Indication	Adjuvant treatment of adult patients who are at significant risk of relapse following resection of Kit (CD117)-positive GIST, as defined by the Miettinen 2006 criteria (based on tumour size, location and mitotic rate).	
Antineoplastic agent		400mg orally once daily with a meal and a large glass of water
		Dispense every 30 days.
		For patients unable to swallow the tablets, disperse in 200ml of still water or apple juice.
Drug Interactions (see SPC for full list)	Drugs that inhibit or induce cytochrome P450 isoenzyme CYP activity could affect imatinib concentrations.	
http://www.medicines.org.uk/emc/medicine/ 15014	The following drugs increase plasma levels of imatinib: clarithromycin, erythromycin, itraconazole.	
	The following drugs decrease plasma levels of imatinib: carbamazepine, dexamethasone and phenytoin.	
	Drugs whose plasma levels may be increased by imatinib: ciclosporin, statins, warfarin.	
	Imatinib may inhibit the metabolism of warfarin; consider using a low molecular weight or unfractionated heparin in patients requiring anticoagulation.	
	Monitor liver function carefully if on concomitant hepatotoxic medications.	
	Grapefruit juice should be avoided; increased imatinib plasma concentration.	
	Aprepitant- potentially elevated plasma levels of imatinib.	
	Levothyroxine- decreased effectiveness, worsening hypothyroidism.	
Length of treatment	Continuous for up to 3 years or until progression of disease.	
Dose modifications	Neutrophils <1.0 x 10 ⁹ or Platelets <50 x 10 ⁹ /L	Stop treatment until neutrophil 1.5 x 10 ⁹ /L and platelets 75 x 10 ⁹ /L. Restart at same dose but if recurrence, repeat delay of treatment and then recommence at 300mg
	Impaired liver function	SELLI M. Colling by Latence and Constitution by
	Impaired renal function	Patients with renal dysfunction or on dialysis should be given the minimum recommended dose of 400mg daily with caution.

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Imatinib for ADJUVANT treatment of gastrointestinal stromal tumour (GIST) UGI-056			
Mandatory tests	FBC, LFT, U&E	Every 4 weeks for the first 3 months and then every 3 months during treatment	
	Hepatitis B serology should be tested before starting treatment.		
Mandatory supportive drugs	None		
Mandatory information	Patients should be supplied with a patient information leaflet, Cancerbackup information sheet and a copy of their treatment plan		

Ratifying consultant:

Justin Waters

Pharmacists:

S Lightfoot & C Waters

DAHOL

Version 2

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