MD Anderson Survivorship – Pancreatic Cancer

Page 1 of 6

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

TABLE OF CONTENTS

Resected Pancreatic Adenocarcinoma	Pages 2-3
Sporadic Neuroendocrine Tumor	Pages 2-3
Resected Duodenal /Peri-ampullary Cancer	Pages 2-3
APPENDIX A: Pancreatoduodenectomy or Total Pancreatectomy Labs	Page 4
APPENDIX B: Bone Mineral Density (BMD) Monitoring	Page 4
Suggested Readings	Page 5
Development Credits	Page 6

Cancer Center

MD Anderson Survivorship — Pancreatic Cancer

Page 2 of 6

Making Cancer History®

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

ELIGIBILITY

CONCURRENT **COMPONENTS** OF VISIT

SURVEILLANCE →

MONITORING

FOR LATE

EFFECTS

RISK REDUCTION/

EARLY

DETECTION

PSYCHOSOCIAL

FUNCTIONING

Resected pancreatic

adenocarcinoma

or

Sporadic neuroendocrine tumor

<u>or</u>

Resected duodenal/periampullary cancer

and

> 3 years posttreatment and no evidence of disease (NED)

Years 3 to 5:

- History and physical every 6-12 months
- CT chest, abdomen, and pelvis with contrast every 6-12 months
- Nutrition evaluation with Registered Dietitian:
- As clinically indicated if status post distal or central pancreatectomy
- o Every 6-12 months if status post pancreatoduodenectomy (PD) or total pancreatectomy (TP)

Years 6 to 10:

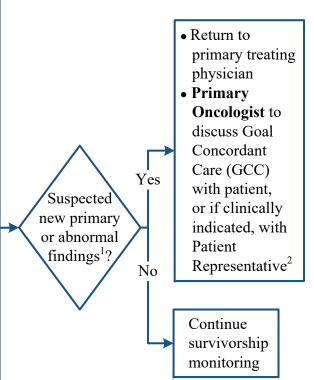
- Annual history and physical
- Annual CT chest, abdomen, and pelvis with contrast or MRI abdomen with and without contrast - MRCP based on age of patient and genetic history
- Nutrition evaluation with Registered Dietitian:
- As clinically indicated if status post distal or central pancreatectomy
- o Annually if status post PD or TP

Years 11 and beyond:

- Annual history and physical
- Annual MRI abdomen with and without contrast - MRCP
- Nutrition evaluation with Registered Dietitian annually or as clinically indicated. If status post PD or TP, additional labs with annual visit (see Appendix A)

- Labs every 6-12 months: CBC with differential, CMP, HbA1c, fasting lipid panel, CA 19-9 and/or CEA
- o If status post PD or TP, will need additional labs annually (see Appendix A)
- Bone mineral density (BMD) baseline at 2 years post-op, then every 2-5 years from baseline or as indicated (see Appendix B)
- Labs annually: CBC with differential, CMP, HbA1c, fasting lipid panel, CA 19-9 and/or CEA o If status post PD or TP, will need additional labs annually (see Appendix A)
- BMD as indicated (see Appendix B)
- For patients with familial PDAC or germline mutations, consider referral to PDAC hereditary High Risk Clinic using the Ambulatory referral to GI High Risk and Genetics order
- Labs annually: CBC with differential, CMP, HbA1c, fasting lipid panel, CA 19-9 and/or CEA
- BMD as indicated (see Appendix B)

DISPOSITION



CMP = complete metabolic panel CA 19-9 = cancer associated antigen 19-9

PDAC = pancreatic ductal adenocarcinoma

CEA = carcinoembryonic antigen

MRCP = magnetic resonancecholangiopancreatography

→ See Page 3

¹CA 19-9 elevated above normal, clinical status decline, patient choice with MD review, abnormal physical exam findings, imaging findings suggestive of: new lesions/lymphadenopathy, stricture, thrombus, fluid collections

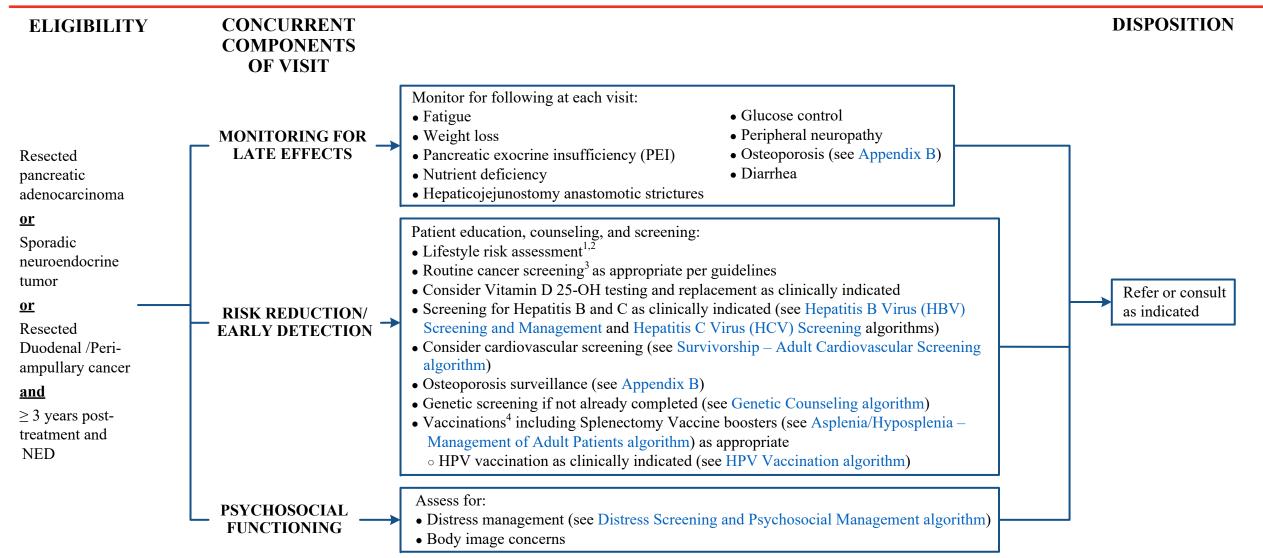
²GCC should be initiated by the **Primary Oncologist**. If Primary Oncologist is unavailable, Primary Team/Attending Physician to initiate GCC discussion and notify Primary Oncologist. Patients, or if clinically indicated, the Patient Representative should be informed of therapeutic and/or palliative options. GCC discussion should be consistent, timely, and re-evaluated as clinically indicated. The Advance Care Planning (ACP) note should be used to document GCC discussion. Refer to GCC home page (for internal use only). Department of Clinical Effectiveness V2

Making Cancer History®

MD Anderson Survivorship — Pancreatic Cancer

Page 3 of 6

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.



¹ See Physical Activity, Nutrition, Obesity Screening and Management, and Tobacco Cessation Treatment algorithms; patient should be encouraged to limit alcohol consumption. Ongoing reassessment of lifestyle risks should be a part of routine clinical practice

² Recommend at least 30 minutes of moderate-intensity activity most days of the week

³ Includes breast, cervical, colorectal, liver, lung, pancreatic, and skin cancer screening

⁴Based on American Society of Clinical Oncology (ASCO) guidelines



Making Cancer History®

MD Anderson Survivorship – Pancreatic Cancer

Page 4 of 6

Cancer Center Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

APPENDIX A: Pancreatoduodenectomy or Total Pancreatectomy Labs

- CBC with differential
- PT with INR
- Copper
- Zinc
- Selenium
- Ferritin
- Iron
- •Transferrin
- Folate
- Vitamin B6
- Vitamin B12
- Methylmalonic acid
- Vitamin A
- CRP
- Vitamin E
- Vitamin D 25-OH
- Albumin
- HbA1C

PT with INR = Prothrombin Time with INR CRP = C-Reactive Protein

APPENDIX B: Bone Mineral Density (BMD) Monitoring

Patient Population	Frequency of Monitoring
Normal bone density	Recheck BMD every 5 years if male or premenopausal; recheck BMD every 2 years if postmenopausal
Osteopenia, ≥ 50 years old	Consider medical therapy or referral to bone health specialist based on FRAX® Calculation¹: if risk of hip fracture is < 3% risk and risk of non-hip fracture is < 20%, recheck BMD in 2 years. If risk of hip fracture is \geq 3% or risk of non-hip fracture is \geq 20%, bone health specialist.
Osteopenia, < 50 years old	Refer to bone health specialist
Osteoporosis	Refer to bone health specialist

PDAC = Pancreatic Ductal Adenocarcinoma

¹ FRAX® - Fracture Risk Assessment Tool at https://frax.shef.ac.uk/FRAX/tool.aspx?country=9

MD Anderson Survivorship — Pancreatic Cancer

Page 5 of 6

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

SUGGESTED READINGS

- Arango, N.P., & Maxwell, J.E. (2023). Pancreatic neuroendocrine tumors and multiple endocrine neoplasia. In BW Feig (Ed). *The MD Anderson surgical oncology handbook* (7th Edition). Philadelphia: Lippincott Williams & Wilkins. Wolters Kluwer.
- Centre for Metabolic Bone Diseases, University of Sheffield. (n.d.). FRAX ® fracture risk assessment tool. Calculation tool. Retrieved from https://www.sheffield.ac.uk/FRAX/tool.aspx
- Gaskill, C.E., Kim, M. P., & Katz. M. K. (2023) Pancreatic ductal adenocarcinoma. In BW Feig (Ed). *The MD Anderson surgical oncology handbook* (7th Edition). Philadelphia: Lippincott Williams & Wilkins. Wolters Kluwer.
- Kamboj, M., Bohlke, K., Baptiste, D. M., Dunleavy, K., Fueger, A., Jones, L., ... Kohn, E. C. (2024). Vaccination of adults with cancer: ASCO guideline. *Journal of Clinical Oncology*, 42(14), 1699-1721. doi:10.1200/JCO.24.00032
- Mason, M.C., & Uppal, A. (2023). Small bowel malignancies. In BW Feig (Ed). *The MD Anderson surgical oncology handbook* (7th Edition). Philadelphia: Lippincott Williams & Wilkins. Wolters Kluwer
- MD Anderson Institutional Policy #CLN1202 Advance Care Planning Policy Advance Care Planning (ACP) Conversation Workflow (ATT1925)
- National Comprehensive Cancer Network. (2024). *Pancreatic Adenocarcinoma* (NCCN Guideline Version 3.2024). Retrieved from https://www.nccn.org/professionals/physician_gls/pdf/pancreatic.pdf
- Petzel, M. Q., & Hoffman, L. (2017). Nutrition implications for long-term survivors of pancreatic cancer surgery. Nutrition in Clinical Practice, 32(5), 588-598. doi:10.1177/0884533617722929



MD Anderson Survivorship – Pancreatic Cancer

Page 6 of 6

Disclaimer: This algorithm has been developed for MD Anderson using a multidisciplinary approach considering circumstances particular to MD Anderson's specific patient population, services and structure, and clinical information. This is not intended to replace the independent medical or professional judgment of physicians or other health care providers in the context of individual clinical circumstances to determine a patient's care. This algorithm should not be used to treat pregnant women.

DEVELOPMENT CREDITS

This survivorship algorithm is based on majority expert opinion of the Pancreatic Survivorship workgroup at the University of Texas MD Anderson Cancer Center. It was developed using a multidisciplinary approach that included input from the following:

Core Development Team Leads

Rae Reynolds, PhD, APRN (Surgical Oncology) Brandon Smaglo, MD (GI Medical Oncology) Rebecca Snyder, MD, MPH (Surgical Oncology) Whittney Thoman, MS (Cancer Survivorship)

Workgroup Members:

Prajnan Das, MD (GI Radiation Oncology)

Olga N. Fleckenstein, BS

Justin Folloder, PA-C (Surgical Oncology)

Katherine Gilmore, MPH (Cancer Survivorship)

Daniel Haldar, MD (GI Medical Oncology)

Emma Holliday, MD (GI Radiation Oncology)

Ryan Huey, MD (GI Medical Oncology)

Naruhiko Ikoma, MD (Surgical Oncology)

Matthew Katz, MD (Surgical Oncology)

Michael Kim, MD (Surgical Oncology)

Eugene Koay, MD, PhD (GI Radiation Oncology)

Albert Koong, MD, PhD (GI Radiation Oncology)

Ethan Ludmir, MD (GI Radiation Oncology)

Jessica Maxwell, MD (Surgical Oncology)

Florencia McAllister, MD (Clinical Cancer Prevention)

Maria Pia Morelli, MD, PhD (GI Medical Oncology)

Sonal Noticewala, MD (GI Radiation Oncology)

Shubham Pant, MD (GI Medical Oncology)

Maria Petzel, RD, CSO, FAND (Clinical Nutrition)

Johnny L. Rollins, MSN, APRN, ANP-C (Cancer Survivorship)

Grace Smith, MD (GI Radiation Oncology)

Hannah Warr, MSN, RN, CPHON

Steven Wei, EdD, PA-C (Surgical Oncology)

Jason Willis, MD, PhD (GI Medical Oncology)

Bob Wolff, MD (GI Medical Oncology)

Dan Zhao, MD (GI Medical Oncology)

^{*}Clinical Effectiveness Development Team