



مركز الكويت لمكافحة السرطان
Kuwait Cancer Control Center

MAI (IFOSFamide / Mesna / ADRIAmycin)



Ministry of Health

Name: _____ **File #:** _____ **Ht (cm):** _____
Nationality: _____ **Civil ID:** _____ **Wt (Kg):** _____
Gender/Age: _____ **DOB:** _____ **BSA (m²):** _____

Indication(s): Advanced Soft Tissue Sarcoma.

Central line: Available NA

Allergies: NKA Yes, specify; _____

Parameters: Initiate treatment only if ANC \geq 1500; HB \geq 80; Plt \geq 100,000; CrCl $>$ 45 ml/min.
 Baseline ECHO and/or MUGA scan before initiation of anthracycline-containing regimen.
 Date of pre-treatment ECHO and/or MUGA scan is _____. LVEF is ____ %.

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

Akynzeo 1 Capsule PO (300 mg NETUpitant/0.5 mg PALONOssetron) on Day 1 and Day 3
 Dexamethasone 10 mg PO/IV

Standard Protocol:

DRUG	DOSE	ADMINISTRATION	DAYS
DOXOrubicin	25 mg/m ²	IV in 100 mL NS over 30 min.	D1, 2, 3
IFOSFamide	3000 mg/m ²	IV in 500 mL NS over 60 min.	D1, 2, 3
Mesna	1800 mg/m ²	In 3 divided doses at 0, 4, & 8 hr from starting IFOSFamide,	D1, 2, 3

To be repeated every 3 weeks for 6 cycles.

Special instructions: The maximum cumulative dose of DOXOrubicin is 450 mg/m² (in normal cardiac function) and 350 mg/m² (in case of cardiac dysfunction or exposed to mediastinal IR).

Treatment Description:

Pre-hydration: 1 Liter NS IV over 2 hrs.

Cycle	Day	Date	DOXOrubicin	IFOSFamide	Mesna
C# __	D1				
	D2				
	D3				

Post-hydration: 1 Liter NS IV over 1 hrs.

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 \geq 7 days? Yes No
 Did it indicate dose reduction? Yes No
 Did it indicate G-CSF support? Yes No

Physician (Stamp and signature)

Consultant (Stamp and signature)