

APIDRYL[®]
▲ **20/80/160 MG**
DOCETAXEL INJECTION, USP

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DOCETAXEL INJECTION, USP

**A CHEMOTHERAPY DRUG USUALLY GIVEN TO TREAT BREAST CANCER,
PROSTATE CANCER AND NON-SMALL CELL LUNG CANCER.**

CLASS - ANTINEOPLASTIC

INDICATIONS AND USAGE

Apidryl[®] DOCETAXEL INJECTION, USP is a microtubule inhibitor indicated for:

Breast Cancer (BC): single agent for locally advanced or metastatic BC after chemotherapy failure; and with doxorubicin and cyclophosphamide as adjuvant treatment of operable node-positive BC

Non-Small Cell Lung Cancer (NSCLC): single agent for locally advanced or metastatic NSCLC after platinum therapy failure; and with cisplatin for unresectable, locally advanced or metastatic untreated NSCLC

Hormone Refractory Prostate Cancer (HRPC): with prednisone in androgen independent (hormone refractory) metastatic prostate cancer.

DOSAGE AND ADMINISTRATION

Administer in a facility equipped to manage possible complications (e.g., anaphylaxis). Administer intravenously over 1 hr every 3 weeks.

PVC equipment is not recommended..

BC locally advanced or metastatic: 60 mg/m² to 100 mg/m² single agent

BC adjuvant: 75 mg/m² administered 1 hour after doxorubicin 50 mg/m² and cyclophosphamide 500 mg/m² every 3 weeks for 6 cycles (2.1)

NSCLC: after platinum therapy failure: 75 mg/m² single agent

NSCLC: chemotherapy-naive: 75 mg/m² followed by cisplatin 75 mg/m²

HRPC: 75 mg/m² with 5 mg prednisone twice a day continuously

For all patients:

Premedicate with oral corticosteroids

Adjust dose as needed

DOSAGE FORMS AND STRENGTHS : Apidryl[®] DOCETAXEL INJECTION, USP

20 mg/2 mL single use vial

80 mg/8 mL multi-use vial

160 mg/16 mL multi-use vial