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:

Durvalumab

lication	lication — Non-small cell lung cancer ns only from a relevant specialist or any relevant practitioner on the recommendation of a relevant specialist. Approvals valid for 4 months. ites (tick boxes where appropriate)
	Patient has histologically or cytologically documented stage III, locally advanced, unresectable non-small cell lung cancer (NSCLC)
	Patient has histologically or cytologically documented stage IIb (T1N2a only), locally advanced, unresectable non-small cell lung cancer (NSCLC)
and	Patient has received two or more cycles of platinum-based chemotherapy concurrently with definitive radiation therapy
	Patient has no disease progression following the second or subsequent cycle of platinum-based chemotherapy with definitive radiation therapy treatment
and	Patient has a ECOG performance status of 0 or 1
and	Patient has completed last radiation dose within 8 weeks of starting treatment with durvalumab
and	Patient must not have received prior PD-1 or PD-L1 inhibitor therapy for this condition
	Durvalumab is to be used at a maximum dose of no greater than 10 mg/kg every 2 weeks
	Durvalumab is to be used at a flat dose of 1500 mg every 4 weeks
and	Treatment with durvalumab to cease upon signs of disease progression

Renewal — Non-small cell lung cancer

Current approval Number (if known):.....

Applications only from a relevant specialist or any relevant practitioner on the recommendation of a relevant specialist. Approvals valid for 4 months. **Prerequisites**(tick boxes where appropriate)

an	The treatment remains clinically appropriate and the patient is benefitting from treatment and		
	or	Durvalumab is to be used at a maximum dose of no greater than 10 mg/kg every 2 weeks	
		Durvalumab is to be used at a flat dose of 1500 mg every 4 weeks	
	and Treatment with durvalumab to cease upon signs of disease progression and		
	Tota	I continuous treatment duration must not exceed 12 months	

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