## {{PANUMCODE}}

{{DISPLAY\_PAGNAME}} {{PACDESCRIPTION}}

This fax machine is located in a secure location as required by HIPAA regulations. Fax complete signed and dated forms to {{COMPANY\_NAME}} at {{CLIENT\_PAG\_FAX}}. Please contact {{COMPANY\_NAME}} at {{CLIENT\_PAG\_PHONE}} with questions regarding the prior authorization process. When conditions are met, we will authorize the coverage of {{DRUGNAME}}.

 Quantity:
 Frequency:
 Strength:

 Route of Administration:
 Expected Length of Therapy:

 Diagnosis:
 <<DIAGNOSIS>>
 ICD Code:
 <<ICD9>>

- What is the diagnosis?
   Colorectal cancer, including appendiceal adenocarcinoma and anal adenocarcinoma
   Other \_\_\_\_\_\_
- 2. What is the ICD-10 code?
- 3. Coverage for the requested drug is provided when the patient has tried and had a treatment failure with at least three of the formulary medications, or all of the formulary alternatives if there are fewer than three. The formulary alternatives for the requested drug are Lonsurf and Stivarga. Can the patient's treatment be switched to a formulary alternative? *If Yes, fax new prescription and no further questions.*□ Yes please specify.
  □ No Continue request for non preferred drug.
- 4. Has the patient tried and had a documented inadequate response or intolerable adverse reaction to at least three of the formulary medications, or all of the formulary alternatives if there are fewer than three? [Note: Formulary medications should be prescribed first unless the patient is unable to use or receive treatment with the alternative.] Formulary alternative(s): Lonsurf and Stivarga. □ Yes □ No

If Yes, indicate the formulary alternative(s) and the reason(s) for treatment failure. and skip to #6.

Drug name: \_\_\_\_\_\_ Reason for treatment failure: \_\_\_\_\_\_

Drug name: Reason for treatment failure:

5. Does the patient have a documented contraindication to all or at least three of the formulary alternatives? Lonsurf and Stivarga. 🗆 Yes 🗅 No

If Yes, specify the formulary alternative(s) the patient is unable to take and describe the contraindication(s).

Drug name:	Contraindication:
Drug name:	_ Contraindication:

- 6. Have chart notes or other documentation supporting the inadequate response, intolerable adverse reaction, or contraindication to at least three of the formulary medications, or all of the formulary alternatives if there are fewer than three been attached? *ACTION REQUIRED: Submit chart note(s) or other documentation indicating prior treatment failure, severity of the adverse event (if any), and dosage and duration of the prior treatment, or contraindication to formulary alternatives.*  $\Box$  Yes  $\Box$  No
- 7. Is the patient currently receiving treatment with the requested medication?  $\Box$  Yes  $\Box$  No If No, skip to #9

- 8. Is there evidence of unacceptable toxicity or disease progression while on the current regimen?
   □ Yes □ No
- 9. What is the clinical setting in which the requested medication will be used?
  □ Advanced disease □ Metastatic disease □ Other
- 10. Is the patient's disease RAS wild type?
  - Yes
  - 🗖 No
  - Unknown
- 11. Is an anti-epidermal growth factor receptor (EGFR) therapy such as Erbitux (cetuximab) or Vectibix (panitumumab) medically appropriate?
  Yes I No
- 12. Has the patient been previously treated with an anti-epidermal growth factor receptor (EGFR) therapy? □ Yes □ No
- 13. Does the patient have a contraindication or intolerance to an anti-epidermal growth factor receptor (EGFR) therapy?
  - $\hfill\square$  Yes contraindication
  - □ Yes intolerance
  - 🛛 No
- 14. Has the patient been previously treated with fluoropyrimidine-, oxaliplatin- and irinotecan-based chemotherapy (with or without bevacizumab)? If Yes, no further questions.  $\Box$  Yes  $\Box$  No
- 15. Does the patient have a contraindication or intolerance to fluoropyrimidine-, oxaliplatin- and irinotecan-based chemotherapy?
  - Yes contraindication
  - □ Yes intolerance
  - 🛛 No

I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.