APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:
Reg No:	First Names:	First Names:
Name:	Surname:	Surname:
Address:	DOB:	Address:
	Address:	
Fax Number:		Fax Number:

Sunitinib

Application	ication — RCC s only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 3 months. tes(tick boxes where appropriate)
and	The patient has metastatic renal cell carcinoma
	 The patient is treatment naive The patient has only received prior cytokine treatment The patient has only received prior treatment with an investigational agent within the confines of a bona fide clinical trial which has Ethics Committee approval The patient has discontinued pazopanib within 3 months of starting treatment due to intolerance The cancer did not progress whilst on pazopanib
and [and [and T	The patient has good performance status (WHO/ECOG grade 0-2) The disease is of predominant clear cell histology The patient has intermediate or poor prognosis defined as:
	Image: Constraint of the constraint
and [Sunitinib to be used for a maximum of 2 cycles
Initial appl	ication — GIST

Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 3 months. **Prerequisites**(tick boxes where appropriate)

a	The patient has unresectable or metastatic malignant gastrointestinal stromal tumour (GIST) and			
		or		The patient's disease has progressed following treatment with imatinib
		or		The patient has documented treatment-limiting intolerance, or toxicity to, imatinib

I confirm the above details are correct and that in signing this form I understand I may be audited.

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Sunitinib - continued

Current approval Number (if known):						
Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 3 months. Prerequisites (tick boxes where appropriate)						
No evidence of disease progression						
The treatment remains appropriate and the patient is benefiting from treatment						
Note: Sunitinib treatment should be stopped if disease progresses. Poor prognosis patients are defined as having at least 3 of criteria 5.1-5.6. Intermediate prognosis patients are defined as having 1 or 2 of criteria 5.1-5.6						
Renewal — GIST						
Current approval Number (if known):						
Applications only from a relevant specialist or medical practitioner on the recommendation of a relevant specialist. Approvals valid for 6 months. Prerequisites (tick boxes where appropriate)						
The patient has responded to treatment or has stable disease as determined by Choi's modified CT response evaluation criteria as follows:						
The patient has had a complete response (disappearance of all lesions and no new lesions)						
The patient has had a partial response (a decrease in size of 10% or more or decrease in tumour density in Hounsfield Units (HU) of 15% or more on CT and no new lesions and no obvious progression of non measurable disease)						
The patient has stable disease (does not meet criteria the two above) and does not have progressive disease and no symptomatic deterioration attributed to tumour progression						
and The treatment remains appropriate and the patient is benefiting from treatment						
Note: It is recommended that response to treatment be assessed using Choi's modified CT response evaluation criteria (J Clin Oncol, 2007, 25:1753-1759) Progressive disease is defined as either: an increase in tumour size of 10% or more and not meeting criteria of partial response (PR) by tumour density (HU) on CT; or: new lesions, or new intratumoral nodules, or increase in the size of the existing intratumoral nodules.						
Renewal — GIST pandemic circumstances						
Current approval Number (if known):						
Applications from any relevant practitioner. Approvals valid for 6 months. Prerequisites (tick boxes where appropriate)						
The patient has unresectable or metastatic malignant gastrointestinal stromal (GIST)						
The patient is clinically benifiting from treatment and continued treatment remains appropriate and						
Sunitinib is to be discontinued at progression and						
The regular Special Authority renewal requirements cannot be met due to COVID-19 constraints on the health sector						

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