Enquiries to Ministry of Health 0800 855 066

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA2496 August 2025

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:						
Trametinib								
Initial application — stage III or IV resected melanoma - adjuvant Applications from any relevant practitioner. Approvals valid for 4 months. Prerequisites(tick boxes where appropriate) The individual is currently on treatment with dabrafenib and trametinib and met all remaining criteria prior to commencing treatment or The individual has resected stage IIIB, IIIC, IIID or IV melanoma (excluding uveal) (see note a) The individual has received neoadjuvant treatment with a PD-1/PD-L1 inhibitor and Adjuvant treatment with trametinib is required and Treatment must be adjuvant to complete surgical resection and Treatment must be initiated within 13 weeks of surgical resection, unless delay is necessary due to post-surgery recovery (see note b) and Treatment must be addivant to combination with dabrafenib and Trametinib must be administered in combination with dabrafenib and Trametinib must be administered in combination with dabrafenib The individual has ECOG performance score 0-2								
a) Stage IIIB, IIIC, IIID or IV melanoma defined as per American Joint Committee on Cancer (AJCC) 8th Edition								
b) Initiating treatment within 13 weeks of complete surgical resection means 13 weeks after resection (primary or lymphadenectomy)								

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

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	Address:						
Fax Number:		Fax Number:					
Renewal — stage III or IV resected melanoma - adjuvant Current approval Number (if known):							
and Trametinib must be admin and Treatment to be discontinu	Trametinib must be administered in combination with dabrafenib and Treatment to be discontinued at signs of disease recurrence or at completion of 12 months' total treatment course, including any systemic neoadjuvant treatment The individual has received adjuvant treatment with a BRAF/MEK inhibitor The individual has metastatic or unresectable melanoma (excluding uveal) stage III or IV and The individual meets initial application criteria for trametinib for unresectable or metastatic melanoma						
and The individual has metast and The individual meets initia							
and The individual has receive							

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Addre	ess:				DOB:	Address:
			Address:			
Fax Number:						Fax Number:
Initia Appli Prere	al applications	ication s from a tes(tick	or	unresectable or metastation relevant practitioner. Approves where appropriate) dividual is currently on treater. The individual has metastation be asseline measurement of oxona the individual has ECOG period to be administed by the individual has bear the individual has bear the individual did not to the individual of the individual did not to the individual did not to the individual did not to the individual did not the individual	ment with dabrafenib and trametinib and met all remate or unresectable melanoma (excluding uveal) stage werall tumour burden is documented clinically and raderformance score 0-2 d BRAF mutation ered in combination with dabrafenib en diagnosed in the metastatic or unresectable stage receive treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment in the adjuvant setting with a BRAF ecceived treatment disease recurrence while on treatment in the experience disease recurrence within six month bitor	III or IV iologically III or IV setting F/MEK inhibitor F/MEK inhibitor ent with that BRAF/MEK inhibitor
				able or metastatic melano ber (if known):		
Appli	ication	s from a	ıny	relevant practitioner. Approves where appropriate)		
	and [] ·	The individual's disease has The individual has stable dis nse to treatment in target le	had a complete response to treatment had a partial response to treatment sease with treatment sions has been determined by comparable radiologic	assessment following the most recent treatment

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