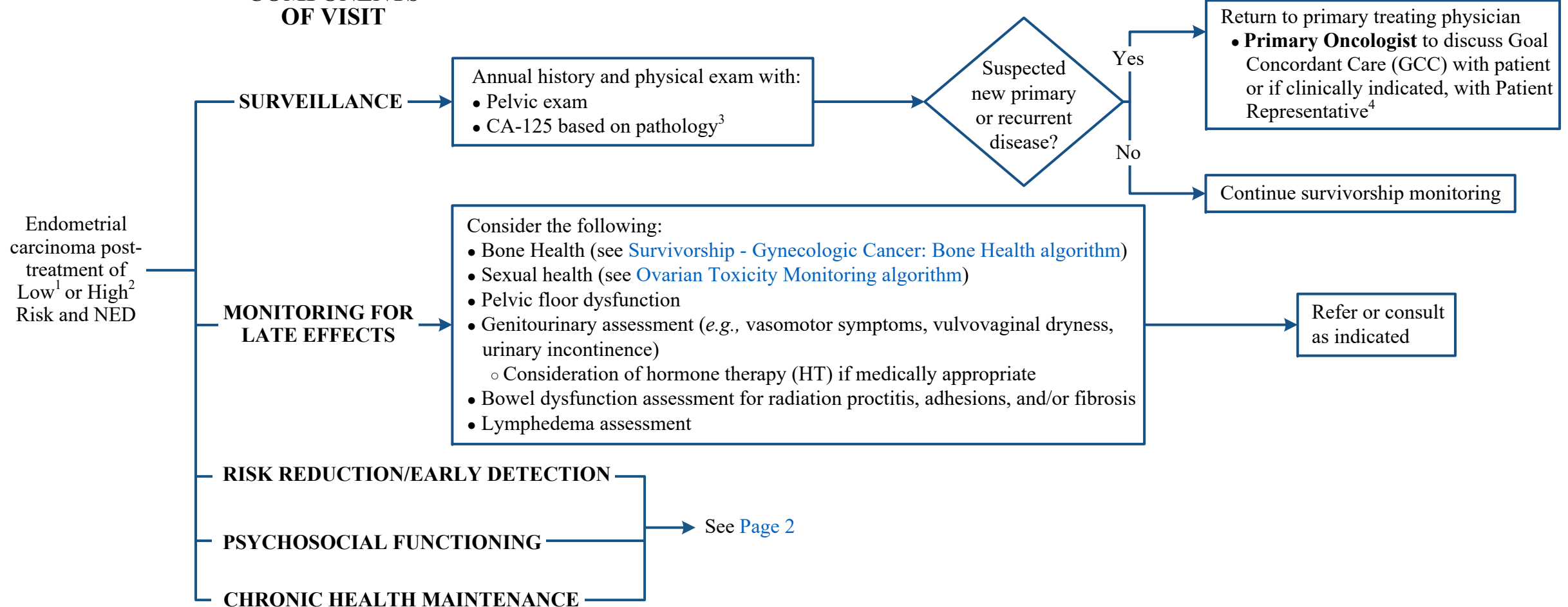


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ELIGIBILITY

CONCURRENT COMPONENTS OF VISIT

DISPOSITION



NED = no evidence of disease

¹ Low risk endometrial cancer is defined as any patient who did not receive chemotherapy or radiotherapy as adjuvant treatment after their initial surgery. Survivorship begins 3 years post-treatment and NED.

² High risk defined as patients who received chemotherapy or radiotherapy as adjuvant treatment after their surgery. Survivorship begins 5 years post-treatment and NED.

³ • Uterine carcinosarcoma – CA-125 annually • High grade, uterine serous types – CA-125 annually

⁴ GCC 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).

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ELIGIBILITY

CONCURRENT
COMPONENTS
OF VISIT

DISPOSITION

Endometrial carcinoma post-treatment of Low¹ or High² Risk and NED
(continued from previous page)

RISK REDUCTION/
EARLY DETECTION

- Patient education, counseling, and screening:
- Patient education regarding symptoms including radiation therapy complications if appropriate
 - Lifestyle risk assessment³
 - Cancer screening⁴
 - Vaccinations⁵ as appropriate
 - HPV vaccination as clinically indicated (see [HPV Vaccination algorithm](#))
 - Screening for Hepatitis B and C as clinically indicated (see [Hepatitis B Virus \(HBV\) Screening and Management](#) and [Hepatitis C Virus \(HCV\) Screening](#) algorithms)
 - Consider cardiovascular screening (see [Survivorship – Adult Cardiovascular Screening algorithm](#))
 - Genetic screening⁶ (see [Genetic Counseling algorithm](#))

PSYCHOSOCIAL
FUNCTIONING

- Assess for:
- Distress management (see [Distress Screening and Psychosocial Management algorithm](#))
 - Social support
 - Financial stressors

CHRONIC HEALTH
MAINTENANCE

- Confirm primary care provider (PCP) or recommend establishing care with a local PCP
- PCP responsible for assessment and management of non-cancer chronic health conditions (hyperlipidemia, diabetes, hypertension, obesity, *etc.*) or refer as indicated

Refer or consult as indicated

¹ Low risk endometrial cancer is defined as any patient who did not receive chemotherapy or radiotherapy as adjuvant treatment after their initial surgery. Survivorship begins 3 years post-treatment and NED.

² High risk defined as patients who received chemotherapy or radiotherapy as adjuvant treatment after their surgery. Survivorship begins 5 years post-treatment and NED.

³ See [Physical Activity](#), [Nutrition](#), [Obesity Screening and Management](#) and [Tobacco Cessation Treatment](#) algorithms; ongoing reassessment of lifestyle risks should be a part of routine clinical practice

⁴ Includes [breast](#), [cervical](#), [colorectal](#), [liver](#), [lung](#), [pancreatic](#), and [skin](#) cancer screening

⁵ Based on [American Society of Clinical Oncology \(ASCO\) guidelines](#)

⁶ Consider genetic counseling if there has been a significant family history change since the last genetic consult, or if the patient has not previously had genetic counseling and has Lynch Syndrome risk factors. Lynch Syndrome risk factors: personal history of colon or rectal cancer; immediate family (first degree relatives such as parent, child, or sibling) with colorectal or endometrial cancer; immediate or extended family (first, second or third degree relatives including parent, child, sibling, aunt, uncle, nieces, nephews, grandparents, and first cousins) diagnosed before age 50 with colon, rectal or uterine cancer; any relatives tested positive for a Lynch Syndrome mutation (EPCAM, MLH1, MSH2, MSH6, PMS2 genes).

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DEVELOPMENT CREDITS

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

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