Molecular analysis of homeostatic iron regulator, transmembrane protease serine-6, and BTB domain-containing protein-9 variants and iron parameters in blood donors

Manal S. Fawzy1,2, Abeer Fakhr-Eldeen3,4, Baraah T. Abu AlSel5 and Eman A. Toraih6,7

1Department of Biochemistry, Faculty of Medicine, Northern Border University, Arar, Saudi Arabia; 2Department of Medical Biochemistry and Molecular Biology, Faculty of Medicine, Suez Canal University, Ismailia, Egypt; 3Department of Pathology, Faculty of Medicine, Northern Border University, Arar, Saudi Arabia; 4Department of Clinical Pathology, Faculty of Medicine, Sohag University, Sohag, Egypt; 5Department of Pathology, Faculty of Medicine, Northern Border University, Arar, Saudi Arabia; 6Department of Surgery, Tulane University, School of Medicine, New Orleans, Louisiana, U.S.A.; 7Genetics Unit, Department of Histology and Cell Biology, Faculty of Medicine, Suez Canal University, Ismailia, Egypt

Correspondence: Manal S. Fawzy (manal2_khashana@ymail.com, manal_mohamed@med.suez.edu.eg)

Genetic variants associated with iron homeostasis have been identified, but their association with iron-related indices and variables among different ethnic populations remains controversial. We aimed to explore the genotype frequency and allelic distribution of three iron-metabolism related variants in homeostatic iron regulator gene (HFE; rs1800562 G/A), transmembrane protease, Serine-6 gene (TMPRSS6; rs855791 A/G), and BTB domain-containing protein-9 gene (BTBD9; rs9357271 C/T) among a sample of the Middle Eastern blood donors and to detect the association of these variants on blood indices, and serum hepcidin/ferritin levels. Real-Time TaqMan genotyping assay for the specified variants was applied for 197 unrelated blood donors. Complete blood picture and serum ferritin/hepcidin levels were assessed. All participants were carriers of rs1800562*G/G genotype for HFE. The frequency of A/A and A/G genotypes of TMPRSS6 rs855791 variant was 55% and 45%, and for C/C, C/T, and T/T of BTBD9 rs9357271, were 15%, 43%, and 42%, respectively. Minor allele frequencies of rs855791*G and rs9357271*C were 0.23 and 0.37. The GGC genotype combination (for HFE/TMPRSS6/BTBD9, respectively) was more frequent in male participants. Higher serum hepcidin and hepcidin/ferritin ratio was observed in TMPRSS6 (A/G) carriers. While subjects with BTBD9 C/T and TT genotypes had lower serum ferritin values and higher levels of hepcidin and hepcidin/ferritin ratio compared with C/C genotype. No significant associations were found with any other blood parameters.

In conclusion, TMPRSS6 rs855791 (A/G) and BTBD9 rs9357271 (C/T) variants were prevalent in the present blood donor population and may influence the serum hepcidin and/or ferritin levels.

Introduction
Numerous biologic processes rely upon adequate iron levels [1]. Over the past decade, many traits/disease-associated single-nucleotide polymorphisms (SNPs) have been reported by genome-wide association studies, which have revolutionized our understanding of different genotype–phenotype associations [2].
Data from these studies have identified multiple SNPs associated with changes in serum iron-related parameters and blood cell phenotypes in the general population [3–5]. This influential effect could be explained in part by the intermediate effect on hepcidin and/or ferritin concentrations, although conflicting results in different populations remain [6,7].

Among these SNPs, the HFE variants in the human homeostatic iron regulator gene (HFE; rs1800562) have been suggested to exert multiple effects on the iron parameters and serum transferrin levels, in part independent of hepcidin [8]. The protein encoded by this gene is an HLA class I-like membrane protein that associates with β2-microglobulin and transferrin receptors on the cell surface [9]. HFE may reduce the affinity of transferrin receptor 1 for transferrin, [10] but its major role is in regulating hepcidin expression, likely via the bone morphogenetic protein (BMP)–SMAD signaling pathway [11].

Another SNP implicated in iron regulation is the transmembrane protease, serine 6 gene (TMPRSS6; rs855791), mapped on chromosome 22q12-13, which encodes matritase-2 (MT-2) [12]. MT-2 is mainly expressed in the liver, where it acts to down-regulate hepcidin expression [13]. It was found to be significantly associated with serum iron, hemoglobin (Hb), the mean corpuscular volume (MCV), and the mean cell hemoglobin (MCH) in genome-wide association studies (GWAS) in some populations [4,14,15].

Accumulating evidence revealed that some variants related to genes not directly implicated in iron homeostasis might also show an association with iron status parameters. For example, rs9357271 SNP in the BTB domain-containing protein-9 gene (BTBD9) was associated with decreased serum ferritin in patients and their relatives; despite the emergence of conflicting results [5,6]. The protein coded by this gene is ubiquitously expressed in the central nervous system and in the periphery, and both during development and adulthood. [16,17] Although the BTBD9 variants may directly influence iron metabolism and may be associated with proteins that participate in iron regulation signaling pathways (Figure 1A), yet the mechanism is not known [5].

Based on the earlier works [18,19], serum hepcidin concentration and serum ferritin showed a strong association in both sexes that withstood adjustment for age, BMI, time of blood sampling, and other measured iron parameters like iron and TIBC, as well as biochemical variables such as ALT, eGFR, and CRP [18]. Great interindividual variation in serum hepcidin concentration was observed [20]. This implies that population-based normalcy values may have limitations when used for the interpretation of hepcidin concentrations. Such it appears that hepcidin values should be interpreted in the context of biochemical tests used to evaluate iron metabolism [21]. Normalization of hepcidin to ferritin levels in the form of hepcidin/ferritin ratio has been evaluated in several genetic studies [22,23]. For example, Dijk et al. found that low serum hepcidin/ferritin ratio was associated with HFE C282Y-homozygosity compared with the wild genotype carriers and those with ferritin levels that remain within reference values. These investigators concluded that ‘this ratio might be a useful index for assessing inadequate responses to iron loading and might be of help in predicting which homozygoates may be at risk of developing clinically significant iron loading’ [22]. Additionally, Heeney et al. reported the clinical utility of this ratio in predicting TMPRSS6 mutation status in patients with chronic iron deficiency [23]. In the case of normal liver function, the hepcidin/ferritin ratio might be a useful indicator of erythropoiesis and iron kinetics, as reported by Lotfi and colleagues [18]. They have realized the importance of this ratio in providing a statistical algorithm to predict the spontaneous recovery from iron loss following blood donation [18].

Taken collectively, based on the literature mentioned above and the limited studies that relate the specified SNPs with serum iron-related parameters in the authors’ region, the present study has been conducted to determine the prevalence of rs1800562 in HFE (p. Cys282Tyr), rs855791 in TMPRSS6 (p. Ala736Val) and the intrinsic variant rs9357271 in BTBD9 among a sample of Middle Eastern population, and to evaluate the association of these variants with the available iron-related parameters, including the serum hepcidin, ferritin, and hepcidin/ferritin ratio.

The criteria and the selection process of the specified variants are detailed in the ‘Materials and Methods’ section of the present study. The identification and characterization of such associations may lead to better risk stratification for iron-related disorders in this population according to their genetic profile and early implementation of the preventive measurements and individualized iron-based therapeutic strategies for selected one.

**Subjects and methods**

**Study participants**

The present cross-sectional study enrolled 197 consecutive unrelated blood donors attending the blood bank at the Sohag University Hospital, Sohag, Egypt. The participants selection criteria were age > 18 years, non-pregnant, have no history or laboratory findings of chronic disease, inflammatory disorders (leukocyte count more than 10 × 10^3/μl),...
Figure 1. In silico data analysis

(A) Gene–gene functional interaction network generated by GeneMANIA. The network nodes have been colored by function (i.e., gene ontology annotation) (data source: https://genemania.org/search/homo-sapiens/HFE/TMPRSS6/BTBD9). (B) Protein–protein interaction using STRING version 11. Each network node represents the protein produced by a single locus. Edges represent protein–protein associations (i.e., contribute to a shared function; this does not necessarily mean they are physically binding to each other). The line colors of the edges indicate the type of interaction evidence that is explained in the figure key. The Network showed a relationship with iron-metabolism related proteins (data source: https://string-db.org). (C–E) Subcellular localization of HFE, TMPRSS6, and BTBD9 proteins. Darker color, according to the provided color key, is indicating more abundance (data source: https://compartments.jensenlab.org). Abbreviations: ANLN, anillin actin-binding protein; ATF2, activating transcription factor 2; B2M, beta-2-microglobulin; BTBD9, BTB domain containing 9; CBR3, carbonyl reductase 3; CDH18, cadherin 18; CFI, complement factor I; FBXO11, F-box protein 11; GALNS, galactosamine (N-acetyl)-6-sulfatase; GALNT2, polypeptide N-acetylgalactosaminyltransferase 2; HAMP, Hepcidin; HFE, homeostatic iron regulator; ID3, inhibitor of DNA binding 3; IGFBP4, insulin-like growth factor binding protein 4; PTPRN, protein tyrosine phosphatase, receptor type N; SMARCB1, SWI/SNF related, matrix associated, actin-dependent regulator of chromatin, subfamily b, member 1; SKA1, spindle and kinetochore associated complex subunit 1; TFR2, transferrin receptor 2; TFRC, transferrin receptor; TMPRSS6, transmembrane protease, serine 6; TPP1, tripeptidyl peptidase 1; USP25, ubiquitin specific peptidase 25; major histocompatibility complex, class I-related, HLA-E, major histocompatibility complex, class I, E; HLH protein.

Criteria for selecting the study variants and in silico data analysis

Literature review and in silico approach were applied for retrieving and selection of common (i.e. minor allele frequency > 0.05) SNPs located in genes coding for proteins related to iron-metabolism. Genomic sequence and variants were analyzed in ensemble.org, and the specified variants were selected for each gene (Table 1). Subcellular localization was identified in the Compartments database (compartments.jensenlab.org). Gene–gene interaction was analyzed using Gene Mania version 3.6.0 (https://genemania.org/search/homo-sapiens/HFE/TMPRSS6/BTBD9). Protein–protein association networks and gene ontology were analyzed using String version 11.0 (string-db.org).
### Table 1 Characteristics of the selected gene variants

<table>
<thead>
<tr>
<th>Locus</th>
<th>Position</th>
<th>Gene</th>
<th>SNP ID</th>
<th>Alleles</th>
<th>MAF</th>
<th>Type</th>
<th>Citations</th>
</tr>
</thead>
<tbody>
<tr>
<td>6p22.2</td>
<td>6:26092913</td>
<td>HFE</td>
<td>rs1800562</td>
<td>G/A</td>
<td>0.06</td>
<td>(A)</td>
<td>missense</td>
</tr>
<tr>
<td>22q12.3</td>
<td>22:37066896</td>
<td>TMPRSS6</td>
<td>rs855791</td>
<td>G/A</td>
<td>0.50</td>
<td>(G)</td>
<td>missense</td>
</tr>
<tr>
<td>6p21.2</td>
<td>6:38398097</td>
<td>BTBD9</td>
<td>rs9357271</td>
<td>T/C</td>
<td>0.50</td>
<td>(T)</td>
<td>intronic</td>
</tr>
</tbody>
</table>

Abbreviations: BTBD9, BTB domain-containing protein-9 gene; HFE, homeostatic iron regulator gene; MAF, minor allele frequency; SNP, single-nucleotide polymorphism; TMPRSS6, transmembrane protease, serine 6 gene. Data are currently mapped to Genome Assembly GRCh38.p13 (Data source: www.ensembl.org)

### Sample collection and the laboratory analysis

Seven milliliters of blood samples were withdrawn from all participants by early day time to minimize the effects of diurnal variation in hepcidin in vacutainer serum separator tubes and ethylenediaminetetraacetic acid (EDTA)-vacutainers. The former tubes were centrifuged at 2500 rpm for 15 min, and the separated sera were subjected to analysis of the routine infectious disease screening according to the local protocols. Ferritin and hepcidin were measured by enzyme-linked Immunosorbent assay Human Ferritin Kit (BioVision, Inc) and Human Hecpidin Quantikine Kit (R&D system) following the manufacturer's instructions, respectively. The EDTA tubes were used for complete blood count (cell Dyne-3700 fully automated cell counter; Abbott Diagnostics, Wiesbaden, Germany) with blood film examination and for subsequent DNA extraction.

### Iron metabolism-related genes genotyping

Genomic DNA was purified from whole blood using the QIAamp DNA Blood Mini kit (Qiagen, Hilden, Germany) following the manufacturer's protocol. The concentration and purity of the extracted genomic DNA were assessed by NanoDrop ND-1000 (NanoDrop Technologies, Inc. Wilmington, DE, U.S.A.). Genotyping for the selected three SNPs in iron metabolism-related genes (HFE; rs1800562; TMPRSS6; rs855791 and BTBD9; rs9357271) were assayed using TaqMan Real-Time polymerase chain reaction (PCR) allelic discrimination assay as described in details in our previous work [24]. PCR reactions were run in a 25-μl final volume containing 40 ng genomic DNA, TaqMan genotyping Master Mix, and TaqMan SNP Genotyping Assay Mix following the standard protocols. Appropriate controls were used in each reaction plate. The authors who ran the PCR reactions were blinded to the identity of the samples. PCR amplification was done using StepOne™ Real-Time PCR System, and allelic discrimination was called by the SDS software (version 1.3.1, Thermo Fisher Scientific Inc., Waltham, MA, U.S.A.). All quality control measures were applied according to the standard protocols, and replicates of 10% of samples were run with a 100% concordance rate.

### Statistical analysis

Categorical variables were presented as frequency counts and compared using the chi-square test. Meanwhile, the continuous data were expressed as mean ± standard deviation and compared using Student’s t-test or Mann–Whitney U tests according to data distribution and variance homogeneity. Genotype frequencies for each selected variant were tested for Hardy–Weinberg equilibrium (HWE). The odds ratio (OR) and the 95% confidence interval (CI) were calculated by logistic regression analysis for each variant [25]. Overall and sex-stratified analyses were run. Univariate analysis was performed to test associations. A bivariate correlation matrix using Spearman’s rank correlation analysis was applied to correlate the laboratory results’ different parameters. A multivariate test using the principal component analysis for data exploration was run to test the possibility of participant clustering according to sex and/or genotyping. Statistical significance was set at P-value < 0.05. Statistical Package for Social Science software version 23.0 was used for the statistical analysis.

### Results

#### Baseline characteristics of the study population

The study enrolled 197 blood donors; 179 men (90.0%) aged 29.1 ± 6.3 years and 18 women (9.1%) aged 30.2 ± 11.6 years old. More than half of the participants (56%) gave blood once for transfusion, 25% donated twice throughout their entire life, while less than a quarter underwent blood donation more than two times. Laboratory results are shown in Supplementary Table S1. No significant differences were found in blood parameters among men and women. In contrast, lower values of MCH and MCHC were observed in young subjects under 30 years old (P=0.003 and 0.018, respectively).
### Table 2 Genetic association models for the study variants

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Female</th>
<th>Male</th>
<th>OR (95% CI)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>rs855791</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Heterozygote comparison</td>
<td>A/A</td>
<td>10 (56)</td>
<td>98 (55)</td>
<td>1.00</td>
</tr>
<tr>
<td></td>
<td>A/G</td>
<td>8 (44)</td>
<td>81 (45)</td>
<td>1.03 (0.39–2.74)</td>
</tr>
<tr>
<td>Allelic model</td>
<td>A</td>
<td>28 (78)</td>
<td>277 (77)</td>
<td>1.00</td>
</tr>
<tr>
<td></td>
<td>G</td>
<td>8 (22)</td>
<td>81 (23)</td>
<td>1.02 (0.44–2.33)</td>
</tr>
<tr>
<td><strong>rs9357271</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Co-dominant model</td>
<td>T/T</td>
<td>10 (55.6)</td>
<td>72 (40.2)</td>
<td>1.00</td>
</tr>
<tr>
<td></td>
<td>C/T</td>
<td>7 (38.9)</td>
<td>78 (43.6)</td>
<td>1.55 (0.56–4.28)</td>
</tr>
<tr>
<td>Dominant model</td>
<td>T/T</td>
<td>10 (55.6)</td>
<td>72 (40.2)</td>
<td>1.00</td>
</tr>
<tr>
<td></td>
<td>C/C</td>
<td>1 (5.6)</td>
<td>29 (16.2)</td>
<td>4.03 (0.49–32.90)</td>
</tr>
<tr>
<td>Recessive model</td>
<td>T/T/C/C</td>
<td>8 (44.4)</td>
<td>107 (59.8)</td>
<td>1.86 (0.70–4.93)</td>
</tr>
<tr>
<td>Over-dominant model</td>
<td>T/T/C/T</td>
<td>17 (94.4)</td>
<td>150 (83.8)</td>
<td>1.00</td>
</tr>
<tr>
<td>Allelic model</td>
<td>T</td>
<td>27 (75)</td>
<td>222 (62)</td>
<td>1.00</td>
</tr>
<tr>
<td></td>
<td>C</td>
<td>9 (25)</td>
<td>136 (38)</td>
<td>1.83 (0.83–4.02)</td>
</tr>
<tr>
<td>Log-additive</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>1.77 (0.82–3.82)</td>
</tr>
</tbody>
</table>

Values are shown as number (percentage). Chi-square or Fisher’s exact tests were used. Abbreviation: OR (95% CI), odds ratio and confidence interval.

### Table 3 Genotype combination frequencies in men and women

<table>
<thead>
<tr>
<th>HFE</th>
<th>TMPRSS6</th>
<th>BTBD9</th>
<th>All</th>
<th>Female</th>
<th>Male</th>
<th>P-value</th>
<th>Crude OR (95% CI)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>G</td>
<td>A</td>
<td>T</td>
<td>0.49</td>
<td>0.52</td>
<td>0.48</td>
<td>0.034</td>
<td>1.00 (reference)</td>
</tr>
<tr>
<td>2</td>
<td>G</td>
<td>A</td>
<td>C</td>
<td>0.28</td>
<td>0.25</td>
<td>0.29</td>
<td>0.043</td>
<td>1.25 (0.64–2.43)</td>
</tr>
<tr>
<td>3</td>
<td>G</td>
<td>G</td>
<td>T</td>
<td>0.14</td>
<td>0.22</td>
<td>0.13</td>
<td>0.043</td>
<td>0.64 (0.29–1.41)</td>
</tr>
<tr>
<td>4</td>
<td>G</td>
<td>G</td>
<td>C</td>
<td>0.08</td>
<td>0.00</td>
<td>0.09</td>
<td>0.008</td>
<td>20.56 (1.16–36.29)</td>
</tr>
</tbody>
</table>

Chi-square or Fisher’s exact tests were used. Abbreviations: BTBD9, BTB domain-containing protein-9 gene; HFE, homeostatic iron regulator gene; OR (95% CI), odds ratio and confidence interval; TMPRSS6, Transmembrane Protease gene, Serine-6. Bold value indicates significance at P<0.05.

### Allelic discrimination analysis

All participants (n=197) had rs1800562*G/G genotype for the HFE gene. Thus, this variant was not included in the further downstream analysis. On the other hand, rs855791 (A/G) of the TMPRSS6 gene showed only two genotypes: A/A and A/G in 55% and 45% of the population, respectively, suggesting that this could be due to the small sample size. This later variant was not consistent with HWE in the overall analysis and the male population when stratified by sex (P>0.001). Whereas BTBD9 rs9357271 (C/T) variant was in agreement with HWE in overall and stratified analyses (P>0.05) and showed three genotypes: C/C, C/T, as well as T/T accounting for 15%, 43%, and 42% of the sample population, respectively. Minor allele frequencies of rs855791*G and rs9357271*C were 0.23 and 0.37 in the study population (Supplementary Table S2).

On comparing genotype and allele frequencies in men and women, there was no significant difference observed for both TMPRSS6 rs855791 and BTBD9 rs9357271 variants (Table 2). Paired loci analysis revealed more frequency of GGC genotype combination in men (Table 3).

### Univariate analysis

Associations of rs855791 TMPRSS6 and rs9357271 BTBD9 variants with biochemical results were depicted in Figure 2 and Figure 3. For TMPRSS6 gene, A/G genotype carriers exhibited higher levels of serum hepcidin (P=0.039) and hepcidin/ferritin ratio (P=0.041). While in BTBD9 gene, subjects with C/T and TT genotypes had lower values of...
Figure 2. Association between rs855791 \textit{TMPRSS6} and laboratory results

(A) Number (No) of donation, (B) Serum hepcidin (pg/ml), (C) Serum ferritin (ng/ml), (D) Serum hepcidin/ ferritin ratio, (E) White blood cells count (10^3/L), (F) Lymphocyte count (10^3/L), (G) Granulocytes count (10^3/L), (H) Red Blood Cells count (106/L), (I) Hemoglobin level (g/dl), (J) Hematocrit concentration (%), (K) Mean cell volume, (L) Mean cell hemoglobin concentration (g/dl), (M) Mean cell hemoglobin (pg), (O) Mean platelet volume (fL).

Box plots represent median and quartile values. Mann–Whitney \textit{U} test was used. \textit{P}<0.05 was set to be statistically significant.

Abbreviations: FER, ferritin; GRA, granulocytes; HCT, hematocrit; HEP, hepcidin; H/F, hepcidin/ferritin ratio; HGB, hemoglobin; LYM, lymphocyte; MCH, mean cell hemoglobin; MCHC, mean cell hemoglobin concentration; MCV, mean red cell volume; MPV, mean platelet volume; PLT, platelets; RBC, red blood cells; RDW, red cell distribution width; WBC, white blood cells.

serum ferritin (\textit{P}=0.007) and higher levels of hepcidin (\textit{P}=0.010) and hepcidin/ferritin ratio (\textit{P}=0.025) compared with C/C genotype. No significant associations were found with any other blood parameters.

\section*{Multivariate analysis}

As illustrated in Figure 4, the principal component analysis did not show discrete clustering of individuals based on their sex or genotypes.

\section*{Discussion}

There is a growing body of literature that recognizes the implication of multiple SNPs in several aspects of iron metabolism through known and yet unknown mechanisms or to be associated with some iron-related regulatory pathways \cite{26}. In this sense, we explored in the present study the association of three iron homeostasis-related SNPs on the iron parameters and serum hepcidin/ferritin levels in a cohort of Middle Eastern donors. We show that the SNPs \textit{TMPRSS6}; rs855791 and \textit{BTBD9}; rs9357271 were potentially associated with variations in serum hepcidin, serum ferritin, and/or hepcidin normalized to ferritin (the hepcidin/ferritin ratio).
Figure 3. Association between rs9357271 BTBD9 and laboratory results

(A) Number (No) of donation, (B) Serum hepcidin (ng/ml), (C) Serum ferritin (ng/ml), (D) Serum hepcidin/ ferritin ratio, (E) White blood cells count (10^3/L), (F) Lymphocyte count (10^3/L), (G) Granulocytes count (10^3/L), (H) Red Blood Cells count (10^6/L), (I) Hemoglobin level (g/dl), (J) Hematocrit concentration (%), (K) Mean cell volume, (L) Mean cell hemoglobin concentration (g/dl), (M) Mean cell hemoglobin concentration (g/dl), (N) Red cell distribution width (%), (O) Platelets count (10^3/L), (P) Mean Platelet Volume (fL).

Box plots represent median and quartile values. Mann–Whitney U test was used. \( P < 0.05 \) was set to be statistically significant. Abbreviations: FER, ferritin; GRA, granulocytes; HCT, hematocrit; HEP, hepcidin; H/F, hepcidin/ferritin ratio; HGB, hemoglobin; LYM, lymphocyte; MCH, mean cell hemoglobin; MCHC, mean cell hemoglobin concentration; MCV, mean red cell volume; MPV, mean platelet volume; PLT, platelets; RBC, red blood cells; RDW, red cell distribution width; WBC, white blood cells.

Although some studies, including the one done by Traglia et al. [8] and replicated by Galesloot et al. [4] in the Italian population, postulated the implication of HFE; rs1800562 in several aspects of iron parameters through hepcidin-dependent and independent mechanisms. Currently, this specified SNP in HFE gene could not be further analyzed because of its monomorphic genotype (GG; 100%) in the present population. However, this finding was in line with previous studies that failed to detect this variant in adult hepatitis C virus-induced liver cirrhosis, hemochromatosis, pediatric lymphoblastic leukemia survivors, or the controls in the same region [27–29]. These studies, including ours, are consistent with that rs1800562 (G/A) frequency, in general, ranged from 0 to 9.9% and seen to be nearly 0% in the North African population. Further large-scale multicenter studies are warranted to confirm this finding.

Several studies have realized the associations of the TMPRSS6 variants with decreased serum iron, [14] serum ferritin, [30], and Hb levels [14,31,32]. The current study showed that A/G genotype carriers of this variant have higher levels of serum hepcidin and hepcidin/ferritin ratio compared with the wild genotype carriers. Although this variant showed a departure from HWE, authors did not reject this SNP from further analysis as they speculated this could be attributed to the missing of the homozygous genotype of this SNP minor allele in the study population,
Figure 4. Principal component analysis

No clear demarcation was found between males and females (A) or subjects with different genotypes (B and C).
In conclusion, the present study demonstrated that TMPRSS6 rs855791 (A/G) and BTBD9 rs9357271 (C/T) variants were associated with serum hepcidin and/or serum ferritin levels in a preliminary sample of Middle Eastern blood donors. However, the present study could be limited by the relatively small sample size and the study design's cross-sectional analysis. Large-scale, multi-central, and prospective follow-up studies are warranted to verify the current conclusions.

Data Availability
All data used were included in the manuscript, and additional results were available in the Supplementary Material.

Competing Interests
The authors declare that there are no competing interests associated with the manuscript.

Funding
This work was supported by a Deanship of Scientific Research, Northern Border University (NBU), Arar, Saudi Arabia [grant number MED-2017-1-B-F-7477].

Author Contribution
M.S.F. and E.A.T. conceived the study, participated in the design of the study, carried out the molecular lab work, and drafted the manuscript. E.A.T. carried out the statistical analysis. A.F-E. collected field samples and data, carried out the hematological lab work, and participated in manuscript drafting. B.T.A. participated in the literature review, and helped draft the manuscript. All authors gave final approval for publication and agreed to be held accountable for the work performed therein.

Acknowledgements
The authors would like to thank the Deanship of Scientific Research, Northern Border University (NBU), Arar, Saudi Arabia, for the financial support of this study.

Abbreviations
BMP, bone morphogenetic protein; BTBD9, BTB domain-containing protein-9 gene; CI, confidence interval; GWAS, genome-wide association studies; Hb, hemoglobin; HWE, Hardy–Weinberg equilibrium; MCH, mean cell hemoglobin; MCV, mean corpuscular volume; MT-2, matriptase-2; OR, odds ratio; SNP, single-nucleotide polymorphism.

References

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