

FLEXIBLE DOSING OF ORAL TREATMENTS IN mCRC TO MANAGE ADVERSE EVENTS¹

REGORAFENIB

Multikinase inhibitor

RECOMMENDED DOSING

WEEK 1	WEEK 2	WEEK 3	WEEK 4
160mg	160mg	160mg	--
ONCE DAILY ORALLY	ONCE DAILY ORALLY	ONCE DAILY ORALLY	DOSE-FREE INTERVAL



REPEAT CYCLE

CYCLE 1
(28 days)

MOST COMMON ADVERSE EVENTS

Pain (including gastrointestinal and abdominal pain), hand-foot skin reaction, asthenia/fatigue, diarrhoea, decreased appetite/food intake, hypertension, infection, dysphonia, hyperbilirubinemia, fever, mucositis, weight loss, rash, and nausea^a

FLEXIBLE DOSING STRATEGIES TO OPTIMISE TREATMENT DURATION^{1,2}

WEEK 1	WEEK 2	WEEK 3	WEEK 4	WEEK 1
80mg	120mg	160mg	--	HIGHEST TOLERATED DOSE FROM CYCLE 1
ONCE DAILY ORALLY	ONCE DAILY ORALLY	ONCE DAILY ORALLY	DOSE-FREE INTERVAL	ONCE DAILY ORALLY

CYCLE 1
(28 days)

CYCLE 2

^a. Most common adverse reactions ≥20%

Stivarga (regorafenib) Prescribing Information Dec 2020

1. Bekaii-Saab TS, et al. Lancet Oncol. 2019; 20:1070–1082 | 2. NCCN Clinical practice guidelines for Colon Cancer V1.2022

mCRC: metastatic colorectal cancer

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FLEXIBLE DOSING OF ORAL TREATMENTS IN **mCRC** TO MANAGE ADVERSE EVENTS

TRIFLURIDINE/TIPIRACIL (TAS-102)

Combination of a nucleoside metabolic inhibitor and a thymidine phosphorylase inhibitor

RECOMMENDED DOSING

35mg/m² dose, orally, twice daily

WEEK 1	WEEK 2	WEEK 3	WEEK 4
35mg/m² TWICE DAILY <i>Day 1-5 treatment</i>	35mg/m² TWICE DAILY <i>Day 8-12 treatment</i>	-- DOSE-FREE INTERVAL	-- DOSE-FREE INTERVAL
<i>Day 6-7 dose-free</i>	<i>Day 13-14 dose-free</i>		



CYCLE 1
(28 days)

MOST COMMON ADVERSE EVENTS

Anaemia, neutropenia, fatigue/asthenia, nausea, thrombocytopenia, decreased appetite, diarrhoea, vomiting, and pyrexia^a

FLEXIBLE DOSING STRATEGIES TO MANAGE ADVERSE EVENTS

Reduce dose by 5mg/m² for adverse events a maximum of 3 times to a minimum of 20mg/m^{2b}



^a. Most common adverse reactions ≥10% | ^b. Do not escalate the TAS-102 dosage after it has been reduced
Lonsurf (trifluridine/tipiracil) Prescribing Information Dec 2019
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FLEXIBLE DOSING OF ORAL TREATMENTS IN mCRC TO MANAGE ADVERSE EVENTS

CAPECITABINE

Nucleoside metabolic inhibitor with antineoplastic activity

RECOMMENDED DOSING

1250 mg/m² dose, orally, twice daily

WEEK 1	WEEK 2	WEEK 3
1250mg/m² TWICE DAILY	1250mg/m² TWICE DAILY	-- DOSE-FREE INTERVAL



REPEAT CYCLE

CYCLE 1
(21 days)

MOST COMMON ADVERSE EVENTS

Diarrhoea, hand-and-foot syndrome, nausea, vomiting, abdominal pain, fatigue/weakness, and hyperbilirubinemia^a

FLEXIBLE DOSING STRATEGIES TO MANAGE ADVERSE EVENTS

Using the NCIC CTC grade reduce dose by occurrence and severity

NCIC GRADE	1st OCCURENCE	2nd OCCURENCE	3rd OCCURENCE	4th OCCURENCE
G1	<i>maintain dose</i>	<i>maintain dose</i>	<i>maintain dose</i>	<i>maintain dose</i>
G2	<i>100% dose</i>	<i>75% dose</i>	<i>50% dose</i>	<i>discontinue</i>
G3	<i>75% dose</i>	<i>50% dose</i>	<i>discontinue</i>	
G4	<i>50% / discontinue</i>			

^a. Most common adverse reactions ≥30%

Xeloda (capecitabine) Prescribing Information May 2021

NCIC CTC: National Cancer Institute of Canada Common Toxicity Criteria | mCRC: metastatic colorectal cancer

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