

**APPLICANT** (stamp or sticker acceptable)      **PATIENT NHI:** .....      **REFERRER** Reg No: .....

Reg No: .....      First Names: .....      First Names: .....

Name: .....      Surname: .....      Surname: .....

Address: .....      DOB: .....      Address: .....

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Fax Number: .....      Fax Number: .....

**Elexacaftor with tezacaftor, ivacaftor and ivacaftor**

**Initial application**

Applications from any relevant practitioner. Approvals valid without further renewal unless notified.

**Prerequisites**(tick boxes where appropriate)

Patient has been diagnosed with cystic fibrosis

**and**

Patient has two cystic fibrosis-causing mutations in the cystic fibrosis transmembrane regulator (CFTR) gene (one from each parental allele)

**or**

Patient has a sweat chloride value of at least 60 mmol/L

**and**

Patient has a heterozygous or homozygous F508del mutation

**or**

Patient has a mutation responsive to elexacaftor/tezacaftor/ivacaftor (see note)

**and**

The treatment must be the sole funded CFTR modulator therapy for this condition

**and**

Treatment with elexacaftor/tezacaftor/ivacaftor must be given concomitantly with standard therapy for this condition

Note: Eligible mutations are listed in the Food and Drug Administration (FDA) Trikafta prescribing information [https://www.accessdata.fda.gov/drugsatfda\\_docs/label/2019/212832Orig1s001.pdf](https://www.accessdata.fda.gov/drugsatfda_docs/label/2019/212832Orig1s001.pdf)

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

Signed: ..... Date: .....

Post application to Health New Zealand, Private Bag 3015, Wanganui – email: [customerservice@health.govt.nz](mailto:customerservice@health.govt.nz)