Enquiries to Ministry of Health 0800 855 066

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA2418 January 2026

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:	
Osimertinib			
Initial application — NSCLC – first line Applications from any relevant practitioner. Approvals valid for 4 months. Prerequisites (tick boxes where appropriate) Patient is currently on treatment with osimertinib and met all remaining criteria prior to commencing treatment or Patient has locally advanced or metastatic, incurable, non-squamous non-small cell lung cancer (NSCLC) and Patient has received prior chemotherapy in the adjuvant setting and/or while awaiting EGFR results or Patient has discontinued gefitinib or erlotinib due to intolerance and The cancer did not progress while on gefitinib or erlotinib and There is documentation confirming that the cancer expresses activating mutations of EGFR and Baseline measurement of overall tumour burden is documented clinically and radiologically Renewal — NSCLC – first line			
Current approval Number (if known):			
Prerequisites(tick box where appropriate)			
Response to or stable disease with treatment in target lesions has been determined by comparable radiologic assessment following the most recent treatment period			

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Page 2 Form SA2418 January 2026

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Address:	DOB:	Address:	
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Fax Number:		Fax Number:	
Osimertinib - continued			
Initial application — NSCLC – second line Applications from any relevant practitioner. Approvals valid for 4 months. Prerequisites(tick boxes where appropriate) Patient is currently on treatment with osimertinib and met all remaining criteria prior to commencing treatment or Patient has locally advanced or metastatic, incurable, non-squamous non-small cell lung cancer (NSCLC) and Patient has an ECOG performance status 0-3 and The patient must have received previous treatment with erlotinib or gefitinib and There is documentation confirming that the cancer expresses T790M mutation of EGFR following progression on or after erlotinib or gefitinib and Baseline measurement of overall tumour burden is documented clinically and radiologically			
Renewal — NSCLC – second line			
Current approval Number (if known):			
Applications from any relevant practitioner. Approx Prerequisites (tick box where appropriate)	vals valid for 6 months.		
Response to treatment in target lesions has been determined by comparable radiologic assessment following the most recent treatment period			