RUNNING HEAD: Candidate gene analysis for pig meat quality

An association analysis for 14 candidate genes mapping to meat quality QTL in a Duroc pig population reveals that the *ATP1A2* genotype is highly associated with muscle electric conductivity

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Summary

In previous genome-wide association studies carried out in a Duroc commercial line (Lipgen population), we detected on pig chromosomes 3, 4 and 14 several QTL for *gluteus medius* muscle redness (GM a*), electric conductivity in the *longissimus dorsi* muscle (LD CE) and vaccenic acid content in the LD muscle (LD C18:1 n-7), respectively. We have genotyped, in the Lipgen population, 19 single nucleotide polymorphisms (SNP) mapping to 14 genes located within these three QTL. Subsequently, association analyses have been performed. After correction for multiple testing, two SNPs in the *TGFBRAP1* (rs321173745) and *SELENOI* (rs330820437) genes were associated with GM a*, while *ACADSB* (rs81449951) and *GPR26* (rs343087568) genotypes displayed significant associations with LD vaccenic content. Moreover, the polymorphism of the *ATP1A2* (rs344748241), *ATP8B2* (rs81382410) and *CREB3L4* (rs321278469 and rs330133789) genes showed significant associations with LD CE. We made a second round of association analyses including the SNPs mentioned above as well as other SNPs located in the chromosomes to which they map to. After performing a correction for multiple testing, the only association that remained significant at the chromosome-wide level was that between the *ATP1A2* genotype and LD CE. From a functional point of view, this association is meaningful because this locus encodes a subunit of the Na^+/K^+-ATPase responsible of maintaining an electrochemical gradient across the plasma membrane.

Keywords: Pig, single nucleotide polymorphism, meat quality, Na^+/K^+-ATPase.

Meat quality traits are of paramount importance for the pig industry because they determine, to a great extent, consumer acceptance and financial profit. Once pigs are slaughtered, there is a decline of the pH of the skeletal muscle due to the production of lactic acid through anaerobic glycolysis (Rosenvold & Andersen 2003). The rate of muscle acidification has a strong effect on meat color and water-holding capacity. In this way, a low ultimate pH (5.4-5.3) is associated with pale, soft and exudative (PSE) meat as well as with an increased electrical conductivity (CE) and elevated drip and cooking losses (Lee *et al.* 2000; Rosenvold & Andersen 2003). In contrast, a high ultimate pH (6.3 or higher) results in dark, firm and dry (DFD) meat with a high water-holding capacity and a lower CE (Lee *et al.* 2000; Kim *et al.* 2016). Adverse effects on meat quality are influenced by both genetic and environmental factors. Recessive and dominant genotypes in the porcine ryanodine receptor 1 (*RYR1*) and the protein kinase AMP-activated non-catalytic subunit γ_3 (*PRKAG3*) genes, respectively, are strong predisposing factors to the occurrence of PSE meats (Fujii *et al.* 1991; Milan *et al.* 2000). On the other hand, there are multiple factors related with pig management and transportation (pre-slaughter stress), stunning method at slaughter, carcass chilling and pelvic suspension of carcasses that influence pork quality (Rosenvold & Andersen 2003). Another important parameter that determines meat quality is intramuscular fat (IMF) composition. In this regard, it is well known that fatty acid composition can have important consequences on the oxidative stability of meat during processing and retail display as well as on fat firmness (Wood *et al.* 2008).

In previous genome-wide association studies, we identified several genomic regions containing quantitative trait loci (QTL) for meat Minolta a* value (redness), CE (González-Prendes *et al.* 2017) and IMF composition (González-Prendes *et al.* 2019) traits measured in the *longissimus dorsi* (LD) and *gluteus medius* (GM) muscle samples of 350 Duroc

barrows (Lipgen population). Details about the rearing of the pigs can be found in Gallardo *et al*. (2009), while a thorough description of QTL mapping methods is reported in González-Prendes *et al*. (2017). The measurement of CE was done 24 hours after slaughter by using a Pork Quality Meter (Intek GmbH), while Minolta a* value was determined with a Minolta Chroma-Meter CR-200 (Konica Minolta) equipment at the same time point. Muscle fatty acid composition was measured as previously described by Quintanilla *et al*. (2011). In the current work, we have selected 14 candidate genes located within QTL regions for GM a* on SSC3, LD CE on SSC4, and LD vaccenic content on SSC14 (**Table 1**). These genes were: phosphorylase kinase catalytic subunit γ 1 (*PHKG1*), transforming growth factor ß receptor associated protein 1 (*TGFBRAP1*), selenoprotein I (*SELENOI*), hydroxyacil-CoA dehydrogenase trifunctional multienzyme (*HADHA*), coatomer protein complex subunit α (*COPA*), proliferation and apoptosis adaptor protein 15 (*PEA15*), calsequestrin 1 (*CASQ1*), ATPase Na^+/K^+ transporting α_2 subunit (*ATP1A2*), ATPase phospholipid transporting 8B2 (*ATP8B2*), cAMP responsive element binding protein 3 like 4 (*CREB3L4*), CREB regulated transcription coactivator 2 (*CRTC2*), acyl-CoA dehydrogenase short/branched chain (*ACADSB*), G protein-coupled receptor 26 (*GPR26*) and C-terminal binding protein 2 (*CTBP2*).

Genes were selected based on bibliographic information about their biological functions which suggested that they could be involved in the determinism of any of the three traits under study (GM a*, LD CE and LD C18:1 n-7). Based on available RNA-Seq (Cardoso *et al.* 2017) and whole-genome data (our unpublished results), we called 19 SNPs mapping to these 14 genes by using the GATK Best Practices workflow for SNP calling (https://software.broadinstitute.org/gatk/best-practices/workflow?id=11145) in accordance with protocols reported by Mármol-Sánchez et al. (2019). Nineteen SNPs were finally

selected because the SnpEff software predicted that they might have functional effects (Cingolani *et al.* 2012), as reported in **Supplementary Table 1**. The 19 selected SNPs (**Table 1**) were genotyped at the Servei Veterinari de Genètica Molecular of the Universitat Autònoma de Barcelona (http://sct.uab.cat/svgm/en) by using a QuantStudio 12K Flex Real-Time PCR System (ThermoFisher Scientific). Association analyses between SNPs and phenotypes were performed with the Genome-wide Efficient Mixed-Model Association (GEMMA) software (Zhou & Stephens 2012). The following statistical model was used:

$$\boldsymbol{y} = \boldsymbol{W\alpha} + \boldsymbol{x\delta} + \boldsymbol{u} + \boldsymbol{\varepsilon}$$

where $\boldsymbol{y}$ is the vector of phenotypic observations for every individual; $\boldsymbol{\alpha}$ corresponds to a vector including the intercept plus the fixed effects, *i.e.* batch effect with 4 categories (all traits), and farm origin effect with 3 categories (all traits). The $\boldsymbol{\alpha}$ vector also contains the regression coefficients of the following covariates: (1) Carcass weight at slaughterhouse for meat quality traits, and (2) IMF content in the LD muscle for LD fatty acid composition; $\boldsymbol{W}$ is the incidence matrix relating phenotypes with the corresponding effects; $\boldsymbol{x}$ is the vector of the genotypes corresponding to the set of selected polymorphisms; $\boldsymbol{\delta}$ is the allele substitution effect for each polymorphism; $\boldsymbol{u}$ is a vector of random individual effects with a n-dimensional multivariate normal distribution $MVN_n\,(0, \lambda\,\tau^{-1}\,K)$, where τ^{-1} is the variance of the residual errors, λ is the ratio between the two variance components and K is a known relatedness matrix derived from the SNPs; and $\boldsymbol{\varepsilon}$ is the vector of residual errors. Results were corrected for multiple testing by using the false discovery rate (FDR) method reported by Benjamini & Hochberg (1995). The correction for multiple testing took into account the number of candidate SNPs (2nd column of **Table 1**) mapping to each one of the

SSC3 GM a* (5 SNPs), SSC4 LD CE (11 SNPs) and SSC14 LD (C18:1) n-7 (3 SNPs) QTL.

Performance of association analyses with the methodology described above revealed the existence of several associations that remained significant even after correction for multiple testing. We have found, for instance, an association between GM Minolta a* value and missense mutations in the *TGFBRAP1* and *SELENOI* genes, which map to two different GM a* QTL on SSC3 (**Table 1**). The inactivation of the *TGFBRAP1* gene results in the suppression of aerobic glycolysis and increased levels of mitochondrial respiration and fatty acid oxidation (Yoshida *et al.* 2013), while *SELENOI* encodes a selenoprotein fundamental for the synthesis of phosphatidylethanolamine, a molecule with important effects on the oxidation of lipid membranes, oxidative phosphorylation and mitochondrial morphology (Tasseva *et al.* 2013; Poyton *et al.* 2016). We have also detected significant associations between LD CE and SNPs in the *ATP1A2*, *ATP8B2* and *CREB3L4* genes, which map to SSC4 LD CE QTL covering two regions spanning from 85.6 to 91 Mb and from 95.2 to 97.8 Mb. These findings are very suggestive because the *ATP1A2* gene, the one showing the most significant association, is preferentially expressed in the skeletal and heart muscle and brain and it encodes the α_2 subunit of the ion pump Na^+/K^+ ATPase (Clausen *et al.* 2017). Noteworthy, Na^+/K^+-ATPases provide the energy necessary for the maintenance of Na^+ and K^+ electrochemical gradients across the plasma membrane by hydrolyzing ATP (Clausen *et al.* 2017; Sampedro *et al.* 2018). These gradients are essential for the preservation of the resting membrane potential as well as for the generation of electrical impulses in the skeletal muscle and nervous system (Clausen *et al.* 2017; Sampedro *et al.* 2018). The ATP8B2 protein is also an ATPase with flippase activity

144 towards phosphatidyl choline, a key component of phospholipid membranes with important
145 effects on the functioning of the sarcoendoplasmic reticulum Ca^{2+}ATPase pumps (Shin &
146 Takatsu 2018; Fajardo *et al.* 2018), while CREB3L4 is a transmembrane bZip transcription
147 factor involved in the modulation of endoplasmic reticulum stress (Kim *et al.* 2014). Our
148 association analysis has also revealed the existence of significant associations between the
149 phenotypic variation of LD vaccenic (C18:1 n-7) content and SSC14 SNPs located in the
150 *ACADSB* gene, which catalyzes the oxidation of branched-chain fatty acids (Porta *et al.*
151 2019) and the *GPR26* gene, whose inactivation leads to hyperphagia, glucose intolerance,
152 hyperinsulinemia, dyslipidemia and obesity in mice (Chen *et al.* 2012).

153 We have made a second round of association analyses in which the SNPs that
154 previously showed evidence of statistical significance were compared against the whole
155 sets of the Porcine SNP60 BeadChip SNPs co-localizing to the same chromosome
156 (chromosome-wide analysis) *i.e.* 3,123 SNPs on SSC3, 3,899 SNPs on SSC4 and 4,203
157 SNPs on SSC14. These 11,225 SNPs were obtained from previously published porcine
158 SNP60 BeadChip data reported by González-Prendes *et al.* (2017). In this case, the
159 correction for multiple testing took into account the number of SNPs mentioned above for
160 each one of the three chromosomes under analysis, *i.e.* 3,128, 3,910 and 4,206 independent
161 tests were taken into consideration when performing association analyses for pig
162 chromosomes SSC3, SSC4 and SSC14. Interestingly, the rs344748241 SNP in the *ATP1A2*
163 gene was the only one that surpassed the chromosome-wide threshold of significance (q-
164 value < 0.05) (**Table 1**, **Figure 1**). Noteworthy, this SNP was not significant when we
165 made an association analysis at the genome-wide level (data not shown). Additionally, we
166 used the *LD* function of *gaston* R package (v1.5.5; Perdry & Dandine-Roulland 2019) to
167 evaluate the presence of linkage disequilibrium among the SNP markers that showed

significant associations with LD CE after correction for multiple testing at the chromosome-wide level (**Supplementary Figure 1**). The amount of linkage disequilibrium was expressed as r^2 in accordance with the definition of Hill & Robertson (1968). As shown in **Supplementary Figure 1**, we observed a high degree of linkage disequilibrium between the rs344748241 (*ATP1A2* gene) and the rs80782100 (*IGSF8* gene) markers. Noteworthy, the rs80782100 SNP, which maps to an intronic position within the immunoglobulin superfamily member 8 gene, displays the highest association with the LD CE phenotype, as described in González-Prendes *et al.* (2017).

As previously discussed, we consider that the *ATP1A2* gene is a strong positional and functional candidate to explain the CE QTL found on SSC4 because Na^+, K^+ ATPases are fundamental to induce an electrochemical gradient across the plasma membrane of cells (Suhail 2010), and their kinetics are modulated by the extracellular pH (Salonikidis *et al.* 2000), a parameter which also displays strong effects on muscle electrical conductivity. In pigs, the *ATP1A2* gene has been sequenced (Henriksen *et al.* 2013) and its polymorphisms have been associated with fat cut percentage (Fontanesi *et al.* 2012). A next step would be to re-sequence the whole gene in Lipgen pigs with alternative genotypes (QQ vs qq) for the LD CE QTL on SSC4, to build a complete catalogue of SNPs with potential effects on protein activity and expression and to investigate their association with CE in the Lipgen population. Subsequently, functional tests should be applied to ascertain whether any of the mutations in the pig *ATP1A2* gene with highly significant *q*-values also have causal effects on muscle conductivity.

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Data availability

These 11,225 SNPs included in this study were obtained from published Porcine SNP60 BeadChip data reported by González-Prendes *et al.* (2017), which can be accessed at the Figshare public repository (https://figshare.com/s/ 2e636697009360986794).

Conflict of interest

The authors declare that they have no conflict of interest.

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329 **Table 1.** An association analysis between 19 SNPs mapping to 14 candidate genes and meat quality traits recorded in a Duroc pig

330 population (significant associations are shown in bold)[1].

Gene	SNP	Type	Trait	P-value	q-value	P-value*	q-value*	δ ± SE	A_1	MAF
PHKG1	rs697732005 (3:16.830 Mb)	Splice region variant (G/A)	**GM a***	0.88661	0.88661	0.68325	0.96577	-0.02 (0.142)	A	0.3443
TGFBRAP1	**rs321173745 (3:49.516 Mb)**	**Missense variant (A/G)**		**0.00361**	**0.00902**	**0.03108**	0.67220	0.549 (0.186)	G	0.1875
SELENOI	**rs330820437 (3:112.635 Mb)**	**Missense variant (A/G)**		**0.00039**	**0.00196**	**0.01307**	0.51778	0.643 (0.181)	G	0.1757
HADHA	rs81215086 (3:112.794 Mb)	Missense variant (G/A)		0.53993	0.67491	0.62966	0.96577	-0.102 (0.169)	A	0.2899
	rs344578723 (3:112.796 Mb)	Missense variant (G/A)		0.53466	0.67491	0.67980	0.96577	-0.104 (0.169)	A	0.2866
COPA	rs340853721 (4:90.163 Mb)	Splice region variant (T/C)	**LD CE**	0.90735	0.95684	0.79005	0.99942	0.014 (0.091)	T	0.4351
	rs333099339 (4:90.183 Mb)	Splice region variant (T/C)		0.87813	0.95684	0.88586	0.99942	0.017 (0.090)	T	0.4381
	rs80949931 (4:90.186 Mb)	Missense variant (A/G)		0.95684	0.95684	0.68990	0.99942	-0.002 (0.091)	A	0.4335
PEA15	rs329681990 (4:90.266 Mb)	Splice region variant (G/A)		0.85666	0.95684	0.58021	0.99942	-0.014 (0.091)	G	0.433
CASQ1	rs334946278 (4:90.280 Mb)	Splice region variant (G/A)		0.95267	0.95684	0.92240	0.99942	0.005 (0.104)	A	0.1304
ATP1A2	**rs344748241 (4:90.356 Mb)**	**Splice region variant (G/A)**		**6.515E-06**	**7.167E-05**	**0.00006**	**0.02518**	-0.325 (0.066)	G	0.497
ATP8B2	**rs81382410**	**Splice region**		**0.00285**	**0.01565**	**0.00256**	0.21113	-0.233	T	0.3345

	(4:95.435 Mb)	**variant (T/C)**						(0.077)		
	rs329686514 (4:95.717 Mb)	Missense variant (C/T)		0.08043	0.17695	0.22592	0.97957	-0.155 (0.088)	T	0.3063
CREB3L4	**rs321278469 (4:95.717 Mb)**	**Missense variant (C/A)**		**0.00639**	**0.01757**	**0.00554**	0.30475	-0.228 0.083)	C	0.3084
	rs330133789 (4:95.721 Mb)	**Missense variant (G/A)**		**0.00493**	**0.01757**	**0.01769**	0.57188	0.254 (0.075)	A	0.3373
CRTC2	rs330198768 (4:95.740 Mb)	Intron variant (C/T)		0.32931	0.60373	0.56310	0.99942	-0.083 (0.085)	T	0.3687
ACADSB	rs81449951 (14:132.588 Mb)	Missense variant (C/A)	**LD (C18:1) n-7**	**0.04036**	0.08073	**0.0424837**	0.8322423	0.093 (0.045)	A	0.2109
GPR26	**rs343087568 (14:133.182 Mb)**	**Splice region variant (A/G)**		**0.00333**	**0.01334**	0.1269422	0.9956111	-0.096 (0.032)	G	0.4632
CTBP2	rs339956077 (14:134.334 Mb)	Splice region variant (G/A)		0.88166	0.88166	0.1269422	0.9956111	0.007 (0.046)	A	0.2094

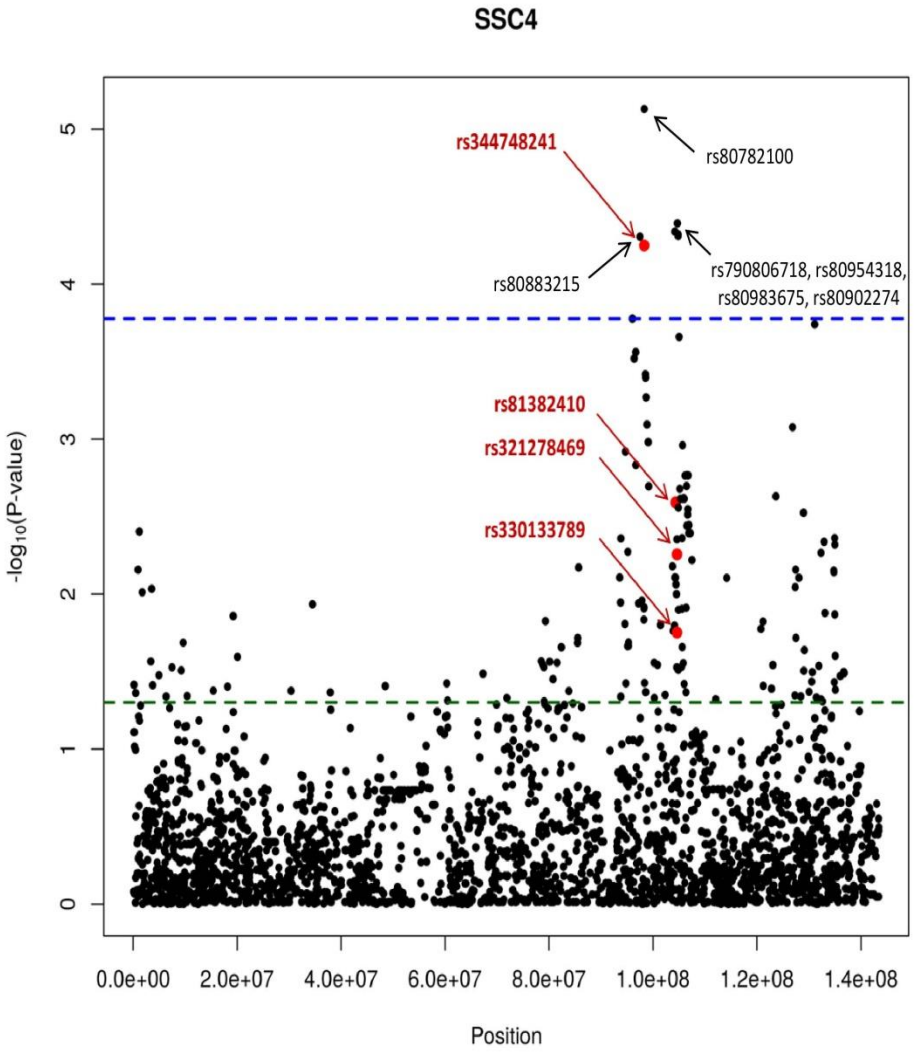
[1] The *P*-value and the *q*-value terms define the statistical significance of the association analysis before and after correcting for multiple testing with a fase discovery rate approach, respectively. The correction for multiple testing took into account the number of candidate SNPs (2nd column of Table 1) mapping to each one of the SSC3 GM a* (5 SNPs), SSC4 CE (11 SNPs) and SSC14 LD (C18:1) n-7 (3 SNPs) QTL. The *P*-value* and the *q-value** terms define the statistical significance of the chromosome-wide association analysis before and after correcting for multiple testing with a false discovery rate approach, respectively. In this case, the correction for multiple testing took into account the number of markers in the Porcine SNP60 BeadChip mapping to pig chromosomes SSC3 (3,123 SNPs), SSC4 (3,899 SNPs) and SSC14 (4,203 SNPs). Other terms that need to be defined are: δ, estimated allele substitution effect and its standard error (SE); A_1, minor allele; MAF, minor allele frequency; GM a*, Minolta a* value (redness) in the

339 *gluteus medius* muscle; LD CE, electric conductivity in the *longissimus dorsi* muscle; and LD (C18:1) n-7, vaccenic acid content in the

340 *longissimus dorsi* muscle.

LEGENDS TO FIGURES

Figure 1: Manhattan plot depicting associations between electrical conductivity in the *longissimus dorsi* muscle and the genotypes of markers in the *ATP1A2* (rs344748241), *ATP8B2* (rs81382410) and *CREB3L4* (rs321278469 and rs330133789) loci plus 3,899 additional SNPs mapping to pig chromosome 4 (SSC4). The positions of these three genes are SSC4: 90.292-90.371 Mb (*ATP1A2*), SSC4: 95.426-95.446 Mb (*ATP8B2*) and SSC4: 95.714-95.723 Mb (*CREB3L4*). The green line represents the nominal *P-value* of significance, while the blue line indicates the *P-value* of significance after correcting for multiple testing with a false discovery rate approach (*q*-value). The rs344748241 SNP in the *ATP1A2* gene is located 23 kb away from the peak of the LD CE QTL, *i.e.* ALGA0026686 (rs80782100; 4:90.378 Mb) SNP, as reported by González-Prendes *et al.* (2017).



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SUPPLEMENTARY DATA

Supplementary Table 1: Additional information about selected SNP and their potential impact and deleteriousness (SIFT).

Supplementary Figure 1: Graph depicting the magnitude of linkage disequilibrium among SNPs that showed significant associations with *longissimus dorsi* electric conductivity after correction for multiple testing at the chromosome-wide level. Here, the amount of linkage disequilibrium is expressed as r^2 as defined by Will & Robertson (1968) and such parameter was calculated with the *LD* function of *gaston* R package.

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SNP_ID	Allele	Consequence	Impact	Symbol	SIFT
rs697732005	A	splice_region_variant,intron_variant	LOW	*PHKG1*	-
rs321173745	G	missense_variant	MODERATE	*TGFBRAP1*	tolerated(0.22)
rs330820437	G	missense_variant	MODERATE	*SELENOI*	tolerated(1)
rs81215086	A	missense_variant	MODERATE	*HADHA*	deleterious(0.04)
rs344578723	A	missense_variant	MODERATE	*HADHA*	tolerated(0.81)
rs340853721	C	splice_region_variant,intron_variant	LOW	*COPA*	-
rs333099339	C	splice_region_variant,intron_variant	LOW	*COPA*	-
rs80949931	G	missense_variant	MODERATE	*COPA*	tolerated(0.15)
rs329681990	A	splice_region_variant,intron_variant	LOW	*PEA15*	-
rs334946278	A	splice_region_variant,synonymous_variant	LOW	*CASQ1*	-
rs344748241	A	splice_region_variant,synonymous_variant	LOW	*ATP1A2*	-
rs81382410	G	splice_region_variant,intron_variant	LOW	*ATP8B2*	-
rs329686514	T	missense_variant	MODERATE	*CREB3L4*	tolerated_low_confidence(0.05)
rs321278469	A	missense_variant	MODERATE	*CREB3L4*	tolerated(0.24)
rs330133789	A	missense_variant	MODERATE	*CREB3L4*	tolerated(0.32)
rs330198768	T	intron_variant	MODIFIER	*CRTC2*	-
rs81449951	A	missense_variant	MODERATE	*ACADSB*	tolerated(0.32)
rs343087568	G	splice_region_variant,intron_variant	LOW	*GPR26*	-
rs339956077	T	splice_region_variant,intron_variant	LOW	*CTBP2*	-

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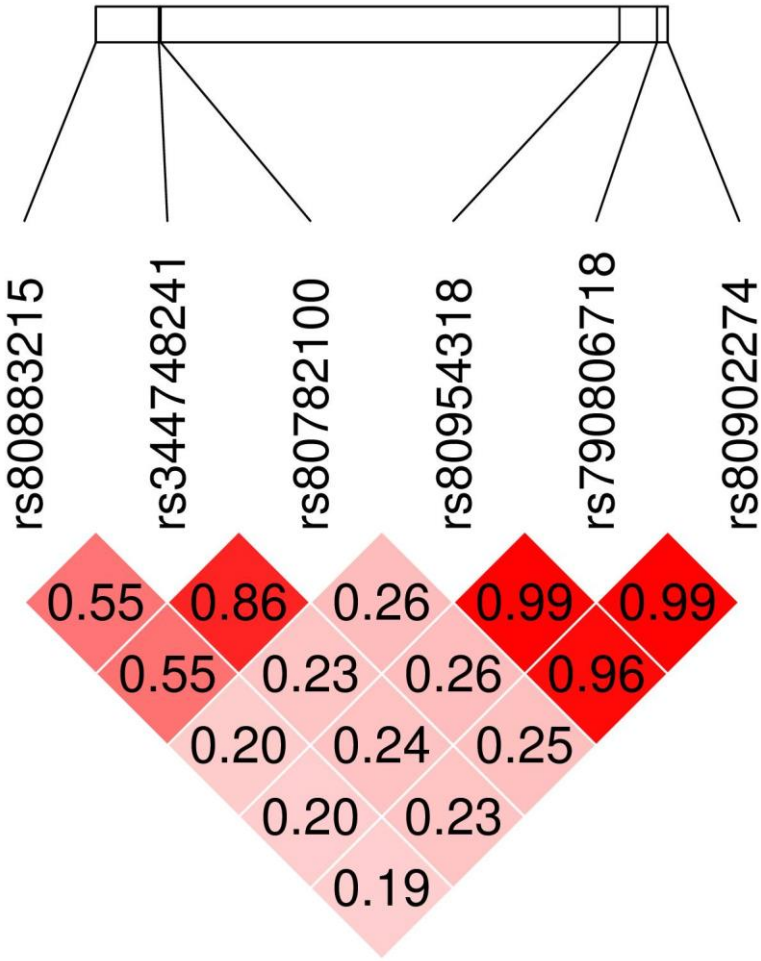
rs80883215
rs3447748241
rs807782100
rs809543118
rs79080671 8
rs809022 74
0.55
0.86
0.26
0.99
0.99
0.55
0.23
0.26
0.96
0.20
0.24
0.25
0.20
0.23
0.19

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