

Use this checklist to determine if a patient meets the restrictions for funding in the **hospital setting**. For more details, refer to [Section H](#) of the Pharmaceutical Schedule. For community funding, see the [Special Authority Criteria](#).

PRESCRIBER

Name:

Ward:

PATIENT:

Name:

NHI:

Osimertinib

INITIATION – NSCLC – first line

Re-assessment required after 4 months

Prerequisites (tick boxes where appropriate)

Patient has locally advanced or metastatic, incurable, non-squamous non-small cell lung cancer (NSCLC)
and

Patient is treatment naïve
or
 Patient has received prior treatment in the adjuvant setting and/or while awaiting EGFR results
or

The 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
 Patient has an ECOG performance status 0-3
and
 Baseline measurement of overall tumour burden is documented clinically and radiologically

CONTINUATION – NSCLC – first line

Re-assessment required after 6 months

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

INITIATION – NSCLC – second line

Re-assessment required after 4 months

Prerequisites (tick boxes where appropriate)

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
 The treatment must be given as monotherapy
and
 Baseline measurement of overall tumour burden is documented clinically and radiologically

CONTINUATION – NSCLC – second line

Re-assessment required after 6 months

Prerequisites (tick box where appropriate)

Response to treatment in target lesions has been determined by comparable radiologic assessment following the most recent treatment period

I confirm that the above details are correct:

Signed: Date: