

*HLA-DRB1*1501* and Spinal Cord Magnetic Resonance Imaging Lesions in Multiple Sclerosis

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eTable. Single-Nucleotide Polymorphisms Included on DNA Chip

| Gene | rs-nr | Chromosome | Polymorphism | HWE ^a | MAF |
|-----------|------------------|------------|--------------|------------------|-------------------|
| ADAMTS14 | rs4747075 | 10q22 | A/G | <i>P</i> < .01 | 0.30 |
| ADAMTS14 | rs7081273 | 10q22 | C/G | NS | 0.34 |
| ADAMTS14 | rs4746060 | 10q22 | C/T | NS | 0.08 |
| Apo I/Fas | rs1800682 | 10q23 | C/T | NS | 0.47 |
| Apo I/Fas | rs3781202 | 10q23 | C/T | <i>P</i> < .01 | 0.40 |
| Apo I/Fas | rs2234978 | 10q23 | C/T | NS | 0.31 |
| BTNL2 | rs2076530 | 6p21.3 | A/G | <i>P</i> < .01 | 0.26 |
| CIITA | rs3087456 | 16p13 | A/G | NS | 0.26 |
| CACNG4 | rs4790896 | 17q24 | A/G | NS | 0.41 |
| CCR5 | rs333 | 3p21 | -/+ | NS | 0.11 |
| CD24 | rs8734 | 6q21 | C | NA | 0.00 ^b |
| GNTF | rs1800169 | 11q12 | A/G | NS | 0.12 |
| CRYAB | rs14133 | 11q21-q23 | C/G | NS | 0.27 |
| CRYAB | rs762550 | 11q21-q23 | A/G | NS | 0.42 |
| CRYAB | rs2234702 | 11q21-q23 | C | NA | 0.00 ^b |
| CTLA4 | rs231775 | 2q33 | A/G | NS | 0.37 |
| CTLA4 | rs5742909 | 2q33 | C/T | NS | 0.09 |
| EBF1 | rs1368297 | 5q34 | A/T | NS | 0.38 |
| GABBR1 | rs1805057 | 6p22 | C | NA | 0.00 ^b |
| HELZ | rs2363846 | 17q24 | C/T | NS | 0.48 |
| HLA | rs2395166 | 6p21.3 | C/T | NS | 0.47 |
| HLA | rs2213584 | 6p21.3 | A/G | NS | 0.40 |
| HLA | rs2227139 | 6p21.3 | C/T | NS | 0.40 |
| HLA | rs3135388 | 6p21.3 | A/G | NS | 0.33 |
| HLA | rs9268458 | 6p21.3 | A/C | NS | 0.20 |
| HLA | rs6457594 | 6p21.3 | A/G | <i>P</i> < .01 | 0.40 |
| HLA-DRA | rs2395182 | 6p21.3 | G/T | NS | 0.38 |
| HLA-DRA | rs2239802 | 6p21.3 | C/G | NS | 0.38 |
| IFNAR1 | rs2257167 | 21q22 | C/G | NS | 0.08 |
| IFNGR2 | rs9808753 | 21q22 | A/G | NS | 0.14 |
| IKBL | rs3130062 | 6p21.3 | C/T | NS | 0.18 |
| IL-10 | rs1800896 | 1q32 | A/G | NS | 0.46 |
| IL-1β | rs1799916 | 2q14 | A | NA | 0.00 ^b |
| IL-1β | rs1143627 | 2q14 | A/G | NS | 0.34 |
| IL-1β | rs1143634 | 2q14 | C/T | NS | 0.23 |
| IL-1RN | rs419598 | 2q12-q14 | C/T | NS | 0.31 |
| IL-1RN | 2073 C/T Intron2 | 2q12-q14 | C/T | NS | 0.30 |
| IL-2 | rs2069763 | 4q26 | G/T | NS | 0.36 |
| IL-2 | rs2069762 | 4q26 | G/T | NS | 0.27 |
| IL-4R | rs1801275 | 16p12 | A/G | NS | 0.20 |
| IL-7R | rs11567685 | 5p13 | C/T | NS | 0.25 |
| IL-7R | rs7718919 | 5p13 | G/T | NS | 0.13 |
| IL-7R | rs11567686 | 5p13 | A/G | NS | 0.34 |
| MC1R | rs1805009 | 16q24 | C/G | NA | 0.01 ^b |
| MC1R | rs1805006 | 16q24 | A/C | NA | 0.00 ^b |
| MEFV | rs28940577 | 16p13.3 | A | NA | 0.00 ^b |
| MGC33887 | rs987931 | 17q24 | G/T | NS | 0.32 |
| MOG | rs3130250 | 6p22 | A/G | NS | 0.19 |
| MOG | rs3130253 | 6p22 | A/G | NS | 0.12 |
| NDUFA7 | rs2288414 | 19p13.2 | C/G | NA | 0.03 ^b |
| NDUFA7 | rs561 | 19p13.2 | A/G | NS | 0.21 |
| NDUFS5 | rs2889683 | 1p34.2 | C/T | NS | 0.31 |
| NDUFS5 | rs6981 | 1p34.2 | A/G | NA | 0.04 ^b |
| NDUFS7 | rs2074897 | 19p13.3 | A/G | <i>P</i> < .01 | 0.47 |
| NOS2A | rs1137933 | 17q11.2 | A/G | NS | 0.25 |
| NOS2A | rs2779248 | 17q11.2 | C/T | NS | 0.39 |
| NOTCH4 | rs367398 | 6p21.3 | A/G | NS | 0.16 |
| PD-1 | rs11568821 | 2q37 | G/A | NS | 0.11 |
| PITPNC1 | rs1318 | 17q24 | A/G | NS | 0.21 |
| PITPNC1 | rs2365403 | 17q24 | C/G | NS | 0.18 |
| PNMT | rs876493 | 17q11-q23 | A/G | NS | 0.39 |
| PRKCA | rs7220007 | 17q24 | A/G | NS | 0.49 |
| PRKCA | rs887797 | 17q24 | C/T | NS | 0.30 |
| PRKCA | rs2078153 | 17q24 | C/G | NS | 0.23 |
| PRKCA | rs3890137 | 17q24 | A/G | NS | 0.37 |
| PTPN22 | rs2476601 | 1p13 | A/G | NS | 0.11 |

| Gene | rs-nr | Chromosome | Polymorphism | HWE ^a | MAF |
|--------------|------------------------|-------------|--------------|------------------|-------------------|
| <i>PTPRC</i> | rs17612648 | 1q31 | C/G | NS | 0.03 ^b |
| <i>PTPRC</i> | rs4915154 | 1q31 | A/G | NS | 0.00 ^b |
| <i>CCL5</i> | rs2280788 | 17q11.2-q12 | C/G | NS | 0.02 ^b |
| <i>CCL5</i> | rs2107538 | 17q11.2-q12 | C/T | NS | 0.18 |
| <i>Spp1</i> | rs1126616 | 4q21 | C/T | NS | 0.23 |
| <i>Spp1</i> | rs1126772 | 4q21 | A/G | NS | 0.18 |
| <i>Spp1</i> | rs2853744 | 4q21 | G/T | NS | 0.05 |
| <i>Spp1</i> | rs9138 | 4q21 | A/C | NS | 0.24 |
| <i>Spp1</i> | rs4754 | 4q21 | C/T | NS | 0.24 |
| <i>TNF</i> | rs1800629 | 6p21.3 | A/G | NS | 0.17 |
| <i>TRAIL</i> | rs1131568 ^c | 3q26 | C/T | NS | 0.32 |
| <i>UCP2</i> | rs659366 | 11q13 | C/T | NS | 0.37 |
| <i>VDR</i> | rs1544410 | 12q13 | A/G | NS | 0.48 |
| <i>VDR</i> | rs731236 | 12q13 | A/G | NS | 0.48 |

Abbreviations: *ADAMTS14*, a disintegrin and metalloproteinase with thrombospondin motif, type 1 motif 14; *Apo I/Fas*, tumor necrosis factor (TNF) receptor superfamily, member 6; *BTNL2*, butyrophilinlike 2; *CIITA*, class II, major histocompatibility complex, transactivator; *CACNG4*, calcium channel, voltage-dependent, gamma subunit 4; *CCL5*, chemokine (C-C motif) ligand 5; *CCR5*, chemokine (C-C motif) receptor 5; *CNTF*, ciliary neurotrophic factor; *CRYAB*, α -B crystallin; *CTLA4*, cytotoxic T-lymphocyte-associated protein 4; *EBF1*, early B-cell factor 1; *GABBR1*, gamma-aminobutyric acid (GABA) B receptor 1; *HELZ*, helicase with zinc finger; *HLA*, human leukocyte antigen; *HLA-DRA*, human leukocyte antigen DR alpha; HWE, Hardy-Weinberg Equilibrium in our sample; *IFNAR1*, interferon (alpha, beta, and omega) receptor 1; *IFNGR2*, interferon gamma receptor 2 (interferon gamma transducer 1); *IKBL*, inhibitory κ -B-like gene; *IL-10*, interleukin 10; *IL-1 β* , interleukin 1 β ; *IL-1RN*, interleukin 1 receptor antagonist; *IL-2*, interleukin 2; *IL-4R*, interleukin 4 receptor; *IL-7R*, interleukin 7 receptor; *MAF*, minor allele frequency in our sample; *MC1R*, melanocortin 1 receptor; *MEFV*, Mediterranean fever; *MGC33887*, coiled-coil domain containing 46; *MOG*, myelin oligodendrocyte glycoprotein; NA, not applicable; *NDUFA7*, NADH dehydrogenase (ubiquinone) 1 α subcomplex, 7; *NDUFS5*, NADH dehydrogenase (ubiquinone) Fe-S protein 5; *NDUFS7*, NADH dehydrogenase (ubiquinone) Fe-S protein 7; *NOS2A*, nitric oxide synthase 2; *NOTCH4*, Notch homolog 4; NS, not significant; *PD-1*, programmed cell death 1; *PITPNC1*, phosphatidylinositol transfer protein, cytoplasmic 1; *PNMT*, phenylethanolamine *N*-methyltransferase; *PRKCA*, protein kinase C, alpha; *PTPN22*, protein tyrosine phosphatase, nonreceptor type 22; *PTPRC*, protein tyrosine phosphatase, receptor type C; rs-nr, RefSNP accession identification; *Spp1*, secreted phosphoprotein 1 (osteopontin); *TRAIL*, TNF-related apoptosis inducing ligand; *UCP2*, uncoupling protein 2; *VDR*, vitamin D (1,25-dihydroxyvitamin D3) receptor.

^a *P* value of HWE significant. *P* < .01 indicates deviation from HWE.

^b Excluded due to minor allele frequency *P* < .05.

^c Previous rs-number: rs9880164.