Polymorphisms and genetic effects of PRLR, MOGAT1, MINPP1 and CHUK genes on milk fatty acid traits in Chinese Holstein

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Abstract

Background: Our initial genome-wide association study (GWAS) identified 20 promising candidate genes for milk fatty acid (FA) traits in a Chinese Holstein population, including PRLR, MOGAT1, MINPP1 and CHUK genes. In this study, we performed whether they had significant genetic effects on milk FA traits in Chinese Holstein.

Results: We re-sequenced the entire exons and 3000 bp of the 5’ and 3’ flanking regions, and identified 11 single nucleotide polymorphisms (SNPs), containing four in PRLR, two in MOGAT1, two in MINPP1, and three in CHUK. The SNP-based association analyses showed that all the 11 SNPs were significantly associated with at least one milk FA trait (P = 0.0456 ~ < 0.0001), and none of them had association with C11:0, C13:0, C15:0 and C16:0 (P > 0.05). By the linkage disequilibrium (LD) analyses, we found two, one, one, and one haplotype blocks in PRLR, MOGAT1, MINPP1, and CHUK, respectively, and each haplotype block was significantly associated with at least one milk FA trait (P = 0.0456 ~ < 0.0001). Further, g.38949011G > A in PRLR, and g.111599360A > G and g.111601747 T > A in MOGAT1 were predicted to alter the transcription factor binding sites (TFBSs). A missense mutation, g.39115344G > A, could change the PRLR protein structure. The g.20966385C > G of CHUK varied the binding sequences for microRNAs. Therefore, we deduced the five SNPs as the potential functional mutations.

Conclusion: In summary, we first detected the genetic effects of PRLR, MOGAT1, MINPP1 and CHUK genes on milk FA traits, and researched the potential functional mutations. These data provided the basis for further investigation on function validation of the four genes in Chinese Holstein.

Keywords: PRLR, MOGAT1, MINPP1, CHUK, Genetic effects, Milk fatty acid traits, Chinese Holstein
(P = 2.25E-05); BTA-111275-no-rs within the \textit{MINPP1} (Multiple inositol-polyphosphate phosphatase 1) was significantly associated with C12:0 (P = 2.39E-05); and Haplmap46411-BTA-15820 within the \textit{CHUK} (Conserved helix-loop-helix ubiquitious kinase) had strong associations with C14:1 (P = 8.29E-06) and C14index (C14index = 100; P = 1.10E-08). Thus, the four genes were considered as the promising candidates for milk FA traits in Chinese Holstein.

\textit{PRLR} during the pregnancy may be associated with mammary development [13], and it can guide and maintain mammary epithelial cells for continuous lactation during milking of dairy cows [14]. \textit{MOGAT1} catalyzes the conversion of monoacylglycerols to diacylglycerols, the precursor of several physiologically important lipids such as phosphatidylethanolamine and triacylglycerol [15]. \textit{MOGAT1} expression is inversely correlated with lipolytic rates, and its suppression increases basal lipolytic activity [16]. \textit{MINPP1} is a stress protein and it can interact with the glucose-1-phosphate to induce apoptosis [17, 18] and human \textit{MINPP1} plays a role in differentiation and apoptosis [19]. \textit{CHUK} gene is also named IKKA gene, and IKKa is normally an activator of the transcription factor nuclear factor-kB, and it leads to potent activation of SREBP-mediated lipogenesis in the context of hepatitis C virus infection [20].

In this study, we further explored the polymorphisms and genetic effects of \textit{PRLR}, \textit{MOGAT1}, \textit{MINPP1} and \textit{CHUK} genes on milk FA traits, and searched the potential functional mutations.

\textbf{Results}

\textbf{SNPs identification}

In this study, we identified four, two, two, and three SNPs in \textit{PRLR}, \textit{MOGAT1}, \textit{MINPP1}, and \textit{CHUK} genes, respectively (Additional file 2: Table S2). For \textit{PRLR}, two SNPs (g.38948871C > T and g.38949011G > A) were in the 5′ flanking region, and two SNPs (g.39115344G > A and g.39115345T > C) were in the exon 4. In \textit{MOGAT1}, both g.111599360A > G and g.111601747T > A were in the 5′ flanking region. The g.9206582C > T and g.9207070A > G were observed in the 3′ UTR and the intron 5 of \textit{MINPP1}, respectively. For \textit{CHUK}, g.21008688G > T was in the 5′ flanking region, and two SNPs (g.20966385C > G and g.20966354C > T) were in the 3′ UTR. Out of these SNPs, g.39115344G > A in \textit{PRLR} was a missense mutation with the substitution of amino acid from serine to asparagine when the allele mutated from G to A.

\textbf{Associations between SNPs/haplotype blocks and milk fatty acids}

We performed the association analyses between the total 11 SNPs and 23 milk FA traits, and the results were shown in Additional file 3: Table S3. For \textit{PRLR} gene, g.38948871C > T was significantly associated with C6:0 (P = 0.0027) and UFA (P = 0.0364); g.38949011G > A was significantly associated with C8:0 (P = 0.0108); g.39115344G > A was significantly associated with C6:0, C8:0, C14:0, C17:1, C17index, SFA, and UFA (P = 0.022 ~ < 0.0001); and g.39115345T > C was significantly associated with C6:0, C8:0, C10:0, C14:0, C18index, C17index, SFA, and UFA (P = 0.0456 ~ < 0.0001). For \textit{MOGAT1} gene, the two SNPs (g.111599360A > G and g.111601747T > A) were significantly associated with C8:0 (P = 0.0001 and P < 0.0001), and g.111599360A > G was also significantly associated with C18:0 (P = 0.0058) and C17index (P = 0.0153). Both g.9206582C > T and g.9207070A > G of \textit{MINPP1} gene had significant associations with C6:0, C8:0, C10:0 and C17:0 (P = 0.0271 ~ < 0.0001), and g.9206582C > T was also significantly associated with C20:0 (P = 0.0436). The association analyses results of three SNPs (g.21008688G > T, g.20966385C > G and g.20966354C > T) of the \textit{CHUK} showed that they were all significantly associated with C8:0, C10:0, C12:0, C14:0, C16:1, C17:0, C17:1, C18:0, C16index, and C17index (P = 0.0423 ~ < 0.0001). In addition, g.21008688G > T was strongly associated with C18index, C20:0, and UFA (P = 0.0099 ~ 0.0014); g.20966385C > G was strongly associated with C14:1 (P = 0.0368) and C14index (P < 0.0001); and g.20966354C > T had significant associations with C6:0, C18:1cis-9, C18index, C20:0, SFA, UFA, and SFA/UFA (P = 0.0114 ~ < 0.0001). Correspondingly, the significances of additive (a), dominant (d) and substitution (s) effects for the 11 SNPs with milk FA traits were shown in Additional file 4: Table S4.

We found five haplotype blocks (D′ = 0.97 ~ 1.00; Fig. 1) in the study, including two in \textit{PRLR}, one in \textit{MOGAT1}, one in \textit{MINPP1}, and one in \textit{CHUK}. By haplotype-based association analyses (Additional file 5: Table S5), we found that the block 1 in \textit{PRLR} was significantly associated with C6:0 (P = 0.0389); block 2 in \textit{PRLR} was significantly associated with C6:0, C14:0, C17:1, C17index, SFA and UFA (P = 0.0392 ~ 0.0001); block 3 in \textit{MOGAT1} had significant associations with C8:0, C16:1, C18:0 and C17index (P = 0.0364 ~ < 0.0001); block 4 in \textit{MINPP1} had strong associations with C6:0, C8:0, C10:0, C18:1cis-9, C20:0, C17index, and UFA (P = 0.0306 ~ < 0.0001); and block 5 in \textit{CHUK} was strongly associated with C6:0, C8:0, C10:0, C14:0, C16:1, C17:0, C18:0, C18:1cis-9, C18index, C20:0, C14index, C16index, C17index, SFA, UFA, and SFA/UFA (P = 0.0256 ~ < 0.0001).

\textbf{Functional variation caused by SNPs of \textit{PRLR}, \textit{MOGAT1}, \textit{MINPP1} and \textit{CHUK}}

We used the Genomatix software to predicted the changes of TFBSs for all the five SNPs in 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that totally three SNPs (g.38949011G > A, g.111599360A > G and g.111601747C > T) were in the 5′ flanking region of the four genes, and found that total...
T > A) could alter the TFBSs (Fig. 2). The allele G of g.38949011G > A created a BS for the transcription factor MYBL1 (V-myb avian myeloblastosis viral oncogene homolog-like 1; MST = 0.8), and the allele A created the BSs for MEL1 (MDS1/EVI1-like gene 1; MST = 1.0) and LTSM (LTSM elements with 8 bp spacer; MST = 0.8). Alleles A and G of g.111599360A > G created the BSs for PAX6 (Pax-6 paired domain binding site; MST = 0.8) and TEAD4 (TEA domain family member 4; MST = 1.0), respectively. The allele A of g.111601747 T > A created three TFBSs, namely, AP4 (Activating enhancer binding protein 4; MST = 1.0), TCF21 (Transcription factor 21; MST = 1.0) and NEUROG (Neurogenin 1 and 3 binding sites; MST = 0.9).

We further used the SOPMA to predict the protein secondary structure variation for the missense mutation (g.38948871C > T and g.38949011G > A in PRLR; haplotype block 2 included g.39115344G > A and g.39115345 T > C in PRLR; haplotype block 3 included g.111599360A > G and g.111601747 T > A in MOGAT1; haplotype block 4 included g.9206582C > T and g.9207070A > G in MINPP1; and haplotype block 5 included g.20966385C > G and g.20966354C > T in CHUK. D’ = the value of D prime between the two loci

In addition, we used the position information of all the SNPs in the 3′ UTR to confirm that whether the SNPs altered the BSs for the microRNAs according to the Tar-gestScanHuman database, and found that the allele C of g.20966385C > G in the 3′ UTR of CHUK was located in the seed region for targeting the specific microRNAs, bta-miR-2392, bta-miR-2434/3602 and bta-miR-2395.

Discussion
The PRLR, MOGAT1, MINPP1 and CHUK genes were considered as the promising candidates for milk FA traits in our previous GWAS [12], and their polymorphisms and genetic effects were determined in this study. In recent years, the association between the gene polymorphism and milk production traits in dairy cattle has become a hotspot [21–23]. In this study, we found that each SNP of the PRLR, MOGAT1, MINPP1, and CHUK genes had significant association with at least one milk FA trait. Haplotypes formed by the SNPs have important implications for identifying associations with complex traits [24]. For the identified SNPs, we estimated their LD, and found that they were highly linked, which might...
be due to the properties of each SNP, such as the frequency and population history [25]. The haplotype-based association analyses showed that each haplotype block was also significantly associated with at least one milk FA trait.

From the KEGG database (https://www.genome.jp/kegg/pathway.html), we found that PRLR gene is involved in PI3K-AKT (ko04151) and Jak-STAT (ko04630) signaling pathways, which were identified to be associated with lipid metabolism [26, 27]. In this study, g.38949011G > A in the 5′ flanking region of PRLR altered the TFBSs, in which, the allele G created a BS for MYBL1, and the allele A created the BSs for MEL1 and LTSM. As we know, transcription factors (TFs) are the sequence-specific DNA-binding proteins that can regulate gene transcription, and genomic locations at which TFs interact with DNA are considered as TFBSs [28]. Some studies showed that TFs can play important roles in gene expression [29, 30]. MYBL1 as a DNA-binding TF can bind the mim-1 promoter and to activate its expression to regulate the oncogenesis [31], and MEL1 can stabilize the inactive Smad3-SKI complex on the promoter of TGF-β target genes to inhibit TGF-β signaling [32]. LTSM can enhance the transcriptional activity of the promoters in dependency of the distance from the transcription start site [33]. Hence, we deduced that these TFs might regulate PRLR gene expression to impact the components of milk FAs. In addition, we found a missense mutation in PRLR gene, g.39115344G > A, altered the protein secondary structure. Proteins are the utmost multi-purpose macromolecules (i.e., main catalysts, structural elements, signaling messengers and molecular machines of biological tissues) [34], which play a crucial function in many aspects of biological processes [35]. The protein structure plays a decisive role in function of protein, for example, a study reports that the β-turn is essential for the structure and function of proteins [36]. The prediction of protein structure from amino acid sequences is one of the most vital problems in molecular biology, and the fundamental elements of the protein secondary structure are α-helix, extended strand, β-turn, and coils [37]. In this study, α-helix, extended strand, β-turn, and random coil were all changed because of the mutation from G to A in g.39115344G > A, indicating that this missense mutation might affect the function of PRLR through changing the protein secondary structure.

MOGAT1 gene is involved in the glycerolipid metabolism (ko00561), and codes the MGAT (monoacylglycerol acyltransferases) enzyme, which is active in human liver and its activity can represent a viable target for pharmacologic intervention to treat nonalcoholic fatty liver disease [38]. A study reported that up-regulation of MOGAT1 gene can mediate hepatic steatosis by increasing intracellular diacylglycerol content [39]. In this study, g.111599360A > G in 5′ flanking region of MOGAT1 was predicted to create the TFBSs for PAX6 with the allele A and TEAD4 with the allele G. PAX6 can respectively down-regulate Sox3 and up-regulate Lhx9 in the Pax6-mutant cortex to exert its effects at the molecular level during murine forebrain neurogenesis [40]. TEAD4 can directly induce Myogenin, CDKN1A and Cavelin3 expression to promote myoblast differentiation [41]. Mutation of TAED4 at either site can decrease its occupancy on the promoter region of target genes, and largely impair the target gene transcription to inhibit the growth and colony formation of gastric cancer cell HGC-27 [42]. By association analyses, the cows with AA genotype of g.111599360A > G had higher C8:0 than that with GG, implicating that PAX6 might enhance the content C8:0 by
regulating the target gene MOGAT1. While, the TEAD4 might regulate MOGAT1 to reduce C8:0 by binding the G allele. In addition, g.111601747 T > A in the 5’ flanking region of MOGAT1, created the BSs for three TFs, namely, AP4, TCF21, and NEUROG. It is reported that AP4 can up-regulate the expression of LAPT4M4 to promote cell growth, migration, invasion, and cisplatin resistance in breast cancer [43]. AP4-geminin complex suppresses the precocious expression of target genes in fetal brain [44]. TCF21 plays a repression role for its target gene to affect the phenotypes [45, 46]. Over-expression of NEUROG can override the pluripotency-specific gene network and force human embryonic stem cells to differentiate into neurons [47]. The cows with AA genotype of g.111601747 T > A had significantly lower C8:0, suggesting that the three TFs (AP4, TCF21 and NEUROG) might work together to regulate the expression of MOGAT1 gene to finally decrease the C8:0.

MINPP1 is involved in the glycolysis/gluconeogenesis (ko00010) and inositol phosphate metabolism (ko00562) signaling pathways. Glycolysis can completely bypass 3-phosphoglycerate through that MINPP1 converts 2,3-bisphosphoglycerate to 2-phosphoglycerate, which activates the AMPK cascade [48]. Furthermore, AMPK can stimulate the fatty acid oxidation [49]. In this study, we identified g.9207070A > G in the intron 5 of the MINPP1 gene. In 1987, the first finding that introns can increase the gene expression was found in maize [50]. Introns include the regulatory regions, that can confer developmental and cell-specific expression of a gene reside [51]. The rs734553 located on the intron 7 of SLC2A9 in human is greatly related with serum uric acid of healthy individuals with normal renal function, thus it is powerful for prediction of chronic kidney disease progression [52]. Our association analyses showed that the cows with GG genotype of g.9207070A > G had significantly higher C6:0, and lower C17:0. We herein concluded that the intron mutation might be able to affect the milk FA traits that deserved the further validation.

CHUK gene is involved in the MAPK (ko04010), mTOR (ko04150), PI3K-Akt (ko04151), and Adipocytokine (ko04920) signaling pathways associated with lipid metabolism. In 2012, the associations between a SNP (rs11597086) of the human CHUK and lipid phenotype were identified [53]. In this study, the allele C of g.20966385C > G in the 3’ UTR of CHUK is in the seed sequences for targeting the microRNAs (bta-miR-2392, bta-miR-2434/3602 and bta-miR-2395). MicroRNA is a class of gene expression regulating factors and plays an important role in maintaining genome stability, regulating growth and development, and other biological processes [54, 55]. Some studies reported that microRNA regulates the fat metabolism and lipid metabolism disorders through the targeting genes, for example, miR-196 may be related to the gene expression of the homologous genes in subcutaneous adipose tissue and lipid distribution [56]. For the three microRNAs, bta-miR-2392, bta-miR-2434/3602 and bta-miR-2395, there have not been studied to reveal the regulatory function in cattle until now. In human, the miR-2392 can regulate its target gene MAML3 and WHSC1 to suppress metastasis and epithelial-mesenchymal transition in gastric cancer [57]. Our association analyses showed that the cows with CC genotype of g.20966385C > G had significantly higher C14:1, C17:0, C18:0 and C14index, and lower C14: 0, C16:1, C16index and C17index, indicating that the three microRNAs might regulate the expression of CHUK to affect the milk FA traits.

In dairy cattle breeding, the integration of DNA marker technology and genomics into the traditional evaluation system decreased generation interval and increased the selection accuracy, so that the cost of progeny testing was reduced [58]. Here, we found significant SNPs for milk FAs in dairy cattle, which could be used as the genetic markers for the genomic selection to improve the accuracy of selection and lower the breeding cost. This study provided the evidence for associations of PRLR, MOGAT1, MINPP1 and CHUK genes with milk FAs, and the in-depth study should be performed to verify the regulatory mechanism of these genes for milk FAs in dairy cattle by biotechnologies, such as RNA interference and gene editing.

Conclusions
Our findings first confirmed the genetic effects of PRLR, MOGAT1, MINPP1 and CHUK genes on milk FAs in Chinese Holstein. Three SNPs, g.38949011G > A of PRLR, g.111599360A > G and g.111601747 T > A of MOGAT1, might be the functional mutations by changing the promoter activities. In addition, the missense mutation in PRLR, g.39115344G > A, changed the protein secondary structure was suggested to be a critical mutation to affect the PRLR protein function. Furthermore, g.20966385C > G in 3’ UTR of CHUK varied the binding sequences for microRNAs that could regulate the gene expression of CHUK. This study provided the basis for further investigation on function validation of PRLR, MOGAT1, MINPP1 and CHUK genes, and the potential functional mutations might serve as genetic markers to apply to the breeding program for milk FA traits in Chinese Holstein.

Methods
Animals and phenotypic data
A total of 1065 Chinese Holstein cows belonging to 44 sire families were collected from 23 dairy farms of Beijing Sanyuanlvhe Dairy Farming Center (Beijing, China), where the standard performance testing for dairy herd improvement (DHI) has been regularly conducted
since 1999. These cows were full blood of Chinese Holstein breed that were originated from cross-breeding between the Chinese Yellow cattle and European Holstein, over the past 100 years, by continuous import of foreign Holstein bulls, semen and embryos, mainly from USA and a few from Canada and Europe, which were directly used to AI or used to cross Chinese Holstein cows through planned mating to generate breeding bulls [59]. All the cows were fed with the same regular total mixed ration composed of concentrated feed and coarse fodder. We obtained 50 ml milk samples during November to December of 2014 to measure the DHI, and then the dairy rations composed of concentrated feed and coarse fodder.

DNA extraction and SNP identification

We used the TIANamp Blood DNA kit (Tiangen, Beijing, China) and salt-out procedure to extract the genomic DNA form the blood samples of 1065 Chinese Holsteins and the semen samples of 44 sires, respectively. Then the quantity and quality of the genomic DNAs were measured by NanoDrop 2000 Spectrophotometer (Thermo Scientific, DE, USA) and gel electrophoresis, respectively.

We designed a total of 85 primers (Additional file 1: Table S1) in the entire exons with their partial adjacent introns, and 3000 bp of 5′ and 3′ flanking regions of the PRLR, MOGAT1, MINPP1, and CHUK genes using the Primer 3 (http://bioinfo.ut.ee/primer3-0.4.0/) according to their bovine genomic sequences (GenBank accession no. AC_000177.1, AC_000159.1, AC_000183.1 and AC_000183.1). The primers were synthesized in Beijing Genomics Institute (BGI, Beijing, China). The 44 semen DNAs were randomly divided into two DNA pools (22 DNA in each pool) for the PCR amplification, and the concentration of each DNA was 50 ng/μL. PCR conditions were as follows: initial denaturation at 94 °C for 5 min, followed by 35 cycles at 94 °C for 30s; 60 °C for 30s; 72 °C for 30s; and a final extension at 72 °C for 7 min. Afterwards, we sequenced the PCR products using ABI3730XL DNA analyzer (Applied Biosystems, Foster, CA, USA), and aligned the sequences with the bovine reference sequence (UMD 3.1) for identifying the potential SNPs.

Genotyping and linkage disequilibrium (LD) analyses

Genotypes of the identified SNPs were obtained from 1065 cows with Sequenom MassArray by matrix-assisted laser desorption/ionization time of flight mass spectrometry (MALDI-TOF MS, Sequenom MassARRAY, Bioyong Technologies Inc., HK). In addition, we used the Haploview 4.1 (Broad Institute, Cambridge, MA, USA) to analyze the LD extent among the identified SNPs.

Association analyses

The association analyses between each SNP/haplotype block and 23 milk FA traits were conducted with SAS9.2 (SAS Institute Inc., Cary, NC, USA) using the following model:

\[ Y_{ijklm} = \mu + G_i + h_j + I_k + a_l + b \times M_m + e_{ijklm} \]

For each trait, \( Y_{ijklm} \) was the phenotypic value of the 1065 cows; \( \mu \) was the overall mean; \( G_i \), \( h_j \), \( I_k \) and \( M_m \) were the fixed effects of the genotypes or haplotypes, farm (23), stage of lactation (4) and calving months (293), respectively; \( a_l \) was the random polygenic effect; \( b \) represented the regression coefficient of covariate \( M \); and \( e_{ijklm} \) was the random residual. We considered a significant association when the \( P \) was less than 0.05/\( N \), where \( N \) was the number of genotypes or haplotype combinations. In addition, we calculated the additive effect (a), dominant effect (d), and substitution effect (s) according to the formulas by Falconer & Mackay [62]:

- Additive effect (a) = \((AA - BB)/2\)
- Dominant effect (d) = \(AB - (AA + BB)/2\)
- Substitution effect (s) = \(a + d(q - p)\), where AA, AB and BB represent the square means of milk FA traits corresponding to the genotypes, \( p \) and \( q \) refer to the allele frequencies of corresponding loci.

Biological function prediction

We used the Genomatix software (http://www.genomatix.de/cgi-bin/welcome/welcome.pl?se=d1b5c9a9015b02bb3b1a806f9c03293f) to predict whether the SNPs in the 5′ flanking region of the PRLR, MOGAT1, MINPP1 and CHUK genes altered the transcription factor binding sites (TFBSs; matrix similarity threshold, MST > 0.8). Furthermore, we explored the changes of protein secondary structure for missense mutation by the SOPMA software (https://prabi.ibcp.fr/htm/site/web/services/secondaryStructurePrediction#SOPMA). For the SNPs in the 3′ untranslated region (UTR), we aligned them to the TargetScanHuman database (http://www.targetscan.org/vert_71/) for researching whether they changed the binding sites (BSs) of seed sequences with the microRNAs.
Additional files

**Additional file 1:** Table S1. Details of PCR primers of PRLR, MOGAT1, MINPP1, and CHUK genes. (XLSX 20 kb)

**Additional file 2:** Table S2. Information about the 11 identified SNPs. (XLSX 10 kb)

**Additional file 3:** Table S3. Associations of 11 SNPs of PRLR, MOGAT1, MINPP1, and CHUK genes with fatty acid traits in Chinese Holstein (LSM ± SE). (XLSX 26 kb)

**Additional file 4:** Table S4. Additive (a), dominant (d), and allele substitution (a) effects of 11 SNPs on milk fatty acid traits in Chinese Holstein cows. (XLSX 18 kb)

**Additional file 5:** Table S5. Associations of the haplotype blocks with milk fatty acid traits in Chinese Holstein (LSM ± SE). (XLSX 19 kb)

Abbreviations

- a: Additive effect; BS: Binding site; CHUK: Conserved helix-loop-helix ubiquitous kinase; d: Dominant effect; DH: Dairy herd improvement; FA: Fatty acids; GWAS: Genome-wide association study; LD: Linkage disequilibrium; MINPP1: Multiple inositol-polyphosphate phosphate 1; MOGAT1: Monoacylglycerol O-acyltransferase 1; MST: Matrix similarity threshold; PRLR: Prolactin receptor; SFA: Saturated fatty acids; SNP: Single nucleotide polymorphism; TF: Transcription factor; TFBS: Transcription factor binding site; UFA: Unsaturated fatty acids; UTR: Untranslated region; α: Substitution effect.

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Authors’ contributions

BH and DS conceived and designed the experiments, LL, ZM and YY prepared the milk, blood and semen samples, LS extracted the DNA for SNP identification and genotyping with the help of, XL, YL, and ZF, XL measured the milk fatty acids, LS analyzed the data, and the manuscript was prepared by LS, DS and BH. All authors read and approved the final manuscript.

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Availability of data and materials

All relevant data are available within the article and its additional files.

Ethics approval and consent to participate

All protocols for collection of the tissues of experimental individuals and phenotypic observations were reviewed and approved by the Institutional Animal Care and Use Committee (IACUC) at China Agricultural University. Tissue samples were collected specifically for this study following standard procedures with the full agreement of the Beijing Sanyuanlvhe Dairy Farming Center who owned the Holstein cows and bulls, respectively.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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