

QC₃: QUALITY OF COMPREHENSIVE CANCER CARE IN SOUTHERN SWITZERLAND



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INTRODUCTION

International studies on Quality of Cancer Care (QoCC) since the 90's showed a constant and continuous improvement of the delivered oncologic care and a consequent spread of the advanced specialist care on the territory. The most of the study were developed on a regional basis as well as our project: this helps to increase the enrolment of the involved physicians and it shares a common basis about the sanitary laws and the territorial characteristics. One of the main challenges in the previous studies was due to the delay from the analysis of the first specimen in pathology and the registration of the incident cases at the cancer registry. The peculiarity of the Ticino Cancer Registry is of being located inside the Cantonal Institute of Pathology, so all the histological verified incident cases are recorded real-time.

The aims of the present study are to produce evidence-based quality indicators (QIs), whom application could allow an immediate change in the diagnostic-treatment process, that could be translated in a short-term benefit for patients.

METHODS

The QC₃ project is a population-based, prospective study, implemented on a three-year time period (2011-2013) on the territory of Canton Ticino. It deals with the QoCC of colorectal, prostatic, ovarian, endometrial and lung cancers. From the Ticino Cancer Registry we extract the patients and the cases regarding the above considered pathologies, treated both in the regional public and private hospitals; we include in the study all the patients > 18 years old, that is to say that all the elderly, usually excluded from randomized clinical trials (RCTs), are here evaluated. In the preliminary phase of the project, the QC₃ Qls derived from a comprehensive literature search on PubMed/MEDLINE of relevant peer-reviewed articles are developed using a 2-step modified Delphi process, involving dedicated working groups (WG) of local health care providers (colorectal WG, lung cancer WG, prostate cancer WG, ovarian/endometrial cancers WG, lung WG) to obtain expert opinions in a systematic, anonymous and individual manner for the validation of both evidence- and expert-based items. Then, for each localization, the list of selected cancer-specific Qls derived from the two Delphi rounds is submitted to an independent international multidisciplinary cancer-specific Advisory Board (AB), in order to get an additional evaluation and to define a final approved list of Qls. The final selected Qls are applied to the regional routine oncologic care, so to evaluate the performance of the currently used pattern of care according to the international guidelines.

RESULTS

Here we present a selection of the preliminary QIs results about the colorectal cancer (CRC) cases incident in 2011 (n=243). The initial evidence-based cancer-specific list of QC₃ QIs (n=149) was proposed to the CRC WG in an in-person meeting. for a preliminary revision and their selection (n=104), then it underwent to a 2-step modified Delphi process, shortening the QIs candidates to 89. The AB revised them and extracted the final 74 QIs (Fig.1). In **Tab. 1** we describe the demographic characteristics of the CRC cases incident in 2011 (n=243), whereas **Tab. 2** reports the CRC cases incident in 2011 undergoing surgery. In **Tab. 3** is represented a selection of the final QIs. For each QI is described its own *denominator*, i.e. the population on whom the QI is calculated, the results expressed in **YES** (QIs satisfied), **NO** (QIs not satisfied) and **MISSING** (data not present in the whole medical documentation examined). Furthermore, for each QI is indicated the literature used to define it (**G**= guidelines; **R**= reviews; **M**= meta-analysis; **PBS**: population-based studies; **CCS**: case-control or cohort studies).



Fig. 1 - QC, QIs SELECTION

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Tab. 1 -	CKCS	INCIDE	NI IN 2	(011

Variabile	Totale n = 243		COLON n = 167 (69%)		RETTO n = 76 (31%)	
Sesso, n (%)						
Uomini	129	(53.1%)	84	(50.3%)	46	(60.5%)
Donne	114	(46.9%)	83	(49.7%)	30	(39.5%)
Età						
media±sd (anni)	72±12		73±11		71±14	
mediana		74	74		74	
Gruppi età-specifici, n (%)						
0.49	10	(4.1%)	- 3	(1.8%)	7	(9.2%)
50,59	25	(10.3%)	15	(9.0%)	10	(13.2%)
60-69	56	(23.0%)	45	(26.9%)	11	(14.5%)
70-79	79	(32.5%)	56	(33.6%)	23	(30.3%)
>80+	73	(30.1%)	48	(28.7%)	25	(32.8%)
Gruppi età specifici, n (%)						
<70	91	(37,4%)	63	(37,7%)	28	(36.8%)
≥70	152	(62,6%)	104	(62,3%)	48	(63.2%)
Localizzazione tumorale ICD O II		(,		(02.0.0)		(
18.00 cieco	29	(11.9%)	29	(17,4%)		
18.20 colon ascendente	33	(13.6%)	33	(19.8%)		
18.30 flessura epatica	7	(2.9%)	7	(4,2%)		
18.40 colon trasverso	18	(7,4%)	18	(10.8%)		
18.50 flessura splenica	10	(4.1%)	10	(6,0%)		
18.60 colon discendente	16	(6.6%)	16	(9.6%)		
18.70 sigma	50	(20.6%)	50	(29.8%)		
18.80 overlapping lesion	2	(0.8%)	2	(1.2%)		
18.90 colon, NOS	2	(0.8%)	2	(1.2%)		
20.90 ampolla rettale. NOS	6	(2.5%)			6	(7.9%)
20.90 ampolia rettale, NOS 20.91 retto distale (4-7.5 cm)	28	(11.5%)			28	(36.8%)
20.92 retto medio (7.5-12 cm)	7	(2.9%)			7	(9.2%)
20.93 retto prossimale (>12 cm)	26	(10.7%)			26	(34.3%)
20.94 retto medio-prossim NOS	3	(1.2%)			3	(3.9%)
20.95 retto medio-distale. NOS	6	(2.5%)			6	(7.9%)
20.33 retto medio-distale, NOS	·	(2.5%)				(2.5%)
Histological type						
adenocarcinoma	225	(92.6%)	153	(91.6%)	72	(94.7%)
carcinoma mucinoso	1	(0.4%)	1	(0.6%)	0	(0.0%)
carcinoma sigillo cellulare	15	(6.2%)	11	(6.6%)	4	(5.3%)
carcinoma NOS	2	(0.8%)	2	(1.2%)	0	(0.0%)

Tab. 2 - OPERATED CRCs INCIDENT IN 2011

Variabile	Totale COLON n = 200 n = 150 (75%)			RETTO n = 50 (25%)		
Sesso, n (%)						
Uomini	108	(54%)	79	(53%)	29	(58%)
Donne	92	(46%)	71	(47%)	21	(42%)
Età						
media±sd (anni)	72±11		72±11		69±12	
mediana	72		73		72	
Gruppi età-specifici, n (%)						
0-49	7	(3.5%)	3	(2%)	4	(8%)
50-59	21	(10.5%)	13	(8.7%)	8	(16%)
60-69	52	(26%)	42	(28%)	10	(20%)
70-79	66	(33%)	49	(32.7%)	17	(34%)
>80+	54	(27%)	43	(28.6%)	11	(22%)
Gruppi età specifici, n (%)						
<70	80	(40%)	58	(38.7%)	22	(44%)
≥70	120	(60%)	92	(61.3%)	28	(56%)
Tipo istologico						
adenocarcinoma	184	(92%)	137	(91,3%)	47	(94%)
carcinoma mucinoso	13	(6.5%)	10	(6.7%)	3	(6%)
carcinoma sigillo cellulare	2	(1%)	2	(1.3%)	0	(0%)
carcinoma NOS	1	(0.5%)	1	(0.7%)	0	(0%)
Classificazione AJCC						
1.0	42	(21%)	34	(22.7%)	8	(16%)
IIA	58	(29%)	45	(30%)	13	(26%)
IIB	12	(6%)	10	(6.7%)	2	(4%)
IIC	3	(1.5%)	2	(1.3%)	1	(2%)
IIIA	6	(3%)	3	(2%)	3	(6%)
IIIB	43	(21.5%)	26	(17.3%)	17	(34%)
IIIC	10	(5%)	9	(6%)	1	(2%)
IV	26	(13%)	21	(14%)	5	(10%)

Tab. 3 – CRCs INCIDENT IN 2011 – QC₃ QIs SELECTION

QUALITY INDICATORS (QI)	DENOMINATORS	RESULTS		LITERATURE	
Proportion of patients evaluated by	Patients with colorectal cancer	N %			CCS, R, G
preoperative colonoscopy	undergoing surgery (n=200)	Yes	172	91 9	
		No	17	9	
		Missing	11		
Proportion of patients with preoperative	Patients with colorectal cancer	www	N	%	G
staging according to the AJCC TNM 7 th ed.	undergoing surgery (n=200)	2000		76	G
staging according to the AJCC TNW 7 ed.	undergoing surgery (n=200)	Yes	126	94 6	
		No	8	6	
		Missing	66		
Proportion of patients undergoing rectal-	Patients with rectal cancer (n=76)		N	%	P, CCS, R, M
sigmoidoscopy /colonoscopy		Yes	68	96	,
		No	3	4	
	+	Missing	5		
		iviissing			
Proportion of patients undergoing biopsy	Patients with rectal cancer (n=76)		N	%	P, CCS, R, M
		Yes	65	93	
		No	5 6	7	
		Missing	6		
Proportion of patients with description of the	Patients with rectal cancer (n=76)		N	%	CCS, R
clinical-endoscopic visit, particularly of the		Yes	60	85	CC3, 1.
tumour localization (distance ab ano)					
tumour localization (distance ab ano)		No	11 5	15	
		Missing			
Proportion of patients with definitive	Patients with colorectal cancer		N	%	CCS, R, G, PBS
pathological report including the number of	undergoing surgery (n=200)	Yes	196	98	
lymph nodes retrieved		No	1	2	
		Missing	2		
Proportion of patients with definitive	Patients with colorectal cancer		N	%	G
pathological report including the margin		26	196	99	ď
	undergoing surgery (n=200)	Yes			
status		No	1	1	
		Missing	3		
Proportion of patients with definitive	Patients with colorectal cancer		N	%	ccs, g
pathological report including the pTNM	undergoing surgery (n=200)	Yes	193	98	7.77
classification		No	4	2	
Classification		Missing	3		
		iviissing			
Proportion of patients operated in emergency	Patients with colorectal cancer		N	%	ccs
	undergoing surgery (n=200)	Yes	29	16	
		No	153	84	
		Missing	18		
Proportion of patients operated on with free	Patients with colorectal cancer		N	%	CCS, G, M
margins	undergoing surgery (n=200)	Yes		96	CC3, G, W
margins	undergoing surgery (11–200)		8	96 4	
		No	188	4	
		Missing	4		
Proportion of patients NOT undergoing neo-	Patients with colon cancer and patients		N	%	CCS, R, G, PBS
adjuvant RT or RT-CT, with a number of	with rectal cancer undergoing primary	Yes	148	82	
resected lymph nodes ≥ 12	surgery (n=183)	No	32	18	
		Missing	3		
Proportion of patients with clinical stage from	Patients with AJCC stage I (from	wiissing	N	%	G
I (T2N0M0) to III (every T,N1-2M0) undergoing			171	99	G
	12 Notivio) - III colorectal cancer (N=1/3)	Yes		99 1	
		No	1	1	
anastomosis		Missing	1		
Proportion of patients with metastases for	Patients with colorectal cancer with		N	%	CCS, R, G
which the first line of systemic therapy was	unresectable metastases undergoing	Yes	12	44	
planned on the basis of molecular factors	chemotherapy (n=28)	No		56	
(KRAS, BRAF, etc)	,	Missing	15 1		
	Patients with colorectal cancer with	iviissifig		0.6	_
Proportion of patients with single pulmonary		Yes	N	%	G
			2	7	
metastasis or hepatic metastases undergoing	hepatic or singular pulmonary				
metastasis or hepatic metastases undergoing immediate/synchronous metastasectomy	hepatic or singular pulmonary metastases (n=30)	No	28	93	
			/		
immediate/synchronous metastasectomy	metastases (n=30)	No		93	ccs
immediate/synchronous metastasectomy Proportion of patients with protective stoma	metastases (n=30) Patients with rectal cancer undergoing	No Missing	/ N	%	ccs
immediate/synchronous metastasectomy	metastases (n=30) Patients with rectal cancer undergoing neo-adjuvant radio-chemotherapy	No Missing Yes	/ N 2	% 11	ccs
immediate/synchronous metastasectomy Proportion of patients with protective stoma	metastases (n=30) Patients with rectal cancer undergoing	No Missing Yes	/ N 2 16	%	ccs
Immediate/synchronous metastasectomy Proportion of patients with protective stoma before neo-adjuvant RT-CT	metastases (n=30) Patients with rectal cancer undergoing neo-adjuvant radio-chemotherapy (n=20)	No Missing Yes	/ N 2 16 2	% 11 89	
immediate/synchronous metastasectomy Proportion of patients with protective stoma before neo-adjuvant RT-CT Proportion of patients with locally advanced	metastases (n=30) Patients with rectal cancer undergoing neo-adjuvant radio-chemotherapy (n=20) Patients with locally advanced rectal	No Missing Yes No Missing	/ N 2 16 2 N	% 11 89 %	CCS, R, G
Immediate/synchronous metastasectomy Proportion of patients with protective stoma before neo-adjuvant RT-CT	metastases (n=30) Patients with rectal cancer undergoing neo-adjuvant radio-chemotherapy (n=20)	No Missing Yes	/ N 2 16 2 N 17	% 11 89	
immediate/synchronous metastasectomy Proportion of patients with protective stoma before neo-adjuvant RT-CT Proportion of patients with locally advanced	metastases (n=30) Patients with rectal cancer undergoing neo-adjuvant radio-chemotherapy (n=20) Patients with locally advanced rectal	No Missing Yes No Missing	/ N 2 16 2 N 17	% 11 89 %	
immediate/synchronous metastasectomy Proportion of patients with protective stoma before neo-adjuvant RT-CT Proportion of patients with locally advanced	metastases (n=30) Patients with rectal cancer undergoing neo-adjuvant radio-chemotherapy (n=20) Patients with locally advanced rectal	No Missing Yes No Missing	/ N 2 16 2 N	% 11 89 % 74	

CONCLUSIONS

The study is instrumental to draw a population-based picture of the QoCC currently in use in the territory of Canton Ticino and to open new perspectives on quality-related issues in oncology. In addition, the systematic trend analysis of QI allows to assess immediate changes and improvements in the diagnostic-therapeutic process that could be translated in a short-term benefit for patient, without waiting for survival analysis typically needed some years to be produced because of the patients follow-up. The prospective design allows the production of up-to-date results, reproducing the currently used pattern of care. The population-based design implies the inclusion of the elderly patients usually excluded from RCTs. The study favours the rationalization of data transmission modalities to Cancer Registries and, furthermore, it increases the expectations of Cancer Registry data system, moving from the static retrospective evaluation of cancer treatment outcomes to dynamic interventions to monitor and to ensure optimal multidisciplinary cancer care. Moreover, in a second step, for each QI the minimum and the target requirement at a regional level will be proposed.