ADULT ORAL Updated: August 16, 2023

Regimen Reference Order - SARC - ripretinib

ARIA: SARC - [ripretinib]

Planned Course: Once daily until disease progression or unacceptable toxicity (1 cycle = 30 days)

Indication for Use: Gastrointestinal Stromal Tumor, Advanced/Metastatic

Proceed with treatment if:

- ANC equal to or greater than $1 \times 10^9/L$ AND Platelets equal to or greater than $75 \times 10^9/L$
- Total bilirubin less than or equal to 1.5 times upper limit of normal
- AST/ALT less than or equal to 3 times upper limit of normal
 - Contact Physician if parameters not met

SEQUENCE OF MEDICATION ADMINISTRATION

Treatment Regimen – SARC – ripretinib			
Drug	Dose	CCMB Administration Guideline	
ripretinib	150 mg	Orally once daily with or without food Swallow whole (Self-administered at home)	
ripretinib (QINLOCK®) a Classification: Cytotoxi	available dosage strength c, Hazardous	n: 50 mg tablet	

REQUIRED MONITORING

Cardiac monitoring

- · LVEF at baseline and as clinically indicated
- Blood pressure at baseline and as clinically indicated

Baseline and throughout treatment (at least every 3 months)

• CBC, serum creatinine, urea, electrolytes, liver enzymes and total bilirubin as per Physician Orders

Recommended Support Medications					
	Drug	Dose	CCMB Administration Guideline		
	None required				

INSTRUCTIONS FOR PATIENT

- Patients should be instructed to report rash or any new or changing skin lesions to their cancer team
- ripretinib has potential for drug-drug interactions. Patients should notify clinic prior to starting any new medication
- Avoid grapefruit and grapefruit juice, Seville oranges (i.e. orange marmalade) and starfruit
- Reinforce applicable safe handling precautions of medication, blood and body fluids for 48 hours after completion of ripretinib



ADULT ORAL SARC – ripretinib

ADDITIONAL INFORMATION

- ripretinib can cause cardiac toxicity including cardiac dysfunction, ischemic events and hypertension
- ripretinib may cause hand-foot syndrome, alopecia and arthralgia/myalgia
- ripretinib may increase the risk of new primary cutaneous malignancies including squamous cell carcinoma and melanoma
- Temporary interruption of ripretinib is recommended in patients undergoing surgical procedures due to the potential for impaired wound healing

