Enquiries to Ministry of Health 0800 855 066

APPLICATION FOR SUBSIDY BY SPECIAL AUTHORITY

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| Reg No: | APPLICANT (stamp or sticker acceptable) | PATIENT NHI: | REFERRER Reg No: | |
|--|---|--------------|------------------|--|
| Name: Surname: Surname: Surname: Surname: Address: DOB: Address: Address: Address: Fax Number: Fax Number: Fax Number: Fax Number: Fax Number: Fax Number: Address: Fax Number: Fax Number | , , | First Names: | C | |
| Address: | _ | | | |
| Fax Number: Fax Number: Casirivimab and imdevimab Initial application — Treatment of profoundly immunocompromised patients Applications from any relevant practitioner. Approvals valid for 2 weeks. Prerequisites(tick boxes where appropriate) Patient has confirmed (or probable) COVID-19 and Patient is in the community with mild to moderate disease severity* and Patient is profoundly immunocompromised** and is at risk of not having mounted an adequate response to vaccination against COVID-19 or is unvaccinated Patient's symptoms started within the last 10 days and Patient is not receiving high flow oxygen or assisted/mechanical ventilation | Name: | Surname: | Surname: | |
| Fax Number: Fax Nu | Address: | DOB: | Address: | |
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| Casirivimab and imdevimab is to be administered at a maximum dose of no greater than 2,400 mg Note: * Mild to moderate disease severity as described on the Ministry of Health Website ** Examples include B-cell depletive illnesses or patients receiving treatment that is B-Cell depleting. | | | | |

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