

BC Cancer Colon Screening Pre-Post Colonoscopy Standards

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Pre-Post Colonoscopy Assessment Standards

Colon Screening Program

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About BC Cancer

BC Cancer, an agency of the Provincial Health Services Authority, provides a comprehensive cancer control program for the people of BC in partnership with regional health authorities. This includes prevention, screening and early detection programs, research and education, and care and treatment.

BC Cancer's mandate is a three-fold mission:

- To reduce the incidence of cancer
- To reduce the mortality rate of people with cancer
- To improve the quality of life of people living with cancer

This mission drives everything we do, including providing screening, diagnosis and care, setting treatment standards, and conducting research into causes of, and cures for cancer.

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1. Introduction

1.1 Colon Screening Program

Colorectal cancer (CRC) is the second most commonly diagnosed cancer and the second leading cause of cancer death in men and third leading cause of cancer death in women¹. The Colon Screening Program seeks to reduce the incidence and mortality of colorectal cancer by providing timely and equitable access to high quality screening and diagnostic services to eligible people. The program is available in all areas of B.C. except Northern Health Authority where local screening processes are used in lieu of the BC Colon Screening Program process.

1.2 Purpose of the Standards

These standards are designed to maximize participant safety and program efficiency and efficacy by ensuring pre and post colonoscopy assessment is carried out in a safe, effective and consistent manner across the province.

1.3 Sources of Information

The Pre-Post Colonoscopy Assessment Standards are based on the experiences of the BC Cancer Colon Check pilot program, the Vancouver Island Health Authority Pilot Program and the NHS Bowel Cancer Screening Programme (UK).

1.4 General Principles

- Maximize follow-up colonoscopy uptake for participants with a positive FIT
- Optimize participant understanding of colonoscopy
- Optimize participant satisfaction
- Minimize colonoscopy related complications
- Optimize follow-up screening and surveillance

1.5 Program Eligibility

Eligible participants are referred to the program by primary care providers. There are three main categories of eligibility:

- 1. Individuals, age 50 to 74 years, without a personal history of pre-cancerous colorectal polyps nor a high-risk family history of colorectal cancer, will be offered FIT every two years.
- 2. Individuals with a personal history of a pre-cancerous colorectal polyp will be offered colonoscopy every 5 years until age 74.
- 3. Individuals with a high-risk family history of colorectal cancer will be offered

colonoscopy every 5 years, commencing at 40 years of age or 10 years younger than the age of colorectal cancer diagnosis of the youngest affected relative, whichever is earliest. A high-risk family history is defined as a single first degree relative (parent, sibling, child) diagnosed with colorectal cancer at less than 60 years of age or two or more first degree relatives diagnosed with colorectal cancer at any age.

Primary care providers are provided with information on the eligibility criteria for the program and it is expected that providers consider and adhere to the criteria. However, some participants will be asked to complete a FIT inappropriately.

If the FIT is abnormal, the Colon Screening Program recommends colonoscopy in all of the following scenarios:

- Participant had a normal FIT recently and is not yet due for repeat FIT.
- Participant is in a colonoscopy surveillance program for a personal history of pre-cancerous colorectal polyps or a high-risk family history of colorectal cancer but has a FIT that is abnormal.
- Participant who had an abnormal FIT followed by a colonoscopy in which neither
 colorectal cancer nor pre-cancerous colorectal polyps were identified and the
 next recommended screening is FIT in 10 years. The participant undergoes FIT
 before they are due and it is abnormal.

Colonoscopy is protective for ten years and previous guidelines based on data using the guaiac fecal occult blood test stated that an abnormal guaiac fecal occult blood test following a negative colonoscopy could be ignored. However, given the improved performance of FIT, more recent guidelines have recommended that colonoscopy be offered to participants with an early FIT that is abnormal. These recommendations were graded as weak and based on low quality evidence. However, a further peer-reviewed publication has demonstrated a risk of post-colonoscopy colorectal cancer in this group. Despite the risk of colorectal cancer for participants with an abnormal FIT who are not yet due for colonoscopy, data does not support the addition of FIT to colonoscopy surveillance in participants with a personal history of neoplastic polyps or a family history of colorectal cancer. There are harms associated with over-screening and the best defense against post-colonoscopy colorectal cancer is ensuring the initial exam is high quality.

Participants not eligible for screening within the Colon Screening Program:

- 1. Personal history of colorectal cancer or inflammatory bowel disease (Crohn's disease or ulcerative colitis)
 - Require individualized screening directed by a colonoscopist.
- 2. Outside the screening age range
 - Participants with abnormal FIT results who are under 50 years old are not referred on for further follow-up. Participants up to age 75.5 are referred for further follow-up to allow for participants who may have been offered FIT prior to their 75th birthday but did not complete the test until after. Those over age

- 75.5 are not referred to the Health Authority for further follow-up.
- If the participant is older than 74 years, then the participant will not be re-called by the Colon Screening Program for further screening or surveillance.
- 3. Symptoms that require a full colonoscopist assessment
 - Local processes should be used in determining whether a participant with symptoms should be assessed and booked through the usual Colon Screening Program follow-up process or if the provider should be notified to refer for follow-up through a different process. In general, the Colon Screening Program is supportive of maintaining participants in the program for follow-up to reduce re-routing referrals and improve follow-up efficiency for participants.
 Participants with significant symptoms should consult with the colonoscopist prior to the procedure.
- 4. Participants with a high-risk family history or a personal history of pre-cancerous colorectal polyps that are referred for colonoscopy but are not yet due
 - Do not complete the colonoscopy and use the Referral Update Form to indicate
 when the participant is due for colonoscopy (see Appendix H). The participant
 will be recalled for colonoscopy when next due. Participants who will be over
 the age of 74 years when due for colonoscopy will not be referred by the
 Program.

2. Hospital and Endoscopy Unit Standards

2.1 Assessment and Participant Education

- Contact referred participants and establish a time to complete assessment. Each
 Regional Health Authority will determine whether the assessment takes place by
 telephone, in person or through group education sessions. Self-reported height and
 weight is acceptable for phone assessments.
- Confirm the participant's primary care provider. A primary care provider is required for participants undergoing colonoscopy to support any follow-up that the participant may need.
- Confirm family history or personal history of pre-cancerous colorectal polyps for those being referred for screening colonoscopy. If the information provided does not meet the program eligibility requirements for screening colonoscopy communicate this back with the participant's primary care provider to ensure appropriate screening is arranged.
- Complete pre-colonoscopy assessment. The elements of a recommended assessment are available in the Assessment Form example (see Appendix A).
 - Identify any high risk factors that require colonoscopist assessment prior to colonoscopy and liaise with colonoscopist as indicated. See Section 3 and Participant Assessment Process document (see Appendix A).
- Identify the presence of a high-risk family history for hereditary colorectal cancer. If a
 high-risk family history is identified, advise the participant to discuss their history with
 their primary care provider.
- Provide education to the participant regarding:
 - o Implications of an abnormal FIT and the reasons for colonoscopy follow-up.
 - Colonoscopy is always indicated after a positive FIT, even if there is a subsequent negative FIT.
 - Bowel preparation and colonoscopy.
 - Explain the risks of colonoscopy.
 - Provide the participant with the Colon Screening Program Colonoscopy Brochure (sample in Appendix B) to inform them about colonoscopy.
 - Give the participant written bowel preparation instructions, based on the assessment and the local practices for selecting bowel preparation type.
- Book participant for colonoscopy:
 - If not proceeding to colonoscopy, advise primary care provider using Not Proceeding to Colonoscopy letter and send the Referral Update Form to the Colon Screening Program (see Appendix C and H).
- Participants who do not proceed to their colonoscopy within 6 months of the assessment should be re-assessed prior to proceeding to colonoscopy.

2.2 Bowel Preparation

Participants should be provided with written preparation instructions as per the Bowel Preparation Algorithm in Appendix G.

Fleet phospho-soda is contraindicated. (Health Canada Reference: www.healthycanadians.gc.ca/recall-alert-rappel-avis/hc-sc/2009/9807r-eng.php)

Split-dose bowel preparations, in which the second dose of the bowel preparation is given 4 to 6 hours prior to the colonoscopy and same-day bowel preparations for afternoon procedures are recommended. Studies have shown that split-dose bowel preparations improve the quality of the bowel preparation as compared to bowel preparations administered the day prior to colonoscopy and this has led to a significant increase in the adenoma detection rate².

Polyethylene glycol (PEG) based regimens are the preferred preparation for:

- Age > 65 years
- Diuretic use
- Renal insufficiency (GFR < 60)
- Diabetes
- Congestive heart failure
- Liver cirrhosis or ascites

If a colonoscopy is incomplete due to a poor bowel preparation, then the colonoscopist should specify the bowel preparation for the next colonoscopy and re-book the participant in a Colon Screening Program slot. After a failed preparation, an individualized bowel preparation will be required. On the Colonoscopy Reporting Form, the colonoscopist will tick the box for "Repeat Colonoscopy". Local processes should be used for re-booking the participant. The colonoscopist is responsible for ensuring the participant is re-booked.

3. Alerts for Colonoscopy

3.1 Pre-Colonoscopy Assessment

A pre-colonoscopy questionnaire is a useful tool to identify participants being considered for colonoscopy and polypectomy who may be at increased risk, see Assessment Form (Appendix A). Two methods of contact, separated by a two week interval, is the minimum requirement for contacting participants for colonoscopy assessment. For example, call the participant, wait two weeks, if no response then mail a letter to client requesting they contact the health authority.

Pre-existing medical conditions and medications may conflict with a safe bowel preparation, medications used for sedation, electrocautery equipment or be associated with increased risk of complications.

Each individual is unique and the clinical circumstances with each participant prevent clear guidelines as to appropriate adjustments required in every circumstance of identified increased risk. When in doubt as to the appropriate action, the participant's family physician and/or the attending colonoscopist should be consulted for clinical direction.

If any of the following conditions exist, then the health authority staff should alert the colonoscopist and the participant may require a consultation prior to colonoscopy. The participant may also see the colonoscopist prior to the colonoscopy at the participant's request.

GI Symptoms

- Rectal bleeding
- Chronic diarrhea
- Persistent change in bowel habits
- Chronic abdominal pain
- Unexplained weight loss

Significant co-morbid medical illnesses

- Cancer
- Dialysis participants
- Insulin-dependent diabetics
- Bleeding disorders and participants on antithrombotics
- Cardiac disease requiring a pacemaker or defibrillator
- Respiratory disease requiring home oxygen or CPAP
- Congestive heart failure

- Current angina or history of a myocardial infarction
- Cirrhosis with ascites
- Morbid obesity (BMI ≥ 40)

Other

Participant who will not consent to blood products (e.g. Jehovah's Witness)

3.2 Antithrombotic Therapy³

Antithrombotic agents are medications that prevent blood clot formation and can be divided into anticoagulants and antiplatelet agents. These medications may increase a participant's risk of bleeding following colonoscopic polypectomy. Recommendations state that polypectomy should not be performed while a participant is on anti-thrombotics; biopsies are permitted. Non-steroidal anti-inflammatory drugs (NSAIDs) such as ibuprofen and naproxen are not prescribed to prevent clot formation but as a side effect they do inhibit platelet function and increase the bleeding time. Prospective studies have concluded that aspirin and NSAIDs can be safely continued for colonoscopy and polypectomy.

Whether a medication is discontinued prior to undergoing colonoscopy involves balancing the risk of bleeding following polypectomy and the risk of clotting if the antithrombotic medication is held. Participants on antiplatelet agents (aside from aspirin and NSAIDs), anti-thrombin agents and anticoagulants should be reviewed by a physician prior to the colonoscopy to decide timing of the colonoscopy, discontinuation of the antithrombotic agent, the need for bridging anticoagulation and when the antithrombotic agent can be restarted. This is the responsibility of the colonoscopist; however, the decisions regarding discontinuation of anti-thrombotics, need for bridging therapy and resumption of anti-thrombotics may be at the recommendation of the participant's primary care provider, cardiologist, neurologist and/or thrombosis clinic.

Two scenarios that have arisen in the Colon Screening Program and recommended actions are below.

- 1. If a participant arrives for their scheduled colonoscopy, prepared, but having neglected to hold the antithrombotic as recommended, the colonoscopy should still be undertaken. If a polyp is discovered, then the procedure will be rescheduled for polypectomy with the anti-thrombotic held. If a mass lesion is discovered, then biopsies can be performed. It is the colonoscopist's responsibility to ensure the participant is re-booked for the colonoscopy.
- 2. If a participant cannot safely discontinue an anti-thrombotic agent as the risk of thrombosis is too high, then the colonoscopy should be undertaken while the participant continues the anti-thrombotic medication. This most commonly occurs following coronary stent placement and the requirement for uninterrupted anti-thrombotics is time-limited. If a polyp is discovered, then the procedure will be re-scheduled for polypectomy with the thrombotic held. If a mass lesion is discovered, biopsies can be performed. It is the colonoscopist's responsibility to ensure the participant is re-booked for the colonoscopy.

The following are examples of anticoagulants and antiplatelet agents with the Canadian brand names in brackets. New antithrombotic agents may be available in the near future so this list should not be considered exclusive:

Anticoagulants

- Warfarin (Coumadin)
- Heparin
- Low-molecular weight heparin
 - Enoxaparin (Lovenox)
 - Dalteparin (Fragmin)
- Fondaparinux (Arixtra)
- Dabigatran (Pradax)
- Rivaroxaban (Xarelto)
- Apixaban (Eliquis)
- Desirudin (Iprivask)

Antiplatelet Agents

- Aspirin
- Cilostazol (Pletal)
- Thienopyridine agents
 - Clopidogrel (Plavix)
 - Ticlopidine (Ticlid)
 - Prasugrel (Effient)
 - Ticagrelor (Brilinta)

3.3 Cardiac Defibrillator

Implantable cardiac defibrillators are increasingly common and may be activated inadvertently during endoscopy if electocautery is used. Most participants with cardiac pacemakers may undergo routine uses of electocautery (e.g. polypectomy) with no alterations in management. Some standard precautions are necessary during the procedure to minimize risk.

In all participants with implanted cardiac devices, determine the type of cardiac device, indication for the device and degree of pacemaker dependence before endoscopy. Most participants carry a wallet card, which identifies the device and contact numbers.

In participants with cardiac defibrillators, consultation with cardiologist is recommended and deactivation of the device by qualified personnel should be considered. Continuous cardiac monitoring during the procedure is recommended. The device should be reprogrammed as soon as possible after the procedure.

3.4 Diabetes

Diabetic participants may experience difficulty with glucose control during the bowel preparation and require fasting prior to colonoscopy. Most participants on oral agents (e.g. Metformin (Glucophage), Glipizide (Glucotrol), Glyburide (Diabeta/Micronase), Pioglitazone (Actos), Rosiglitazone (Avandia), Acorbose (Precose) or Miglito (Glyset)) can safely continue the medications until their usual diet is interrupted. During fasting and the bowel preparation time, the drugs should be held. Drugs should be restarted when normal oral intake is resumed after the procedure.

Participants requiring insulin will need to reduce the insulin dosage during fasting for the bowel preparation and day of the colonoscopy procedure. Most participants on insulin have been educated on how to adjust their own insulin during periods of fasting. Participants should be asked to consult with their physician ahead of the procedure.

Participants with diabetes are at increased risk of renal disease and should be questioned as to any pre-existing renal impairment, as this would impact the type of bowel preparation that would be recommended.

3.5 Iron Tablets

Oral iron compounds interact with colonic mucous and dietary compounds and impair the effect of bowel preparations. Participants should be advised to discontinue oral iron preparations 7 days prior to the procedure. Even oral vitamins containing iron are best discontinued to improve colonoscopy quality.

3.6 Glaucoma

Glaucoma (an optic neuropathy due to increased intro-ocular pressure) is present in ~1-8% of individuals over 40 and more common in diabetics. Participants with increased intraocular pressure or glaucoma are often treated with topical eye drop medications. Glaucoma can be aggravated by anti-cholinergic drugs, which are occasionally used during endoscopic procedures to reduce smooth muscle spasm. Glaucoma is usually well controlled with topical medications, which should be continued, and does not interfere with colonoscopy or polypectomy. Anti-spasmodic drugs should be avoided during the procedure.

3.7 Renal Insufficiency/Dialysis

Participants with impairment of renal function can be adversely affected by the dehydrating potential of colonoscopy bowel preparations. Participants with significant kidney disease (e.g. eGFR of less than 60ml/min) should be offered an electrolyte solution containing polyethylene glycol (PEG) for bowel cleansing.

Participants receiving dialysis who require colonoscopy present challenges for safe, effective bowel preparation that does not seriously affect their fluid balance. Colonoscopy is best scheduled in consultation with the participant's nephrologists to discuss bowel preparation and appropriate timing of the procedure in relation to the participant's dialysis times.

Routine antibiotic prophylaxis is not recommended prior to colonoscopy. Antibiotic prophylaxis prior to colonoscopy is recommended for participants undergoing continuous peritoneal dialysis to prevent peritonitis. A single dose of ampicillin plus an aminoglycoside may be given intravenously just prior to the colonoscopy. Intraperitoneal antibiotics the night prior to colonoscopy is an alternative strategy. The abdomen should be emptied of fluid prior to colonoscopy ^{4, 5}.

3.8 Congestive Heart Failure (CHF)

Participants with congestive heart failure may be at increased risk of complications related to colonoscopy bowel preparation and should be offered the PEG based bowel preparations. Participants with severe congestive heart failure, which causes shortness of breath on exertion or significantly limits activity, require a medical consult before colonoscopy should be considered.

4. Informed Consent

Requirements for written informed consent will differ according to the institution. The Colon Screening Program "What is a Colonoscopy?" brochure provides information on the risks of colonoscopy. This must be provided to each participant, in addition to any institution specific consent requirements. It's important that the participant be given time to process the consent information and ask questions. The health authority staff will provide the participant with the information necessary to give informed consent. The colonoscopist will obtain consent prior to the procedure.

Colonoscopy has a 5/1000 risk of a serious complication⁶. This includes the following:

- Reaction to the bowel preparation
- Reaction to the medication used for sedation
- Cardiopulmonary event
- Infection
- Bleeding
- Perforation (<1/1000)

The chance of death from colonoscopy is $1/14,000^7$.

The chance of a significant abnormality being missed is $1/10^8$.

Additional information to answer participant's questions is provided below.

- Cardiopulmonary event refers to desaturation, low blood pressure and rarely angina or myocardial infarction.
- Infection refers to phlebitis related to the IV, pneumonia (aspiration), and diverticulitis. Infection can be transmitted by the colonoscope between participants or from a contaminated water supply. If infection is transmitted between participants, it indicates an error has occurred in the colonoscope cleaning.
- Bleeding is almost always at the site of a polyp removal. It is usually self-limited but will occasionally require hospital admission with a repeat colonoscopy, blood transfusion, radiologic intervention or surgery.
- Perforation is usually at the site of a polyp removal. It almost always requires surgery.

5. Post Colonoscopy Assessment

All participants with colonoscopy information recorded on colonoscopy reporting forms – whether assessed and booked by Health Authority staff or by a colonoscopist only – require post-colonoscopy assessment to monitor for unplanned events and to ensure that the program has information on file to recall participants as needed.

5.1 Telephone Follow-up at 14 Days

Fourteen days after the procedure, the health authority staff will contact the participant. Two methods of contact, separated by a two week interval, is the minimum requirement for contacting participants for post-colonoscopy follow-up. For example, call the participant, wait two weeks, if no response then mail a letter to the participant requesting they contact the health authority. The purpose of the 14-day telephone interview is to:

- assess for any unplanned events following colonoscopy and
- recommend the next re-screening or surveillance interval

5.2 Unplanned Events

Any unplanned event occurring the day before or following colonoscopy should be recorded using the Unplanned Event Form (Appendix D). A serious adverse event is an adverse event that results in a hospitalization, blood transfusion, interventional radiology procedure, other intervention, surgery, or death.

5.3 Re-screening and Surveillance Guidelines

Re-screening and surveillance intervals are based on the findings at colonoscopy, see the Colonoscopy Standards document for current program standards. The health authority staff should review the participant's pathology report and the recommendations in the colonoscopist's Procedure Report. If recommendations differ from the re-screening or surveillance guidelines outlined in the Colonoscopy Standards document, then the next recommended screening type and interval should be discussed with the colonoscopist. There is a Colonoscopy Follow-up Reference Guide that can be used to help determine the appropriate follow-up interval for participants based on their history and pathology findings (Appendix I).

Complete the Follow-Up Form (Appendix E) based on the guidelines and colonoscopist's recommendations and fax the form to the Colon Screening Program. The Program will generate a letter outlining re-screening/surveillance recommendations to be sent to the family physician, colonoscopist and health authority staff who completed the assessment.

Deviations in the recommendations are appropriate under certain circumstances. Examples are in the Colonoscopy Standards.

Where multiple colonoscopies are needed to complete a screening interval, the final follow-up recommendations should consider the outcomes of all procedures and document the next recommended screening as needed. This can be managed through a deviation if the standard intervals do not apply (e.g. participant needs to return in 30 months from the second procedure when a second was completed to assess a piecemeal resection of a high risk lesion six months after the first procedure).

The only reasons for a participant to leave the Colon Screening Program are for age > 74 years, a diagnosis of colorectal cancer and a diagnosis of ulcerative or Crohn's colitis. A diagnosis of ulcerative or Crohn's colitis cannot be determined from a pathology report alone and needs to be determined in discussion with the colonoscopist regarding other clinical findings. Individuals with Lynch Syndrome or Attenuated Familial Adenomatous Polyposis require screening for other malignancies and should also be managed outside the Colon Screening Program by a colonoscopist with expertise in hereditary colon cancer syndromes. All other participants should continue to be screened in the Colon Screening Program and if their screening needs to be individualized, then this can be done by citing a deviation and explanation on the Follow-up Form.

While there may be an indication to do a colonoscopy at an earlier interval, there is never an indication to do a FIT at an earlier interval. If a colonoscopy is not high quality the participant should have a repeat colonoscopy as soon as possible and certainly within 1 year.

If the colonoscopist disagrees with the Colon Screening Program's recommendations and decides upon a different FIT follow-up interval for a participant who has undergone a high quality colonoscopy, then this will need to be arranged by the colonoscopist outside of the Colon Screening Program. Unfortunately, the primary care provider will receive two different recommendations - those in the colonoscopy report and those from the program.

Regarding participants with an abnormal FIT and a colonoscopy without a cancer or polyp, the participant will be recalled to undergo repeat FIT in 10 years. Follow-up Forms received by the program that indicate a deviation with FIT prior to the 10 year recall will not have the deviation entered and the follow-up letter to the colonoscopist, health authority staff and primary care provider will indicate rescreening or surveillance based on current guidelines.

Colonoscopies performed within the Colon Screening Program may reveal significant findings beyond the scope of the program. For instance, participants diagnosed with anal intraepithelial neoplasia or squamous cell carcinoma of the anus, carcinoid tumors, gastrointestinal stromal tumors, or Peutz-Jehger polyps. In this situation, the colonoscopist should either arrange follow-up or guide the primary care provider in the appropriate management. These participants will remain in the Colon Screening Program and be re-called at the appropriate interval for re-screening or surveillance as outlined in the Colonoscopy Standards.

6. Quality Assurance

6.1 Data Collection

Each colonoscopy unit will need a quality program in place. The Colon Screening Program has a central database where the performance indicators will be maintained and reported back to Health Authorities. By providing complete and accurate information on the relevant forms, health authority staff will help with appropriate data collection for performance indicator and participant outcome monitoring.

6.2 Pre-Post Colonoscopy Assessment Performance Indicators

- Number of participants not proceeding to colonoscopy due to poor medical fitness
- Compliance with follow-up colonoscopy
- Time from positive FIT to colonoscopy
- Time from referral to colonoscopy for surveillance procedures
- Number of participants deemed medically unfit by colonoscopist at time of colonoscopy (i.e. prepped for procedure but medically unfit)
- Bowel preparation quality
- Participant, primary care provider, colonoscopist satisfaction with pre-post colonoscopy assessment

7. Medical Record Retention Policy

The Health Authority is the primary record holder for documentation pertaining to pre and post colonoscopy assessment. Health Authorities follow their own policies with respect to record retention and documentation. The Colon Screening Program is a secondary user of the forms and records that are completed for program participants. Participants and providers requesting copies of screening records will be directed to obtain copies from the facility where the interaction occurred.

8. References

- 1. Canadian Cancer Society (CCS). Canadian Cancer Statistics 2015, http://www.cancer.ca/en/cancer-information/cancer-101/canadian-cancer-statistics-publication/?region=bc
- 2. Johnson, David A., et al. "Optimizing adequacy of bowel cleansing for colonoscopy: recommendations from the US multi-society task force on colorectal cancer." Gastroenterology 147.4 (2014): 543-562.
- 3. Baron TH et al. Management of antithrombotic therapy in patients undergoing invasive procedures. N Engl J Med 2013; 368; 2113-24.
- 4. Piraino, B et al. ISPD position statement on reducing the risk of peritoneal dialysis related infections. Peritoneal Dialysis International 2011;Vol. 31:614-630.
- 5. Antibiotic prophylaxis for GI Endoscopy. ASGE Standards of Practice Committee. Gastrointestinal Endoscopy 2015; In press.
- 6. Levin et al. Complications of Colonoscopy in an Integrated Health Care Delivery System. Annals of Internal Medicine 2006:145(12); 880-887.
- 7. Rabeneck L, et al. Bleeding and perforation after outpatient colonoscopy. Gastroenterology 2008. Rex et al. American Journal of Gastroenterology 2002:97(6); 1296-1308.
- 8. Heresbach D, et al., Miss rate for colorectal neoplastic polyps: a prospective, multicenter study of back-to-back video colonoscopies. Endoscopy 2008;40(4): 284-290.

Appendix A - Assessment Form

| BC CAN CER COLON CER SCREENING Provincial Health Services Authority | sse | ssm | ent | Form | | | Affix Label Here | | |
|--|------------------------------------|---------------|--------|--|----------|-------|---|------|-------------|
| 1st CONTACTED DATE (YYYYMMDD) | COMPLE | TED DATE (YYY | YMMDD) | PATIENT NAME LAST | | Pr | KTIENT NAME FIRST | _ | |
| HEALTH AUTHORITY SERVICE CENTRE | AMENDE | D DATE (YYYY | MMDD) | PHN | | D | ATE OF BIRTH (YYYYMMDD) | SEX | (F/M/X) |
| | | | | PRIMARY CARE PROVIDE | (MSC) | н | UMARY PROVIDER LAST, FIRST | _ | |
| Alerts for Colonoscopy Anticoagulation Antiplatelet ag Defibrillator/P Diabetic insulication Sleep Apnea Comments: Reason for Colonoscopy | on gent acemake n/tablets | | | Iron tablets (stop 7 Glaucoma COPD CHF Contact Precautio | n (speci | | Significant co-morbid ii Allergies/sensitivities No blood transfusions Renal insufficiency/dia | | - - - |
| Medication | Dose | Freq. | Medica | ation | Dose | Freq. | Medication | Dose | Freq. |
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| Allergies: NKA | | | | | | | | | |
| Symptoms (within las | + C m and | thal Na | Vac | Comments | | | | | |
| BM Frequency (specify) | t o mon | uis) ive | 163 | Comments | | | | | |
| Recent changes in boy | vel hahit | ·c | | | | | | | \dashv |
| Diarrhea | vernabit | - | + | | | | | | \dashv |
| Constipation | | | | | | | | | \neg |
| Rectal bleeding | | | | | | | | | \neg |
| Bowel urgency | | | | | | | | | \neg |
| Unexplained weight lo | oss | | | | | | | | \neg |
| Abdominal pain | | | | | | | | | |
| Upper GI Symptoms (e swallowing difficulties, GER | | | | | | | | | |
| Comments: | | | | | | | | | |
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| COLON SCREENING PROC | | ver, BC \ | | 1 of 4 ■ 1-877-70-COLON | | | VERSION: 20APRIL2021 | | |



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| PATIENT NAME LAST | PATIENT | NAME F | IRST PHN | DATE OF BIRTH (YYYYMMDD) |
|--|----------|--------|------------|---|
| Medical History | No | Yes | Comments | |
| Gastrointestinal (eg.Ulcers, | | | | |
| arrets, Hiatus hernia, Diverticular disease) | | | | |
| x colonoscopy or flexible | | | | |
| igmoidoscopy | +- | | | |
| urgery | | | | |
| g.Abdominal and other) | +- | | | |
| ardiac eg.A. Fib, Pacemaker, ICD, CHF) | | | | |
| ед.н. гів, Расетакет, ІСИ, СПГ) | +- | | | |
| lypertension | | | | |
| espiratory | | | | |
| g.Sleep apnea, asthma, COPD) | | | | |
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| iver | | | | |
| enal (eg.document eGFR <60ml/min, | | | | |
| reatinine >100umol/L, if known) | + | | | |
| iabetes (eg.Type 1/2, Insulin, oral | | | | |
| /poglycaemic) | +- | | | |
| laucoma | | | | |
| eurological (e.g. Epilepsy, Stroke, MS, | + | | | |
| orkinson's, Alzheimer's, dementia, etc.) | | | | |
| arkinson's, Alzheimer's, dementid, etc.) | 1 | | | |
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| leeding disorder | | | | |
| lood transfusion concerns | | | | |
| g.Jehovah's witness) | + | | | |
| roblems with sedation or | | | | |
| naesthesia | | | | |
| omments / Other Medical Concer | ns: | | | |
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| Patient lives: Alone | Mish (o. | | | |
| Do you consider yourself to h | | | | |
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| Blind/partially blind | Doof/L | | Other (it) | ssive disability (eg MS) Learning disab |
| | | | | te (approximate): |
| EtOH: No Yes | | | | te (approximate). |
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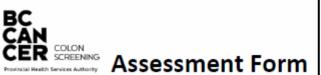




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Assessment Form NOT REQUIRED TO FAX TO BC CANCER

| T REGOINED TO 17 | IN TO BE CHINEEN | | | |
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| 1st CONTACTED DATE (YYYMMDD) | COMPLETED DATE (YYYYMMDD) | PATIENT NAME LAST | PATIENT NAME FIRST | _ |
| | | | | |
| HEALTH AUTHORITY SERVICE CENTRE | AMENDED DATE (YYYMMDD) | PHN | DATE OF BIRTH (YYYYMMDD) | SEX (F/M/) |
| | | PRIMARY CARE PROVIDER (MSC) | PRIMARY PROVIDER LAST, FIRST | _ |
| Assessment In Pers | on By Phone Pat | tient Not Contacted | | |
| FOR ALL PATIENTS: Family H | listory | | | |
| FDR diagnosed CRC: No | Yes More than 3 F | DR Any relatives with | HNPCC connected Cancers? | Yes |
| Relative: | Age at Diagnosis | Specify: | | |
| Relative: | Age at | | | |
| Relative: | Diagnosis Age at | | | |
| | Diagnosis | · | | |
| _ | | | | |
| Patient proceeding to co | lonoscopy as part of the Colo | on Screening Program | | |
| | | | | |
| 1st available date (YY | YMMDD) Booked date (YY | (YYMMDD) Procedure Locat | tion | |
| Patient teaching | Patient instru | ictions (if applicable) | | |
| Appointment details pro | | discontinue iron 7 days prior | | |
| Procedure explained | | • | or specialist regarding fasting & medic | |
| ☐ Bowel prep explained | _ | botics - patient aware to discuss r - ensure hospital protocols are: | with GP/specialist when to stop medic | ations |
| | | er - ensure nospital protocols are | met for these patients | |
| ☐ Sedation options discuss ☐ Risks/complications disc | | | Teachingdate/time: | |
| _ | | | Teaching Coordinator: | |
| La Transportation nome dis | cussed, ride to be provided b | у | Teaching Coordinator. | |
| ☐ Patient NOT proceeding | to colonoscopy as part of the | e Colon Screening Program (plea | ase specify): | |
| Communication provide | d to GP/NP | | | |
| Crohn's or ulcerative | colitis | ■ Not due for colonoscop | y screening/surveillance/follow-up: | |
| Colorectal cancer his | tory | | FIT Colonoscopy | |
| Symptomatic, GP/NP | to refer to specialist | (specify future date) (YYY | YMM) | |
| Outside the target ag | e | Patient declined | | |
| ☐ Medically unfit | | Unable to contact patie | ent | |
| Family history does n | ot meet colonoscopy eligibili | ty Other (specify): | | |
| Patient is not proceeding | g at this time but a future rec | call is required - future date (YYY | YMM): FIT | Colonoscopy |
| | | | | |
| Colonosconist consult re | united. | - | HCP Referral: | , |
| Colonoscopist consult re | quired: | | HCP Referral: | , |
| | quired: | | HCP Referral: | |
| | quired: | [| HCP Referral: | |
| Comments: | | | | |
| | Patien | t Coordinator Signature | HCP Referral: Location VERSION:20APRIL2021 | 211 |



|--|

| PATIEN | T NAME LAST | PATIENT NAME FIRST | PHN | DATE OF BIRTH (YYYYMMDD) |
|------------------------|--|--|---|-------------------------------|
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| COLON 50 801- 686 \ | REENING PROGRAM West Broadway Vanco | Page 4 of 4 uver, BC V5Z 1G1 🖀 1-877- | FORM: 21100 70-COLON www.screening | VERSION: 20APRIL2021 bc.ca |

Appendix B - Colonoscopy Brochure



Are there any risks with colonoscopy?

As with any medical procedure, colonoscopy has a small risk of complications.

Approximately 5/1,000 people will have a serious complication. Complications can include a reaction to the bowel preparation or medication used for sedation, heart or lung problems, an infection, bleeding from the colon and/or perforation of the colon (hole in the colon).

If a complication occurs, treatment including If a complication occurs, treatment including antibiotics, blood transfusion, hospitalization, repeat colonoscopy or surgery may be required. The risk of dying from colonoscopy is less than 1/14,000. There is also a risk of missing a significant abnormality. This occurs in less than 1/10 cases.

Certain cancers may never cause any symptoms or affect life expectancy or quality of life. However, research shows that most colon cancers are harmful and that colon cancer should be detected and treated as early as possible.





Colonoscopy

Who should get a colonoscopy?

A personal reconcerous polyp; or,

One first degree relative (parent, sibling or child) with colon cancer diagnosed under the age of 60; or.*

Contact Us

BC Cancer Colon Screening 801-686 West Broadway Vancouver, BC V5Z 1G1

Phone: 1-877-702-6566 Email: screening@bccancer.bc.ca Web: www.screeningbc.ca/colon

Version: June 2021



Before the colonoscopy

- Expect to be at the hospital for two to three
- You will be asked to change into a gown.
- A nurse will complete your admission history and measure your vital signs.
- · You will be asked to provide a list of your
- A nurse will start an intravenous (IV) to administer sedation and pain medication.

What happens during a colonoscopy?

- · A colonoscopist inserts the colonoscope into the rectum and advances it along the length of the
- · Air is sent through the colonoscope to expand the colon for better viewing. It is normal throughout the procedure to feel slight pressure or experience
- · Images of the lining of the rectum and colon are sent to a video monitor where the colonoscopist will look for anything unusual, like a polyp. A polyp is a small growth of tissue on the wall of the
- Polyps can grow very slowly, and some can become cancerous. It may be necessary to take a sample (biopsy) or remove the polyp (polypectomy). This is painless.
- . The biopsy or polyp is then sent to a lab for

What happens after a colonoscopy?

- Have an adult accompany you home. You cannot drive until the following day.
- You may be sleepy after you arrive home from the procedure. It is recommended that you do not operate equipment, sign legal papers or drink alcohol until the following day.
- You will be able to resume your regular diet and medications after your colonoscopy, unless otherwise directed by the health care team in your community.
- The air inside your colon may cause you to feel bloated and/or have cramping after the procedure. It is important to relax and pass the air as soon as possible. If this discomfort increases or is unrelieved, go to the emergency department and advise them that you had a colonoscopy.

You will be closely monitored before, during and after the procedure.

What do I need to know about my colonoscopy results?

You will be given preliminary results before you leave the hospital. Then, approximately two weeks after your procedure, the health care team in your community will inform you of your complete results and answer your questions during the follow up call. Your doctor will also receive your results.

If your colonoscopy is normal, your family history will determine when you will be re-screened. The health care team in your community will advise you of your next screening date.

If your colonoscopy is abnormal, further procedures or more regular surveillance may be necessary. The health care team in your community, or your doctor will explain the process for further appointments and next steps

Appendix C – Sample Not Proceeding to Colonoscopy Letter

| ar Dr | Fax # | | Date |
|---|--|--|---|
| tient Name | | PHN | DOB |
| Your patient was refer | ed for pre-colonoscopy | assessment on | (date) due to: |
| Abnormal FIT | □ Family History | □ Surveillan | ce Requirement |
| Your patient has <u>NOT</u> t | een booked for a colon | oscopy procedure o | lue to: |
| | | | |
| | | | |
| | | irectly to a specialist | for assessment. The patient will |
| Program will recall the patier | nt at that time. If no date is | indicated, the patier | t will not be recalled by the Colon |
| | | | |
| | | | |
| please send the Program a (| Colonoscopy Referral For | m. | |
| patient to advise that they ha Colon Screening Program. If | ave not been booked for a the patient wishes to par | colonoscopy. The pa | atient will not be recalled by the |
| Patient is required to be scop | ped outside of the Colon S | Screening Program. | |
| | | | |
| | | | |
| ncerely, | | | |
| LON SCREENING PROGRAM | Fax: | | APRIL 2021 |
| | Your patient was reference. Abnormal FIT Your patient has NOT I Patient has a history of inflar his/her specialist for ongoing Program. Patient has a history of color monitoring. The patient will repatient will recalled by the Colon Medically unfit for colonosco Program will recall the patier Screening Program. The patient Screening Program. The patient patient patient of colonoscopy. Your patient does not meet a The patient will be recalled by Patient declined proceeding please send the Program a Colonoscopy. We were unable to reach your patient to advise that they had Colonoscopy Referral Form. Patient is required to be scopeding required to be scopeding. | Your patient was referred for pre-colonoscopy Abnormal FIT | Your patient was referred for pre-colonoscopy assessment on |

Appendix D – Unplanned Events Form

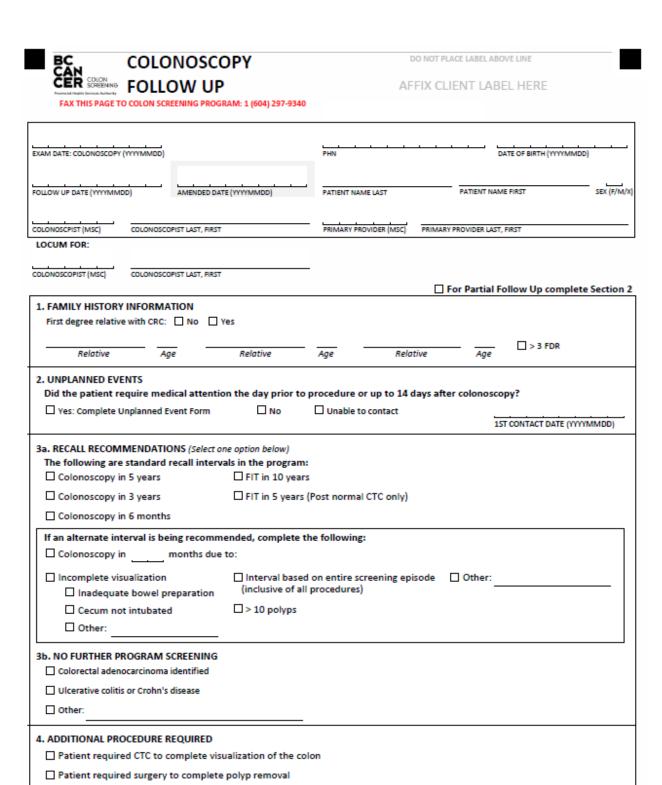
Pre/Post Colonoscopy CER SCHEDNIG TOUGH Unplanned Event FAX THIS PAGE TO COLON SCREENING PROGRAM: 1 (604) 297-9340 Pre/Post Colonoscopy

DO NOT PLACE LABEL ABOVE LINE

AFFIX CLIENT LABEL HERE

| XAM DATE: COLONOSCOPY (YYYYMMDD) | | PHN | DATE OF BIRTH (Y) | YYMMDD) |
|---------------------------------------|----------------------|-----------------------------|-----------------------------|------------|
| OLLOW UP DATE (YYYYMMDD) AMEND | DED DATE (YYYYMMDD) | PATIENT NAME LAST | PATIENT NAME FIRST | SEX (F/M/X |
| OLONOSCOPIST [MSC] COLONOSCOPIST LAST | r, FIRST | PRIMARY PROVIDER (MSC) | RIMARY PROVIDER LAST, FIRST | |
| DATE OF ONSET SYMPTOMS (YYYYMMDD) | Symptoms ongoing? (| O No O Yes | UTION (YYYYMMDD) | |
| The day prior to, or within 14 da | ays after undergoing | g a colonoscopy, this patie | ent had these uplanned ev | ent(s): |
| ☐ Bowel prep complication | | ☐ Perforation | | |
| ☐ Rectal bleeding → Anticoa | gulation: O No O | es Respiratory | | |
| ☐ Infection | | ☐ Cardiac | | |
| Death: (YYY | | Other: | | |
| Cause of death: | YMMDD) | | | |
| Comments: | | | | |
| | | | | |
| Patient first obtained medical a | ttention: | | | |
| Patient required the following i | | | | |
| ☐ Blood transfusion | ☐ Additiona | al Colonoscopy: | (manus et app) | |
| | Other: | | - (YYYYMMDD) - | |
| Antibiotics | | | | |
| ☐ Antibiotics ☐ Surgery: | Hospital : | admission: | (YYYMMDD) to | (YYYYMMDD) |
| □ Surgery: (YY | | | | |
| □ Surgery: (YY | | | (YYYMMDD) to | |
| □ Surgery: (YY | | | | |
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| Surgery:(YYY | | | | |
| Surgery:(YYY | | | | |

Appendix E - Follow-Up Form



PATIENT COORDINATOR

PATIENT COORDINATOR SIGNATURE

20410



Appendix F – Colonoscopy Reporting Form

| В | BC COLONOSCOPY | | | | | | | | DO I | OT PLACE | LABEL ABO | VE LINE | | | |
|---|---|-----------------------|------------------------------|--|---------------|----------------|---------------|----------------------------|--------------------------------------|------------------------|--------------------------------|----------------------------------|----------------------------|-----------|----------|
| PRESS FIRMLY TO ENSURE LEGIBILITY FOR MULTIPLE COPIES FAX TOP COPY TO COLON SCREENING PROGRAM: 1 (604) 297 9340 GREY SECTIONS TO BE COMPLETED AS REQUIRED | | | | | | | | AFFI: | K CLIEN | IT LABE | L HERE | | | | |
| EXAM DATE (YYYYMMDD) START TIME (HRS) | | | | | PHI | N . | | | DA | ATE OF BIRTH | (YYYYMMDD) |) | | | |
| FACILITY N | AME | | _ | AMENI | DED DATE | (YYYYN | IMDD) | PAT | PATIENT NAME LAST PATIENT NAME FIRST | | | | SEX (M/F/X) | | |
| COLONOS | COPIST (MSC) | COLO | NOSCOPI | ST LAST | , FIRST | | | PRII | MARY PROVIDE | R (MSC) | RIMARY PRO | VIDER LAST, | FIRST | | |
| l _ | for Colon | | | | illance | = 🗆 | Deviation | | son Colono No Show fo | • • | | | • | day of pr | ocedure |
| ☐ Exc | L PREPARA ellent r (adequat or (inadeq | ☐ Good te to visua | alize a | | | | n) | | CALINTUB Yes → PI No □ | oto docum Uncertair | entation? | No □ Fle | ☐ Yes xible Sign | noidoscop | py |
| ☐ Per ☐ Blee ☐ Car | foration eding diovascula piratory | ar [| Admi Reve Deat Othe | it to h rsal a h | gents | | | 5. 00 | OMMENTS 1 | TO PATHOL | OGIST: | | | (M | linutes) |
| | Specimen Type | Location | <u><</u> 5 | Size 6-9 | (mm) 10-19 | <u>></u> 20 | Morphology | Primary Removal Mode | Submucosal Injection (Y/N) | Plecemeal (Y/N) | Complete Removal (Y/N/U) | Complete Retrieval (Y/N/U) | Specimen Sent (Y/N#) | Time | Initials |
| Example | Р | T | | ✓ | | | Р | HS | Y | Y | Y | Y | Υ | 14:00 | AB |
| 1/A | | | | | | | | | | | | | ļ | | |
| 2/B | | | | | | | | | - | | | | | | |
| 3/C | | | | | | | | | - | | | | | | |
| 4/D | | | - | | | | <u> </u> | | - | | | | ļ | | |
| 6. ☐ Additional specimens recorded on Page 2 7. ☐ Repeat Colonoscopy Required COMPLETE COLONOSCOPY REPORTING FORM FOR NEXT SCOPE | | | | Specimen Type B = biopales P = polypectomy P = yes N = no U = uncertain Specimen Type A = ascending colon C = cecum D = descending I = lieum L = left colon O = other/irandom U = uncertain D = descending N = nectum S = sigmoid T = transverse colon Morphology F = fat M = mass C = cold snare D = pedunculated B = hot snare B = hot snare B = hot snare B = hot snare B = sossile B = sossile | | | | | y forceps snare lopsy forceps | | | | | | |
| MD NAM | ME: | | s | IGNAT | URE: | | | | RN NAME | <u> </u> | | SIGNA | TURE: | | |
| 1. BC | COPIES OF Cancer Co #: 1 (604) | lon Screen | ing | ORT T | | ry Provi | ider (Name 8 | k MSC#) | _ 3. c | Other (Name | & MSC#) | 4 Oti | her (Name 8 | & MSC#) | |
| Spe | cimen track | ing required | | | | | imples sent t | | _ | _ | TIALS | | | | _ |
| | | , | | | | | imples trans | | _ | _ | TIALS | _ | TE: TE: | | _ |

PATHOLOGY COPY | FAX THIS COPY TO 1 (604) 297 9340

INFORMATION ON THIS FORM IS CONFIDENTIAL. IF YOU RECEIVE THIS IN ERROR PLEASE FAX TO QUALITY DEPT: 1 (604) 675 7223





Appendix G - Bowel Preparation Algorithm

Colon Screening Program

Bowel Preparation Guidelines



Bowel Preparations

High Volume (4L PEG)

Consider for:

- · Constipation
- · Previous poor preparation
- · Narcotic use
- · Poor mobility
- · Morbid obesity

Examples:

- CoLyte
- GoLYTELY
- PegLyte

Low Volume (PEG / 2L PEG)

Examples:

- Bi-PegLyte (do not take Bisacodyl)
- MoviPrep

Low Volume (Hyperosmolar)

Examples:

- Picoflo
- PicoSalax
- · Purg-Odan

Split-dose regimens are preferred.

PEG-based regimens are the preferred preparation for:

- · Age > 65 years
- Diuretic use
- Renal insufficiency (GFR< 60)
- Diabetes
- · Congestive heart disease
- · Liver cirrhosis or ascites

Adjuncts (bisacodyl, magnesium citrate, enemas) are not recommended for standard bowel preparations.

Participants requiring a repeat colonoscopy due to a poor preparation should have their preparation directed by the colonoscopist.

References:

Optimizing adequacy of bowel cleansing for colonoscopy: recommendations from the US Multi-Society Task Force on Colorectal Cancer. Gastrointestinal Endoscopy 2014;80:543-562.

Appendix H – Referral Update Form

| | DC | NOT PLACE LABEL ABOVE LINE | • |
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| CER SOLON CERSON REFERRAL UPDATE FORM | AFF | X CLIENT LABEL HERE | |
| PRESS FIRMLY TO ENSURE LEGIBILITY | | | |
| FAX TOP COPY TO COLON SCREENING PROGRAM: 1 (604) 297-9340 | | | |
| IST CONTACTED DATE (YYYYMMDD) COMPLETED DATE (YYYYMMDD) | PHN | DATE OF BIRTH (YYY | VMMDD) |
| COMPLETED DATE (TTYTMINDD) | | Sile of Sillin (File | ······································ |
| HEALTH AUTHORITY SERVICE CENTRE AMENDED DATE (YYYYMMDD) | PATIENT NAME LAST | PATIENT NAME FIRST | SEX (F/M/ |
| | PRIMARY PROVIDER (MSC) | RIMARY PROVIDER LAST, FIRST | |
| COMPLETE ONLY ONE SECTION BELOW | , | | |
| COMPLETE ONLY ONE SECTION BELOW | | | |
| SECTION A: TRANSFER REQUEST Complete only if referral r | equires a transfer to another s | ervice centre. | |
| Transfer Request To: | | | |
| (Name of Hospital or City) | | | |
| Transfer Request ☐ Medical Reason ☐ Patient Reason: | Preference Par | ient Address Related | |
| Other (Please specify): | | | |
| | | | |
| SECTION B: PATIENT NOT PROCEEDING Complete only if p Please ensure the patient's primary provider has been not | tified if the patient is not go | oing to proceed. | tre. |
| Please ensure the patient's primary provider has been not | tified if the patient is not go | | tre. |
| Please ensure the patient's primary provider has been not | tified if the patient is not go | oing to proceed. | tre. |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up | tified if the patient is not go | rectal cancer history n's or ulcerative colitis | tre. |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: FIT Colonoscopy | tified if the patient is not go Patient has colo | rectal cancer history n's or ulcerative colitis | tre. |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: | Patient has colo Patient has Croh Patient has Croh Patient is deceas | rectal cancer history n's or ulcerative colitis | |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: | Patient has colo Patient has Croh Patient has Croh Patient is decea: Patient moved o | rectal cancer history n's or ulcerative colitis sed ut of province | |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: | Patient has colo Patient has Croh Patient has Croh Patient is decea: Patient moved o Patient family hi | oing to proceed. rectal cancer history n's or ulcerative colitis sed ut of province story does not meet colonosco | py eligibility |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: | Patient has colo Patient has Croh Patient has Croh Patient is decea: Patient moved o Patient family hi | rectal cancer history n's or ulcerative colitis sed ut of province story does not meet colonosco | py eligibility |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: | Patient has colo Patient has Croh Patient is deceas Patient moved of Patient family hi Patient is medic Patient is Sympt Other: | rectal cancer history n's or ulcerative colitis sed ut of province story does not meet colonosco ally unfit for follow up pmatic, provider to refer to spe | py eligibility |
| Please ensure the patient's primary provider has been not Patient not due for screening/surveillance/follow up Recall for: | Patient has colo Patient has Croh Patient is deceas Patient moved of Patient family hi Patient is medic | rectal cancer history n's or ulcerative colitis sed ut of province story does not meet colonosco | py eligibility |

INFORMATION ON THIS FORM IS CONFIDENTIAL IF YOU RECEIVE THIS IN ERROR PLEASE FAX TO QUALITY DEPT: 1 (604) 675-7223



Appendix I – Follow-Up Reference Guide



Colonoscopy Follow-up Reference Guide

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| Standard Standard Standard Standard Standard Standard Standard Standard Standard Standard Average risk patient with at teast one high risk lession completely the excision site more low risk lession removed. The excision site more low risk months to ensure complete removal. Sor more low risk months to ensure complete removal. Sor more low risk months to ensure complete removal. Adenomas with: Adenomas with: Adenomas with: High Risk Lesions High-grade dysplasia; High-grade dysplasia; | Recall Recommendations | Alternate Interval | If patient's circumstance does not match a Standard Interval/Reason, use the Alternate Interval section to indicate when patient should return for colonoscopy (3 months mininum*) and provide reason. Incomplete visualization Inadequate bowel preparation Cecum not intubated Other visual issue Interval based on entire screening epidsode (inclusive of all procedures) | Colonoscopy in months (Select one of the above reasons on the follow-up form) | Family Histories that require ongoing colonoscopy screening (NOT considered average risk) 1 first degree relative (parent, sibling, child) diagnosed with colon cancer under age 60 2 or more first degree relatives diagnosed at any age | | | |
|--|------------------------|--------------------|--|--|---|---|--------------------------|--|
| t one high sisten with: stelely d at time onoscopy; oscopy in YEARS High | | Standard | Average risk patient who required CT colonography to complete visualization of the colon and had a negative CT colonography. | | FIT in 5 YEARS (Post normal CT colonography only) | Family Histories that reongoing colonoscopy (NOT considered average risk) tirst degree relative (parent, siblin with colon cancer under age 60 a 2 or more first degree relatives dia | | |
| t one high sisten with: stelely d at time onoscopy; oscopy in YEARS High | | | Average risk patient who did not have any precancerous lesions removed. | | FIT in 10 YEARS | ysplasia s he cecum, on that are ≥ 10 mm | | |
| t one high sisten with: stelely d at time onoscopy; oscopy in YEARS High | | | | | Colonoscopy in 6 MONTHS | igh Risk Lesions Adenomas with: □ Villous features; □ High-grade dysplasia; □ 26-ssile serrated lesions with dysplasia Sessile serrated adenomas Hyperplastic polyps found in the cecum, ascending and transverse colon that are ≥ 10 mm | ending and transverse co | |
| tient with: I or 2 low risk esions removed; and who did not have brecancerous esions removed; a personal history of adenomas and who did not have any precancerous esions removed. K Lesions enomas with low-gra hat are smaller than 1 ated lesions with no | | | | | Colonoscopy in 3 YEARS | I | | |
| Ris aradó | | | Patient with: 1 or 2 low risk lesions removed; a family history and who did not have precancerous lesions removed; a personal history of adenomas and who did not have any precancerous lesions removed; | | | Low Risk Lesions Tubular adenomas with low-grade dysplasia that are smaller than 10 m Sessile serrated lesions with no dys that are smaller than 10 mm | | |
| Recall Reason Interval | | | Keason | Recall Interval | LOW Tubul dyspla Sessile that a | | | |

^{*} Alternate recall intervals of <3 months should be booked internally using local workflow processes rather than relying on another referral to come from the program.
** This refers to patients who have more than 10 pre-cancerous polyps (adenomas, sessile serrated lesions, traditional serrated adenomas) removed requiring a more frequent colonoscopy follow-up. Once all polyps have been removed from the colon, the patient should return for surveillance colonoscopy in one year.

| | Pre-Post Colonoscopy Assessment Standards Change Log Revision History | | | | | | |
|---------|---|----------|----------------------------|---|--|--|--|
| Version | Date | Action | Pages Affected | Details | | | |
| 1.0 | May 2014 | | | | | | |
| | May 2015 | | | | | | |
| | March 2016 | | | | | | |
| 1.1 | November 2017 | Updated | ALL | Format updated based on the Colonoscopy Standards Title of document from "Patient Coordinator Standards" to "Pre-Post Colonoscopy Assessment Standards" Page numbers added to the Table of Contents. Titles and section numbers updated Dr. Telford Updated Standards. (p. 4-7, 13, 15, 16) References and Appendices matched and added based on the updated standards. | | | |
| 1.2 | January 2018 | Addition | 6,17 | Added statement on confirming participants PCP Added Medical Records Retention policy | | | |
| 1.3 | March 2018 | Updated | All | New Logo/Branding | | | |
| 1.4 | April 2018 | Updated | 19-24 | Updated Appendices | | | |
| 1.5 | July 2019 | Addition | 14 | Added requirement for two methods of contact for follow up phone call to participant with time interval, and example | | | |
| | August 2019 | Updated | Section 3.1 Section 5.1 | Remove above addition and incorporate minimum contact for assessment standard. | | | |
| | September 2019 | Updated | Section 5.1 | Added requirement for two methods of contact for follow up phone call to participant with time interval, and example | | | |
| | April 2020 | Updated | Sections 1.5, 2.2, 3.2, | Clarify eligibility and when colonoscopy should proceed. Assessment when on Antithrombotic Therapy updated. | | | |
| | October 2021 | Updated | Appendices H, I | Clarify eligibility and update process for Referral Update Form and new Follow-up Form. Added two appendices. | | | |