Enquiries to Ministry of Health 0800 855 066

## APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

Page 1 Form SA2418 November 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:	
Osimertinib			
Patient has locally advance and  Patient is treatment in  Patient has received  or  Patient has received  The patient has and The cancer did  and Patient has an ECOG perform and Baseline measurement of or	rith osimertinib and met all remaining criteria prior to or door metastatic, incurable, non-squamous non-small or arive prior chemotherapy in the adjuvant setting and/or which discontinued gefitinib or erlotinib due to intolerance not progress while on gefitinib or erlotinib	cell lung cancer (NSCLC)  le awaiting EGFR results  s of EGFR	
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			

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

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APPLICANT (stamp or sticker acceptable)	PATIENT NHI:	REFERRER Reg No:	
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Address:	DOB:	Address:	
	Address:		
Fax Number:		Fax Number:	
Osimertinib - continued			
Patient has locally advanced and Patient has an ECOG perfor and The patient must have received and There is documentation conferiorinib or gefitinib  and The treatment must be given	th osimertinib and met all remaining criteria prior to color metastatic, incurable, non-squamous non-small commance status 0-3  yed previous treatment with erlotinib or gefitinib  firming that the cancer expresses T790M mutation of	ell lung cancer (NSCLC)  EGFR following progression on or after	
Renewal — NSCLC – second line			
Current approval Number (if known):			
Applications from any relevant practitioner. Approv <b>Prerequisites</b> (tick box where appropriate)	als valid for 6 months.		
Response to treatment in target lesions h	has been determined by comparable radiologic asses	sment following the most recent treatment period	

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