

Supplementary Table 2. Association between secretor MBL2 polymorphisms and disease-specific survival, stratified by study population*

Genotype	Case patients from case-control study				Case-only series			
	White patients		African American patients		White patients		African American patients	
	No.	RR of death† (95% CI)	No.	RR of death (95% CI)	No.	RR of death† (95% CI)	No.	RR of death† (95% CI)
Promoter SNP								
mbl2_11 (<i>rs11003125</i>)								
C/C (<i>L/L</i>)	140	1.0 (referent)	89	1.0 (referent)	75	1.0 (referent)	31	1.0 (referent)
C/G (<i>L/H</i>)	159	1.10 (0.75 to 1.61)	35	1.08 (0.60 to 1.95)	79	0.72 (0.45 to 1.15)	9	0.64 (0.19 to 2.20)
G/G (<i>H/H</i>)	49	1.12 (0.67 to 1.89)	4	2.11 (0.63 to 7.04)	21	1.70 (0.89 to 3.26)	0	—
mbl2_12 (<i>rs7096206</i>)								
G/G (<i>Y/Y</i>)	219	1.0 (referent)	93	1.0 (referent)	110	1.0 (referent)	31	1.0 (referent)
G/C (<i>Y/X</i>)	120	0.65 (0.45 to 0.93)	31	1.07 (0.59 to 1.93)	60	0.56 (0.34 to 0.92)	10	1.69 (0.57 to 5.01)
C/C (<i>X/X</i>)	15	0.59 (0.24 to 1.48)	3	1.81 (0.41 to 7.99)	9	0.55 (0.17 to 1.83)	1	—
G/C + C/C	135	0.64 (0.45 to 0.91)	34	1.13 (0.64 to 1.99)	69	0.56 (0.35 to 0.90)	11	1.90 (0.68 to 5.32)
Exon 1 SNP								
mbl2_03 (<i>rs5030737</i>)								
C/C (<i>Non-D/Non-D</i>)	292	1.0 (referent)	122	1.0 (referent)	155	1.0 (referent)	38	1.0 (referent)
C/T (<i>Non-D/D</i>)	56	0.74 (0.47 to 1.16)	5	0.43 (0.10 to 1.81)	19	1.72 (0.94 to 3.13)	1	—
T/T (<i>D/D</i>)	3	2.82 (0.38 to 20.72)	0	—	1	—	0	—
mbl2_01 (<i>rs1800450</i>)								
G/G (<i>Non-B/Non-B</i>)	264	1.0 (referent)	117	1.0 (referent)	127	1.0 (referent)	35	1.0 (referent)
G/A (<i>Non-B/B</i>)	80	1.00 (0.67 to 1.51)	7	1.30 (0.29 to 5.81)	40	0.80 (0.48 to 1.34)	6	1.55 (0.49 to 4.92)
A/A (<i>B/B</i>)	3	3.19 (0.43 to 23.60)	1	—	4	1.20 (0.28 to 5.03)	0	—
mbl2_02 (<i>rs1800451</i>)								
G/G (<i>Non-C/Non-C</i>)	337	1.0 (referent)	73	1.0 (referent)	42	1.0 (referent)	18	1.0 (referent)
G/A (<i>Non-C/C</i>)	8	1.38 (0.50 to 3.77)	42	0.92 (0.53 to 1.59)	6	0.65 (0.09 to 4.95)	16	1.08 (0.37 to 3.09)
A/A (<i>C/C</i>)	1	—	6	0.59 (0.14 to 2.53)	0	—	4	7.42 (1.16 to 47.60)
Haplotype								
GGCGG (<i>HYA</i>)	192	1.0 (referent)	38	1.0 (referent)	102	1.0 (referent)	8	1.0 (referent)
CGCGG (<i>LYA</i>)	186	1.00 (0.73 to 1.37)	107	0.65 (0.38 to 1.11)	94	1.38 (0.94 to 2.05)	28	0.93 (0.33 to 2.65)

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CCCGG (<i>LXA</i>)	142	0.69 (0.48 to 0.99)	34	0.72 (0.38 to 1.35)	78	0.70 (0.44 to 1.12)	12	2.31 (0.73 to 7.34)
CGCAG (<i>LYB</i>)	84	1.03 (0.69 to 1.54)	9	0.56 (0.13 to 2.48)	51	0.96 (0.59 to 1.55)	6	1.67 (0.43 to 6.48)
CGCGA (<i>LYC</i>)	10	1.17 (0.42 to 3.20)	53	0.58 (0.32 to 1.06)	6	0.52 (0.07 to 3.82)	23	1.22 (0.43 to 3.44)
GGTGG (<i>HYD</i>)	60	0.73 (0.46 to 1.16)	5	0.31 (0.07 to 1.37)	21	1.85 (0.99 to 3.34)	1	—
Diplotype								
YA/YA	98	1.0	40	1.0	61	1.0	6	1.0
YA/XA	85	0.72 (0.45 to 1.16)	25	1.30 (0.64 to 2.64)	39	0.65 (0.35 to 1.21)	6	1.30 (0.64 to 2.64)
XA/XA	14	0.61 (0.23 to 1.59)	2	—	9	0.56 (0.16 to 1.90)	1	—
YA/B	52	1.27 (0.77 to 2.10)	4	0.96 (0.12 to 7.61)	26	0.81 (0.43 to 1.54)	3	0.96 (0.12 to 7.61)
XA/B	19	0.30 (0.09 to 0.99)	0	—	11	0.37 (0.13 to 1.06)	2	—
YA/C	6	1.57 (0.48 to 5.18)	33	1.15 (0.59 to 2.25)	2	—	14	1.15 (0.59 to 2.25)
YA/D	39	0.65 (0.36 to 1.17)	3	0.26 (0.03 to 1.99)	7	3.03 (1.22 to 7.52)	1	—
XA/D	9	0.59 (0.21 to 1.67)	0	—	8	0.58 (0.17 to 1.91)	0	—
XA/C	1	—	5	0.32 (0.07 to 1.46)	2	—	2	—
O/O	14	0.80 (0.31 to 2.10)	11	0.62 (0.18 to 2.17)	11	1.36 (0.60 to 3.08)	4	0.62 (0.18 to 2.17)

*RR = risk ratio; CI = confidence interval; SNP = single-nucleotide polymorphism; — = insufficient data. Statistically significant associations are in boldface type.

†Risk ratio of death was adjusted for sex, stage (III–IV vs. I–II), age at diagnosis, current smoking status, and pack-years of smoking.