

Genetic Appendix

Carter Tinkler

ALLERGY					
SNP	rsID	Minor Allele	Genotype	Phenotype	
C11ORF30	rs2155219	G	TT	-/-	
HLA-DQB1	rs7775228	C	TT	-/-	
CLEFT LIP/CLEFT PALATE					
SNP	rsID	Minor Allele	Genotype	Phenotype	
IRF6	rs861020	A	GG	-/-	
INTERGENIC	rs987525	A	CC	-/-	
CLOTTING FACTORS					
SNP	rsID	Minor Allele	Genotype	Phenotype	
CETP	rs1800775	C	AC	+/-	Increased risk of high cholesterol
CYP4V2	rs13146272	C	AC	+/-	Increased sensitivity to dietary fats
F10	rs3211719	G	AG	+/-	Increased risk of deep vein thrombosis
F11	rs2036914	T	CC	-/-	
F11	rs2289252	T	CT	+/-	Increased risk of deep vein thrombosis
F12	rs1801020	A	GG	-/-	
F12	rs2731672	T	CC	-/-	
F5	rs6025	T	CC	-/-	
F7	rs6046	A	GG	-/-	
F9	rs6048	G	AA	-/-	
GP6	rs1613662	G	AA	-/-	
ITGB3	rs5918	C	CT	+/-	Increased risk of clotting disorders
NR1H2	rs1523127	C	CC	+/+	Increased risk of clotting disorders
SERPINC1	rs2227589	T	CC	-/-	
DETOX					
SNP	rsID	Minor Allele	Genotype	Phenotype	
CTH	rs1021737	T	GG	-/-	
CYP1A1	rs1048943	C	TT	-/-	
CYP1A1	rs1799814	T	GT	+/-	Decreased estrogen metabolism
CYP1A1	rs4986883	C	TT	-/-	
CYP1A2	rs762551	C	AA	-/-	Increased caffeine metabolism
CYP1B1	rs1056836	C	CG	+/-	Decreased estrogen metabolism
CYP1B1	rs1800440	C	TT	-/-	
CYP2A6	rs1801272	T	AA	-/-	
CYP2C19	rs12248560	T	CT	+/-	Increased testosterone to estrogen conversion
CYP2C9	rs1057910	C	AA	-/-	
CYP2C9	rs1799853	T	CT	+/-	Altered estrogen and drug metabolism
CYP2E1	rs2070676	G	CC	-/-	
CYP2E1	rs55897648	A	GG	-/-	
CYP2E1	rs6413419	A	GG	-/-	
CYP3A4	rs12721627	C	GG	-/-	
CYP3A4	rs2740574	C	TT	-/-	
CYP3A4	rs55785340	G	AA	-/-	
GPX3	rs8177412	C	CT	+/-	Decreased metal detoxification
GSTM1	rs2239892	G	AA	-/-	
GSTP1	rs1138272	T	CC	-/-	
GSTP1	rs1695	G	AA	-/-	
NAT1	rs4986782	A	GG	-/-	
NAT2	rs1208	G	AG	+/-	
NAT2	rs1799930	A	GG	-/-	
NAT2	rs1799931	A	AG	+/-	
NAT2	rs1801279	A	GG	-/-	
NAT2	rs1801280	C	CT	+/-	
GLUTEN INTOLERANCE					
SNP	rsID	Minor Allele	Genotype	Phenotype	
HLA-DQA1	rs2187668	T	CC	-/-	
HLA-DQA2	rs2858331	G	AA	-/-	

IGA				
SNP	rsID	Minor Allele	Genotype	Phenotype
CFH	rs6677604	A	AG	+/-
HLA-DPB2	rs1883414	A	GG	-/-
HLA-DQA2	rs9275224	A	AG	+/-
HLA-DRB1	rs9271366	G	AA	-/-
HORMAD2	rs2412971	G	AG	+/-
IFIH1	rs1990760	T	CT	+/-
IGF1R	rs2229765	A	AA	+/+
IRF5	rs4728142	A	GG	-/-
MTC03P1	rs2856717	A	AG	+/-
PSMB8	rs9357155	A	GG	-/-
TRAF1	rs3761847	G	AG	+/-
IGE				
SNP	rsID	Minor Allele	Genotype	Phenotype
ACKR1	rs2814778	C	TT	-/-
FCER1A	rs2251746	C	CC	+/+
FCER1A	rs2427837	A	AA	+/+
FCER1A	rs2494262	A	AA	+/+
IL5	rs2069812	G	AG	+/-
RAD50	rs17772565	T	CC	-/-
RAD50	rs17772583	G	AG	+/-
RAD50	rs2040704	G	AA	-/-
RAG1	rs3740955	A	AG	+/-
IGG				
SNP	rsID	Minor Allele	Genotype	Phenotype
FCGR2A	rs1801274	G	AG	+/-
GSTM3	rs7483	T	CT	+/-
MUC21	rs1634731	G	AA	-/-
TNFRSF13B	rs4792800	G	AG	+/-
METHYLATION				
SNP	rsID	Minor Allele	Genotype	Phenotype
ACAT1	rs3741049	A	AG	+/-
ACE	rs4343	G	AG	+/-
AGT	rs699	A	AG	+/-
BHMT	rs3733890	A	GG	-/-
BHMT-02	rs567754	T	CT	+/-
BHMT-04	rs617219	C	AC	+/-
C10RF167	rs4846048	G	AA	-/-
CBS	rs2851391	T	TT	+/+
CBS	rs4920037	A	AG	+/-
CBS A360A	rs1801181	A	GG	-/-
CBS C699T	rs234706	A	AG	+/-
CBS N212N	rs2298758	A	GG	-/-
CLCN6	rs13306560	T	CC	-/-
CLCN6	rs13306561	G	AA	-/-
CLCN6	rs3737964	T	CC	-/-
COMT	rs6269	G	AG	+/-
COMT H62H	rs4633	T	CT	+/-
COMT P199P	rs769224	A	AG	+/-
COMT V158M	rs4680	A	AG	+/-
DAO	rs2070586	A	AG	+/-
DAO	rs2111902	G	GG	+/+
FOLR1	rs2071010	A	GG	-/-
FOLR2	rs651933	G	AG	+/-
FOLR3	rs7925545	G	AA	-/-
FOLR3	rs7926875	A	CC	-/-
FUT2	rs492602	G	AG	+/-
FUT2	rs601338	A	AG	+/-
FUT2	rs602662	A	AG	+/-
G6PD	rs1050828	T	CC	-/-
G6PD	rs1050829	C	TT	-/-

GAD1	rs2241165	C	CT	+/-	Increased risk of anxiety/altered gut function
GAD1	rs3749034	A	AG	+/-	Increased risk of anxiety/altered gut function
GAD1	rs3791850	A	AG	+/-	Increased risk of anxiety/altered gut function
GAD1	rs3828275	T	CC	-/-	
GAD1	rs701492	T	CT	+/-	Increased risk of anxiety/altered gut function
GAD1	rs769395	G	AG	+/-	Increased risk of anxiety/altered gut function
GAD2	rs1805398	T	GT	+/-	Increased risk of anxiety/altered gut function
GAMT	rs17851582	A	GG	-/-	
MAOA	rs6323	G	TT	-/-	Decreased breakdown of Serotonin/Adrenaline
MAOB	rs1799836	C	TT	-/-	Decreased breakdown of Dopamine/Histamine
MTHFD1	rs2236225	A	AA	+/+	Increased need for choline
MTHFD1L	rs11754661	A	GG	-/-	
MTHFD1L	rs17349743	C	CT	+/-	Increased need for choline
MTHFD1L	rs803422	A	AG	+/-	Increased need for choline
MTHFR	rs1476413	T	CC	-/-	
MTHFR	rs17037390	A	GG	-/-	
MTHFR	rs17037396	T	CC	-/-	
MTHFR	rs17367504	G	AA	-/-	
MTHFR	rs2066470	A	GG	-/-	
MTHFR	rs2274976	T	CC	-/-	
MTHFR A1298C	rs1801131	G	TT	-/-	
MTHFR C677T	rs1801133	A	AG	+/-	Decreased methylaton potential
MTHFS	rs6495446	T	CC	-/-	
MTR A2756G	rs1805087	G	AG	+/-	Increased need for B12
MTRR	rs10380	T	CC	-/-	
MTRR	rs1801394	G	GG	+/+	Increased need for B12
MTRR A664A	rs1802059	A	AA	+/+	Increased need for B12
MTRR K350A	rs162036	G	AA	-/-	
MTRR R415T	rs2287780	T	CC	-/-	
NOS2	rs2297518	A	GG	-/-	
NOS3	rs1800779	G	GG	+/+	Decreased antioxidant potential
NOS3	rs2070744	C	CC	+/+	Decreased antioxidant potential
NOS3	rs7830	T	GT	+/-	Decreased antioxidant potential
PEMT	rs4244593	T	GG	-/-	
SOD2	rs2758331	A	AC	+/-	Decreased antioxidant potential
SOD2	rs4880	G	AG	+/-	Decreased antioxidant potential
SOD3	rs2855262	C	CT	+/-	Decreased antioxidant potential
TCN1	rs526934	G	AA	-/-	
TCN2	rs1801198	G	CC	-/-	
TYMSOS	rs502396	T	TT	+/+	Increased need for folate
VDR BSM	rs1544410	T	TT	+/+	Increased need for Vitamin D
VDR TAQ	rs731236	G	GG	+/+	Increased need for Vitamin D

MITOCHONDRIAL FUNCTION					
SNP	rsID	Minor Allele	Genotype	Phenotype	
ATP5C1	rs1244414	T	CC	-/-	
ATP5C1	rs1244422	T	CT	+/-	Increased need for mitochondrial support especially B vitamins
ATP5C1	rs2778475	A	AG	+/-	Increased need for mitochondrial support especially B vitamins
CCL2	rs1024611	G	AA	-/-	
COX6C	rs4626565	C	TT	-/-	
NDUFS7	rs1142530	T	CT	+/-	Increased need for mitochondrial support especially CoQ10
NDUFS7	rs11666067	A	AC	+/-	Increased need for mitochondrial support especially CoQ10
NDUFS7	rs7254913	G	AG	+/-	Increased need for mitochondrial support especially CoQ10
NDUFS8	rs1104739	C	AC	+/-	Increased need for mitochondrial support especially CoQ10
NDUFS8	rs2075626	C	CT	+/-	
NDUFS8	rs4147776	C	AA	-/-	
SLC19A1	rs1051266	C	CC	+/+	Increased need for folate
UQCRC2	rs6497563	C	TT	-/-	

OTHER IMMUNE FACTORS

SNP	rsID	Minor Allele	Genotype	Phenotype
ADD1	rs4961	T	GG	-/-
ATG16L1	rs10210302	T	TT	+/+
HLA-DRB1	rs660895	G	AG	+/-
IL13	rs20541	A	GG	-/-
IL4R	rs1801275	G	AA	-/-
KIAA1109	rs6822844	T	GG	-/-
MEFV	rs11466023	A	GG	-/-
MEFV	rs3743930	G	CC	-/-
TNF	rs1800629	A	GG	-/-
TNF	rs361525	A	GG	-/-
TYR	rs28940879	C	GG	-/-

Increased risk of inflammation and immune dysfunction

SULFOTRANSFERASE

SNP	rsID	Minor Allele	Genotype	Phenotype
SULT2A1	rs2547231	C	AC	+/-
SULT2A1	rs2910393	T	CT	+/-
SULT2A1	rs4149449	T	CC	-/-
SULT2A1	rs4149452	T	CC	-/-

Decreased phase 2 detoxification

Decreased phase 2 detoxification

THYROID

SNP	rsID	Minor Allele	Genotype	Phenotype
CTLA4	rs231775	G	AA	-/-
FOXE1	rs10984009	A	GG	-/-

BIOHACKING DIET

SNP	rsID	Minor Allele	Genotype	Phenotype
ABCG8	rs4299376	G	TT	-/-
ABCG8	rs6544713	T	CC	-/-
ACE DEL16	rs4343	G	AG	+/-
ADIPOQ	rs17366568	A	AG	+/-
ADRB2	rs1042713	A	AG	+/-
ADRB2	rs1042714	G	CC	-/-
ADRB3	rs4994	G	AA	-/-
ANKK1	rs1800497	A	AG	+/-
APOA2	rs5082	G	AG	+/-
APOA5	rs662799	G	AA	-/-
APOB	rs1367117	A	GG	-/-
APOE	rs7412	T	CC	-/-
FABP2	rs11724758	A	AG	+/-
FTO	rs1121980	A	AG	+/-
FTO	rs9939609	A	AT	+/-
FTO	rs9940128	A	AG	+/-
GCKR	rs780094	T	CC	-/-
IGF1	rs35767	A	GG	-/-
MCM6	rs182549	T	CT	+/-
MCM6	rs4988235	A	AG	+/-
MIR4761 (COMT V1)	rs4680	A	AG	+/-
PPARD	rs2267668	G	AA	-/-
PPARG	rs1801282	G	CC	-/-
PPARGC1A	rs8192678	T	CT	+/-
SLC2A2	rs5400	A	GG	-/-
SNP?	rs17782313	C	TT	-/-
SNP?	rs2943634	A	AC	+/-
TAS2R38	rs1726866	A	AG	+/-
TNF	rs1800629	A	GG	-/-

Decreased blood sugar regulation

Increased risk of Diabetes, Obesity and Athsma

Increased dopamine due to lack of inhibitory receptor activity

Increased risk of high cholesterol

Sensitivity to polyunsaturated plant oils

Decreased blood sugar regulation

Decreased blood sugar regulation

Decreased blood sugar regulation

Likely lactose tolerant

Likely lactose tolerant

Decreased blood sugar regulation

Increased risk of stroke and CVD

Increased desire for sweet foods

IRON DISREGULATION				
SNP	rsID	Minor Allele	Genotype	Phenotype
G6PD	rs1050828	T	CC	-/-
G6PD	rs1050829	C	TT	-/-
G6PD	rs2230037	A	AA	+/+
G6PD	rs5030868	A	GG	-/-
G6PD	rs72554664	C	CC	+/+
HFE	rs1799945	G	CC	-/-
HFE	rs1800562	A	GG	-/-
HFE	rs2071303	C	TT	-/-
HFE	rs2794719	G	GT	+/-
HFE	rs9366637	T	CC	-/-
SNP?	rs235756	G	GG	+/+
SOD2	rs4880	G	AG	+/-

No genetic hemochromatosis

No genetic hemochromatosis

GLUCURONIDATION				
SNP	rsID	Minor Allele	Genotype	Phenotype
UGT1A10	rs4148323	A	GG	-/-
UGT1A7	rs887829	T	CT	+/-
UGT1A8	rs4148325	T	CT	+/-
UGT1A8	rs6742078	T	GT	+/-

Decreased phase 2 detoxification

Decreased phase 2 detoxification

Decreased phase 2 detoxification

DOPAMINE BETA HYDROXYLASE				
SNP	rsID	Minor Allele	Genotype	Phenotype
DBH	rs1108580	G	AG	+/-
DBH	rs1611115	T	CT	+/-
DBH	rs4531	T	GG	-/-
DBH	rs77905	A	AG	+/-

Decreased conversion from Dopamine to Adrenaline

Decreased conversion from Dopamine to Adrenaline

Decreased conversion from Dopamine to Adrenaline

PON1				
SNP	rsID	Minor Allele	Genotype	Phenotype
PON1	rs662	T	TT	+/+

Decreased ability to detoxify Organophosphates