

HOW TO TREAT PATIENTS AFTER FAILURE ON TRIPLET THERAPY +/- BEV?

By Dr. Chiara Cremolini, Dr. Erika Martinelli and Dr. Dominik Modest

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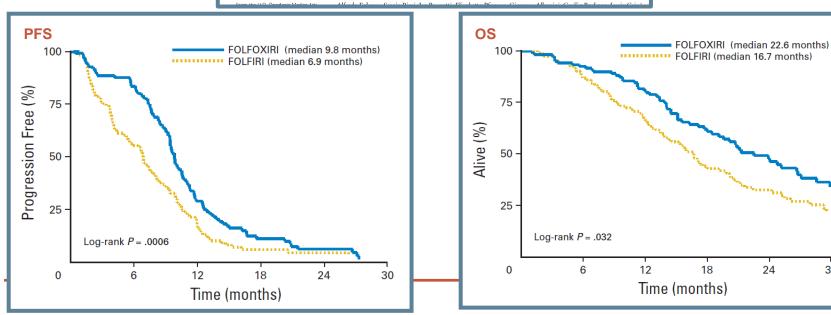
THE UPFRONT USE OF THE THREE ACTIVE CYTOTOXICS MAY RAISE CONCERNS ABOUT THE FEASIBILITY AND EFFICACY OF GIFURTHER TREATMENTS, BUT...



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1) OS results of trials investigating the triplet +/- bevacizumab are reassuring

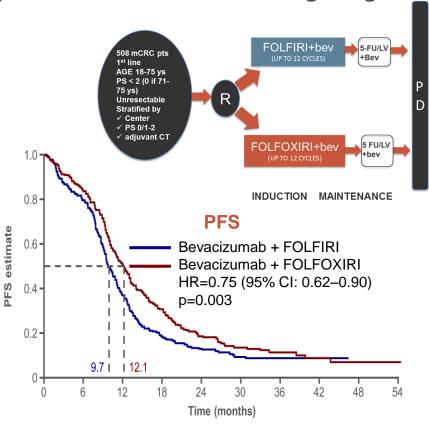


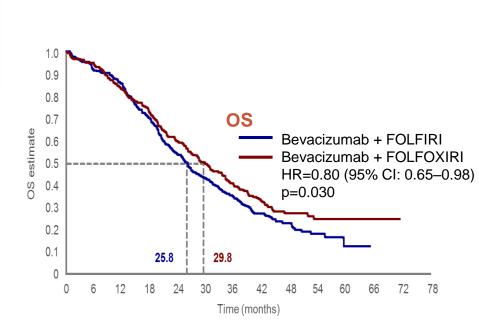


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Consistent results are reported in PFS and OS. If the use of the triplet impaired the feasibility and efficacy of further treatments, the advantage in PFS would hardly translate into an OS benefit

2) 2nd- and further-lines treatments seem feasible

TRIBE: 2ND-LINE TREATMENTS



	FOLFIRI + bev	FOLFOXIRI + bev
	Arm A	Arm B
Potential candidates to 2 nd line	92%	92%
	(236/256)	(232/252)
Any 2 nd -line therapy	76%	76%
	(180/236)	(176/232)
2 nd -line agents	N=180	N=176
Fluoropyrimidine, %	93%	81%
Oxaliplatin, %	67%	29%
Irinotecan, %	31%	63%
Bevacizumab, %	31%	32%
Anti-EGFR mAB, %	15%	29%
3 rd -line Anti-EGFR mAB, %	21%	13%

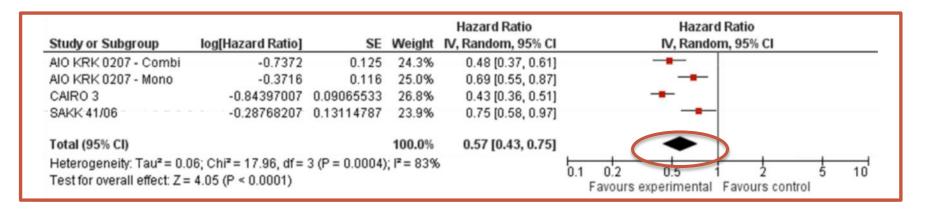
WHICH TREATMENT AFTER PROGRESSION?



The first-line treatment with FOLFOXIRI +/- bev should not be continued until disease progression

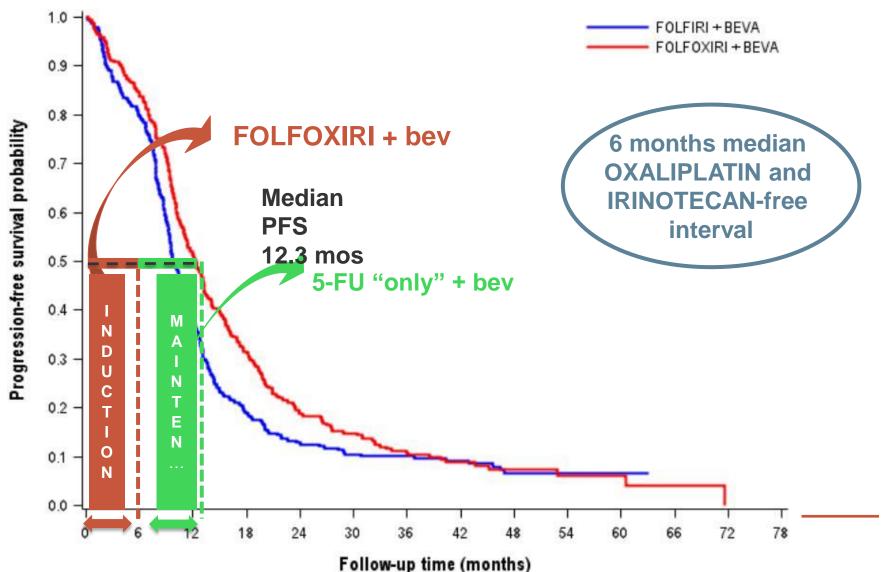


The role of maintenance is crucial to prolong the oxaliplatin-and irinotecan-free interval



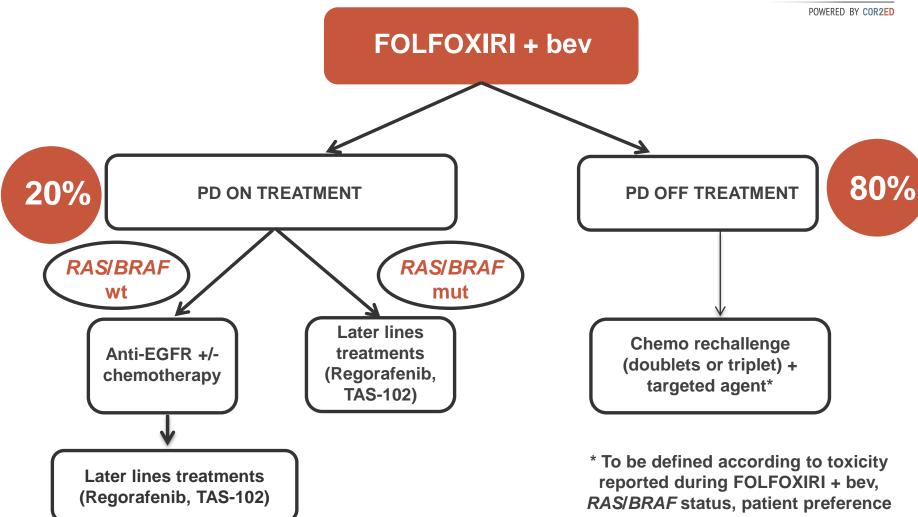
FOLFOXIRI PLUS BEV: WHAT TO DO AFTER PROGRESSION?





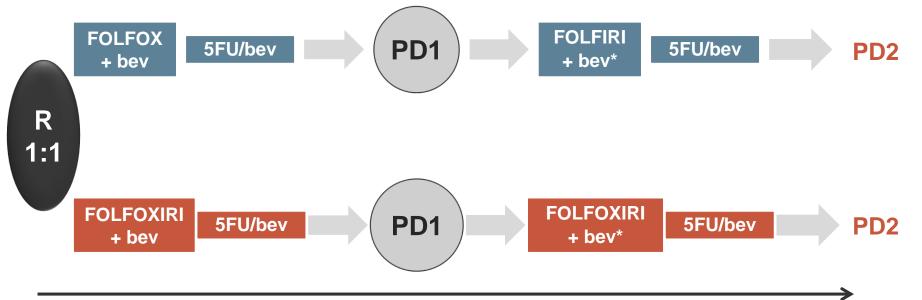
SUMMARIZING... A POTENTIAL ALGORITHM





CURRENTLY ONGOING...TRIBE-2





Primary endpoint:

Progression Free Survival 2

*all repeated for 8 cycles (4 months) followed by maintenance with 5FU/bev until PD

Target accrual: 654 patients in 60 Italian centers







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