

The second ESGAR consensus statement on CT colonography

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Abstract

Objective To update quality standards for CT colonography based on consensus among opinion leaders within the European Society of Gastrointestinal and Abdominal Radiology (ESGAR).

Material and methods A multinational European panel of nine members of the ESGAR CT colonography Working Group (representing six EU countries) used a modified Delphi process to rate their level of agreement on a variety of statements pertaining to the acquisition, interpretation and implementation of CT colonography. Four Delphi rounds were conducted, each at 2 months interval.

Results The panel elaborated 86 statements.

In the final round the panelists achieved complete consensus in 71 of 86 statements (82 %). Categories including the highest proportion of statements with excellent Cronbach's internal reliability were colon distension, scan parameters, use of intravenous contrast agents, general guidelines on patient preparation, role of CAD and lesion measurement.

Lower internal reliability was achieved for the use of a rectal tube, spasmolytics, decubitus positioning and number of CT data acquisitions, faecal tagging, 2D vs. 3D reading, and reporting.

Conclusion The recommendations of the consensus should be useful for both the radiologist who is starting a CTC service and for those who have already implemented the technique but whose practice may need updating.

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Key Points

- *Computed tomographic colonography is the optimal radiological method of assessing the colon*
- *This article reviews ESGAR quality standards for CT colonography*
- *This article is aimed to provide CT-colonography guidelines for practising radiologists*
- *The recommendations should help radiologists who are starting/updating their CTC services*

Keywords CT colonography · Guidelines · Computed tomography · Colon · Polyps

Introduction

Since its introduction (in 1994) [1], clinical implementation of computed tomography (CT) colonography has been governed by advances in CT technology, improvements in dedicated analysis software, development of patient preparation regimens and local diagnostic policies.

In 2007 the European Society of Gastrointestinal and Abdominal Radiology (ESGAR) consensus statement on CT colonography was published, detailing how best to conduct and interpret the examination [2]. That document was based on collective experience up to the beginning of 2006, and the authors represented the EU countries in which CTC underwent consistent clinical implementation (UK, Italy, Belgium and The Netherlands). Over the last 5 years expansion of the CT colonography literature has continued and several important studies, including multicentre studies, have been published [3–5]. These new data have provided further insight regarding optimisation of the CT colonography technique, interpretation and diagnostic capabilities. Indeed CT colonography is now recommended for colorectal cancer screening by several international groupings and is widely used to investigate patients with symptoms suggestive of colorectal cancer [6, 7]. Although recent review articles provide some guidance regarding the optimal CT colonography technique, given the evolving data [8–11] there is a current need to update the ESGAR consensus document.

The purpose of this article is therefore to update quality standards for CT colonography based on examination of the existing literature and expert opinion from key opinion-leaders within the European Society of Gastrointestinal and Abdominal Radiology.

Materials and methods

Consensus panel

A multinational European panel of nine members of the ESGAR CTC Working Group (comprising J.S., S.H., S.T.,

P.L., T.M., D.R., M.H., A.L., E.N., and representing six EU countries: Austria, Belgium, Italy, The Netherlands, Sweden and the UK) used a modified Delphi process [12, 13]. The Delphi process consists of a survey conducted in two or more rounds; the answers (or statements) collected in the first survey are modified in the second, the third, etc., to reach the maximum consensus among the experts. We rated the level of agreement among the experts on a variety of statements pertaining to the acquisition, interpretation and implementation of CT colonography. Four Delphi rounds were conducted, each at 2 months interval.

One of the panellists was chosen as the facilitator (E.N.).

In the first round the facilitator emailed a questionnaire with 22 items pertaining to panel members' personal approaches to CTC, including items on patient preparation, data acquisition technique, image interpretation and clinical implementation (Table 1). Responses collected from all panellists were merged into a unique datasheet that served to identify areas of agreement and conflict in panellist opinion.

In the second round, the panellists attended a 1-day, face-to-face meeting, and, on the basis of their main areas of research and expertise, were divided into four working groups (WG) as follows: bowel preparation and tagging (WG 1), insufflation and scanning protocols (WG 2), reading paradigm (WG 3) and reporting (WG 4). Each WG independently drafted a cluster of statements pertaining to their allocated subject (Table 2). Each statement was built on the basis of panellists' expertise and available indexed literature. Each WG then presented their proposed statements to the whole panel for consideration and subsequent discussion, during which time the content and wording of statements were modified until a general consensus emerged.

In the third and fourth rounds, copies of the latest statements were sent by email to panellists, who then indicated independently their level of agreement with each individual statement using a 5-point scale, as follows: 1, strongly disagree with the statement; 2, disagree somewhat with the statement; 3, undecided; 4, agree somewhat with the statement; 5, strongly agree with the statement.

After the third round the facilitator collected panellists' ratings and determined the agreement score for each statement. If the mean score for an individual item was lower than four (maximum possible=five) the facilitator asked panellists to review the statement and attempt to reach a consensus in the fourth round.

Statistical analysis

To measure the internal consistency of panellist's ratings for each statement, a quality analysis was performed using Cronbach's α correlation coefficient and SPSS (SPSS, Chicago, Ill.) [14]. Cronbach's α was determined after each round.

Table 1 Second ESGAR CT colonography consensus. Survey of the first Delphi round

Please rate your suggestions as follows (in some questions)

1 2 3 4

Do not suggest Could suggest if no other choice Regularly suggest Strongly suggest

Preparation and tagging. Which would you suggest? Cathartic and no tagging Cathartic and fecal tagging Reduced laxative and fecal tagging No laxative and fecal tagging

Cleansing and tagging agents: PEG / MACROGOL Phosphosoda Gastrografin Barium alone Iodine alone Barium and iodine

Cleansing regimen (when to start) (1-4)

1 day before exam 2 days before exam 3 days before exam

Tagging regimen (when to start) (1-4)

Same day of the exam 1 day before exam 2 days before exam 3 days before exam

Insufflation and scanning protocol

Which distension agent? Air CO₂

Who should perform the insufflation? Medical doctor Radiographer Nurse

In which situations you would suggest the use of spasmolytics? (check all that apply):

regularly, in all patients in patients where at scout view the colon appears poorly distended

if patient is finding insufflation unduly uncomfortable other known diverticular disease not used

Which spasmolytics (1-4) Buscopan Glucagon

Which rectal tube (1-4) Flexible with balloon Flexible without balloon Rigid with balloon Rigid without balloon

CT scanner type (minimum number of rows) 1-2 4 8 16 32 64 more rows

CT Low dose protocol: never use use only in asymptomatic patients to use in both asymptomatic and symptomatic patients

CT Normal dose protocol: never use use only in symptomatic patients to use in both asymptomatic and symptomatic patients

Intravenous contrast: not used used in all symptomatic patients used in patients with known cancer or suspected after first series acquisition always used

Decubitus (1-4): Supine first and after prone Prone first and after supine Additional lateral decubitus

Who could read and report the exam?

Radiologist alone Radiographer alone By radiographer (preliminary read) then by radiologist (verification) By resident (preliminary read) then by radiologist (verification) Other (gastroenterologist)

Which is your preferred reading paradigm?

primary 2D + 3D as problem solving primary 3D + 2D as problem solving virtual dissection

Which is your preferred CAD reading paradigm? 2nd reader 1st reader concurrent reading

Who should write the report? radiologist radiographer other (gastroenterologist, etc...)

Do you follow C-RADS (CT Colonography Reporting and Data System) recommendations?

yes no in some occasions (please specify)

When a <6 mm polyp is detected at CTC, what is your suggested policy?

never report report only in symptomatic or high risk patients report it, but don't advise a polypectomy

report it, and advise a polypectomy report it, and advise a follow up

When a 6-9 mm polyp is detected at CTC, what is your suggested policy?

never report report only in symptomatic or high risk patients report but don't advise a polypectomy

report and advise a polypectomy report and advise a follow up

Informed consent should be obtained? yes no

Table 2 Statements elaborated by the panellists in the second Delphi round, and discussed in the third and fourth to reach the maximum consensus and Cronbach's internal reliability. Statements with score

between 4 and 5 are highlighted to show the situations in which all panellists agreed on the statement but the level of support differed (i.e. “agree somewhat” versus “agree strongly”)

Cluster	Stat. #	Second round Statements	Third round		Cronbach's alpha	Fourth round		Cronbach's alpha
			mean scores	std dev		mean scores	std dev	
Rectal tube	1	The use of thin and flexible rectal tubes is recommended (1). Rigid catheters should not be used. Inflation of a small balloon with air is optional.	5.00	.000	0,074	5.00	.000	0,53
	2	If an inflated rectal catheter balloon is used, it should be deflated in one scan acquisition to reduce the risk of masking a distal lesion.	4.78	.441		5.00	.000	
	3	Placement of the rectal tube must be performed by a practitioner specifically trained in the technique. This may be a radiologist, resident, radiographic technician or radiographic nurse, depending on local practice.	5.00	.000		5.00	.000	
Spasmolytics	4	A digital rectal examination is not mandatory. If inflation of a rectal balloon is performed consideration should be given to digital rectal examination.	4.56	.527		4.56	.527	
	5	Use of Spasmolytics (Hyoscin-N-Buthylbromide (Buscopan®)) is preferable prior colonic distension, noting specific contraindications. However, decision to administer spasmolytics should be also based on patient's history (diverticular disease, previous incomplete colonoscopy for strictures, etc).	4.78	.667	0,124	4.78	.667	0,25
	6	Hyoscine-N-butylbromide (buscopan) is the spasmolytic of choice.	5.00	.000		5.00	.000	
	7	If spasmolytics are used, administration should be before commencing insufflation.	4.89	.333		4.89	.333	
	8	If buscopan is contraindicated Glucagon (1 mg) may be used.	4.56	.726		4.67	.500	
Colon distension	9	Automatic distension with CO2 is the method of choice, to optimize colonic distension and to maximise patient comfort	5.00	.000	1	5.00	.000	1
Method of colon distension	10	Manual distension with carbon dioxide or room air is an acceptable alternative if automated insufflations facilities are not available.	5.00	.000		5.00	.000	
	11	Colonic insufflation must be performed by a practitioner specifically trained in the technique. This may be a radiologist, resident, radiographic technician or radiographic nurse, depending on local practice	5.00	.000		5.00	.000	
Quality of colonic distension	12	Colonic distension should be sufficient such that all segments are fully visualized in at least one patient position and ideally in both.	5.00	.000	1	5.00	.000	1
	13	The volume of gas administered does not alone indicate adequate distension.	5.00	.000		5.00	.000	
	14	The optimum insufflated gas volume differs between individuals and should be judged taking into consideration colonic pressure (if measured), patient tolerance and the appearance on the scout image.	5.00	.000		5.00	.000	
	15	The degree and completeness of colonic distension should be checked by inspection of a scout image acquired in both patient positions prior to each full CT data acquisition.	5.00	.000		5.00	.000	
Image acquisition	16	A combination of supine and prone positioning is standard.	5.00	.000		5.00	.000	
Decubitus and number of CT scans	17	If the patient is unable to lie in the prone position, a lateral decubitus scan is recommended.	5.00	.000	N/A	5.00	.000	0,3
	18	If segments are inadequately visualized due to poor distension, an additional scan is indicated with further insufflation and/or in a different position.	4.78	.441		5.00	.000	
	19	There is little evidence that the order of patient positioning (ie supine or prone position first) influences distension quality.	5.00	.000		5.00	.000	
Scan parameters	20	Use of multidetector row CT scanners (= and >4 rows) is a prerequisite for CTC given the requirement to achieve a CT scan of the whole abdomen with a narrow collimation within one breath-hold.	5.00	.000	1	5.00	.000	1
	21	Maximum collimation influences colonic lesion detection and should be no more than 2.5mm, although newer generation CT-Scanners allow routine acquisition of thinner slices which is preferable.	5.00	.000		5.00	.000	
	22	Images should be reconstructed with an overlap. (20-30% overlap).	5.00	.000		5.00	.000	
	23	CT scans should be performed in cranio-caudal direction to minimize breathing artifacts.	5.00	.000		5.00	.000	
	24	Low radiation dose protocols without IV contrast should be used for screening CTC.	5.00	.000		5.00	.000	
	25	120 kV should be used for both supine and prone acquisitions, but lower kV may be acceptable in specific situations.	5.00	.000		5.00	.000	
	26	When IV contrast is not administered ≤ 50 mAs is preferable for prone and supine positions, excepting overweight patients.	5.00	.000		5.00	.000	
	27	Dose modulation and iterative reconstruction should be applied if available.	5.00	.000		5.00	.000	

Table 2 (continued)

<i>IV Contrast</i>	28	IV contrast is not required for colonic evaluation but improves evaluation of extra-colonic organs.	4.89	.333	0,38	5.00	.000	1
	29	Oral tagging agents do not preclude the use of IV contrast.	5.00	.000		5.00	.000	
	30	IV contrast should be administered in all patients with known colorectal cancer (unless contraindicated) to facilitate staging.	5.00	.000		5.00	.000	
	31	In symptomatic patients without known colorectal cancer, routine administration of IV contrast depends on the clinical indication and requirement to fully evaluate the extracolonic organs, especially if an abnormality has been seen on the unenhanced scan.	5.00	.000		5.00	.000	
	32	If IV contrast is administered, acquisition should be in the portal venous phase.	4.89	.333		5.00	.000	
	33	If IV contrast is administered a standard radiation dose protocol should be applied, although a reduced mA acquisition ≤ 50 mAs should be utilized during the unenhanced acquisition	5.00	.000		5.00	.000	
	34	If intravenous contrast media is administered, it is preferable to do so in the supine position	4.89	.333		4.89	.333	
<i>Precautions before and after CT scan</i>	35	Before the patient leaves the CT table, the quality of the examination should ideally be assessed by a practitioner specifically trained in the technique. Specific attention should also be made for the presence of perforation.	4.89	.333	0,75	4.89	.333	0,75
	36	If colonic perforation is a possibility, for example following difficult optical colonoscopy, this should be excluded via acquisition of "low dose" abdominal CT, before starting CTC.	4.89	.333		4.89	.333	
	37	If polypectomy has recently been performed there is a case for delaying CTC depending on the type of biopsy. There is no clear evidence regarding the interval.	5.00	.000		5.00	.000	
Patient preparation	38	General patient preparation for CTC is mandatory for proper detection of polyps and CRC in both symptomatic and asymptomatic individual. This may include dietary restriction, oral contrast agent and bowel purgation.	5.00	.000	1	5.00	.000	1
<i>General guidelines</i>	39	The general patient preparation scheme, including bowel purgation if used, should be straightforward and simple.	5.00	.000		5.00	.000	
	40	An information leaflet with detailed description of the preparation scheme is advised.	5.00	.000		5.00	.000	
<i>Aggressiveness of preparation</i>	41	Tagging regimens should be restricted to no more than 24 hours.	4.67	.707	0,57	4.78	.667	0,6
	42	Aggressive catharsis (purgation) should be restricted to 24 hours or less.	5.00	.000		5.00	.000	
	43	Bowel preparation should include dietary restrictions (e.g. Low fibre diet), to reduce faecal volume and faecal heterogeneity.	5.00	.000		5.00	.000	
	44	The bowel preparation for CTC should normally include laxative agents.	5.00	.000		5.00	.000	
	45	A trade-off between the patient burden and the required image quality to detect the target lesion should be considered when choosing a laxative agent.	5.00	.000		5.00	.000	
	46	CTC without laxative, but with tagging, may be considered in frail and elderly patients where CRC is the diagnostic target.	5.00	.000		5.00	.000	
	47	Sodium phosphate is efficient but not recommended at double dose since this may cause serum electrolyte disturbances, phosphate nephropathy.	4.67	1.000		4.67	1.000	
	48	Magnesium citrate has less side effects and should therefore be preferred over sodium phosphate. However it has restricted availability in Europe.	4.67	1.000		4.67	1.000	
	49	Polyethylene glycol preparations avoid many electrolyte disturbances, but may result in excess colonic fluid.	4.89	.333		4.89	.333	
	50	Iodinated contrast media are used for tagging and also may have a laxative effect.	5.00	.000		5.00	.000	
Faecal tagging	51	Faecal tagging is mandatory.	4.89	.333	0,378	5.00	.000	0,4
	52	Faecal tagging can be achieved with either iodine or barium or both.	4.78	.667		5.00	.000	
	53	Insufficient scientific evidence exists to favour one tagging agent over the other.	4.33	1.118		4.78	.441	
	54	Iodine results in homogeneous tagging which may facilitates interpretation.	5.00	.000		5.00	.000	
	55	Hyperosmolar iodine based preparations have a laxative effect, which should be taken into account.	5.00	.000		5.00	.000	
	56	Caution is necessary when prescribing iodine-based preparations in cases of known iodine-contrast medium allergy.	5.00	.000		5.00	.000	
	57	Barium is inert and consequently has no cathartic effect but may cause constipation.	5.00	.000		5.00	.000	
	58	Barium may produce heterogeneous tagging of stool and fluid.	5.00	.000		5.00	.000	
	59	Barium suspensions may impair same-day colonoscopy.	5.00	.000		5.00	.000	

Table 2 (continued)

	60	Barium AND iodine combine stool and fluid tagging which may be desirable, but this more complex preparation scheme may reduce patient compliance.	5.00	.000		5.00	.000	
	61	There is a wide variability in patient preparation schemes between experienced centres.	5.00	.000		5.00	.000	
Reading paradigm	62	Interpretation of CT colonography should incorporate both 2D and 3D visualization (i.e. fly-through).	4.89	.333	N/A	4.89	.333	N/A
<i>2D and 3D reading</i>	63	Initial interpretation using either primary 2D or primary 3D methods are acceptable depending on personal preference and on WS availability.	5.00	.000		5.00	.000	
	64	On average the primary 2D interpretation is likely to be faster.	5.00	.000		5.00	.000	
	65	Other 3D visualization options (e.g. virtual dissection, panoramic view, file view, ecc) are viable alternatives provided that the reader is fully trained in conventional 2D and 3D visualisation displays, and is aware that other data display may introduce distortion.	5.00	.000		5.00	.000	
<i>CAD</i>	66	2nd read CAD is recommended because it increases sensitivity for polyp detection without an unacceptable decrease in specificity.	5.00	.000	1	5.00	.000	1
	67	Readers should be aware that it is possible to reject true positive CAD prompts in error.	5.00	.000		5.00	.000	
	68	CAD should be adopted by radiologists only after they have been adequately trained in unassisted interpretation of CT colonography and the use of CAD.	5.00	.000		5.00	.000	
	69	CAD is an adjunct to unassisted interpretation and its implementation will depend on local factors including costs, personal preference and algorithm and/or WS availability.	5.00	.000		5.00	.000	
	70	CAD is less likely to be useful in situations where there are multiple false positive prompts, for example a poorly prepared colon.	5.00	.000		5.00	.000	
	71	CAD algorithms have been developed primarily for polyp detection although they may also detect cancer.	5.00	.000		5.00	.000	
Lesion measurement	72	The maximal diameter of a lesion should be measured on the plane that best demonstrates this dimension, excluding any stalk if present, and its segmental location reported.	5.00	.000	1	5.00	.000	1
	73	Diameter may be estimated using 2D and or 3D methods but readers should be aware that 3D estimates may occasionally be unreliable.	5.00	.000		5.00	.000	
	74	Readers should be aware that there is frequent disagreement between CT and the endoscopic measurements, and this may influence management when patients are defined by polyp size categories.	5.00	.000		5.00	.000	
	75	Readers should be aware that neither endoscopic nor CT estimates are wholly accurate and both are affected by the way the measurement is made (e.g. CT window level and width). Narrow windows should be avoided.	5.00	.000		5.00	.000	
<i>Flat lesions</i>	76	The precise definition of a flat lesion is variable and controversial at the present time. Lesion height above the surrounding mucosa should be reported when flat lesions are encountered. An increasingly acceptable definition of a flat lesion on CT colonography is one where the elevation of a lesion of 6mm or larger above the surrounding mucosa is 3 mm or less.	4.89	.333	N/A	5.00	.000	1
	77	Readers should be aware that CT colonography is less sensitive for flat lesions than for other polyp morphologies.	5.00	.000		5.00	.000	
	78	The likelihood of cancer increases in line with lesion diameter. There is no exclusive threshold that defines cancer at CT colonography. When the morphology of the lesion strongly suggests a cancer this terminology should be used and alternatives such as "mass" should be avoided.	5.00	.000		5.00	.000	
	79	Occasionally factors other than the maximal diameter of a single lesion may be useful to indicate the clinical importance, for example fat attenuation which indicates a lipoma.	5.00	.000		5.00	.000	
Reporting	80	A report should include the Clinical information (under pinning the request should be included in the report, along with personal and family history), the Technical data (low or normal dose protocol, intravenous contrast) and, if desired, preparation and tagging (laxative agent), tagging (tagging regimen), insufflation (air or CO ₂), spasmolytics (used or not used), the effective dose in mSv.	4.44	1.014	0,4	4.44	1.014	0.4
	81	The reported colonic findings should be: colonic anatomy (normal or abnormal.), polyps and cancer (size, shape, maximum diameter, infiltration of the extracolonic fat, location and other colonic (e.g. wall thickening, strictures, diverticula, extrinsic compressions, post-surgical variations).	5.00	.000		5.00	.000	
	82	The extracolonic organs should be interrogated and abnormalities reported, noting the limitations if an unenhanced and or low dose technique was used.	4.89	.333		4.89	.333	
	83	CT colonography should be reported by a radiologist, specifically trained in the technique.	5.00	.000		5.00	.000	

Table 2 (continued)

84	CT Colonography reading requires specific training and expertise in abdominal cross-sectional imaging, which is only conveyed by radiological training	5.00	.000	5.00	.000
85	CT colonography has limited diagnostic value for lesions less than 6 mm. However, if detected with high confidence such lesions might be reported (particularly if ≥ 3), in both asymptomatic and symptomatic patients.	5.00	.000	5.00	.000
86	All polyps of 6mm or larger should be reported in both asymptomatic and symptomatic.	5.00	.000	5.00	.000

Cronbach's α reliability coefficient normally ranges between 0 and 1. The closer the Cronbach's α coefficient is to 1.0, the greater the internal consistency of the item. An α coefficient >0.9 was considered excellent, $\alpha >0.8$ good, $\alpha >0.7$ acceptable, $\alpha >0.6$ questionable, $\alpha >0.5$ poor and $\alpha <0.5$ unacceptable. For the iterations, an α of 0.8 was considered a reasonable goal for internal reliability. All panellist ratings for each statement were also analysed with descriptive statistics, estimating the mean, maximum and minimum score, and their standard deviation.

A mean score of 4 was considered to represent "good" agreement between panellists, a score of 5 "complete" agreement.

Results

Based on the questionnaire provided by the facilitator, the panel elaborated 86 statements that were collected by the facilitator and organised into nine groups, as follows: (1) rectal tube, (2) spasmolytics, (3) colon distension, (4) image acquisition, (5) patient preparation, (6) faecal tagging, (7) reading paradigm, (8) lesion measurement and (9) reporting (Table 2).

In the third round the panelists achieved complete consensus (i.e. mean score 5) in 64 of 86 statements (75 %), which improved to 71 (82 %) in the fourth round (Table 2).

Categories including the highest proportion of statements achieving excellent internal reliability (i.e. Cronbach's α value >0.7) in the final round were colon distension, scan parameters, use of intravenous contrast medium, general guidelines on patient preparation, role of CAD and lesion measurement.

Lower internal reliability was achieved for statements regarding the use of a rectal tube, spasmolytics, decubitus positioning and number of CT data acquisitions, faecal tagging, 2D vs. 3D reading and reporting. However, in the last round, no panellist scored their individual statements as less than 4 on the 5-point rating scale. This indicates that all panellists agreed on the statement but the level of support differed (i.e. "agree somewhat" versus "agree strongly").

Discussion

Full consensus was reached by our expert panel in 82 % of the statements. In the remaining statements, full consensus

was not reached but all panellists achieved a "good" level of agreement. In total, the panellists completed four rounds; the first and second rounds served to elaborate the basic statements. The third and fourth rounds contained the core of the discussion and were necessary to reach the maximum consensus possible, so creating an optimised, homogeneous opinion for each statement.

All panellists exhibited a high level of agreement for the technical performance of CTC, with clear recommendations regarding colon distension, CT parameters, use of intravenous contrast agents and patient preparation. Full agreement was also reached regarding the role of CAD and lesion measurement. These data reflect a general homogeneity of approach between panel members despite their wide geographical spread. All panel members are regular tutors on the ESGAR CTC course, which may have increased their level of agreement; there is a tendency to promote a common message during panel discussions occurring during the ESGAR CTC courses [15, 16]. Furthermore, in these areas the indexed literature is relatively mature and stable; for example available data supporting the use of automated CO² for optimal colonic distension is relatively consistent [17–20].

However, certain aspects of practice achieved less than "full" agreement. In particular, a digital rectal examination, before insertion of the rectal tube (if rectal examination had not been performed previously), was not standard practice in many centres, but was nevertheless recommended by some panellists (with a mean score 4.56). This difference could be explained by the practice to perform a digital rectal examination before CTC amongst a few of the experts involved in the consensus. Similarly, practice differed regarding the use of intravenous spasmolytics, with many administering such agents to all patients, whereas some (in Italy) only used it in selected individuals [21, 22]. Accordingly, use of spasmolytics is recommended by the majority but is not considered mandatory.

There were minor variations in recommended CT parameters between panellists but all recommended data acquisition in at least two patient positions, without any overall preference regarding the order of acquisitions (i.e. supine or prone first). The differences in CT protocols included the need for additional CT data acquisition and insufflation in cases of poor colonic distension; a minority of experts did not consider this mandatory although they agreed it should be recommended. An additional decubitus acquisition was recommended, if required, to improve the diagnostic quality of the examination [23, 24].

Although available CT technology differed among panellists, all agreed that 2.5-mm collimation was the maximum permissible (although thinner collimation is recommended when available) and use of low radiation dose protocols is to be employed when the overriding purpose of the study is the evaluation of the colonic lumen, for example as in screening [25, 26]. A low radiation dose should be considered a study in which the median effective dose is lower than 5.7 mSv, according to the results of the survey by Leidenbaum et al. [26]. For the staging of patients with known malignancy all the panellists agreed upon the use of standard-dose protocols and intravenous contrast medium [27, 28].

Substantial agreement was reached between panelists regarding the reading methods for interpretation of CT colonography. A combination of 2D and 3D reading was emphasised. Most of the panel were primary 2D readers but all recognised the importance of 3D integration, noting the range of different three-dimensional approaches available. The need for the reader to be adequately trained before interpreting CT colonography was emphasised and is strongly supported by the indexed literature [29–33].

Computer-aided diagnosis was acknowledged by all panellists as a potentially useful tool for CTC interpretation, if employed in a second reader paradigm. Accordingly, the use of CAD was recommended provided that readers have already undergone adequate training in general CT colonography interpretation so that they can discriminate between true- and false-positive CAD marks appropriately [34–42].

Panellists acknowledged that accurate polyp measurement is problematic for both CTC and endoscopy, with some evidence that CTC may be the superior technique [43, 44]. Despite this advantage, it is still uncertain whether a 2D or a 3D measurement should be made from CT. Moreover, the accuracy of such measurements has important clinical implications for the correct classification and risk stratification of lesions, influencing subsequent recommendations for patient management [45–50]. The panel concluded that the maximal diameter of lesions should be primarily estimated using axial and MPR 2D views (which were considered to be the most reliable), avoiding a narrow CT window. Some caution should be exercised when measurements are taken using 3D perspectives given the potential for distortion generated by the three-dimensional endoluminal rendering [51–55].

All panellists agreed that CTC should only be reported by a radiologist, and then only after adequate training [56–59]. Motivations behind this recommendation are mainly the medico-legal implications of non-radiologists reporting CTC in EU countries. In all EU countries the radiological report is definitively validated by the radiologist despite, in a few centres, a preliminary reading being performed by a radiographer. Adequate training means having interpreted a minimum amount of colonoscopy-verified cases. Although the precise number has not yet been clearly defined, the

literature shows that 175 is even not sufficient for several individuals [60, 61].

It was acknowledged that diagnostic accuracy is lower for polyps with a maximal diameter less than 6 mm [3, 4] but if detected with high confidence, and particularly if more than three in number, such polyps should still be reported. This contrasts with recommendations from the CT Colonography Reporting and Data System (C-RADS), authored by Zalis et al., where lesions less than 6 mm are considered diminutive and the recommendation is that they should not be reported [45]. The panel agreed that the patient's risk (age, family history of colorectal cancer, previous polypectomy, etc.), as well as the number of diminutive lesions detected, should be considered in the decision to report them or not.

There was little disagreement between panellists regarding the need to calibrate the laxative effect of bowel preparation/purgation to the individual patient and potential target lesion. All panellists agreed that faecal tagging should be used routinely. Different preferences for specific laxative and tagging agents were expressed (for example sodium phosphate, magnesium citrate, polyethylene glycol for cleansing, and barium, iodine or a combination of both agents for tagging), reflecting local practice [62–75].

In summary, the panel covered all important aspects regarding the practice of CTC and reached full agreement on most statements. The Consensus has been structured to give clear guidelines for the practice of CT colonography. The recommendations should be useful for both the radiologist who is starting a CTC service and for those who have already implemented the technique but whose practice may need updating in the light of recent developments.

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References

1. Vining DJ, Celfand DW, Bechtold RE, Scharling ES, Grishaw EK, Shifrin RY (1994) Technical feasibility of colon imaging with helical CT and virtual reality. *AJR* 162:104
2. Taylor SA, Laghi A, Lefere P, Halligan S, Stoker J (2007) European Society of Gastrointestinal and Abdominal Radiology (ESGAR): consensus statement on CT colonography. *Eur Radiol* 17:575–579
3. Johnson CD, Chen MH, Toledano AY et al (2008) Accuracy of CT colonography for detection of large adenomas and cancers. *N Engl J Med* 359:1207–1217
4. Regge D, Laudi C, Galatola G et al (2009) Diagnostic accuracy of computed tomographic colonography for the detection of

- advanced neoplasia in individuals at increased risk of colorectal cancer. *JAMA* 17:2453–2461
5. Graser A, Stieber P, Nagel D et al (2009) Comparison of CT colonography, colonoscopy, sigmoidoscopy and faecal occult blood tests for the detection of advanced adenoma in an average risk population. *Gut* 58:241–248
 6. Levin B, Lieberman DA, McFarland B et al (2008) American Cancer Society Colorectal Cancer Advisory Group; US Multi-Society Task Force; American College of Radiology Colon Cancer Committee. Screening and surveillance for the early detection of colorectal cancer and adenomatous polyps, 2008: a joint guideline from the American Cancer Society, the US Multi-Society Task Force on Colorectal Cancer, and the American College of Radiology. *CA Cancer J Clin* 58:130–160
 7. McFarland EG, Fletcher JG, Pickhardt P et al (2009) American College of Radiology. ACR Colon Cancer Committee white paper: status of CT colonography 2009. *J Am Coll Radiol* 6:756–772
 8. Mang T, Graser A, Schima W, Maier A (2007) CT colonography: techniques, indications, findings. *Eur J Radiol* 61:388–399
 9. Yee J, Rosen MP, Blake MA (2010) ACR Appropriateness Criteria on colorectal cancer screening. *J Am Coll Radiol* 7:670–678
 10. Cash BD (2010) Establishing a CT colonography service. *Gastrointest Endosc Clin N Am* 20:379–398
 11. Mang T, Schima W, Brownstone E et al (2011) Consensus statement of the Austrian Society of Radiology, the Austrian Society of Gastroenterology and Hepatology and the Austrian Society of Surgery on CT colonography (Virtual Colonoscopy). *Rofo* 183:177–184
 12. Graham B, Regehr G, Wright JG (2003) Delphi as a method to establish consensus for diagnostic criteria. *J Clin Epidemiol* 56:1150–1156
 13. Vakil N (2011) Editorial: consensus guidelines: method or madness? *Am J Gastroenterol* 106:225–227
 14. Cronbach LJ (1951) Coefficient alpha and the internal structure of tests. *Psychometrika* 16:3
 15. Burling D (2010) International Collaboration for CT Colonography Standards. CT colonography standards. *Clin Radiol* 65:474–480
 16. Boone D, Halligan S, Frost R et al (2011) CT colonography: who attends training? A survey of participants at educational workshops. *Clin Radiol* 66:510–516
 17. Shinnars TJ, Pickhardt PJ, Taylor AJ, Jones DA, Olsen CH (2006) Patient-controlled room air insufflation versus automated carbon dioxide delivery for CT colonography. *AJR Am J Roentgenol* 186:1491–1496
 18. Burling D, Taylor SA, Halligan S et al (2006) Automated insufflation of carbon dioxide for MDCT colonography: distension and patient experience compared with manual insufflation. *AJR Am J Roentgenol* 186:96–103
 19. Kim SY, Park SH, Choi EK et al (2008) Automated carbon dioxide insufflation for CT colonography: effectiveness of colonic distension in cancer patients with severe luminal narrowing. *AJR Am J Roentgenol* 190:698–706
 20. Neri E, Laghi A, Regge D (2008) Re: Colonic perforation during screening CT colonography using automated CO₂ insufflation in an asymptomatic adult. *Abdom Imaging* 33:748–749
 21. Taylor SA, Halligan S, Goh V et al (2003) Optimizing colonic distension for multi-detector row CT colonography: effect of hyoscine butylbromide and rectal balloon catheter. *Radiology* 229:99–108
 22. Rogalla P, Lembcke A, Rückert JC et al (2005) Spasmolysis at CT colonography: butyl scopolamine versus glucagon. *Radiology* 236:184–188
 23. Gryspeerdt SS, Herman MJ, Baekelandt MA, van Holsbeeck BG, Lefere PA (2004) Supine/left decubitus scanning: a valuable alternative to supine/prone scanning in CT colonography. *Eur Radiol* 14:768–777
 24. Buchach CM, Kim DH, Pickhardt PJ (2011) Performing an additional decubitus series at CT colonography. *Abdom Imaging* 36:538–544
 25. Graser A, Wintersperger BJ, Suess C, Reiser MF, Becker CR (2006) Dose reduction and image quality in MDCT colonography using tube current modulation. *AJR Am J Roentgenol* 187:695–701
 26. Liedenbaum MH, Venema HW, Stoker J (2008) Radiation dose in CT colonography—trends in time and differences between daily practice and screening protocols. *Eur Radiol* 18:2222–2230
 27. Filippone A, Ambrosini R, Fuschi M, Marinelli T, Genovesi D, Bonomo L (2004) Preoperative T and N staging of colorectal cancer: accuracy of contrast-enhanced multi-detector row CT colonography—initial experience. *Radiology* 231:83–90
 28. Mainenti PP, Cirillo LC, Camera L et al (2006) Accuracy of single phase contrast enhanced multidetector CT colonography in the preoperative staging of colo-rectal cancer. *Eur J Radiol* 60:453–459
 29. Taylor SA, Halligan S, Slater A et al (2006) Polyp detection with CT colonography: primary 3D endoluminal analysis versus primary 2D transverse analysis with computer-assisted reader software. *Radiology* 239:759–767
 30. Neri E, Vannozzi F, Vagli P, Bardine A, Bartolozzi C (2006) Time efficiency of CT colonography: 2D vs 3D visualization. *Comput Med Imaging Graph* 30:175–180
 31. Mang T, Schaefer-Prokop C, Schima W, Maier A et al (2009) Comparison of axial, coronal, and primary 3D review in MDCT colonography for the detection of small polyps: a phantom study. *Eur J Radiol* 70:86–93
 32. Mang T, Kolligs FT, Schaefer C, Reiser MF, Graser A (2011) Comparison of diagnostic accuracy and interpretation times for a standard and an advanced 3D visualisation technique in CT colonography. *Eur Radiol* 21:653–662
 33. Lostumbo A, Wanamaker C, Tsai J, Suzuki K, Dachman AH (2010) Comparison of 2D and 3D views for evaluation of flat lesions in CT colonography. *Acad Radiol* 17:39–47
 34. Baker ME, Bogoni L, Obuchowski NA et al (2007) Computer-aided detection of colorectal polyps: can it improve sensitivity of less-experienced readers? Preliminary findings. *Radiology* 245:140–149
 35. Taylor SA, Burling D, Roddie M et al (2008) Computer-aided detection for CT colonography: incremental benefit of observer training. *Br J Radiol* 81:180–186
 36. Petrick N, Haider M, Summers RM et al (2008) CT colonography with computer-aided detection as a second reader: observer performance study. *Radiology* 246:148–156
 37. Taylor SA, Charman SC, Lefere P et al (2008) CT colonography: investigation of the optimum reader paradigm by using computer-aided detection software. *Radiology* 246:463–471
 38. Regge D, Hassan C, Pickhardt PJ et al (2009) Impact of computer-aided detection on the cost-effectiveness of CT colonography. *Radiology* 250:488–497
 39. de Vries AH, Jensch S, Liedenbaum MH et al (2009) Does a computer-aided detection algorithm in a second read paradigm enhance the performance of experienced computed tomography colonography readers in a population of increased risk? *Eur Radiol* 19:941–950
 40. Fischella VA, Jäderling F, Horvath S et al (2009) Computer-aided detection (CAD) as a second reader using perspective file view at CT colonography: effect on performance of inexperienced readers. *Clin Radiol* 64:972–982
 41. Halligan S, Mallett S, Altman DG et al (2011) Incremental benefit of computer-aided detection when used as a second and concurrent reader of CT colonographic data: multiobserver study. *Radiology* 258:469–476
 42. Dachman AH, Obuchowski NA, Hoffmeister JW et al (2010) Effect of computer-aided detection for CT colonography in a multireader, multicase trial. *Radiology* 256:827–835

43. Punwani S, Halligan S, Irving P et al (2008) Measurement of colonic polyps by radiologists and endoscopists: who is most accurate? *Eur Radiol* 18:874–881
44. Jeong JY, Kim MJ, Kim SS (2008) Manual and automated polyp measurement comparison of CT colonography with optical colonoscopy. *Acad Radiol* 15:231–239
45. Zalis ME, Barish MA, Choi JR et al (2005) Working Group on Virtual Colonoscopy. CT colonography reporting and data system: a consensus proposal. *Radiology* 236:3–9
46. Kim DH, Pickhardt PJ, Taylor AJ (2007) Characteristics of advanced adenomas detected at CT colonographic screening: implications for appropriate polyp size thresholds for polypectomy versus surveillance. *AJR Am J Roentgenol* 188:940–944
47. Pickhardt PJ, Hassan C, Laghi A et al (2008) Clinical management of small (6- to 9-mm) polyps detected at screening CT colonography: a cost-effectiveness analysis. *AJR Am J Roentgenol* 191:1509–1516
48. Shah JP, Hynan LS, Rockey DC (2009) Management of small polyps detected by screening CT colonography: patient and physician preferences. *Am J Med* 122:687–689
49. Heresbach D, Chauvin P, Hess-Migliorretti A, Riou F, Grolier J, Josselin JM (2010) Cost-effectiveness of colorectal cancer screening with computed tomography colonography according to a polyp size threshold for polypectomy. *Eur J Gastroenterol Hepatol* 22:716–723
50. Neri E, Faggioni L, Vagli P et al (2011) Patients' preferences about follow-up of medium size polyps detected at screening CT colonography. *Abdom Imaging* 36:713–717
51. Pickhardt PJ, Lee AD, McFarland EG, Taylor AJ (2005) Linear polyp measurement at CT colonography: in vitro and in vivo comparison of two-dimensional and three-dimensional displays. *Radiology* 236:872–878
52. Burling D, Halligan S, Altman DG et al (2006) Polyp measurement and size categorisation by CT colonography: effect of observer experience in a multi-centre setting. *Eur Radiol* 16:1737–1744
53. Park SH, Choi EK, Lee SS et al (2007) Polyp measurement reliability, accuracy, and discrepancy: optical colonoscopy versus CT colonography with pig colonic specimens. *Radiology* 244:157–164
54. Park SH, Choi EK, Lee SS et al (2008) Linear polyp measurement at CT colonography: 3D endoluminal measurement with optimized surface-rendering threshold value and automated measurement. *Radiology* 246:157–167
55. Bethea E, Nwawka OK, Dachman AH (2009) Comparison of polyp size and volume at CT colonography: implications for follow-up CT colonography. *AJR Am J Roentgenol* 193:1561–1567
56. Pickhardt PJ (2009) Editorial: CTC interpretation by gastroenterologists: feasible but largely impractical, undesirable, and misguided. *Am J Gastroenterol* 104:2932–2934
57. Carpenter S (2010) Gastroenterologists should read CT colonography. *Gastrointest Endosc Clin N Am* 20:271–277
58. Kim DH, Pickhardt PJ (2010) Radiologists should read CT colonography. *Gastrointest Endosc Clin N Am* 20:259–69
59. Fletcher JG, Chen MH, Herman BA et al (2010) Can radiologist training and testing ensure high performance in CT colonography? Lessons From the National CT Colonography Trial. *AJR Am J Roentgenol* 195:117–125
60. Heresbach D, Djabbari M, Riou F et al (2011) Accuracy of computed tomographic colonography in a nationwide multicentre trial, and its relation to radiologist expertise. *Gut* 60:658–665
61. Liednbaum MH, Bipat S, Bossuyt PM et al (2011) Evaluation of a standardized CT colonography training program for novice readers. *Radiology* 258:477–487
62. Lefere PA, Gryspeerdt SS, Dewyspelaere J, Baekelandt M, Van Holsbeeck BG (2002) Dietary fecal tagging as a cleansing method before CT colonography: initial results polyp detection and patient acceptance. *Radiology* 224:393–403
63. Iannaccone R, Laghi A, Catalano C et al (2004) Computed tomographic colonography without cathartic preparation for the detection of colorectal polyps. *Gastroenterology* 127:1300–1311
64. Gryspeerdt S, Lefere P, Herman M et al (2005) CT colonography with fecal tagging after incomplete colonoscopy. *Eur Radiol* 15:1192–1202
65. Lefere P, Gryspeerdt S, Marrannes J, Baekelandt M, Van Holsbeeck B (2005) CT colonography after fecal tagging with a reduced cathartic cleansing and a reduced volume of barium. *AJR Am J Roentgenol* 184:1836–1842
66. Zalis ME, Perumpillichira JJ, Magee C, Kohlberg G, Hahn PF (2006) Tagging-based, electronically cleansed CT colonography: evaluation of patient comfort and image readability. *Radiology* 239:149–159
67. Taylor SA, Slater A, Burling DN et al (2008) CT colonography: optimisation, diagnostic performance and patient acceptability of reduced-laxative regimens using barium-based faecal tagging. *Eur Radiol* 18:32–42
68. Slater A, Planner A, Bungay HK, Bose P, Milburn S (2009) Three-day regimen improves faecal tagging for minimal preparation CT examination of the colon. *Br J Radiol* 82:545–548
69. Neri E, Turini F, Cerri F, Vagli P, Bartolozzi C (2009) CT colonography: same-day tagging regimen with iodixanol and reduced cathartic preparation. *Abdom Imaging* 34:642–647
70. Liednbaum MH, de Vries AH, Gouw CI et al (2010) CT colonography with minimal bowel preparation: evaluation of tagging quality, patient acceptance and diagnostic accuracy in two iodine-based preparation schemes. *Eur Radiol* 20:367–376
71. Campanella D, Morra L, Delsanto S et al (2010) Comparison of three different iodine-based bowel regimens for CT colonography. *Eur Radiol* 20:348–358
72. Zueco Zueco C, Sobrido Sampedro C, Corroto JD, Rodriguez Fernández P, Fontanillo Fontanillo M (2012) CT colonography without cathartic preparation: positive predictive value and patient experience in clinical practice. *Eur Radiol* 22:1195–204
73. Liednbaum MH, Denters MJ, de Vries AH et al (2010) Low-fiber diet in limited bowel preparation for CT colonography: Influence on image quality and patient acceptance. *AJR Am J Roentgenol* 195:31–37
74. Davis W, Nisbet P, Hare C, Cooke P, Taylor SA (2011) Non-laxative CT colonography with barium-based faecal tagging: is additional phosphate enema beneficial and well tolerated? *Br J Radiol* 84:120–125
75. Liednbaum MH, Denters MJ, Zijta FM et al (2011) Reducing the oral contrast dose in CT colonography: evaluation of faecal tagging quality and patient acceptance. *Clin Radiol* 66:30–37