

Acute Lymphoblastic Leukemia and Lymphoblastic Lymphoma (ALL) – Adult¹

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.*

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¹ Age ≥ 18 years old

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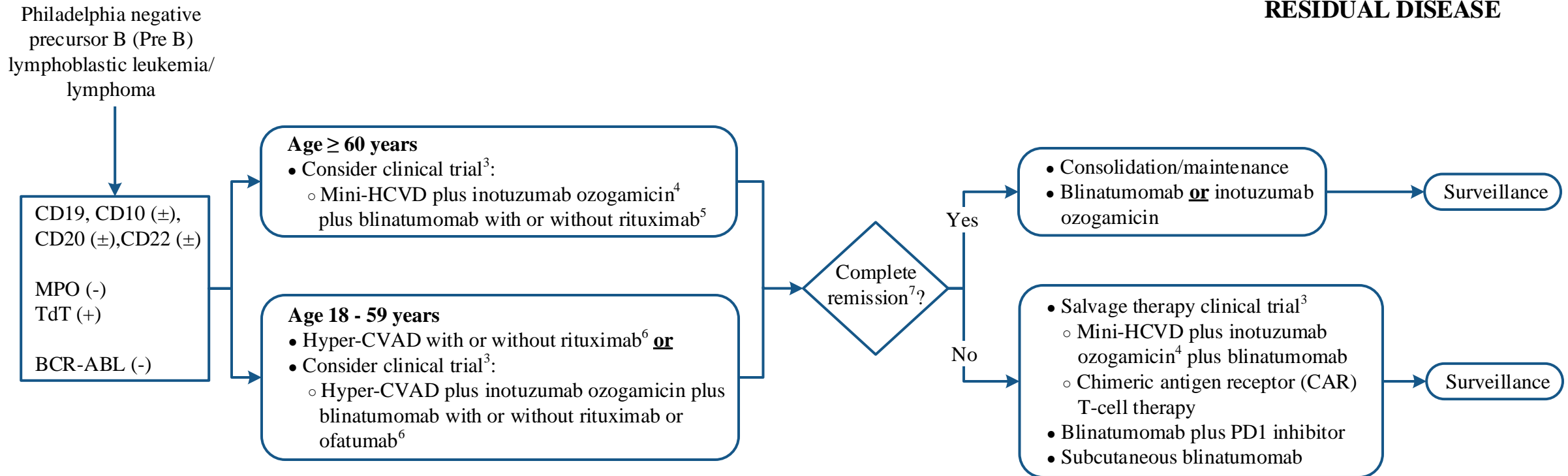
Note: Consider clinical trials as treatment options for eligible patients. Stem Cell Transplant (SCT) guidelines are not included with this algorithm. Leukemia patients should be referred and treated at a Comprehensive Cancer Center.

PATIENT PRESENTATION^{1,2}

TREATMENT

ASSESSMENT OF RESPONSE

POST-REMISSION THERAPY/MINIMAL RESIDUAL DISEASE



¹ See [Physical Activity](#), [Nutrition](#), and [Tobacco Cessation](#) algorithms; ongoing reassessment of lifestyle risks should be a part of routine clinical practice

² Consider MD Anderson approved biomarkers

³ See [Leukemia Clinical Trials](#)

⁴ Mini-HCVD (cyclophosphamide and dexamethasone at 50% dose reduction, no anthracycline, methotrexate at 75% dose reduction, cytarabine at 0.5 g/m² for 4 doses) plus inotuzumab ozogamicin

⁵ Rituximab if CD20 ≥ 20%

⁶ Hyper-CVAD (hyper-fractionated cyclophosphamide, vincristine, doxorubicin, dexamethasone); rituximab if CD20 ≥ 20%

Hyper-CVAD (hyper-fractionated cyclophosphamide, vincristine, doxorubicin, dexamethasone); ofatumumab if CD20 ≥ 1%

⁷ Failure after induction with hyper-CVAD based regimen means no response after 2 cycles of chemotherapy

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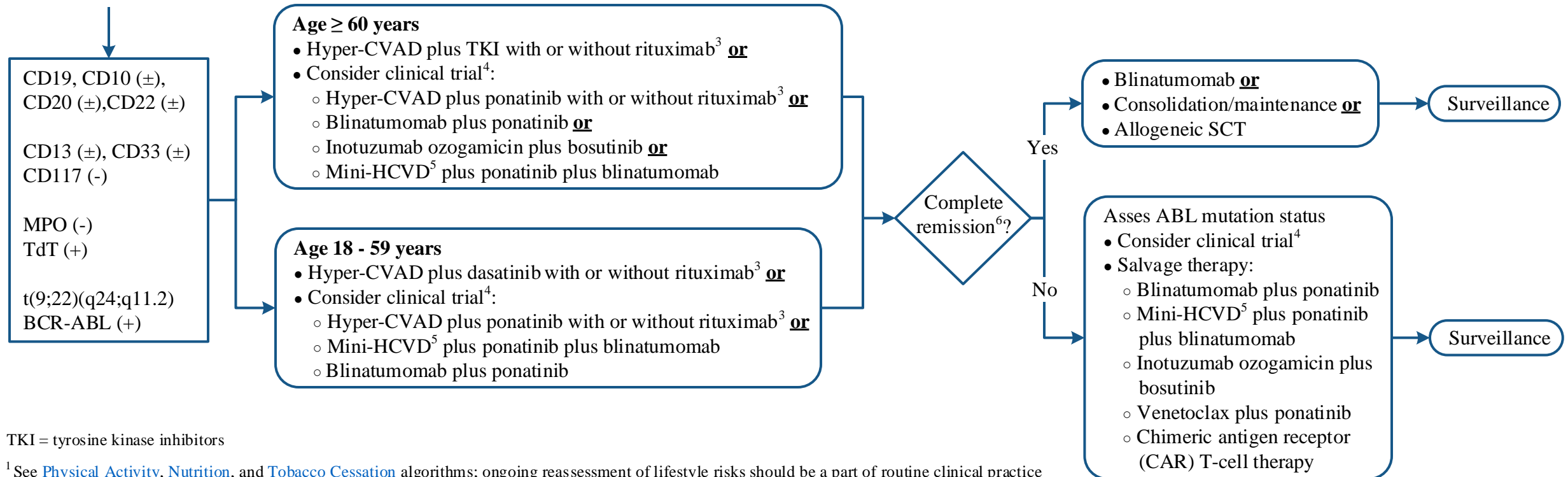
PATIENT PRESENTATION^{1,2}

TREATMENT

ASSESSMENT OF RESPONSE

POST-REMISSION THERAPY

Philadelphia chromosome (Ph) positive acute lymphoblastic leukemia



TKI = tyrosine kinase inhibitors

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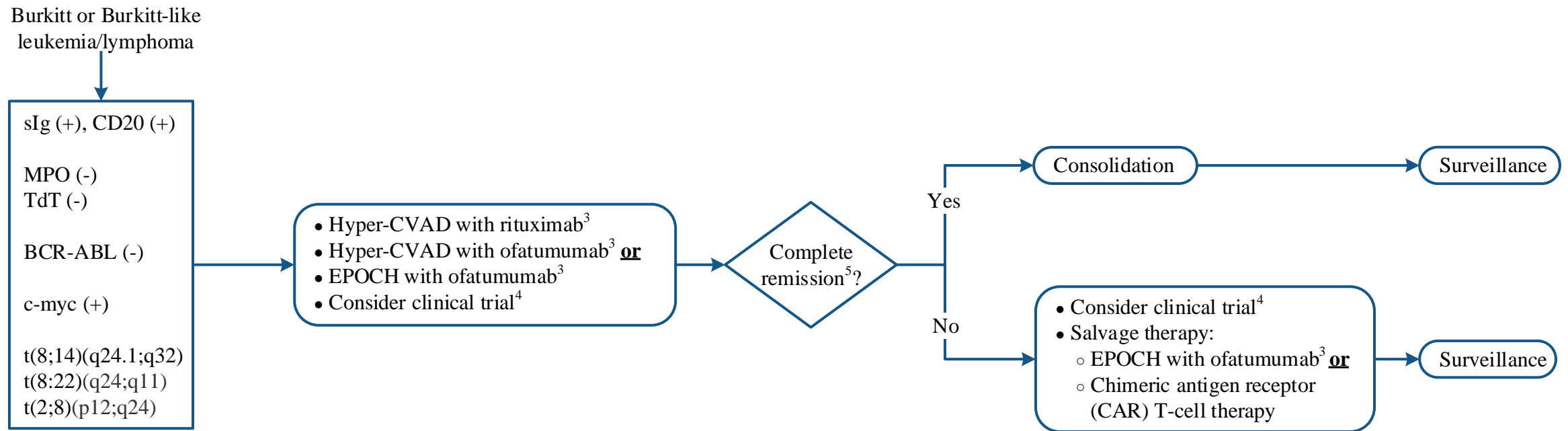
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PATIENT PRESENTATION^{1,2}

TREATMENT

ASSESSMENT OF RESPONSE

POST-REMISSION THERAPY



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² Consider MD Anderson approved biomarkers

³ Hyper-CVAD (hyper-fractionated cyclophosphamide, vincristine, doxorubicin, dexamethasone) plus rituximab
 Hyper-CVAD (hyper-fractionated cyclophosphamide, vincristine, doxorubicin, dexamethasone) plus ofatumumab
 EPOCH (etoposide, prednisone, vincristine, cyclophosphamide, doxorubicin) plus ofatumumab

⁴ See [Leukemia Clinical Trials](#)

⁵ Failure after induction with hyper-CVAD based regimen means no response after 2 cycles of chemotherapy

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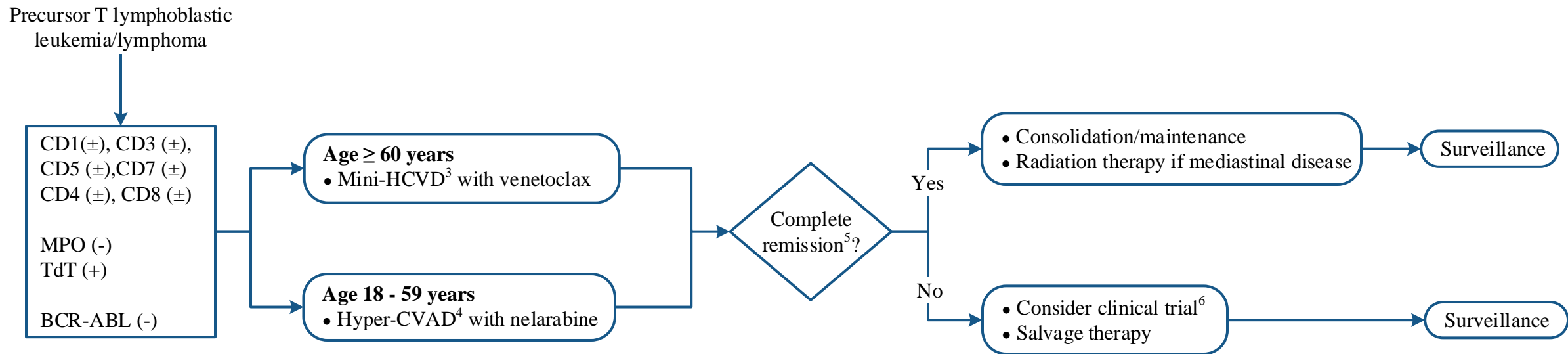
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TREATMENT

ASSESSMENT OF RESPONSE

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DEVELOPMENT CREDITS

This practice algorithm is based on majority expert opinion of the Leukemia Center Faculty 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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