



مركز الكويت لمكافحة السرطان
Kuwait Cancer Control Center

pemBROLizumab (Keytruda®)



Ministry of Health

Name:

File #:

Ht (cm):

Nationality:

Civil ID:

Wt (Kg):

Gender/Age:

DOB:

BSA (m²):

Indication(s): Multiple malignancies, specify: _____

Central line: Available NA

Allergies: NKA Yes, specify; _____

Parameters: Initiate treatment only if ANC ≥ 1000; HB ≥ 80; Plt ≥ 100,000; CrCl > 45 ml/min.

Pre-treatment Medications: (30-60 min before starting treatment)

Chlorphenamine 10 mg PO/IV

Standard Protocol:

| DRUG | DOSE | ADMINISTRATION | DAYS |
|--|--------|------------------------------|------|
| pemBROLizumab | 200 mg | IV in 250 mL NS over 30 min. | D1 |
| To be repeated every 3 weeks until disease progression or intolerable toxicity. | | | |

Treatment Description:

| Cycle | Date | pemBROLizumab | Physician | Consultant |
|-------|------|---------------|-----------|------------|
| C# __ | | | | |
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| C# __ | | | | |
| C# __ | | | | |

Important Notes:

Reported grade 3/4 toxicities: None Hematological Non-Hematological

If yes; Did it indicate hospitalization? Yes No

Did it indicate chemo-delay for ≥ 7 days? Yes No

Did it indicate dose reduction? Yes No

Did it indicate G-CSF support? Yes No