Note: The submission erroneously copied the OS Kaplan Meier curve for the PFS results. This was corrected in the evaluation.

* 1. Table 5 presents the subgroup results for the pre-specified KN590 primary endpoints discussed above, as well as additional exploratory analyses compiled by the FDA (including PD-L1 CPS >/< 1, and OAC patients by CPS status).

Table 5: Subgroup analyses from KN590 prespecified analyses and FDA analyses from TGA Delegate’s Overview

| **Population** | **N** | **OS HR (CI)** | **P** |
| --- | --- | --- | --- |
| OSCC CPS ≥10 | 286 | 0.57 (0.43, 0.75) | Significant |
| OSCC all CPS | 548 | 0.72 (0.60, 0,88) | Significant |
| All cancer types, CPS ≥10 | 383 | 0.62 (0.49, 0.78) | Significant |
| All patients | 749 | 0.73 (0.62, 0.86) | Significant |
| All cancer types, CPS <10 a | 347 | 0.86 (0.68, 1.10) | Not tested |
| OSCC, CPS <10 a | 247 | 0.99 (0.74, 1.32) | Not tested |
| OAC, CPS <10 a | 100 | 0.66 (0.42, 1.04) | Not tested |
| CPS <1 a | ''''''' | '''''''''''' '''''''''''''' '''''''''''' | ''''''' '''''''''''''''' |
| CPS >1 a | ''''''''' | ''''''''''' '''''''''''''' ''''''''''''' | ''''''''' ''''''''''''' |

Source: Page 27 of TGA Delegate’s Overview

CI; = confidence interval; CPS = combined positive score; HR = hazard ratio; OAC = oesophageal adenocarcinoma; OS = overall survival; OSCC = oesophageal squamous cell carcinoma

*a Note that these results presented in Table 5 were not part of the pre-specified statistical plan for KN590. Interpretation of the results and their application should therefore be limited to seeking to understand the basis for the PBAC outcome and should not be used for any other purpose.*

* 1. The KN590 trial included results of the European Organisation for Research and Treatment of Cancer Quality of Care (EORTC-QLQ) C30 questionnaire and the Oesophageal cancer module (OES18) as well as the European Quality of life 5 dimensions Visual Analogue Scale (EQ-5D VAS) as exploratory analyses. No statistically significant differences were observed in these quality of life measures.
  2. The PBAC noted the objective response rate (ORR) for patients treated with pembrolizumab in combination with chemotherapy was 45.0% with a median duration of response of 8.3 months compared to a 29.2% ORR and a median duration of response of 6.0 months for patients treated with chemotherapy.

## Comparative harms

* 1. The analysis of safety data in this study were based on the All Subjects as Treated (ASaT) population which consists of all randomised participants who received at least one dose of study treatment. Table 6 presents a summary of adverse events (AEs) in the KN590 trial.

Table 6: **Summary of key adverse events in KN590 (ASaT population)**

| **Trial** | **Pembrolizumab plus chemo**  **n (%)** | **Chemo alone**  **n (%)** | | **RD (95% CI)** |
| --- | --- | --- | --- | --- |
| Subjects in population (N) | 370 | 370 | | -- |
| One or more AEs | 370 (100) | 368 (99.5) | | 0.01 (0.00, 0.02) |
| Drug-related† AEs | 364 (98.4) | 360 (97.3) | | 0.01 (-0.01, 0.03) |
| Toxicity grade 3-5 AEs | 318 (85.9) | 308 (83.2) | | 0.03 (-0.02, 0.08) |
| Toxicity grade 3-5 drug-related AEs | 266 (71.9) | 250 (67.6) | | 0.04 (-0.02, 0.11) |
| Non-serious AEs | 368 (99.5) | 367 (99.2) | | 0 (-0.01, 0.01) |
| Serious AEs | 205 (55.4) | 204 (55.1) | | 0 (-0.07, 0.07) |
| Serious drug-related AEs | 117 (31.6) | 97 (26.2) | | 0.05 (-0.01, 0.12) |
| Died | 28 (7.6) | 38 (10.3) | | -0.03 (-0.07, 0.01) |
| Died due to a drug-related AE | 9 (2.4) | 5 (1.4) | | 0.01 (-0.01, 0.03) |
| Discontinued drug due to an AE | 90 (24.3) | 74 (20.0) | | 0.04 (-0.02, 0.1) |
| Discontinued drug due to a drug-related AE | 72 (19.5) | 43 (11.6) | | 0.08 (0.03, 0.13) |
| Discontinued drug due to a serious AE | 58 (15.7) | 47 (12.7) | | 0.03 (-0.02, 0.08) |
| Discontinued drug due to a serious drug-related AEs | 38 (10.3) | 17 (4.6) | | 0.06 (0.02, 0.09) |
| AEOSIa | 95 (25.7) | | 43 (11.6) | 0.14 (0.09, 0.20 |
| Hypothyroidism | 38 (10.3) | | 22 (5.9) | 0.04 (0.00, 0.08) |
| Hyperthyroidism | 19 (5.1) | | 2 (0.0.5) | 0.05 (0.02, 0.07) |
| Pneumonitis | 20 (5.4) | | 0 | 0.05 (0.03, 0.08) |

Source: Table 2.5-12, p41, Table 2.5.14, p43, and Table 2.5.17, p47 of the submission

AE = adverse event; AEOSI = adverse event of special interest CI = confidence interval; n = number of participants reporting data; N = total participants in group; RD = risk difference; RR = relative risk

a Comprises adrenal insufficiency, colitis, hepatitis, hyperthyroidism, hypophysitis, hypothyroidism, infusion reactions, myositis nephritis, pancreatitis, pneumonitis, severe skin reactions thyroiditis, and type 1 diabetes mellitus.

Risk difference calculated post-hoc during the evaluation.

* 1. The submission noted that the most frequently reported AEs (defined as ≥40%) in the pembrolizumab + chemotherapy and in the chemotherapy alone arms were nausea, decreased appetite, fatigue, and constipation. These events were predominantly Grade 1 or 2 and manageable.
  2. The submission stated that the drug-related AEs observed for participants treated with pembrolizumab + chemotherapy was generally consistent with the known safety profiles of pembrolizumab monotherapy and 5-FU/cisplatin chemotherapy.
  3. The TGA Delegate considered that (p25-26 of Delegate’s Overview):
* “Safety results from KEYNOTE-590 demonstrate that pembrolizumab in combination with 5FU/cisplatin has a tolerable safety profile that reflects the adverse reaction profiles of the components, with no new safety concerns identified. The addition of pembrolizumab does not appear to significantly increase the toxicity of the chemotherapy backbone, taking into consideration the different lengths of treatment exposure.”
* “Some patients can however experience severe and/or serious toxicity from pembrolizumab, generally due to immune related AEs.”
  + “Specifically, the incidence of death was similar in both treatment arms, (pneumonia being he commonest cause of death). The incidence of serious AEs was also similar in both arms, except for pneumonitis which was more common in combination arm (3.2% versus 0.3%); of note, pneumonitis is an expected immune related AE for pembrolizumab.”
* “More serious drug-related AEs and discontinuations were seen for those in the pembrolizumab plus chemotherapy arm compared to the placebo plus chemotherapy arm. The incidence of AEOSI was higher in the pembrolizumab plus chemotherapy arm (compared to the placebo plus chemotherapy arm) as would be expected, with the AEOSI generally consistent with the known pembrolizumab safety profile.”
* “The risks of pembrolizumab are therefore largely manageable with patient surveillance, treatment delays and supportive care in most patients, and are considered acceptable given the life-threatening nature of metastatic or locally advanced oesophageal carcinoma.”

## Benefits/harms

* 1. A summary of the comparative benefits and harms for pembrolizumab plus chemotherapy versus chemotherapy is presented in Table 7.

Table 7: Summary of comparative benefits and harms for pembrolizumab plus chemotherapy versus chemotherapy

|  | Pembro plus chemo  n/N (%) | | Chemo alone  n/N (%) | Absolute difference | | HR (95% CI) |
| --- | --- | --- | --- | --- | --- | --- |
| Progression-free survival | | | | | | |
| Progressed, n (%) | 297 (79.6) | | 333 (88.6) | *-* | | - |
| Median PFS, months (95% CI) | 6.3 (6.2, 6.9) | | 5.8 (5.0, 6.0) | 0.5 months | | **0.65 (0.55, 0.76)**  p-value <0.0001 |
| Proportion progression free at 12 months (95% CI) | 62.4 (57.1, 67.3) | | 48.7 (43.4, 53.7) | 13.7% | | **-** |
| Overall survival | | | | | | |
| Deaths, n/N (%) | 262 (70.2) | | 309 (82.2) | *-* | | - |
| Median months OS (95% CI) | 12.4 (10.5, 14.0) | | 9.8 (8.8, 10.8) | 2.6 months | | **0.73 (0.62, 0.86)**  p-value < 0.0001 |
| Proportion alive at 12 months (95% CI) | 50.6 (45.4, 55.6) | | 39.4 (34.4, 44.3) | 11.2% | | **-** |
| Harms | | | | | | |
| **Adverse event** | **Number of patients (%)** | | | **Event rate/ 100 patientsa** | | **Risk difference**  **(95% CI)** |
|  | **Pembro plus chemo (N=782)** | **Chemo alone (N=767)** | | **Pembro plus chemo** | **Chemo alone** |  |
| Serious drug-related AEs | 117 (31.6) | 97 (26.2) | | 32 | 26 | 0.05 (-0.01, 0.12) |
| Discontinuation drug due to a serious drug-related AE | 38 (10.3) | 17 (4.6) | | 10 | 5 | 0.06 (0.02, 0.09) |
| AEOSI | 95 (25.7) | 43 (11.6) | | 26 | 12 | 0.14 (0.09, 0.20) |
| Hypothyroidism | 38 (10.3) | 22 (5.9) | | 10 | 6 | 0.04 (0.00, 0.08) |
| Hyperthyroidism | 19 (5.1) | 2 (0.5) | | 5 | 1 | 0.05 (0.02, 0.07) |
| Pneumonitis | 20 (5.4) | 0 | | 5 | 0 | 0.05 (0.03, 0.08) |

Source: Table 2.5.2, p30, Table 2.5.3, p32 Table 2.5-12, p41, Table 2.5.14, p43, and Table 2.5.17, p47 of the submission

AE = adverse events; AEOSI = adverse event of special interest; Chemo = chemotherapy, CI = confidence interval; HR = hazard ratio; N = total participants in group; Pembro = pembrolizumab, SAE = serious adverse events

Note: Median survival was estimated through Kaplan Meier product-limit method of censored data

Note: Hazard ratio was based on Cox regression model with Efron’s method of tie handling with treatment as a covariate stratified by geographic region (Asia versus Rest of the World) and ECOG performance status (0 versus 1).

Note: The one-sided p-value based on log-rank test stratified by geographic region (Asia versus Rest of the World) and ECOG performance status (0 versus 1)

a Median duration of follow-up: of 12.6 months in pembrolizumab plus chemo arm and 9.8 months in chemotherapy alone arm.

Values in bold indicates statistically significant differences

* 1. On the basis of the KN590 trial presented by the submission, for every 100 patients treated with pembrolizumab plus chemotherapy (for a median duration of 12.6 months) instead of chemotherapy alone (for a median duration of 9.8 months):
* 12 more patients were alive at 12 months.
  1. On the basis of the KN590 trial presented by the submission, for every 100 patients treated with pembrolizumab plus chemotherapy (for a median duration of 12.6 months) instead of chemotherapy alone (for a median duration of 9.8 months):
* Approximately 5 additional patients would experience a serious drug related AE;
* Approximately 4 additional patients would experience hypothyroidism;
* Approximately 5 additional patients would experience hyperthyroidism;
* Approximately 5 additional patients would experience pneumonitis; and
* Approximately 6 additional patients would experience discontinuation of treatment due to a serious drug related AE.

## Clinical claim

* 1. The submission described pembrolizumab plus chemotherapy as superior in terms of effectiveness compared with chemotherapy alone and inferior in terms of safety compared to chemotherapy alone.
  2. The submission stated that the claim of superior effectiveness is supported by:
* A 27% reduction in the risk of death compared to treatment with a platinum- and 5-FU based chemotherapy (OS HR=0.73 (0.62, 0.86)).
* A 35% reduction in the risk of death or disease progression compared to chemotherapy (PFS HR=0.65 (0.55, 0.76)).
* The maintenance of stable QoL in patients and manageable tolerability with AEs generally consistent and similar to those previously reported with either pembrolizumab or chemotherapy when used separately.
  1. The evaluation and the ESC considered that overall, the claim of clinical superiority is reasonable in specific subgroups (OSCC; PD-L1 CPS ≥10), but it is likely that the efficacy of pembrolizumab is reduced in patients with PD-L1 CPS <10 when compared to those whose tumours express CPS ≥10. The proportion of PD-L1 CPS <10/≥10 is unknown for the Australian population, which may impact on the effectiveness of pembrolizumab in clinical practice.
  2. The PSCR acknowledged that the proportion of patients with PD-L1 CPS <10/≥10 is currently unknown in the Australian population due to testing in this indication not being routine. The KN590 study was not designed and powered to compare the efficacy in participants with PD-L1 CPS <10 and was not stratified by CPS score. The PSCR stated that it is possible that the same CPS distribution is apparent in the local population as in the KN590 trial, and as the CPS <10 patients are encapsulated within the ITT population, the overall HR is representative of the potential effect and can ensure equity of access. The pre-PBAC response noted that the KN590 trial included 5 Australian sites, and maintained the ITT results would be applicable to Australian practice.
  3. The proportion of OAC/OSCC in the KN590 trial (OAC in 15% of OC cases, OSCC in 73% of OC cases, paragraph 6.7) is different to the Australian target population, with the OAC trial patients being substantially under-represented compared to Australian patients (OAC in 69.6% of OC cases, OSCC in 24.8% of OC cases, paragraph 4.2). The KN590 trial did not report pre-specified analyses for patients with OAC, and in the exploratory analysis, the difference between arms for OS did not reach statistical significance (HR = 0.74 (95% CI: 0.54, 1.02); noting the trial was not powered to assess statistical significance in this subgroup). The ESC noted that the proportion of patients with PD-L1 CPS ≥10 did not differ significantly between OAC (48%) and OSCC (52%) patients, and the point estimates of benefit are similar between histologies, albeit not powered for significance (Table 5: HR point estimate = 0.73 [ITT]; 0.72 [OSCC]; 0.74 [OAC]). The PBAC agreed with the ESC that differing proportions of histologies in the Australian population are unlikely to make a substantial difference to the magnitude of benefit.
  4. The PSCR stated that an increase in median OS in the ITT population of 2.6 months (12.4 months vs. 9.8 months) is a valuable extension for patients with such a poor prognosis. The ESC and the PBAC noted the significant clinical benefit and agreed with the PSCR regarding the clinical significance of the increase in median OS.
  5. The PBAC considered that the claim of superior comparative effectiveness was reasonable based on the statistically and clinically significant benefits of pembrolizumab with respect to OS and PFS in the KN590 trial. The PBAC noted that the treatment benefit of pembrolizumab was less in patients with PD-L1 CPS <10 (HR 0.86) compared to patients with CPS≥10 (HR 0.62) and, while the study was not powered for this comparison, there are biologically plausible reasons for this observation. The PBAC considered the magnitude of the clinical benefit in the Australian population that would be treated with pembrolizumab is uncertain due to the proportion of patients with PD-L1 CPS <10/≥10 being unknown for the Australian population.
  6. The submission stated that, in terms of safety, analysis of the relative risk and risk difference has demonstrated that pembrolizumab in combination with chemotherapy has a higher incidence of grade 3‑5 AEs than chemotherapy indicating that the safety profile is inferior to that of chemotherapy. The submission quoted the TGA Delegate’s Overview: “The risks of pembrolizumab are therefore largely manageable with patient surveillance, treatment delays and supportive care in most patients, and are considered acceptable given the life-threatening nature of metastatic or locally advanced oesophageal carcinoma.”
  7. The PBAC agreed with the ESC, that the claim of inferior safety was reasonable and considered the additional toxicity (hyper-/hypothyroidism and pneumonitis) was manageable.

## Economic analysis

* 1. The submission presented a cost-utility analysis comparing pembrolizumab plus chemotherapy versus chemotherapy alone (cisplatin plus 5-FU). The submission did not present a stepped economic analysis and did not justify this omission. A summary of the model structure in the submission’s economic evaluation is presented in Table 8.

Table 8: Key components of the economic evaluation

| **Component** | **Description** |
| --- | --- |
| Type of analysis | Cost effectiveness analysis (Cost-utility analysis base case) |
| Outcomes | Quality-adjusted life years (QALY; base case), Life-years (LY; sensitivity analysis) gained |
| Time horizon | 7.5 years in the model base case based on a median follow up of 9.8 and 12.6 months in the chemotherapy and pembrolizumab plus chemotherapy arms of the KN590 trial, respectively. |
| Methods used to generate results | Cohort analysis of partitioned survival (i.e., area under the curve) |
| Health states | Three health states are included in the model: PFS, progressed disease and death |
| Cycle length | 1 week |
| Transition probabilities | Health state allocation over time was determined by PFS and OS curves from the KN590 trial (partitioned survival analysis) |
| Extrapolation method | Piecewise (models for all extrapolations)  *OS:* KM data up to 40 weeks, and then log-logistic for pembrolizumab plus chemo arm and log-normal for chemotherapy alone arm.  *PFS:* KM data up to 37 weeks, then exponential for pembrolizumab plus chemo and for chemotherapy alone  KM data only for time on treatment as all treatment curves reached zero before end of trial.  The model is highly sensitive to choice of extrapolation. |
| Health related quality of life | Progression free (0.811) and progressed (0.696) based on KN590 trial based EQ-5D responses.a |
| Inclusion of terminal care costs | $77,018.71 based on Goldsbury (2018) and adjusted for 2020 prices. May not be reflective of requested population, highly linked to extrapolation of OS benefit. May not be reasonable to include terminal care costs where survival is not expected to surpass the modelled time horizon. The PSCR provided a revised terminal care cost of $38,057. |

Source: Table 3.1-1, p 61 of the submission.

LY life year; OS = overall survival; PFS = progression free survival; QALY = quality-adjusted life year

a The Pre-Sub-Committee Response (PSCR) provided additional information on the derivation of the health utilities. The utilities were first cross walked to EQ-5D-3L and then mapped to Australia specific utilities.

* 1. The ESC noted a key limitation of the model is the applicability of the KN590 data on which it is based, as it was unclear how applicable the trial was to the Australian population (as discussed in paragraph 6.33).
  2. The model’s extrapolation method was explicitly selected to demonstrate a survival plateau effect of pembrolizumab. The evaluation noted the Advisory Committee Medicines (ACM) spoke positively of the long tail in survival for a subset of patients (ACM Minutes).The submission stated that single-piece models generated poorly fitting extrapolations of the data and that this is consistent with the findings of previous independent immuno-oncology (IO) validation studies that have shown that single-piece parametric functions underestimate empirically measured survival data for these therapies (Bullement 2019; Phan 2020; Ouwens 2019) as they inadequately represent the complex pattern of hazard functions or underlying mechanism of action of IO treatments (Ouwens 2019).
  3. The submission considered that the use of more flexible approaches to extrapolation is in line with current academic thinking both in analogous HTA markets (NICE DSU paper 21 2020) and Australian academics (Gray J, Sullivan T, Latimer NR et al 2021). It has also been suggested by the ESC in recent appraisals (darolutamide in July 2020; pembrolizumab in November 2020). The ESC noted that the sources referenced specifically refer to the use of flexible parametric models and cure models (where an assumption of a meaningful cured proportion is supported by the clinical evidence). The ESC noted an assumption underpinning the use of piecewise models is that the data within each section is sufficient for robust survival modelling (NICE DSU paper 21, 2020). The ESC noted that a piecewise extrapolation with a 40 week cut-off results in additional weight being attached to the tail of the Kaplan-Meier data i.e., the extrapolation is more sensitive to survival data informed by low patient numbers.
  4. The submission stated a piecewise approach was selected to represent an assumed plateau in survival. However, ESC considered modelling a survival plateau may not be supported by the clinical evidence presented in the submission. The ESC noted the assumption of a plateau in survival was based on observation of the KM curve (Figure 2)when only a small number of patients remain at risk. The ESC considered the piecewise approach may not be reasonable as it modelled a survival plateau which may not be realised in clinical practice. The piecewise approach in the base case (40 week cut-off) estimated a larger increment in overall survival (0.57 life years, undiscounted) compared to the one-piece approach (0.46 life years, undiscounted). The ESC considered the one-piece approach, in which survival models are fitted to the full data and a parametric model is used to extrapolate from Week 98, was a more appropriate, conservative method. The ESC noted using a one-piece approach, extrapolating from Week 98 with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemotherapy alone for OS and progression free survival [PFS]), rather than the piecewise model, increased the ICER from $95,000 to < $115,000/QALY in the base case to $115,000 to < $135,000/QALY. The ESC noted the revised extrapolation predicted survival differences at 7.5 years, which may still be optimistic.
  5. The PSCR justified using a cut-off point of 40 weeks for piecewise model fitting as being in line with the peak of the Chow test for OS in the pembrolizumab + chemotherapy arm. The PSCR stated that while the peak for the OS Chow test for the chemotherapy arm occurs at a later time point (i.e. after week 100); the Kaplan-Meier event data after this time point are too sparse to be used for parametric fitting.
  6. The pre-PBAC response acknowledged that while the Sponsor considered the piecewise extrapolation method to be the most appropriate approach to accurately capture the long-term benefits of immunotherapies, other extrapolation methods such as a one-piece approach, extrapolating from Week 98 with appropriate functions applied, may also account for the survival plateau seen with immunotherapies.
  7. The submission applied terminal care costs of approximately $77,000 (Goldsbury 2018) to the death state in the model. The estimate was specific to terminal phase of head and neck cancers, which was the highest of all of the 11 cancer types of terminal care cost estimates. No costs were included specific to oesophageal cancer. Goldsbury (2018) defined the cost of terminal phase care as all costs in the final year up to and including the death date. This, consequently, would likely constitute mostly double counted costs, as median survival in KN590 ITT population was 12.4 months for pembrolizumab plus chemotherapy and 9.8 months in the chemotherapy alone arm. The PSCR proposed a revised terminal care cost of $38,057 based on an estimate from Langton 2016 converted to 2021 prices. The ESC noted the Langton paper estimated that elderly patients with a variety of different types of cancers who died of cancer accrued $30,001 of costs in the last six months of life. The PSCR response reduced the price of pembrolizumab (from $'''''''''''''''/vial to $'''''''''''''''''/vial) to maintain an ICER of $95,000 to < $115,000/ QALY using the revised terminal care cost.
  8. Given that few patients would be expected to survive beyond 7.5 years, and there is no indication-specific clinical evidence to support that pembrolizumab plus chemotherapy would improve survival in the long term, and the issues with the cost estimate itself, it is likely more reasonable to assume that there would be no difference in terminal care costs between the treatment arms. Moreover, it is likely that treatment with pembrolizumab plus chemotherapy delays, rather than prevents, incurring terminal care costs in the proposed PBS population. Removing the cost of terminal care increased the ICER substantially. The ESC noted the impact of including terminal care costs on the ICER is driven by the difference in surviving proportions at the end of the model time horizon but ultimately this cost should accrue to all patients in both treatment arms. Additionally, the ESC noted the terminal care cost was based on the 12 months prior to death and considered this may result in double counting as some of these costs would have accrued in the progressed health state. The ESC noted excluding terminal care costs increased the ICER from $95,000 to < $115,000/QALY gained to $95,000 to < $115,000/QALY gained (based on the price proposed in the submission).
  9. Key drivers of the model are presented in Table 9.

Table 9: Key drivers of the model

| **Description** | **Method/Value** | **Impact (Base case ICER: ''''''''''''''''1)** |
| --- | --- | --- |
| Extrapolation | Piecewise extrapolations.  OS: KM data up to 40 weeks, and then log-logistic for pembrolizumab plus chemo arm and log-normal for chemotherapy alone arm.  PFS: KM data up to 37 weeks, then exponential for pembrolizumab plus chemo and for chemotherapy alone | High, favours pembrolizumab.  Using KM data to Week 98, then a one-piece with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemotherapy alone for OS and PFS) rather than piecewise models, leads to an ICER of '''''''''''''''''''''''2. |
| Time horizon | 7.5 years | High, likely favours pembrolizumab. Reducing time horizon to 5 years increases ICER to ''''''''''''''''''''''2. The ESC considered a 7.5 year time horizon may be optimistic for this population. |
| Terminal care costs | Cost of $77,019 added to death state based on Goldsbury (2018). The PSCR provided a revised terminal care cost of $38,057. | High, favours pembrolizumab. Removing terminal care costs increases the ICER to '''''''''''''''''''''''1. Using the revised terminal care cost resulted in an ICER of ''''''''''''''''''''''1. |

Source: pp99-102 of the submission.

ICER = incremental cost effectiveness ratio; KM = Kaplan-Meier; OS = overall survival; PFS = progression free survival; PSCR = Pre-Sub-Committee Response.

*The redacted values correspond to the following ranges:*

*1 $95,000 to < $115,0002 $115,000 to < $135,000*

* 1. Model results are presented in Table 10.

Table 10: Results of the economic evaluation

| **Component** | **Pembrolizumab plus chemo** | **Chemotherapy alone** | **Increment** |
| --- | --- | --- | --- |
| Costs ($) | '''''''''''''''''''''' | $79,712 | '''''''''''''''''' |
| Lys | 1.7102 | 1.2245 | 0.4857 |
| QALYs | 1.2858 | 0.9147 | 0.3711 |
| **Incremental cost/extra LY gained** | | | **'''''''''''''''''1** |
| **Incremental cost/extra QALY gained** | | | **'''''''''''''''2** |

Source: Table 3.8-3, p98 and table 3.8-4, p99 of the submission.

Chemo = chemotherapy; LY = life year; QALY = quality adjusted life year.

*The redacted values correspond to the following ranges:*

*1 $55,000 to < $75,000*

*2 $95,000 to < $115,000*

* 1. Sensitivity analyses for the economic evaluation are presented in Table 11.

Table 11: Results of sensitivity analyses (based on pembrolizumab price proposed in the submission)

| **Analyses** | | | | **Incremental cost ($)** | **Incremental QALY** | **ICER** |
| --- | --- | --- | --- | --- | --- | --- |
| Base case | | | | ''''''''''''''''' | 0.37 | ''''''''''''''''''''1 |
| Trial population (ITT in BC) | OSCC | | | '''''''''''''''''' | 0.41 | '''''''''''''''''''''2 |
|  | PD-L1+ (CPS ≥10) | | | ''''''''''''''''''' | 0.53 | ''''''''''''''''''3 |
|  | OSCC and PD-L1+ (CPS ≥10) | | | ''''''''''''''''''''' | 0.61 | ''''''''''''''''''''3 |
| Discount rate (5% in BC) | 0% | | | '''''''''''''''''' | 0.43 | '''''''''''''''''''''2 |
| 3.5% | | | '''''''''''''''''''' | 0.39 | '''''''''''''''''''2 |
| Time horizon (7.5 years in BC) | 5 years | | | '''''''''''''''''''' | 0.30 | '''''''''''''''''''''''5 |
| 10 years | | | ''''''''''''''''''' | 0.42 | ''''''''''''''''''2 |
| Terminal care costs | Removed | | | '''''''''''''''''' | 0.37 | ''''''''''''''''''''1 |
| Revised cost provided in PSCR | | | '''''''''''''''''' | 0.37 | '''''''''''''''''''''4 |
| No relative dose intensity applied for 1L treatments | | | | '''''''''''''''''''' | 0.37 | '''''''''''''''''''''''4 |
| Time to death utility approach rather than progression based | | | | '''''''''''''''''''' | 0.42 | ''''''''''''''''''''2 |
| Utility increment for treatment | | | | '''''''''''''''''' | 0.40 | ''''''''''''''''''''2 |
| Market share for blended comparator (100% 5-FU + cisplatin in BC) | Assumed equal market share (12.5%) across chemotherapy arm and blended chemotherapy comparators | | | '''''''''''''''''' | 0.37 | '''''''''''''''''2 |
| Assumed 50%/50% CAPOX/FOLFOX | | | '''''''''''''''''' | 0.37 | '''''''''''''''''2 |
| Assumed 80%/20% FOLFOX/ CAPOX | | | ''''''''''''''''' | 0.37 | '''''''''''''''''''''2 |
| Apply OS waning effect for pembrolizumab + chemotherapy (OS converges) | | | | ''''''''''''''''' | 0.37 | '''''''''''''''''''''4 |
| **Extrapolation** | | | | | | |
| Pembrolizumab + chemotherapy and chemo alone: Use Week 32 piecewise with the best fitting function (log-logistic for pembrolizumab + chemotherapy and Gompertz for chemo alone according to AIC and BIC) for OS rather than Week 40 piecewise | | | | ''''''''''''''''''''' | 0.35 | ''''''''''''''''''''''''4 |
| Pembrolizumab + chemotherapy and chemo alone for OS and PFS: Use Week 98 one-piece with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemo alone for OS and PFS) rather than piecewise models | | | | '''''''''''''''''''' | 0.31 | ''''''''''''''''''''''''5 |
| **Multivariate analyses** | | | | | | |
| Use Week 98 one-piece with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemo alone for OS and PFS) | | | And remove terminal care costs | '''''''''''''''''' | 0.31 | '''''''''''''''''''''''5 |
| Use Week 98 one-piece with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemo alone for OS and PFS) | | | Revised terminal care costs in PSCR | '''''''''''''''''' | 0.31 | '''''''''''''''''''''''5 |
| Use Week 98 one-piece with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemo alone for OS and PFS) | | And remove terminal care costs | 5 year time horizon | '''''''''''''''''' | 0.27 | ''''''''''''''''''''6 |

Source: 3.9-1, pp101-102 of the submission.

AIC = Akaike Information Criteria; BC = base case; BIC = Bayesian Information Criteria; CPS = combined positive score; 5-FU = 5‑fluorouracil; ICER = incremental cost effectiveness ratio; ITT = intention-to-treat; OSCC = oesophageal squamous cell carcinoma; OS = overall survival; PD-L1 = programmed death-ligand 1; PFS = progression free survival; PSCR = pre-sub-committee response; QALY = *quality adjusted life year.*

*The redacted values correspond to the following ranges:*

*1 $95,000 to < $115,000*

*2 $75,000 to < $95,000*

*3 $55,000 to < $75,000*

*4 $95,000 to < $115,000*

*5 $115,000 to < $135,000*

*6 $135,000 to < $155,000*

* 1. The ESC noted the multivariate analysis that (i) used the Week 98 one-piece with the best fitting function (log-logistic according to AIC and BIC for pembrolizumab + chemotherapy and chemo alone for OS and PFS) and (ii) removed the terminal care costs resulted in an ICER of $115,000 to < $135,000/QALY gained. The ESC considered this scenario may provide a more reasonable base case, noting the predicted survival differences at 7.5 years may still be optimistic. The ESC noted applying a 5 year time horizon increased the ICER from $115,000 to < $135,000/QALY to $135,000 to < $155,000/QALY. The PBAC noted that applying the revised price proposed in the PSCR resulted in ICERs of $115,000 to < $135,000/QALY (7.5 year time horizon) and $135,000 to < $155,000/QALY (5 year time horizon).
  2. The pre-PBAC response stated that:
* The 7.5 year time horizon within the alternative ESC base case adequately captures the benefits in those patients who have a complete response and who have a longer-term benefit.
* The terminal care costs should not be excluded . The Sponsor maintained that the revised costs based on the publication by Langton et al. 2016 should be used in the economic model.
* The application of a Week 98 one-piece curve may be appropriate as a more conservative method, provided the appropriate survival functions are applied to each arm . The Sponsor agreed with the ESC advice of applying a log-logistic function to the pembrolizumab + chemotherapy arm, as it is representative of the survival plateau seen with immunotherapies in oesophageal cancer and other cancer types, but commented that the application of a log-logistic function to the standard of care arm would result in a greater proportion of patients alive than is clinically plausible. The Sponsor noted that only a very small number of patients receiving chemotherapy would typically be alive at 2-3 years, with none surviving greater than 5 years.

## Drug cost/patient/course $''''''''''''' to $''''''''''''''

* 1. The submission calculated the average cost per patient of pembrolizumab accrued in the economic evaluation. This was calculated by multiplying the DPMQ of pembrolizumab times the KN590 relative dose intensity and estimating the total cost per treatment using the time on treatment survival analysis. During the evaluation, this method of calculating treatment cost was applied to the chemotherapy components of the proposed regimen, cisplatin plus 5-FU and that of the comparator as well.
  2. Drug costs per patient are presented in Table 12.

Table 12: Drug cost per patient for pembrolizumab and chemotherapy

|  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Drug and dose** | **Cost per cycle** | | | **Number of cycles** | | | **Total cost per course** | | |
| **Modela** | **KN590a** | **Financials** | **Modelb** | **KN590** | **Financial** | **Model** | **KN590** | **Financials** |
| **Pembrolizumab plus cisplatin plus 5-FU** | | | | | | | | | |
| Pembrolizumab | $''''''''''''''''''''''' | | $'''''''''''''''''''''' | 11.7 | 11.0 | 10.96 | $''''''''''''''''''''' | $''''''''''''''''''''''' | $'''''''''''''''''''''' |
| Cisplatin | $83.62 | | Not included | 4.79 | 4.7 | Not included | $400.63 | $393.01 | Not included |
| 5-FU | $115.41 | | 8.73 | 8.0 | $1,007.99 | $923.28 |
| **Total** | | | | | | | **$''''''''''''''''''** | **$'''''''''''''''''** | $'''''''''''''''''''''' |
| **Chemotherapy alone** | | | | | | | | | |
| Cisplatin | $93.68 | | Not included | 4.69 | 4.7 | Not included | $439.26 | $440.30 | Not included |
| 5-FU | $123.61 | | 7.60 | 7.1 | $938.83 | $877.63 |
| **Total** | | | | | | | **$1,378.10** | **$1,317.93** |

Source: Table 14.1-35, p339 of the KN590 CSR; ‘Drug Costs’ worksheet of “Sec 3 Workbook’’ attached to submission.

Note: small discrepancies between presented results and products of their presented factors are due to rounding.

a cost per cycle includes adjustment with trial based relative dose intensity: as follows, 93.4% for pembrolizumab, 71.8% for 5-FU in the treatment arm, and 58.2% for cisplatin in the treatment arm, 76.9% for 5-FU in the control arm, and 65.2% for cisplatin in the control arm.

b Cycles of treatment back-calculated in the model by dividing undiscounted total cost of treatment by the dose adjusted cost per cycle of treatment

## Estimated PBS usage & financial implications

* 1. This submission was considered by DUSC.
  2. The submission took an epidemiological approach to estimate use and financial impact.
  3. A summary of inputs and their data sources are presented in Table 13.

Table 13: Data sources and inputs in the financial estimates

| **Data** | **Value** | **Source** | **Comment** |
| --- | --- | --- | --- |
| **Eligible population** | | | |
| Incident patients with oesophageal cancer | '''''''''''''1 in Year 1 | AIHW (2018)  With growth based on average historical growth rate between 2010 and 2021 | Yearly growth of 1.4%. The DUSC noted GOJ patients would not be included in this estimate and considered that Siewert Type I GOJ AC patients should be explicitly included in the financial estimates |
| Prevalent patients with oesophageal cancer | '''''''''''''1 in Year 1 | AIHW (2018) | Yearly growth of 1.4%. The DUSC noted GOJ patients would not be included in this estimate and considered that Siewert Type I GOJ AC patients should be explicitly included in the financial estimates. |
| Proportion of incidence cases that is adenocarcinoma | 74% | Nguyen (2019) | Reasonable. |
| Proportion of patients with Siewert type I AC of the GOJ who are HER2+ | 5% | Based on Expert opinion | The submission only stated that this was based on “KSL advice” and no further details of this opinion were presented. |
| Stage | Stage I: 16%  Stage II/III: 57%  Stage IV: 27% | Nguyen (2019) | Reasonable, although this groups Stages II/III together, which may overestimate the incident population. |
| Proportion resectable | 64% | Nguyen 2019: Proportion of patients treated with curative intent |  |
| ECOG PS 0-1 | 80% | Expert opinion | The submission only stated that this was based on “KSL advice” and no further details of this opinion were presented. |
| Uptake | 90% | Assumption | Could potentially be lower in the Australian treatment population. |
| **Treatment utilisation** | | | |
| Pembrolizumab administrations per patient dispensed | 10.96 per patient | 11.73 x 93.4% dose intensity | Consistent with economic model. |
| **Costs** | | | |
| Chemotherapy | Not included in submission | -- | The submission stated that because pembrolizumab will be used adjunctively to platinum + 5‑FU, the current comparator, no estimated change in use of other therapies was considered and so this was not included.  If the claim of improved OS and PFS is accepted, this would lead to longer concomitant use of chemotherapy and thus the omission of these costs underestimates costs associated with listing pembrolizumab. |

Source: pp102-113. AC = adenocarcinoma; AIHW = Australian Institute for Health and Welfare; ECOG PS = Eastern Cooperative Oncology Group Performance Status; 5-FU = 5-fluorouracil; GOJ = gastro-oesophageal junction; HER2+ = human epidermal growth factor receptor 2 positive; MBS = Medicare Benefits Schedule; OS = overall survival; PFS = progression free survival.

*The redacted values correspond to the following ranges:*

*­1 500 < 5,000*

* 1. The submission estimated the expected population by calculating patients in four separate treatment pathways:
* Incident patients Stage III unresectable/ Stage IV patients at diagnosis.
* Patients recurrent from Stage I and II/III resectable.
* Prevalent patients.
* Grandfathered patients.
  1. The DUSC considered that, overall, the approach to estimating the number of eligible patients using the four separate treatment pathways was reasonable. The DUSC noted that the epidemiological estimate of the eligible population excluded Siewert Type I GOJ AC patients, resulting in an underestimate of usage and financial implications.
  2. The pre-PBAC response accepted the advice by the DUSC that the epidemiological approach to the financial estimates only included the incidence of oesophageal cancer (ICD-10 code C15) and neglected to include the incidence of GOJ adenocarcinoma which is captured under gastric cancer (ICD-10 code C16). The pre-PBAC response stated there is a lack of published data that identifies the proportion of GOJ adenocarcinoma, and the Siewert type classification is not consistent across clinical practice with some clinicians referring to Siewert type 1 as distal oesophageal cancer, and Siewert type II-III as gastric cancer. The pre-PBAC response presented revised estimates to capture the incidence of GOJ AC, based on the following assumptions:
* GOJ patients account for approximately 50% of all gastric cancer patients, based on clinician advice.
* 23-40% of GOJ adenocarcinoma patients are Siewert type 1, based on published literature (midpoint of 30% used, based on clinician advice).
* The proportion of HER2-negative patients was appropriately applied to the GOJ AC population (rather than to the oesophageal population as applied in the submission).
  1. Estimated use and financial implications, including the revised estimates, are presented in Table 14.

Table 14: Estimated use and financial implications

|  | **Year 1** | **Year 2** | **Year 3** | **Year 4** | **Year 5** | **Year 6** |
| --- | --- | --- | --- | --- | --- | --- |
| **Estimated extent of use** | | | | | | |
| Number of patients treateda | ''''''''''1 | '''''''''1 | '''''''''1 | ''''''''''1 | ''''''''''1 | ''''''''1 |
| Number of scripts dispensedb | ''''''''''''''2 | ''''''''''''2 | ''''''''''''''2 | ''''''''''''2 | ''''''''''''2 | ''''''''''''''2 |
| **Estimated financial implications for other medicines: None.** | | | | | | |
| **Net financial implications** | | | | | | |
| Net cost to PBS/RPBS | **''''''''''''''''''''''''''**3 | **''''''''''''''''''''''''''**3 | **'''''''''''''''''''''''''** 3 | **''''''''''''''''''''''** 3 | **'''''''''''''''''''''**3 | **'''''''''''''''''''''''''** 3 |
| Net cost to MBSc | '''''''''''''''''''''5 | ''''''''''''''''''''5 | ''''''''''''''''''''''5 | '''''''''''''''''''''5 | '''''''''''''''''''''5 | ''''''''''''''''''''5 |
| Net cost to Government | **''''''''''''''''''''''''**3 | **'''''''''''''''''''''**3 | **'''''''''''''''''''''''**3 | **''''''''''''''''''''''''''**3 | **'''''''''''''''''''''**3 | **''''''''''''''''''''''''**3 |
| **Revised estimates in the pre-PBAC response** | | | | | | |
| Number of patients treated | '''''''''1 | '''''''''1 | '''''''''1 | '''''''''1 | ''''''''1 | ''''''''1 |
| Net cost to PBS/RPBSd | **'''''''''''''''''''''''''**4 | **'''''''''''''''''''''''''**4 | **''''''''''''''''''''''''**4 | **''''''''''''''''''''''''**4 | **''''''''''''''''''''''**4 | **''''''''''''''''''''''**4 |

a Estimate includes < 500 grandfathered patients.

b Assuming 10.96 scripts per year as estimated by the submission.

c MBS 13950 applied.

d The effective vial price for pembrolizumab reduced from $'''''''''''''''''''/vial to $''''''''''''''''''''''/vial in the PSCR. The new price was applied to the revised estimates in the pre-PBAC response.

Source: Table 4.2-7, p107, Table 4.2-9, p109, Table 4.2-16, pp110-112, and Table 4.4-1, p112, and Table 4.5-3-p113 of the submission. MBS = Medicare Benefits Schedule; PBS = Pharmaceutical Benefits Scheme; RPBS = Repatriation Pharmaceutical Benefits Scheme; PSCR = Pre-Sub-Committee Response.

*The redacted values correspond to the following ranges:*

*1 500 to < 5,000*

*2 5,000 to < 10,000*

*3 $20 million to < $30 million*

*4 $30 million to < $40 million*

*5 $0 to < $10 million*

* 1. The total net cost to the PBS/RPBS was estimated at $20M to < $30M in Year 1, decreasing to $20M to < $30M in Year 2, before increasing back to $20M to < $30M in Year 6. The Sponsor’s revised total net cost to the PBS/RPBS, capturing GOJ AC patients, was estimated at $30M to < $40M in Year 1, decreasing to $30M to < $40M in Year 2, before increasing to $30M to < $40M in Year 6.
  2. The PBAC considered there was a high risk of pembrolizumab being used in patients with Siewert Type II or Type III AC of the GOJ and gastric cancer. Additionally, the PBAC noted there would be an impact on utilisation and financial estimates should nivolumab be recommended for PBS listing in a similar population.

## Quality Use of Medicines

* 1. The submission outlined a number of activities intended to promote safe and effective use of pembrolizumab in clinical practice. These include the development of materials about how to identify and manage potential treatment-related AEs in particular immune-related AEs, as well as education activities and a 1-800 medical information service.

## Financial Management – Risk Sharing Arrangements

* 1. The submission stated that the Sponsor is willing to enter into an RSA with the Commonwealth on sharing the costs of the Commonwealth subsidy for supply of pembrolizumab for the treatment of recurrent or metastatic OC and GOJA, in order to manage any risk to the overall cost to the PBS.
  2. For the purpose of the RSA, the sponsor agrees to reimburse the Commonwealth with a proportion of the treatment costs of pembrolizumab, should use exceed the subsidisation cap in that year. Following a positive recommendation, the submission stated that the sponsor is committed to working with the Commonwealth to finalise the specific parameters for inclusion in the RSA including the rebate proportion to be paid.

*For more detail on PBAC’s view, see section 7 PBAC outcome.*

1. PBAC Outcome
   1. The PBAC did not recommend the listing of pembrolizumab in combination with chemotherapy for the treatment of locally advanced (Stage III) or metastatic (Stage IV) oesophageal adenocarcinoma (OAC) or oesophageal squamous cell carcinoma (OSCC), or human epidermal growth factor receptor 2 (HER2)-negative adenocarcinoma (AC) of the gastro-oesophageal junction (GOJ). The PBAC noted that the clinical need for effective treatments in this therapeutic area is high, given the poor prognosis for patients and the poor efficacy and high toxicity of current treatments. The PBAC considered the evidence presented demonstrated treatment with pembrolizumab resulted in a clinically meaningful improvement in progression free survival (PFS) and overall survival (OS). However, the PBAC considered the incremental cost-effectiveness ratio (ICER) in this setting at the proposed price was high and moderately uncertain.
   2. The PBAC noted the consumer input supported the high clinical need for additional effective treatment options for this population.
   3. The submission nominated chemotherapy alone, represented by cisplatin + 5-FU- based regimens, as the comparator. The PBAC noted nivolumab plus chemotherapy was considered at the same PBAC meeting and agreed with the submission that it is a near-market comparator given the overlap in patient populations.
   4. The PBAC was satisfied that pembrolizumab provided, for some patients, a significant improvement in efficacy over the nominated comparator of chemotherapy (cisplatin + 5-FU). The PBAC noted that this combination could be considered representative of all platinum- and 5-FU-based regimens, including the commonly used FOLFOX and XELOX/CAPOX regimens.
   5. The PBAC noted that the TGA indication and trial population for pembrolizumab reflected a subset of the full patient population likely to be considered clinically appropriate for checkpoint inhibitor treatment. The PBAC indicated its preference for an aligned, simpler restriction for checkpoint inhibitors that reflected likely clinical practice across gastric, GOJ and oesophageal cancers, while noting the different TGA indications.
   6. The submission was based on one head-to-head randomised, phase 3, double-blind multi-centre trial (Keynote 590 [KN590]; N=749) comparing pembrolizumab + chemotherapy to cisplatin + 5-FU, in patients with locally advanced unresectable or metastatic OAC or OSCC or advanced/metastatic Siewert type 1 AC of the GOJ. The PBAC noted that there was a statistically significant OS benefit associated with pembrolizumab (HR=0.73, 95% CI: 0.62, 0.86) with the median survival improving from 9.8 to 12.4 months and an improvement in PFS (5.8 months to 6.3 months, HR=0.65, 95% CI: 0.55, 0.76). The PBAC considered that the extent of OS benefit was clinically meaningful in the context of the poor prognosis of this patient population. The PBAC noted the OS benefit in patients treated with pembrolizumab compared to chemotherapy alone was supported by a higher ORR, longer duration of response and maintenance of QoL.
   7. While the PBAC acknowledged the clinically relevant improvement in PFS and OS with pembrolizumab in the KN590 trial, it considered the magnitude of the benefit in the Australian population to be uncertain. It noted the benefit of pembrolizumab is less in patients with PD-L1 CPS <10 and the proportion of patients with CPS<10 in the Australian population was unknown. The PBAC noted that while the proportions of the OAC/OSCC histologies differed in the KN590 trial compared with the Australian population, it was unlikely to make a substantial difference to the magnitude of the benefit from pembrolizumab.
   8. The PBAC noted that pembrolizumab in combination with chemotherapy had a higher incidence of grade 3‑5 AEs than chemotherapy alone, indicating that the safety profile is inferior to that of chemotherapy. The PBAC considered that the claim of inferior safety was reasonable, and that the additional toxicity (hyper-/hypothyroidism and pneumonitis) was manageable.
   9. The PBAC noted the base case ICER presented in the submission was $95,000 to < $115,000 per QALY gained. The PBAC considered this ICER to be high and moderately uncertain, with the economic model sensitive to time horizon, method of extrapolation and the inclusion of terminal care costs.
   10. The time horizon in the base case economic model in the submission was 7.5 years. The PBAC considered a 7.5 year time horizon was optimistic and a 5 year time horizon would be preferable given the median duration of follow up in KN590 was 9.8 months and 12.6 months in the chemotherapy and pembrolizumab arms, respectively, and the poor prognosis of the patient population. However, the PBAC noted it had previously recommended a 5 year time horizon in the second-line treatment setting for a similar patient population and considered a 7.5 year time horizon may be reasonable, given the earlier treatment setting, if other model assumptions were conservative.
   11. The economic model in the submission extrapolated OS and PFS using a two-piece extrapolation method using KM data up to 40 weeks (OS) or 37 weeks (PFS) and then a log-logistic function for pembrolizumab and log-normal for chemotherapy alone for OS, and an exponential function for both pembrolizumab and chemotherapy alone for PFS. The PBAC agreed with the ESC that the use of a two-piece extrapolation method resulted in additional weight being attached to the tail of the Kaplan-Meier data (where patient numbers are low and the data is less reliable) and that a one-piece extrapolation method fitted to the full KM curve was more reliable. The PBAC noted the ESC preferred approach was a one-piece extrapolation from Week 98 with the best fitting function (log-logistic for pembrolizumab and chemotherapy alone for OS and PFS). The PBAC noted the pre-PBAC response accepted this approach was appropriate for the pembrolizumab arm as it resulted in some plateauing of survival but did not consider it appropriate for the chemotherapy arm as it may overestimate survival. The PBAC agreed with the pre-PBAC response but noted the pre-PBAC response did not propose an alternative extrapolation method for the chemotherapy arm.
   12. The economic model in the submission applied a terminal care cost of $77,000 (reduced to $38,057 in the PSCR) to the death health state. The PBAC noted the ESC consideration of the inclusion of terminal care costs (paragraphs 6.43 and 6.44) and agreed with the ESC that terminal care costs should be excluded from the economic model.
   13. The PBAC noted the revised financial estimates provided with the pre-PBAC response addressed the exclusion of patients with AC of the GOJ in the financial estimates (which was the main issue raised by the DUSC). However, the PBAC considered it was uncertain whether the assumptions applied were reasonable (in particular, whether GOJ cancer accounts for 50% of gastric cancer and whether 30% of GOJ cancer patients are Type I Siewert). The PBAC considered this could be further addressed in a resubmission, noting the PBAC preference for an aligned restriction across all gastric, GOJ and oesophageal cancers (see paragraph 7.6), the high risk of use of pembrolizumab outside of Type 1 Siewert GOJ (see paragraph 3.7) and the likely overlap in patient populations for checkpoint inhibitors (see paragraph 5.3).
   14. The PBAC noted that the Sponsor is willing to enter into an RSA with the Commonwealth on sharing the costs in order to manage any risk to the overall cost to the PBS.
   15. The PBAC considered the outstanding issues could be easily resolved in a simple resubmission for pembrolizumab. The PBAC also considered pembrolizumab addresses a high and urgent unmet clinical need and was expected to provide a substantial and clinically relevant improvement in efficacy over any alternative therapies. Therefore, the PBAC considered an early resolution pathway would be acceptable. If the sponsor accepts this pathway, the following changes may address these outstanding issues without requiring further re-evaluation.

* Provide an economic model based on (i) a 7.5 year time horizon (ii) one-piece extrapolation from Week 98 using the log-logistic function for OS and PFS for the pembrolizumab treatment arm and an appropriate function for the chemotherapy alone treatment arm and (iii) exclusion of terminal care costs.
* Propose a price reduction to achieve an ICER of $55,000 to < $75,000 to $75,000 to < $95,000/ QALY;
* Provide revised financial estimates based on the Section 4 model included with the pre-PBAC response, incorporating the revised price and addressing the issues raised in paragraph 7.14 ;
* Propose an appropriate RSA to manage the risk of use outside the proposed patient population included in the resubmission.

The early resolution resubmission must be lodged by week 7 of the current PBAC cycle or the next cycle. If the issues cannot be addressed by the sponsor in a simple resubmission and the early resolution timing is not acceptable, a standard re-entry pathway is available

* 1. The PBAC noted that this submission is eligible for an Independent Review.

**Outcome:**

Not recommended

1. Context for Decision

The PBAC helps decide whether and, if so, how medicines should be subsidised through the Pharmaceutical Benefits Scheme (PBS) in Australia. It considers applications regarding the listing of medicines on the PBS and provides advice about other matters relating to the operation of the PBS in this context. A PBAC decision in relation to PBS listings does not necessarily represent a final PBAC view about the merits of the medicine or the circumstances in which it should be made available through the PBS. The PBAC welcomes applications containing new information at any time.

1. Sponsor’s Comment

MSD is pleased that the PBAC has recognised the high added therapeutic value that pembrolizumab offers patients with oesophageal/GOJ cancer. MSD is looking forward to working with the department to provide access to pembrolizumab for this patient group as early as possible through the early resolution pathway.

1. Saltzman JR. Clinical manifestations, diagnosis, and staging of esophageal cancer. UptoDate. 2021 [last updated July 23, 2021].

   https://www.uptodate.com/contents/clinical-manifestations-diagnosis-and-staging-of-esophageal-cancer?search=esophageal%20cancer&source=search\_result&selectedTitle=1~150&usage\_type=default&display\_rank=1 [↑](#footnote-ref-1)
2. Nguyen T-MN; Hummel R; Bright T; Thompson SK; Tornqvist B; Watson DI. Pattern of care for cancer of the oesophagus in a western population. ANZ J Surg 89 (2019) E15–E19. [↑](#footnote-ref-2)
3. Cherny NI, Dafni U, Bogaerts J, et al: ESMO-Magnitude of Clinical Benefit Scale version 1.1. Annals of Oncology 28:2340-2366, 2017. [↑](#footnote-ref-3)