# The cost of survival gain in metastatic colorectal cancer (mCRC) in France

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# BACKGROUND

- Chemotherapies are commonly used for the treatment of mCRC. Over the past 10 years, a number of targeted therapies (bevacizumab, cetuximab, panitumumab, aflibercept, and regorafenib) have been approved for the treatment of mCRC1-5
- Targeted therapies are more expensive than chemotherapies; therefore, their clinical benefit is associated with economic implications
- Traditionally, the added value of a new therapy has been estimated through the use of cost-utility analyses
- However, clinical trial data are often incomplete and incremental cost-effectiveness ratios (ICERs) are highly dependent on the assumptions used to extrapolate or adjust trial results. In the UK, ICERs for second-line cetuximab plus irinotecan versus irinotecan alone range from £45,237 to £370,044 depending on the assumptions used<sup>6</sup>
- Outside of cost-effectiveness analyses, the relative cost of overall survival (OS) gain has not been examined extensively in published literature. Thus, a basic cost-effectiveness analysis of targeted therapies in mCRC was conducted from the perspective of a French national payer, using only observed data

### **OBJECTIVE**

 To estimate the incremental cost per month of median OS gained with the use of approved targeted therapies, in addition to chemotherapy or best supportive care (BSC) alone, for first-, second-, and third-line treatment of mCRC

## **METHODS**

 A review was conducted of product labels of bevacizumab, cetuximab, panitumumab, aflibercept, and regorafenib to identify pivotal phase 3 clinical trials. Studies were included if they demonstrated statistically significant improvement in median OS (Table 1)

Table 1: Summary of clinical trials assessing targeted therapies

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Regimen/source	Targeted therapies	Median treatment duration, months	Median PFS, months	Median OS, months	
First-line					
IFL	Bevacizumab	9.3	10.6	20.3	
Hurwitz 2004 <sup>i,9</sup>		6.4	6.2	15.6	
FOLFIRI	Cetuximab	NR	9.9	23.5	
Van Cutsem 2011 <sup>10</sup>		NR	8.4	20.0	
FOLFOX4	Panitumumab	NR	10.0	23.9	
Douillard 2014 <sup>ii,11</sup>		NR	8.6	19.7	
Second-line					
FOLFOX4	Bevacizumab	4.6	7.3	12.9	
Giantonio 2007 <sup>12</sup>		3.2	4.7	10.8	
Oxaliplatin- or	Bevacizumab	4.2	5.7	11.2	
irinotecan-based chemotherapy Bennouna 2013 <sup>7</sup> *		3.2	4.1	9.8	
FOLFIRI	Aflibercept	4.9	6.9	13.5	
Van Cutsem 2012 <sup>13</sup>		4.2	4.7	12.1	
Third-line					
BSC	Cetuximab	NR	3.7	9.5	
Karapetis 2008iii.14		NR	1.9	4.8	
BSC	Regorafenib	1.7	1.9	6.4	
Grothey 2013 <sup>15</sup>		1.6	1.7	5.0	

 IFL is no longer standard of care. ii. An updated exploratory analysis.

the study.8

- A retrospective analysis; tumor samples were not available for all patients. BSC, best supportive care; NR, not reported
- \*Bennouna et all presented the average results for bevacizumab added to a variety of oxaliplatin- or irinotecan-based chemotherapies. The dosing regimens for the most common three regimens were used; these dosing regimens were taken from an ASCO presentation of
- Treatment duration was not consistently reported across all trials. For this analysis, median PFS was used as a proxy for treatment duration, which should be reasonable because most patients discontinue therapy due to progression or death.15 The model considers the drug and administration cost over the treatment duration
- Adverse event management and disease management costs are not included in the analysis

### **Drug costs estimation**

- The acquisition costs for targeted therapies and chemotherapies were based on the 2015 public price including VAT from l'Assurance Maladie de la Sécurité Sociale (Table 2)16
- The recommended dosing regimens were based on clinical trials identified in Table 1. and were used to estimate the costs in Table 3. Body surface area (men: 1.93 ± 0.19 m<sup>2</sup>, women: 1.68 ± 1.8 m<sup>2</sup>) and weight (men: 79.8 ± 15 kg, women: 65.3 ± 14 kg) are based on a study of cancer patients in the UK17
- The analysis assumed that the population is 48% male, based on 2012 OECD population statistics<sup>18</sup>
- The drug cost per month of a treatment regimen was based on the recommended dose of each drug, the number of vials/capsules required to achieve dose, the cycle length, and the number of doses per cycle
- The analysis assumed that vial sharing was not allowed (i.e., after administration, the remaining drug in a vial would be discarded)
- The number of vials/tablets required to achieve the specified dose were based on methods described by Sacco et al, 2010.17 A sample calculation for cetuximab is presented in Table 4; similar calculations were performed for other drugs

Table 2: Drug acquisition cost

Drug	Formulation	Cost per vial/tablet, €			
	500 mg vial	2.13			
5-FU	1000 mg vial	3.73			
	5000 mg vial	9.27			
Aflibercept	100 mg vial	307.50			
Allibercept	200 mg vial	614.99			
Revacizumah	100 mg vial	272.61			
Bevacizumab	400 mg vial	1003.16			
Capecitabine	150 mg tablet	0.49			
Сареспарине	500 mg tablet	1.43			
Cetuximab	100 mg vial	168.24			
Cetuximab	500 mg vial	841.22			
	40 mg vial	64.94			
Irinotecan	100 mg vial	161.65			
IIIIotecan	300 mg vial	487.02			
	500 mg vial	811.70			
	25 mg vial	3.73			
Leucovorin	100 mg vial	15.51			
	175 mg vial	24.98			
	50 mg vial	156.21			
Oxaliplatin	100 mg vial	312.43			
Oxalipiatiii	150 mg vial	468.64			
	200 mg vial	624.85			
	100 mg vial	373.18			
Panitumumab	200 mg vial	746.35			
	400 mg vial	1492.70			
Regorafenib	40 mg tablets, 84 tablets per package	30.40			

Table 3: Drug costs estimation

Regimen/ source	Drug	Drug cost Dose per dose, €		Doses per cycle	Cycle length, weeks	Drug cost per month, €
	Irinotecan	125 mg/m <sup>2</sup>	397.65	4	6	565.91
IFL +	Fluorouracil	500 mg/m <sup>2</sup>	4.14	4	6	5.89
bevacizumab Hurwitz 2004 <sup>9</sup>	Leucovorin	20 mg/m <sup>2</sup>	7.45	4	6	10.60
11d1Witz 2001	Bevacizumab	5 mg/ kg	1054.66	1	2	2390.87
	Irinotecan	180 mg/m <sup>2</sup>	681.52	1	2	1480.68
FOLFIRI + cetuximab	Leucovorin	200 mg/m <sup>2</sup>	53.77	1	2	116.83
Van Cutsem	Fluorouracil*	400 mg/m <sup>2</sup>	3.72	1	2	8.09
201110	Fluorouracil**	2400 mg/m <sup>2</sup>	13.72	1	2	59.64
	Cetuximab	400 mg/m <sup>2</sup>	1296.11	Initial loading dose	NA	NA
		250 mg/m <sup>2</sup>	841.90	1	1	3658.25
FOLFOY4:	Oxaliplatin	85 mg/m <sup>2</sup>	556.24	1	2	1208.51
FOLFOX4 + panitumumab	Leucovorin	200 mg/m <sup>2</sup>	53.77	2	2	120.54
Douillard	Fluorouracil*	400 mg/m <sup>2</sup>	3.72	2	2	16.18
201411	Fluorouracil**	600 mg/m <sup>2</sup>	5.26	2	2	22.85
	Panitumumab	6 mg/kg	1805.97	1	2	3923.69
	Oxaliplatin	85 mg/m <sup>2</sup>	556.24	1	2	1208.51
FOLFOX4 +	Leucovorin	200 mg/m <sup>2</sup>	53.77	2	2	120.54
bevacizumab	Fluorouracil*	400 mg/m <sup>2</sup>	3.72	2	2	16.18
Giantonio 2007 <sup>12</sup>	Fluorouracil**	600 mg/m <sup>2</sup>	5.26	2	2	22.85
2007	Bevacizumab	10 mg/kg	1971.42	1	2	4283.15
	Oxaliplatin	100 mg/m <sup>2</sup>	556.24	2	4	1173.08
FOLFOVA:	Leucovorin	400 mg/m <sup>2</sup>	105.36	2	4	228.91
FOLFOX6 + bevacizumab	Fluorouracil*	400 mg/m <sup>2</sup>	3.72	2	4	8.09
Arnold 2012 <sup>8</sup>	Fluorouracil**	2400 mg/m <sup>2</sup>	13.72	2	4	59.64
Arnoid 2012°	Bevacizumab	5 mg/kg	1054.66	2	4	2291.37
	Capecitabine	1000 mg/m <sup>2</sup>	5.55	28	3	224.95
XELOX +	Oxaliplatin	130 mg/m <sup>2</sup>	809.91	1	3	1173.08
bevacizumab Arnold 2012 <sup>8</sup>	Bevacizumab		1520.22	1	3	2201.91
Amoid 2012		7.5 mg/kg		4		
	Irinotecan	180 mg/m <sup>2</sup>	681.52	1	2	1480.68
FOLFIRI +	Leucovorin	400 mg/m <sup>2</sup>	105.36	1	2	228.91
bevacizumab Arnold 2012 <sup>8</sup>	Fluorouracil*	400 mg/m <sup>2</sup>	3.72	1	2	8.09
Amoid 2012	Fluorouracil**	2400 mg/m <sup>2</sup>	13.72	1	2	59.64
	Bevacizumab	5 mg/kg	1054.66	1	2	2291.37
FOI FIRM	Irinotecan	180 mg/m <sup>2</sup>	681.52	1	2	1480.68
FOLFIRI + aflibercept	Leucovorin	400 mg/m <sup>2</sup>	105.36	1	2	228.91
Van Cutsem	Fluorouracil*	400 mg/m <sup>2</sup>	3.72	1	2	8.09
2012 <sup>13</sup>	Fluorouracil**	2400 mg/m <sup>2</sup>	13.72	1	2	59.64
	Aflibercept	4 mg/kg	1071.53	1	2	2328.03
Cetuximab + BSC	Cetuximab	400 mg/m <sup>2</sup>	1296.11	Initial loading dose	NA	NA
Karapetis 2008 <sup>14</sup>		250 mg/m <sup>2</sup>	841.90	1	1	3658.25
Regorafenib + BSC Grothey 2013 <sup>15</sup>	Regorafenib	160 mg	121.59	21	4	2773.75

\*Bolus. \*\*Continuous infusion.

Table 4: Estimation of number of units: sample calculation for cetuximab 400 mg/m<sup>2</sup>

Number of 100 mg vials	Body surface area, m²	Males				Females			
		Proportion of patients, %		Number of vials per dose		Proportion of patients, %		Number of vials per dose	
		Cumulative	Per dose	100 mg	500 mg	Cumulative	Per dose	100 mg	500 mg
1	0.3	0	0	1	0	0	0	1	0
2	0.5	0	0	2	0	0	0	2	0
3	0.8	0	0	3	0	0	0	3	0
4	1.0	0	0	4	0	0	0	4	0
5	1.3	0	0	0	1	1	1	0	1
6	1.5	1	1	1	1	16	15	1	1
7	1.8	17	16	2	1	65	49	2	1
8	2.0	64	47	3	1	96	31	3	1
9	2.3	95	31	4	1	100	4	4	1
10	2.5	100	5	0	2	100	0	0	2
		Mean vials per dose		2.99	1.05	Mean vials per dose		2.22	1.00
		Cost per vial, €		168.24	841.22	Cost per vial, €		168.24	841.22
		Mean cost per dose, €		1382.72		Mean cost per dose, €		1214.71	
		Weighte average		1296.11					

### Administration costs estimation

- The unit cost of chemotherapy administration is €395, which is based on code 28Z07Z/9606: "Chimiothérapie pour tumeur, en séances". 19
- · For each regimen, the administration cost per month was based on the unit cost of administration per visit, the cycle length, and the number of doses per cycle

- Table 5 reported the results incremental median OS (mOS) gain and incremental cost associated with introducing targeted therapies over chemotherapies
- In first-line, targeted agents were associated with 3.5–4.7 months of mOS gain with an additional cost of €7009 to €13,986/month
- In second-line, the 1.4–2.1 months of mOS gain had an additional cost of €11,217 to €18,848/month
- In third-line, the cost per month of mOS gain was the lowest, ranging from €3764 to €4328, with mOS gains of 1.4–4.7 months

### Table 5: Analysis results

Trial/source	Intervention	Median PFS, months	Median OS, months	Incremental mOS, months	Drug cost, €	Administration cost, €	Incremental cost, €	Incremental cost, € incremental month mOS
First-line								
Hurwitz	IFL + bevacizumab	10.6	20.3	4.7	30,258	15,161	32,941	7009
2004°	IFL	6.2	15.6		3611	8868		
Van Cutsem 2011 <sup>10</sup>	FOLFIRI + cetuximab	9.9	23.5	3.5	53,157	16,992	48,952	13,986
	FOLFIRI	8.4	20.0		13,988	7209		
Douillard 2014 <sup>11</sup>	FOLFOX4 + panitumumab	10.0	23.9	4.2	54,049	17,164	43,714	10,408
2014	FOLFOX4	8.6	19.7		12,738	14,761		
Second-line								
Giantonio 2007 <sup>12</sup>	FOLFOX4 + bevacizumab	7.3	12.9	2.1	42,080	12,529	39,581	18,848
2007	FOLFOX4	4.7	10.8		6962	8067		
Arnold 2012 <sup>s</sup>	FOLFOX6 + bevacizumab (5 mg/kg)	5.7	11.2	1.4	21,640	4892	16,842	12,030
	FOLFOX6	4.1	9.8		6171	3519		
Arnold 2012 <sup>8</sup>	XELOX + bevacizumab	5.7	11.2	1.4	20,520	3261	15,703	11,217
2012	XELOX	4.1	9.8		5732	2346		
Arnold 2012 <sup>8</sup>	sFOLFIRI + bevacizumab	5.7	11.2	1.4	23,192	4892	17,278	12,341
2012	sFOLFIRI	4.1	9.8		7287	3519		
Bennouna 2013 <sup>7</sup>	Fluoropyrimidine + oxaliplatin or irinotecan + bevacizumab	5.7	11.2	1.4	21,784 *	4348 *	16,608	11,863
	Fluoropyrimidine + oxaliplatin or irinotecan	4.1	9.8		6397 *	3128 *		
Van Cutsem	FOLFIRI + aflibercept	6.9	13.5	1.4	28,327	5921	21,941	15,237
201213	FOLFIRI	4.7	12.1		8300	4008		
Third-line								
Karapetis 2008 <sup>14</sup>	Cetuximab (wild-type <i>KRAS</i> )	3.7	9.5	4.7	13,990	6351	20,340	4328
	BSC (wild-type KRAS)	1.9	4.8		0	0		
Grothey	Regorafenib	1.9	6.4	1.4	5270	0	5270	3764
201315	Placebo	1.7	5.0		0	0		

# CONCLUSIONS

- Based on this analysis, which considered only observed data without extrapolation, the incremental cost per month of mOS gain varies greatly in France, both by treatment and by line
- The survival gain on targeted therapies is the highest in first-line treatment compared with second- or third-line treatment. The addition of a targeted agent gives the highest additional cost per month of OS gain in second-line treatment, followed by first-line treatment, with the lowest cost per month of mOS gain provided by third-line treatment. Regorafenib was the most cost-effective treatment in this analysis
- The impact of this analysis on the management of targeted agents in France should be explored further, and future analyses should consider other treatment-related costs, such as adverse event management and disease management costs, as well as dose adjustments to manage toxicities

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