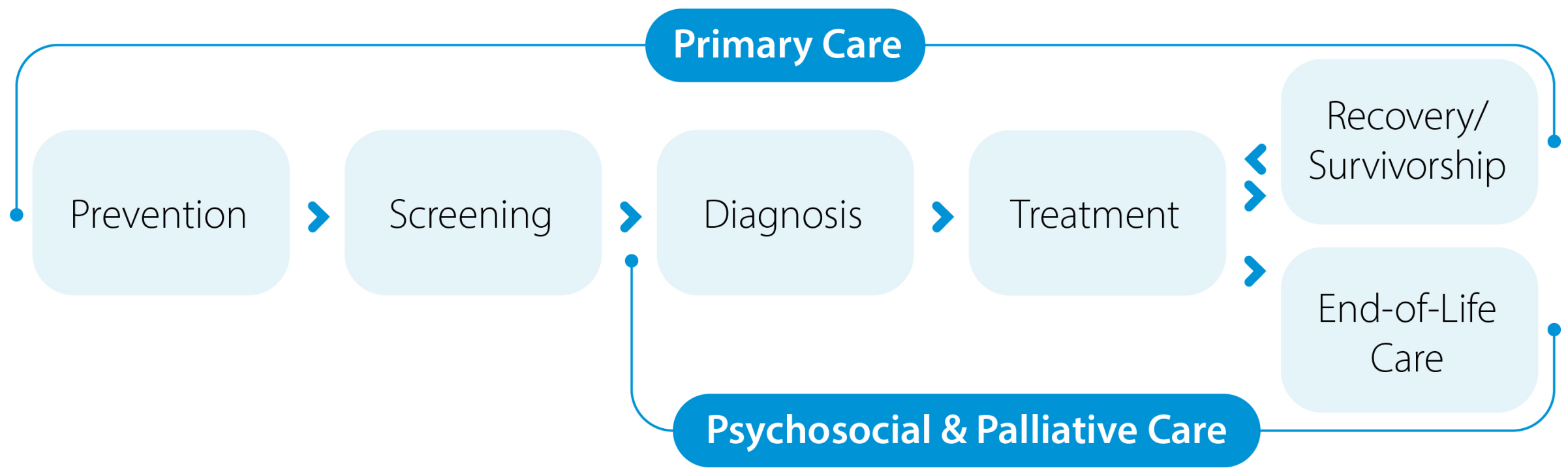


Skin Cancers Pathway Map

Version 2022.10



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


















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Pathway Map Considerations

- Primary care providers play an important role in the cancer journey and should be informed of relevant tests and consultations. Ongoing care with a primary care provider is assumed to be part of the pathway map. For patients who do not have a primary care provider, [Health Care Connect](#) is a government resource that helps patients find a doctor or nurse practitioner.
- Throughout the pathway map, a shared decision-making model should be implemented to enable and encourage patients to play an active role in the management of their care. For more information see [Person-Centered Care Guideline](#) and [EBS #19-2 Provider-Patient Communication.*](#)
- Hyperlinks are used throughout the pathway map to provide information about relevant CCO tools, resources and guidance documents.
- The term ‘healthcare provider’, used throughout the pathway map, includes primary care providers and specialists, e.g. family doctors, nurse practitioners, and emergency physicians.
- Complex skin cancers should be seen at either a cancer centre or a Mohs Centre, as appropriate (see Page 3).
- Physicians may work outside of a cancer centre but should participate in multidisciplinary care.
- For more information on Multidisciplinary Cancer Conferences, visit [MCC Tools](#).
- For more information on wait time prioritization, visit [Surgery](#).
- Clinical trials should be considered for all phases of the pathway map.
- Sexual health should be considered throughout the care continuum. Healthcare providers should discuss sexual health with patients before, during and after treatment as part of informed decision-making and symptom management. See [Psychosocial Oncology Guidelines Resources](#).
- Currently, we are not aware of the effect of systemic agents on skin cancer patients’ fertility. Healthcare providers should discuss potential effects on fertility with patients and arrange referral to a fertility specialist if appropriate. See [Ontario Fertility Program](#).
- Psychosocial oncology (PSO) is the interprofessional specialty concerned with understanding and treating the social, practical, psychological, emotional, spiritual and functional needs and quality-of-life impact that cancer has on patients and their families. Psychosocial care should be considered an integral and standardized part of cancer care for patients and their families at all stages of the illness trajectory. For more information, visit [EBS #19-3.*](#)
- The following should be considered when weighing the treatment options described in this pathway map for patients with potentially life-limiting illness:
 - Palliative care may be of benefit at any stage of the cancer journey, and may enhance other types of care – including restorative or rehabilitative care – or may become the total focus of care.
 - Ongoing discussions regarding goals of care is central to palliative care, and is an important part of the decision-making process. Goals of care discussions include the type, extent and goal of a treatment or care plan, where care will be provided, which health care providers will provide the care, and the patient’s overall approach to care.

* **Note:** [EBS #19-2](#) and [EBS #19-3](#) are older than 3 years and are currently listed as ‘For Education and Information Purposes’. This means that the recommendations will no longer be maintained but may still be useful for academic or other information purposes.

Pathway Map Legend

Colour Guide	Shape Guide	Line Guide
 Primary Care	 Intervention	 Required
 Palliative Care	 Decision or assessment point	 Possible
 Pathology	 Patient (disease) characteristics	
 Surgery	 Consultation with specialist	
 Radiation Oncology	 Exit pathway	
 Medical Oncology	 Off-page reference	
 Radiology	 Referral	
 Multidisciplinary Cancer Conference (MCC)		
 Dermatology		
 Psychosocial Oncology (PSO)		

Pathway Map Disclaimer

This pathway map is a resource that provides an overview of the treatment that an individual in the Ontario cancer system may receive.

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Ontario Health (Cancer Care Ontario) and the pathway map’s content providers (including the physicians who contributed to the information in the pathway map) shall have no liability, whether direct, indirect, consequential, contingent, special, or incidental, related to or arising from the information in the pathway map or its use thereof, whether based on breach of contract or tort (including negligence), and even if advised of the possibility thereof. Anyone using the information in the pathway map does so at his or her own risk, and by using such information, agrees to indemnify Ontario Health (Cancer Care Ontario) and its content providers from any and all liability, loss, damages, costs and expenses (including legal fees and expenses) arising from such person’s use of the information in the pathway map.

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The following definitions of **complex skin cancers** and **non-complex skin cancers** have been created based on clinical consensus from Ontario Health (Cancer Care Ontario)’s Skin Cancers Advisory Committee members. These definitions were created after conducting a literature search for classifications of skin cancers. When addressing clinical service organization and delivery of care for patients, the Advisory Committee felt the terms ‘complex skin cancers’ and ‘non-complex skin cancers’ best addressed this matter. This Advisory Committee includes clinicians from across the province and a variety of disciplines, including primary care, pathology, radiology, general surgery, plastics, dermatology, surgical oncology, radiation oncology, and medical oncology.

Complex Skin Cancers

General factors applicable to all types of skin cancer

Patient factors:

- Inoperable (patient or tumour factors)
- Initial assessment for skin cancers associated with genetic mutations (example: Gorlin’s syndrome)

Tumour factors:

- Node-positive (micro and macro)
- Locally advanced skin cancers (e.g. involving muscle or bone)
- Metastatic
- Subtypes: mucosal melanoma, ocular melanoma
- Cancers that developed in a scar, burn or a site previously treated
- In-transit, satellite disease, or recurrent disease

Treatment factors:

Any patient that needs:

- Surgical treatment including lymph node dissection (modified or radical neck dissection, axillary level 1- 3 dissection, superficial and deep groin dissection), resection of metastatic disease
- A medical oncologist opinion
- A radiation oncologist opinion
- Multidisciplinary care
- Consideration for clinical trials
- Mohs Micrographic surgery at Mohs centre as per Mohs guideline

[Patient Indications for Mohs Micrographic Surgery](#)

Complex Skin Cancers, continued

Factors specific to certain types of skin cancer

Melanoma

- See general factors, Stage IIB-IV

Merkel Cell Carcinoma

- All Merkel cell carcinomas

Squamous Cell Carcinoma

- Squamous cell carcinomas that show rapid growth (i.e. within weeks)
- Histologic Features: Any of depth > 6mm, perineural invasion ≥0.1mm, sensory or motor deficits, poorly differentiated, level IV/V invasion (muscle/bone invasion)

Basal Cell Carcinoma

- Basal cell carcinomas that show rapid growth (i.e. within weeks)
- Histologic features: Perineural invasion, sensory or motor deficits, level IV/V invasion (muscle/bone invasion)

Any other skin cancer histology

- Due to their rare occurrence, any skin cancer that is non-melanoma, non-basal cell carcinoma, non-squamous cell carcinoma (i.e. sebaceous carcinoma, adnexal carcinoma, etc.) is considered complex

Considerations for genetic testing

[Hereditary cancer testing](#) should be considered in patients with:

- ≥3 invasive melanomas (page 14)
- Melanoma, especially if diagnosed ≤40 years of age and have a family history of melanoma and/or pancreatic cancer (page 14)
- ≥5 basal cell carcinomas <30 years of age or with other features of Gorlin syndrome/Nevoid Basal Cell Carcinoma Syndrome (page 21)
- MMR IHC deficient sebaceous neoplasm/carcinoma (page 11)
- melanomas identified as having germline relevant variants in tumour tissue (e.g. CDKN2A, BAP1) (page 8)

For individuals with a hereditary cancer syndrome associated with an increased risk of skin cancer, a cancer genetics clinic will advise on appropriate management and surveillance recommendations.

Non-Complex Skin Cancers

Melanoma

- Stage IA, IB, IIA cutaneous melanoma

Merkel Cell Carcinoma

- None

Squamous Cell Carcinoma (SCC)

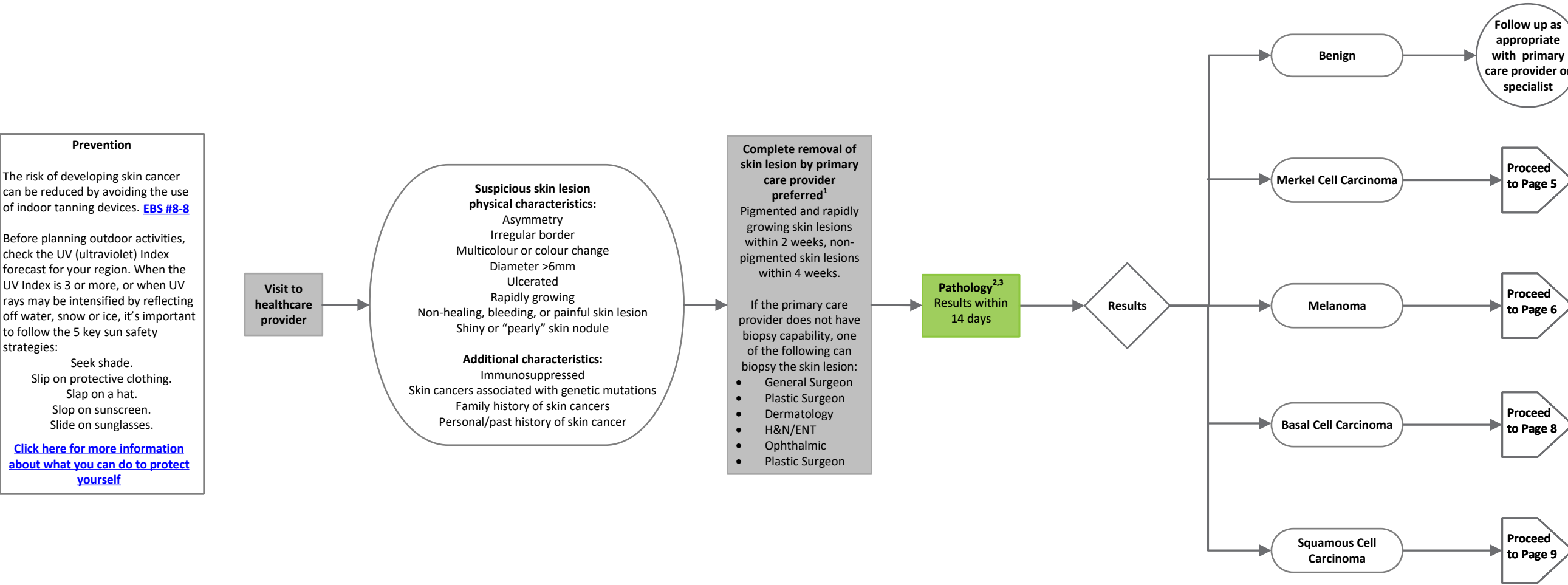
- Any other SCC features not indicated in Complex SCC characteristics

Basal Cell Carcinoma (BCC)

- Any other BCC features not indicated in Complex BCC characteristics

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¹ Biopsy can include punch biopsy, excisional biopsy, shave biopsy or incisional biopsy. For a pigmented lesion, the depth of the biopsy should be at least to deep dermis or subcutaneous tissue to ensure adequate sampling and depth assessment.

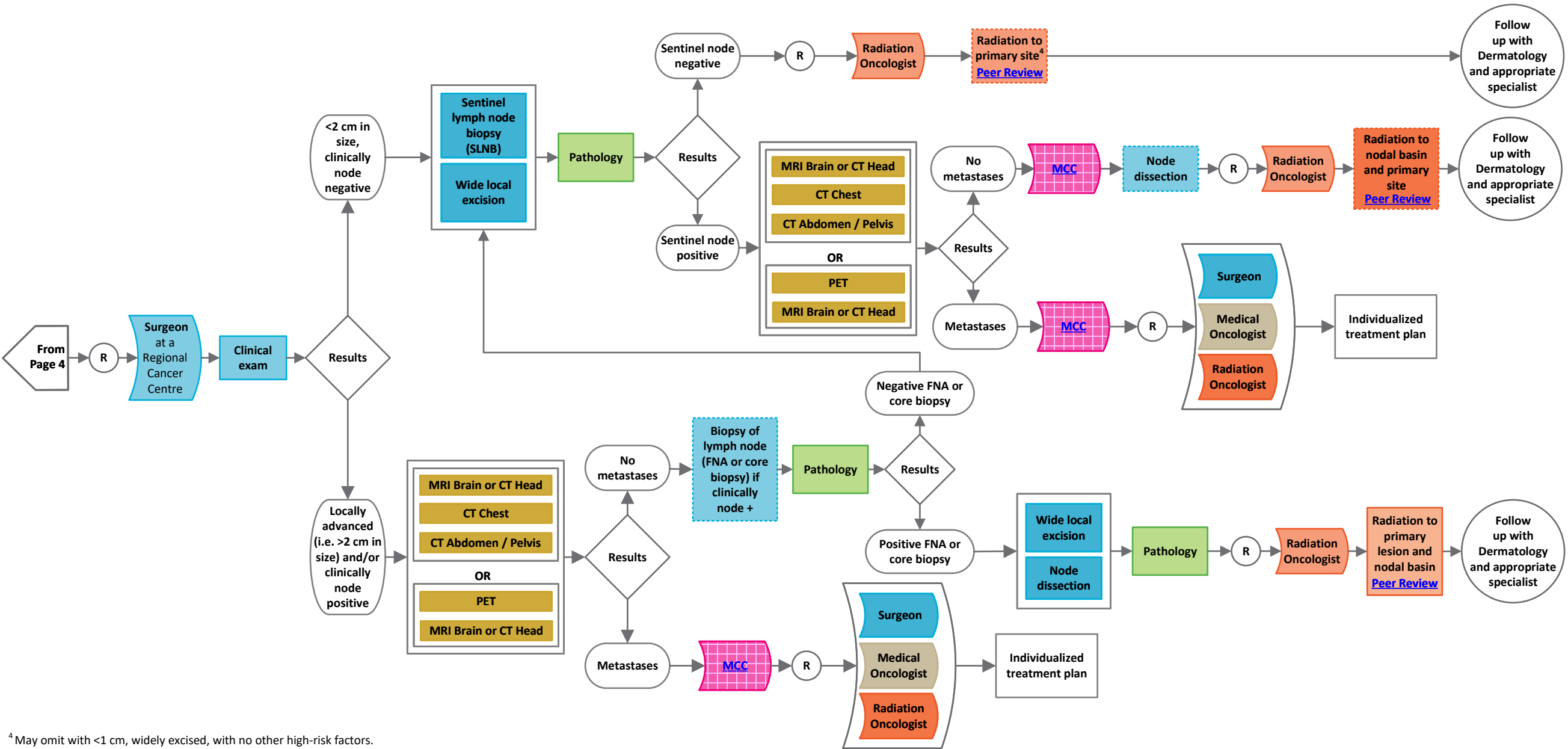
² If >4mm and/or node positive, send specimen for molecular testing.

³ The Ontario Health (Cancer Care Ontario) pathology post-surgical turn-around time indicator targets 85% within 14 days.

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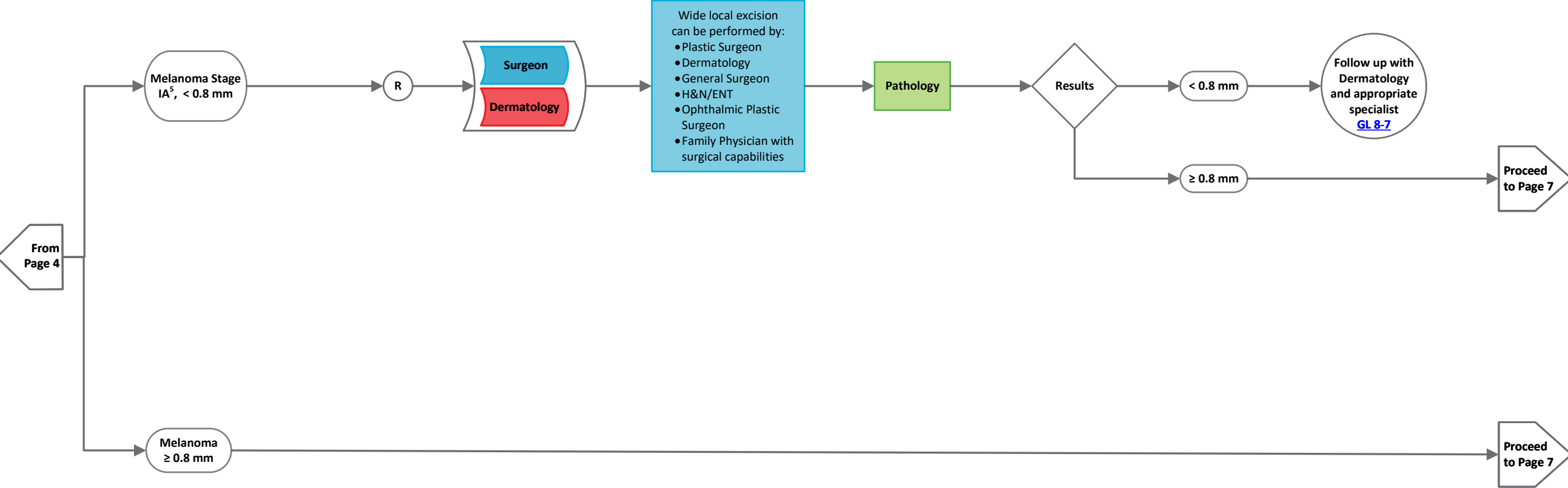


⁴ May omit with <1 cm, widely excised, with no other high-risk factors.

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⁵ <0.8mm and ulcerated or multiple mitoses may require a referral and assessment by a surgeon for sentinel lymph node biopsy.

Skin Cancers Pathway Map

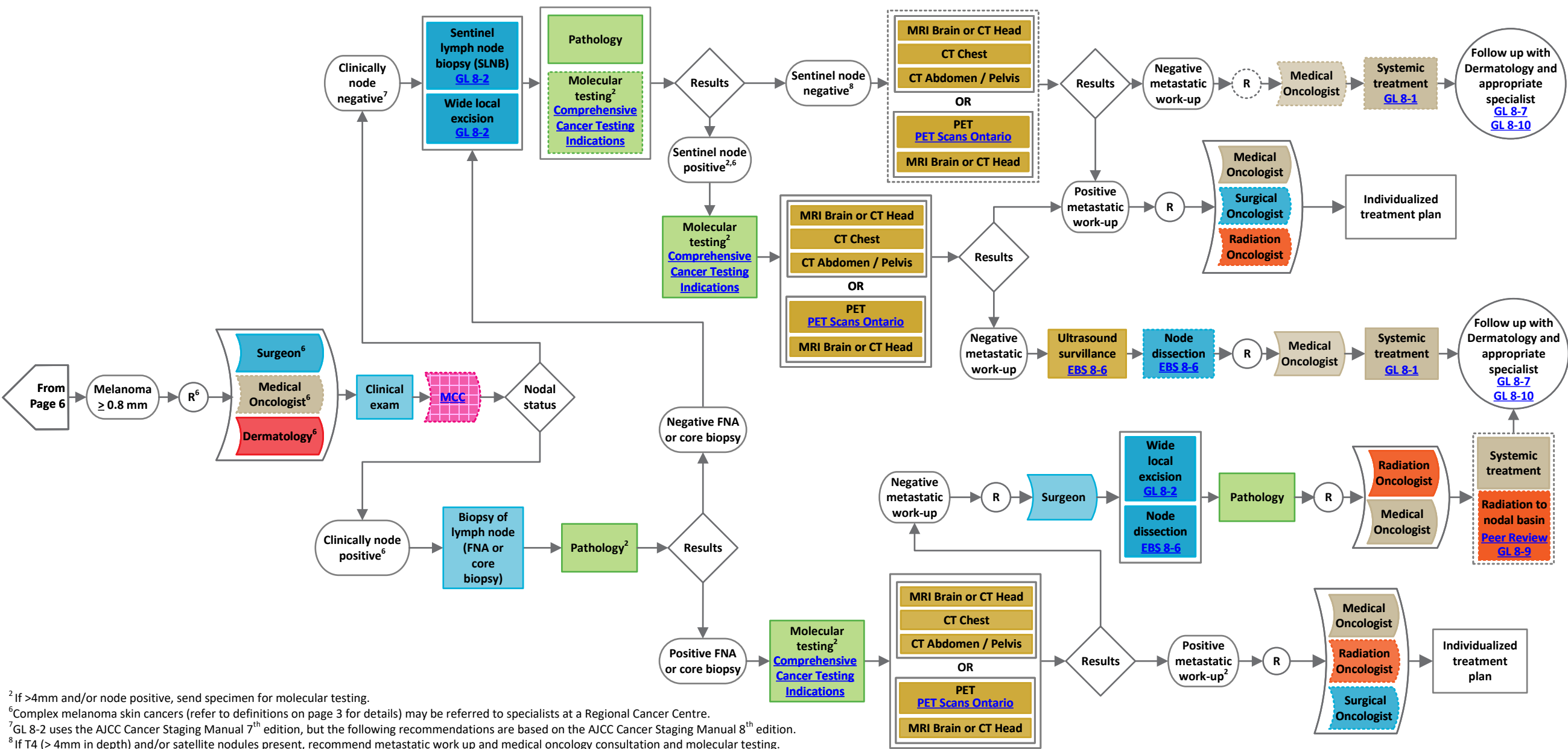
Melanoma - 0.8mm or Deeper

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² If >4mm and/or node positive, send specimen for molecular testing.

⁶ Complex melanoma skin cancers (refer to definitions on page 3 for details) may be referred to specialists at a Regional Cancer Centre.

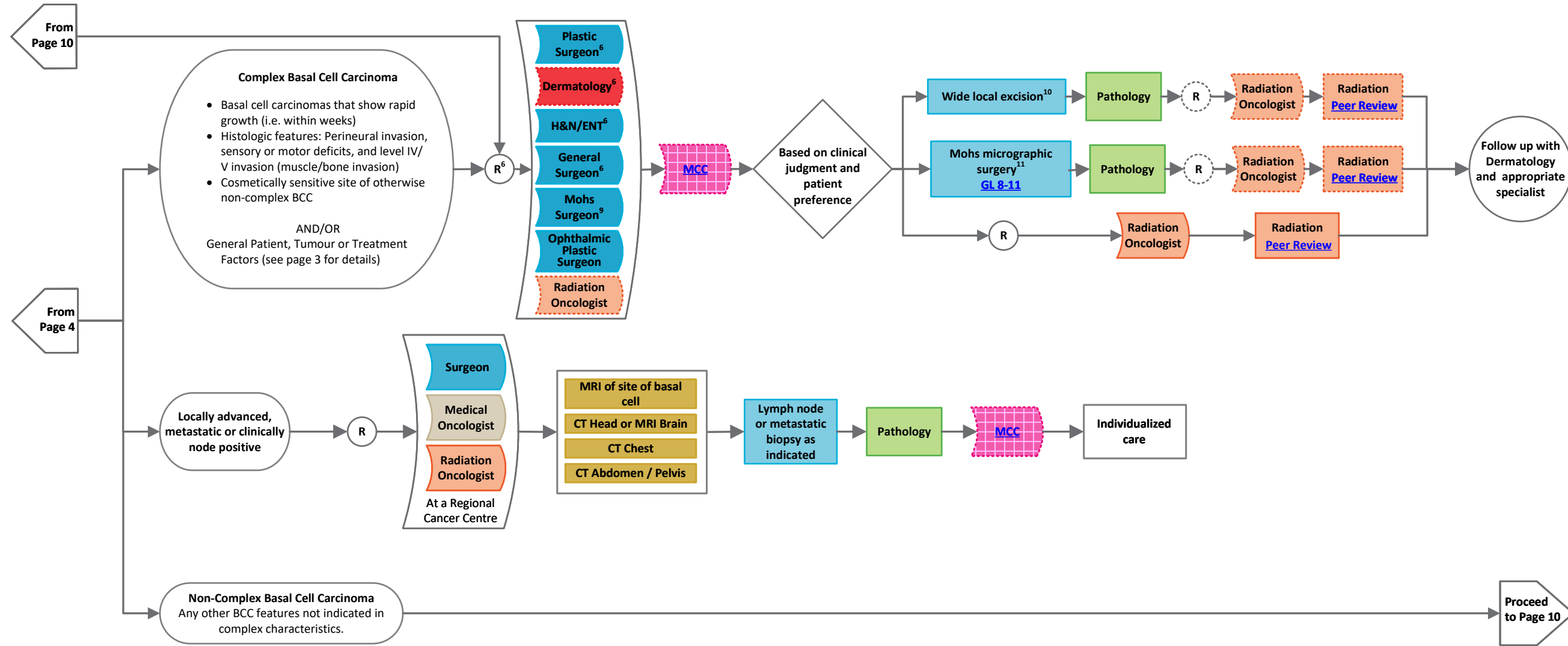
⁷ GL 8-2 uses the AJCC Cancer Staging Manual 7th edition, but the following recommendations are based on the AJCC Cancer Staging Manual 8th edition.

⁸ If T4 (> 4mm in depth) and/or satellite nodules present, recommend metastatic work up and medical oncology consultation and molecular testing.

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⁶ Complex Basal Cell Carcinomas (refer to definitions on page 3 for details) may be referred to specialists at a Regional Cancer Centre.

⁹ Complex BCC or SCC (refer to definitions on page 3 for details) may be referred to specialists at a Regional Cancer Centre or a Mohs centre if eligible for Mohs surgery. [Patient Indications for Mohs Micrographic Surgery](#)

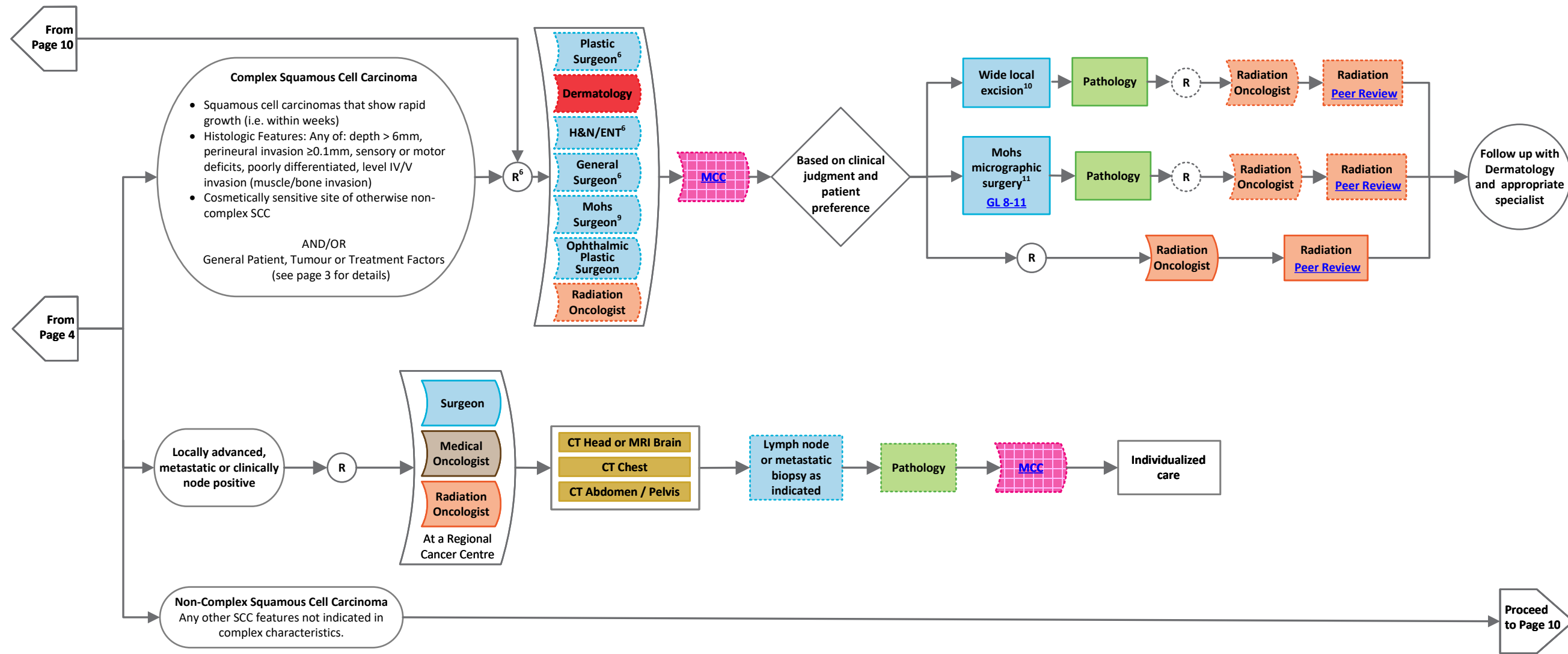
¹⁰ If positive margin, patient should be considered complex and potentially sent to Regional Cancer Centre or Mohs centre.

¹¹ Indications for Mohs micrographic surgery: histologically confirmed recurrent BCC of face, primary BCC of face >1cm, aggressive histology or location on the H zone of the face. Mohs surgery is recommended for SCC in some cases (as per guideline).

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⁶ Complex squamous cell skin cancers (refer to definitions on page 3 for details) may be referred to specialists at a Regional Cancer Centre.

⁹ Complex BCC or SCC (refer to definitions on page 3 for details) may be referred to specialists at a Regional Cancer Centre or a Mohs centre if eligible for Mohs surgery. [Patient Indications for Mohs Micrographic Surgery](#)

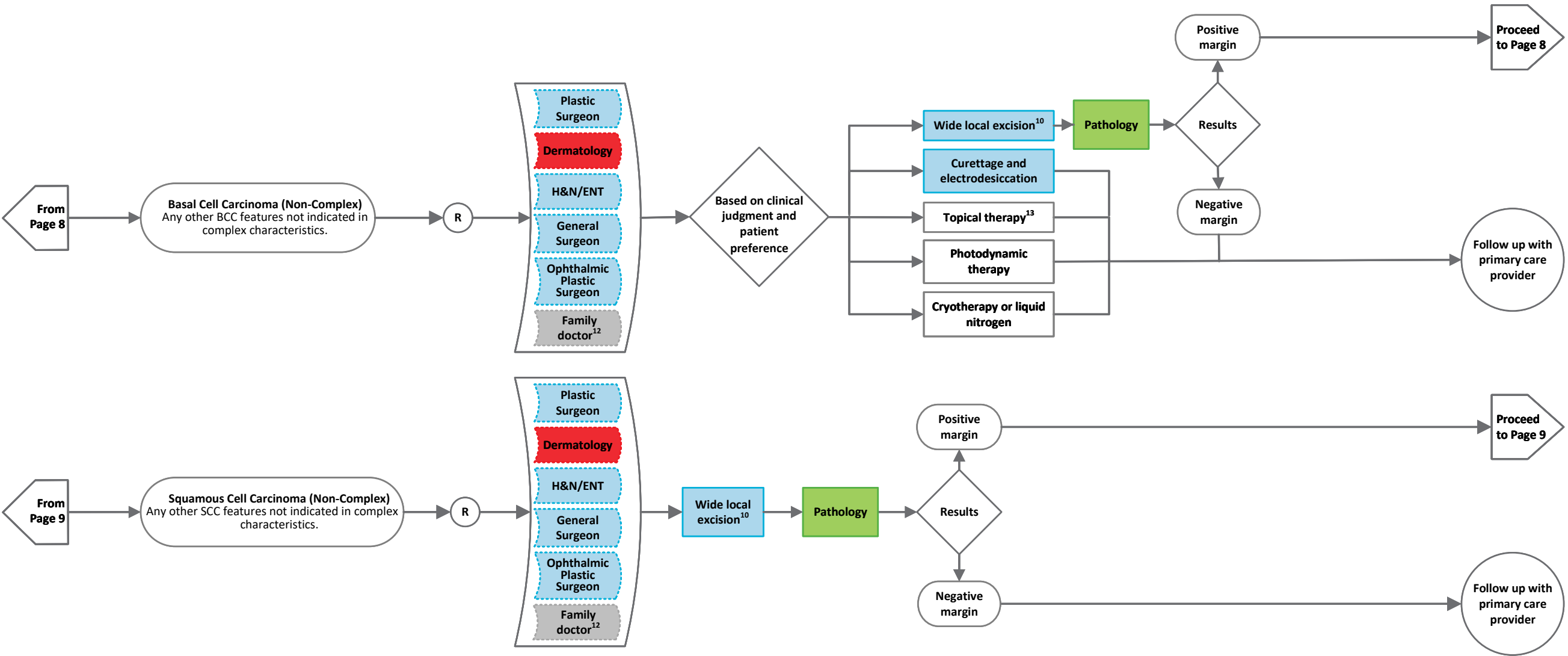
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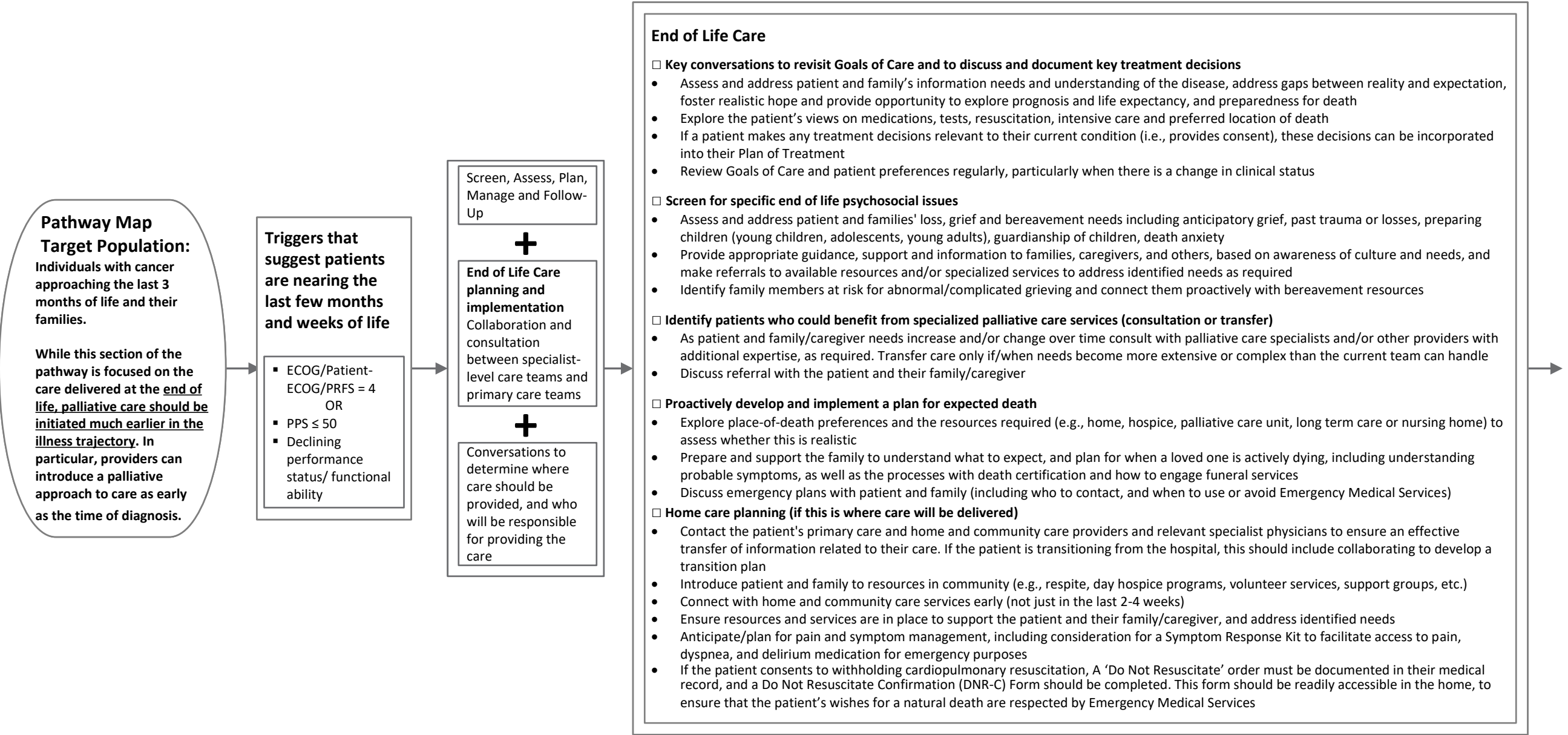
¹² With special interest in surgical procedures.

¹³ Superficial tumours only.

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