

SUPPLEMENTAL MATERIALS

Supplementary Table S1. Regimens considered as first-line therapy for MCL

Abbreviated regimen name	Full regimen name
BR	Bendamustine, rituximab [Rituxan]
CHOP	Cyclophosphamide, doxorubicin, vincristine, prednisone
R-CHOP	Rituximab, cyclophosphamide, doxorubicin [Hydroxydaunorubicin], vincristine [Oncovin], prednisone
R-DHAP*	Rituximab, dexamethasone, high-dose cytarabine (Ara-C), cisplatin
CVP	Cyclophosphamide, vincristine, prednisone
R-CVP	Rituximab, cyclophosphamide, vincristine, prednisone
Hyper-CVAD	Cyclophosphamide, vincristine, doxorubicin [Adriamycin], dexamethasone, methotrexate, cytarabine
R-HyperCVAD*	Rituximab, cyclophosphamide, vincristine, doxorubicin, dexamethasone, methotrexate, cytarabine
CHLO	Chlorambucil
CHLO-R	Chlorambucil, rituximab
CYCL(PO)	Cyclophosphamide (oral)
FC	Fludarabine, cyclophosphamide
FC-R	Fludarabine, cyclophosphamide, rituximab
FR	Fludarabine, rituximab
FLUD-CYCLO-MITOXAN	Fludarabine, cyclophosphamide, mitoxan
VR-CAP*	Bortezomib, rituximab, cyclophosphamide, doxorubicin, prednisone
LR*	Lenalidomide, rituximab
RBAC500*	Rituximab, bendamustine, cytarabine
Nordic regimen*	Dose-intensified induction with rituximab plus cyclophosphamide, vincristine, doxorubicin, prednisone alternating with rituximab plus high-dose cytarabine

*Regimens not identified in the MCL cohort but listed as recommended 1st LoT for MCL in guidelines

Supplemental Table S2. Patient selection into MCL cohort, Ontario, Canada, 2013 – 2016

Cohort Selection Criteria		Remaining Patients		Excluded
		No.	%	patients
MCL cohort inclusion criteria	Note			
Diagnosis prior to December 31 st , 2016 with MCL (Morphology code = 96733)		2,100		
Exclusion Criteria				
Invalid OHIP card number and age<65		1,270		
Invalid or incomplete records (missing age, sex, LHIN, income quintile, and rurality, or death before MCL diagnosis)		1,225		
Age ≥105 years at MCL diagnosis		1,225		
Non-Ontario resident		1,225		
Undetermined MCL diagnosis (cancers other than MCL diagnoses on the same date)		1,212		
No OHIP eligibility at diagnosis		1,206		
Death date recorded on the date of index diagnosis		1,206		
Excluding those having MCL diagnosis date before January 1, 2013		325		
Excluding those having SCT any time before the end of the Analysis Period (including)		313		
Pool of MCL Patients		313	100.0%	
MCL ≥ 65 in 1LoT subgroup inclusion criteria		313		
Patients in the MCL ≥ 65 cohort who has record in ALR within 3 years from index date and up to 31Dec2019		213	68.1%	100
Excluding patients who participated in a clinical trial	(n=*1-5)			*1-5
Patients received confirmed 1st LoT	Excluded patients including those who participated in a trial	183	58.5%	30
Excluding patients who had refractory disease or treatment intolerance		161	51.4%	22
MCL cohort		161		
Matching to 4 general population controls		159	50.8%	2
Matched MCL Patients		159		

*Values were suppressed as per ICES reporting standards to reduce the risk of patient re-identification

Supplemental Table S3. First line therapy received by the MCL cohort^

Lines of Therapy		n (%)
Regimen category	Overall	161^ (100%)
	BR	125 (78.1%)
	R-CHOP	21 (13.1%)
	CHOP	0
	R-CVP	*9-13
	CVP	0
	Hyper-CVAD	0
	CHLO	*1-5
	CHLO-R	0
	CYCL(PO)	0
	FC	0
	FC-R	*1-5
	FR	0
	FLUD-CYCLO-MITOXAN	0
R-maintenance for 1 st LoT, n (%)		126 (78.3%)
Time to next therapy (TTNT), days	n (%)	38 (23.6%)
(from initiation of 1 st LoT to the initiation of 2 nd LoT)	Mean, SD	465.6, 220.6
	Median (IQR)	462.5 (297-628)
	Min - Max	29-988

^MCL cohort prior to general population matching

*Double suppression was conducted according to ICES reporting standards to reduce the risk of patient re-identification

Supplemental Table S4. Demographic and clinical characteristics for matched MCL patients, by LoT categories

Demographic and Clinical Characteristics		BR (n=123)	Other (n=36)
Sex			
	Female	35 (28.5%)	16 (44.4%)
	Male	88 (71.5%)	20 (55.6%)
Age			
	Mean, SD	75.0 ± 6.0	76.1 ± 6.7
	Median (IQR)	74 (70 - 80)	76 (72 - 81)
	Min - Max	65 – 92	65 – 91
	65 - 74 years	63 (51.2%)	14 (38.9%)
	75+ years	60 (48.8%)	22 (61.1%)
Rural residence			
	Urban	*100 - 104	*31 - 35
	Rural	*19 - 23	*1 - 5
Income quintile			
	Q1, lowest	*19 - 23	*1 – 5
	Q2	*26 - 30	*3 – 7
	Q3	23 (18.7%)	6 (16.7%)
	Q4	23 (18.7%)	10 (27.8%)
	Q5, highest	29 (23.6%)	11 (30.6%)
New Ontario Resident at diagnosis		*1 - 5	*1 – 5
Ontario Health Region			
	Central	40 (32.5%)	17 (47.2%)
	East	*31 - 35	*7 – 11

	West	36 (29.3%)	7 (19.4%)
	North	*12 - 16	*1 - 5
Year of MCL cancer diagnosis			
	2013	*14 - 18	*16 - 20
	2014	33 (26.8%)	8 (22.2%)
	2015	35 (28.5%)	7 (19.4%)
	2016	*37 - 41	*1 - 5
Cancer stage at diagnosis			
	Stage I	0 (0%)	*1 - 5
	Stage II	*1 - 5	0 (0%)
	Stage III	*16 - 20	*1 - 5
	Stage IV	*25 - 29	*4 - 8
	Missing	74 (60.2%)	26 (72.2%)
Cancer history (assessed within 5-year lookback)^		*10 - 14	*1 - 5
Charlson's co-morbidity index (assessed within 2-year lookback)			
	0	*18 - 22	*1 - 5
	1	*1 - 5	*1 - 5
	2	*1 - 5	*1 - 5
	3+	*1 - 5	*1 - 5
	Missing	93 (75.6%)	26 (72.2%)
Comorbidities, any time before index date (ICES-derived cohorts)			
	Diabetes mellitus	28 (22.8%)	10 (27.8%)

	Congestive heart failure (CHF)	*3 – 7	*1 – 5
	Chronic obstructive pulmonary disease (COPD)	29 (23.6%)	6 (16.7%)
	Rheumatoid arthritis (RA)	*1 - 5	*1 – 5
	Crohn's disease / Ulcerative colitis	*1 - 5	0 (0%)
Time from index date to end of follow-up			
	Mean, SD	947.2 ± 296.9	806.4 ± 379.0
	Median (IQR)	1,095 (1,025 - 1,095)	1,095 (503 - 1,095)
	Min - Max	28 - 1,095	45 - 1,095

^Due to small cell sizes, the types of previous cancers could not be reported

*Double suppression was conducted according to ICES reporting standards to reduce the risk of patient re-identification