Role of Vitamin-D Receptor (VDR) single nucleotide polymorphisms in gestational hypertension development: A case-control study

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Abstract

Background
Recent literature data have highlighted the important role of hypovitaminosis D in pregnancy complications and prenatal/perinatal health. Vitamin D action takes place through vitamin D receptor (VDR) activation. Two single nucleotide polymorphisms of VDR gene, FokI and BsmI, have been reported to affect VDR molecular signaling and be associated with several disorders, including hypertension.

Methods
We carried out a case-control study aimed to assess vitamin D serum levels together with the distribution of VDR FokI and BsmI in a population of 116 pregnant women with gestational hypertension (GH) and 69 normotensive pregnant women (CTR).

Results
Hypovitaminosis D was largely prevalent both in GH (81%) and CTR (69%) pregnant women. Vitamin D insufficiency (10–30 ng/ml) had a similar frequency in both cohorts (GH 60% vs CTR 58%), while vitamin D deficiency (<10 ng/ml) was more frequent in GH cohort than in CTR one (21% vs 11%). Regression analysis showed that GH was significantly (p = 0.031) linked to vitamin D status. Vitamin D deficiency was associated with a threefold-increased risk of developing GH, while a normal vitamin D status was protective against this pregnancy disorder. The VDR FF/bB haplotype was the most frequent in GH cohort, and resulted to increase by two folds the risk for GH. Notably, hypovitaminosis D was found in 92% of FF/bB GH pregnant women, 27% of which had deficient vitamin D levels compared with 11% of their normotensive counterparts.
Conclusions
Despite being preliminary, these findings suggest that genotyping of pregnant women for VDR polymorphisms may be useful for a tailored vitamin D supplementation strategy.

Introduction
Gestational hypertension (GH), preeclampsia (PE) and eclampsia, are among the main complications related to hypertensive disorders of pregnancy, and represent approximately 14% of maternal mortality [1]. Pregnancy-induced hypertension, defined as blood pressure greater than 140/90 mmHg, usually develops after 20 weeks of gestation [2] and complicates 2% to 8% of pregnancies, thus being, along with PE, a major cause of maternal and perinatal morbidity/mortality [3]. The onset of these pathologies involves several maternal constitutional factors (obesity, genetics, diet), that, in combination with an altered inflammatory state and the mobilization of endothelial progenitor cells and natural killer cells, can lead directly to endothelial dysfunction [4], increasing the risk for cardiovascular disorder [5, 6]. Moreover, it has been suggested that epigenetic alterations may play a role in the abnormal development of placenta, caused by an impairment in angiogenesis and trophoblast invasion [7]. In the last years, the discovery of vitamin D-specific receptors and metabolites in the placenta and decidua, along with CYP27B1, the enzyme converting the 25(OH)D3 to 1,25(OH)2D3, has highlighted an important role for vitamin D in fertility as well as prenatal and perinatal health, outside of its established positive effects on skeletal mineralization, i.e. in the early modulation of trophoblast functions [8–10]. Furthermore, metabolic homeostasis of vitamin D-related proteins has been shown to be significantly altered in placental tissue from pregnancies complicated with PE compared with controls [11, 12]. Hypovitaminosis D, that is defined as a vitamin D level below the normal range (30–80 ng/ml, 75–200 nmol/L), is a common feature in pregnant women, and has been associated with a higher risk of pregnancy complications, i.e. gestational diabetes mellitus, PE, and other disorders, and negative effects for the fetus, including increased risk of preterm and low body weight birth, and increased susceptibility for various disorders in adult life [12–16].

Calcitriol preserves vascular system functions through several mechanisms: (1) inhibition of vascular smooth-muscle cells proliferation, intima-media thickening and metalloproteinase activity; (2) enhancement of vasodilatory action of endothelial nitric oxide synthase; (3) regulation of angiogenesis through modulation of the expression of the vascular endothelial growth factor (VEGF) gene [17, 18]. Moreover, vitamin D influences blood pressure through modulation of the renin-angiotensin-aldosterone system (RAAS) [17, 19, 20], which is also presents in placenta. During pregnancy, RAS normally undergoes major changes, stimulated by the increase of renin circulating levels- due to extra-renal local release by ovaries and maternal decidua- angiotensinogen and angiotensin II [21]. Plasma levels of active renin and angiotensin II type 1 receptor autoantibodies have been found to be higher in PE and GH than in normotensive women [22]. Vitamin D metabolites may reduce autoantibodies to the Angiotensin II type I receptor and suppress renin gene transcription by a vitamin D receptor (VDR)-dependent pathway [23].

The VDR gene consists of two promoter regions, eight coding exons (namely 2–9), and six untranslated exons (1A–1F). Among the many described single nucleotide polymorphisms (SNPs) of VDR, two di-allelic polymorphisms have been reported to affect VDR molecular signaling, namely FokI (C>T, rs2228750) within the first coding exon, and BsmI (A>G, rs1544410) within the third coding exon.
rs1544410) lying in the last intron [24]. The VDR FokI F mutated allele has been shown to increase VDR protein transcriptional activity, while the BsmI B mutated allele affects VDR mRNA stability leading to a reduction of VDR protein amount in tissues. It has been shown that an unfavourable VDR genetic background can significantly reduce the effectiveness of vitamin D action [24], thus contributing to the development of several disorders; in particular, the BsmI B allele has been strongly associated with hypertension [25, 26].

In this study, we assessed the distribution of VDR FokI and BsmI SNPs, as well as the circulating levels of vitamin D, in a population of 116 pregnant women with GH and 69 normotensive pregnant women, in order to evaluate the influence of vitamin D levels and VDR SNPs, as well as their interplay, on the risk for GH.

Materials and methods

Ethics

The study was conducted in accordance with the Declaration of Helsinki, and the study protocol was approved (protocol n. 51/17, approved on date July 17th 2017) by the local Ethics Committee, namely Comitato Etico Messina (website https://www.polime.it/page_view.php?pageid=162). Informed written consent was obtained from all participants.

Study cohorts

Pregnant women with GH, admitted to Operative Unit of Gynecology and Obstetrics, were enrolled for this study at the time of hospitalization for childbirth, from January 1st, 2018 to December 31st, 2019. Women with physiological pregnancies were considered as reference controls.

Data were collected from medical records and inclusion criteria were the following: pregnant women affected by hypertension, not affected by pathologies of the fetus. We excluded all the women who did not meet the inclusion criteria. Healthy pregnant women with the same characteristics as the hypertensive group were subsequently recruited to constitute a homogeneous sample for the control group.

The following anamnestic and obstetric data of all women included in the study were collected: age, height, pre-pregnancy weight, weight at pregnancy term, parity, type of delivery, gestational age at delivery, weight of the newborn, Apgar at 1 and 5 minutes, presence of gestational hypertension, pre-gravid and gravid pathologies, anti-hypertensive therapy if performed, any other therapies carried out during pregnancy.

Blood samples were collected from each participant and routinely tested for the following biochemical parameters: total hemoglobin concentrations, platelet count, prothrombin time (PT), partial thromboplastin time (PTT), fibrinogen levels, glucose levels, urea levels, sodium levels, potassium levels, total protein content, uric acid levels, iron levels, lactic dehydrogenase (LDH) activity, cholinesterase activity, creatine phospho-kinase (CPK) activity, serum total bilirubin, glutamate-pyruvate transaminase (GPT), glutamate-oxaloacetate transaminase (GOT), and serum creatinine. Urine samples were also collected in order to test for the presence of proteinuria.

Assessment of 25(OH)vitamin D3 serum levels

Plasma was obtained after blood centrifugation. Aliquots of whole blood and plasma were stored at -20°C until analysis.

The quantitative determination of 25(OH)Vitamin D3 plasma levels was performed by reverse-phase high-performance liquid chromatography (RP-HPLC) with a commercially
available kit for the determination of 25-OH Vitamin D3/D2 (#195–6529, Bio-Rad, Milan, Italy) using an Agilent 1200 Series HPLC system (Agilent Technologies Italia, Cernusco sul Naviglio, Milan, Italy).

**Genotyping for VDR gene SNPs**

Genomic DNA (gDNA) was isolated from leukocytes of peripheral whole-blood samples collected in EDTA test tubes, using the PUREGENE-DNA purification system (#158389 GENTRA, QIAGEN, Milan, Italy), according to manufacturer’s instructions. The DNA was quantified by spectrophotometric measurement at 260 nm using a BioPhotometer (Eppendorf AG, Hamburg, Germany). DNA quality was considered acceptable for samples having a 260/280 ratio ≥ 1.6. DNA integrity and the presence of contaminant RNA were checked by electrophoresis on 0.8% agarose gel, and subsequent UV detection of DNA bands using a gel documentation system (VilberLourmat, Collégien, France).

The screening for the presence of VDR SNPs FokI (C>T) rs2228570 and BsmI (A>G) rs1544410 was performed by Real-time PCR-based allelic discrimination, using two pre-designed TaqMan-based Genotyping Assays available from Applied Biosystems (assay ID: C_8716062_10, assay ID: C_12060045_20, respectively; ThermoFisher, Monza, Italy). PCR reactions were set up in a 96-well plate on a 7900HT Fast Real-Time PCR System (Applied Biosystems, Foster City, CA), and were carried out in a final volume of 20 μL containing 1x TaqMan™ Genotyping Master Mix (#4381656, Applied Biosystems, Thermo Fisher, Monza, Italy), 1x Mix Primers/TaqMan Probes, and 20 ng gDNA, using thermal cycling conditions suggested by manufacturer’s protocols.

**Statistical analysis**

The numerical data were expressed as mean and standard deviation and the categorical variables as absolute frequency and percentage.

Examined variables were not normally distributed, such as verified by Kolmogorov Smirnov test; consequently, the non-parametric approach has been used.

For each parameter, we performed statistical comparisons between two groups in exam, using Chi Square test for categorical variables (genotypes, haplotypes, haplotypes subgroups, 25(OH)vitamin D3 classes etc….) and Mann-Whitney test for numerical variables.

Moreover, Kruskal-Wallis test was applied to compare 25(OH)vitamin D3 levels among three genotype groups and haplotype subgroups.

For each group, we applied the Spearman correlation test, in order to evaluate the existence of significant correlation between 25(OH)vitamin D3 levels and other numerical parameters. Point biserial correlation was applied to assess the existence of significant inter-dependence between a dichotomous variable, such as binary 25(OH)vitamin D3 levels lower or higher than 30 ng/ml (75 nmol/L), as well as 25(OH)vitamin D3 lower than 10 ng/ml (25 nmol/L) or ranging between 10 and 30 ng/ml (25–75 nmol/L), and numerical variables.

Finally, a binary logistic regression model was applied in order to evaluate a possible significant dependence of the hypertension from 25(OH)vitamin D3 levels and from haplotypes subgroups.

Statistical analyses were performed using SPSS 22.0 for Window package. A P-value lower than 0.050 was considered to be statistically significant.

**Results**

2780 patients gave birth at the U.O.C. of Gynecology and Obstetrics during the study period. 185 women were enrolled for our study (see flow diagram in Fig 1). Among them, 116 patients
were affected by GH (Group 1, GH; 33.0 ± 6.2 years, height 163.7 ± 6.8 cm; pre-pregnancy weight 69.9 ± 12.3 kg, post-pregnancy weight 81.0 ± 11.3 kg), while 69 had normotensive blood pressure during pregnancy and were included as reference control group (Group 2, CTR; age: 33.0 ± 5.9 years; height 163.4 ± 6.3 cm; pre-pregnancy weight 63.3 ± 11.8 kg, post-pregnancy weight 75.9 ± 12.9 kg). Anamnestic data revealed that 22.4% of women with GH and 11.6% of women with normotensive pregnancies had previous abortions, 30.2% of women with GH and 26% of women with normotensive pregnancies had pre-pregnancy pathologies, 20.7% of women with GH and 8.7% of women with normotensive pregnancies had gestational diabetes. Women with GH gave birth after 35.9 (± 3.2 SD) weeks, and their newborns had mean weight of 2,928 (± 760.8 SD) g, 1-minute Apgar score of 8.8 (± 1.2 DS), and 5-minutes Apgar score of 9.5 (± 0.8 SD), while women with normotensive pregnancies gave birth after 36.6 (± 7.9 SD) weeks, to newborns with mean weight of 3,176 (± 472.1 SD) g, 1-minute Apgar score of 9.0 ± 0.9, and 5-minutes Apgar score of 9.7 ± 0.5. However, no significant differences between the two groups were found with regard to age, height, health status, ongoing pharmacological
treatment, and anamnestic-obstetric data, except for birth weeks (p = 0.009), and post-pregnancy weight (p < 0.001).

The biochemical features of the two groups of pregnant women are shown in Table 1. No significant differences were found between GH group and CTR group with regard to almost all biochemical parameters examined, except for blood pressure (systolic, diastolic), circulating fibrinogen levels, blood urea and uric acid levels, GPT, blood creatinine levels, and LDH.

Analysis of 25(OH)vitamin D3 level variability in GH and normotensive pregnant women

The assessment of 25(OH)vitamin D3 circulating levels showed that hypovitaminosis D, defined as a 25(OH)vitamin D3 serum level below the normal reference range of 30–80 ng/ml, was a common condition in pregnant women recruited for this study. Indeed, pregnant women with GH (n = 98) had mean 25(OH)vitamin D3 levels of 21.2 ± 11.8 ng/ml, and normotensive pregnant women (n = 58) had mean 25(OH)vitamin D3 levels of 24.1 ± 12.3 ng/ml (Table 1).

Given that hypovitaminosis D may be classified as insufficiency (serum 25(OH)vitamin D3 10–30 ng/ml) and deficiency (25(OH)vitamin D3 <10 ng/ml), both study cohorts were stratified in three subgroups to divide pregnant women that had normal, insufficient, and deficient 25(OH)vitamin D3 levels.

Table 1. Blood pressure and blood biochemical markers of GH and normotensive pregnant women (CTR) recruited for this study.

|                                | GH          | CTR         | P-value  |
|--------------------------------|-------------|-------------|----------|
| Systolic blood pressure (mmHg) | 149.4 ± 18.4| 116.2 ± 10.7| <0.001   |
| Diastolic blood pressure (mmHg)| 92.2 ± 11.8 | 70.2 ± 8.9 | <0.001   |
| Hemoglobin                     | 11.3 ± 1.3  | 11.1 ± 1.6  | 0.835    |
| PT ratio (%)                   | 107.6 ± 16.4| 111.5 ± 15.3| 0.312    |
| PTT (sec)                      | 27.7 ± 3.9  | 27.3 ± 2.5  | 0.210    |
| Fibrinogen (mg/dL)             | 514.4 ± 105.4| 475.1 ± 82.4| 0.019    |
| Platelet count (cells/mm³)     | 241,327 ± 91,420| 240,266 ± 73,018| 0.759   |
| Glucose (mg/dL)                | 81.7 ± 19.5 | 75.6 ± 9.6  | 0.162    |
| Urea (mg/dL)                   | 20.9 ± 7.4  | 18.4 ± 5.5  | 0.028    |
| Uric acid (mg/dL)              | 5.2 ± 1.5   | 4.1 ± 0.9   | 0.001    |
| Total proteins (g/dL)          | 6.0 ± 0.7   | 6.2 ± 0.5   | 0.105    |
| Bilirubin (mg/dL)              | 0.36 ± 0.26 | 0.36 ± 0.22 | 0.541    |
| GOT (U/L)                      | 31.9 ± 94.3 | 23.1 ± 34.2 | 0.418    |
| GPT (U/L)                      | 29.2 ± 69.0 | 14.8 ± 12.8 | 0.039    |
| CPK (U/L)                      | 79.1 ± 71.4 | 77.9 ± 78.7 | 0.594    |
| Sodium (mmol/L)                | 137.5 ± 3.0 | 138.6 ± 3.5 | 0.099    |
| Potassium (mmol/L)             | 4.3 ± 0.4   | 4.2 ± 0.3   | 0.169    |
| Iron (mcg/dL)                  | 79.4 ± 43.7 | 74.6 ± 49.1 | 0.362    |
| Cholinesterase (U/L)           | 5967.4 ± 1360.1| 6040.3 ± 1185.6| 0.884   |
| LDH (U/L)                      | 414.5 ± 240.6| 353.9 ± 127.8| 0.004    |
| Creatinin (mg/dL)              | 0.53 ± 0.14 | 0.46 ± 0.11 | 0.004    |
| 25(OH)vitamin D3 (ng/ml)       | 21.2 ±11.9  | 24.0 ± 12.3 | 0.181    |

All data values are shown as mean ± SD. CPK, creatine phosphokinase; GOT, glutamate-oxalacetate transaminase; GPT, glutamate-pyruvate transaminase; LDH, Lactate dehydrogenase; PT, Prothrombin time; PTT, partial thromboplastin time.
Among GH pregnant women only 19.4% had normal 25(OH) vitamin D3 levels (39.9 ± 8.3 ng/ml), while 59.2% had insufficient 25(OH) vitamin D3 levels (19.9 ± 5.9 ng/ml), and 21.4% had deficient 25(OH) vitamin D3 levels (7.9 ± 1.6 ng/ml). Instead, control normotensive pregnant women having normal 25(OH) vitamin D3 levels (39.1 ± 8.0 ng/ml) were 30.6%, those with insufficient levels (19.1 ± 5.0 ng/ml) were 57.6%, and those with deficient levels (7.3 ± 0.2 ng/ml) were 10.8%.

No significant differences were found when comparing 25(OH) vitamin D3 concentrations in the three subgroups between the two study cohorts.

Logistic regression analysis showed that the variable hypertension was significantly (p = 0.031) linked to vitamin D status. In particular, pregnant women having 25(OH) vitamin D3 levels below 10 ng/ml (deficiency) had a higher risk of developing hypertension in comparison with those having normal 25(OH) vitamin D3 levels (30–80 ng/ml). In particular, the Odds Ratio calculation showed that vitamin D deficiency was associated with a near three-fold increased risk of hypertension during pregnancy (O.R. = 2.71; C.I. 95% 1.02–7.2; p = 0.04). On the contrary, a normal vitamin D status was protective against the development of GH (O.R. = 0.4; 95% C.I. = 0.22–0.95; p = 0.037).

Distribution of VDR FokI and BsmI SNPs in the study cohorts

The screening of VDR FokI (rs2228570) SNP distribution among women with GH (GH group) and normotensive pregnant women (CTR group) showed that genotype frequencies were in agreement with expectations derived from Hardy-Weinberg equilibrium (HWE) both in GH group (chi-square: 3.54, p > 0.05) and CTR group (chi-square: 1.46, p > 0.05).

The mutated F allele was more frequent than the f wild-type allele in both GH and CTR groups (GH 0.659 vs 0.341, CTR 0.645 vs 0.355), and was more represented in homozygous than in heterozygous state. Indeed, the frequency of FF genotype was higher in GH group than in CTR group (47.4% vs 44.9%), but no statistically significant differences were observed. The homozygous wild-type ff and the heterozygous fF genotypes showed similar frequencies in both groups (ff: GH 15.5% vs CTR 15.9%; fF: GH 37.1% vs CTR 39.2%).

The results obtained by genotyping of GH and CTR pregnant women for VDR BsmI (rs1544410) showed the concordance of genotype distributions in both populations with HWE (GH chi-square: 1.748, p > 0.05; CTR chi-square: 0.344, p > 0.05). The wild-type b allele was more frequent than the B mutated allele both in GH group (0.52 vs 0.48) and CTR group (0.58 vs 0.42), and was largely prevalent in heterozygous state. In particular, the homozygous wild-type bb genotype was more frequent in CTR group than in patients (31.9% vs 24.1%). Instead, the bb and BB mutated genotypes were more frequent among GH pregnant women than in CTR ones (bb: 56.1% vs 52.2%; BB: 19.8% vs 15.9%). However, no statistically significant differences were found between the two groups with regard to all genotype frequencies.

Distribution of VDR FokI/BsmI haplotypes in the study cohorts

Given the reported reduction of VDR signaling efficacy associated with the presence of either the FokI f allele or the BsmI B allele or both [19–21], we aimed to assess whether the combination of these VDR polymorphic alleles can be considered as a risk factor for the development of GH.

The analysis of VDR FokI/BsmI haplotype distribution showed that the FF/Bb haplotype was the most frequent among GH pregnant women, being present in one third of GH population, and was significantly more frequent in GH cohort than in CTR cohort (Table 2).
Notably, the Odds Ratio calculation indicated that women bearing the VDR FF/bB haplotype had a twofold-increased risk for GH compared with those having other haplotypes (OR = 2.18, 95% CI = 1.02–4.66, p = 0.043).

The fF/bB haplotype was the most represented among control pregnant women, being found in one fourth of CTR population, and was more frequent in CTR group than in GH group, even if this difference was not statistically significant (Table 2).

The double heterozygosity condition represented by the fF/bB haplotype was also very common among GH pregnant women, being observed in one fifth of GH population (Table 2).

It’s interesting to note that also the FF/bb haplotype was more frequent in normotensive CTR pregnant women in comparison with GH pregnant women, and this difference tended to be statistically significant (Table 2). Likely, a statistically significant result was not achieved due to the small number of subjects included in both subgroups. This result may suggest a protective effect of FF/bb haplotype against GH.

No statistically significant differences were found when comparing GH pregnant women with CTR pregnant women having other haplotype combinations (Table 2).

### Analysis of 25(OH)vitamin D3 levels in pregnant women with different VDR haplotypes

We also examined the variability of mean 25(OH)vitamin D3 levels in GH and normotensive pregnant women having different VDR FokI/BsmI haplotypes.

After stratification of the study cohorts in nine subgroups according to different VDR haplotypes, we observed that both GH and normotensive pregnant women with fF/BB haplotype had normal 25(OH)vitamin D3 levels, as well as normotensive pregnant women with either fF/BB or fF/bb haplotype (Table 3). GH women with fF/BB haplotype, as well as pregnant women having different haplotypes in both study cohorts, had insufficient 25(OH)vitamin D3 levels (Table 3).

The lowest 25(OH)vitamin D3 levels were found in GH pregnant women with FF/bb haplotype (Table 3). Notably, hypovitaminosis D was largely predominant among FF/bb GH pregnant women (92%) than in normotensive CTR pregnant women having the same haplotype (77%). Vitamin D insufficiency was found at a similar frequency (GH 65% vs CTR 66%), while a deficient vitamin D status was only observed among GH women (26.9%). However, significant differences between the two groups were not found, likely due to the very small number of FF/bb subjects within the two cohorts stratified according to vitamin D status.

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**Table 2. Distribution of VDR FokI/BsmI haplotypes among GH and normotensive pregnant women (CTR) recruited for this study.**

| VDR FokI/BsmI haplotype | GH (n = 116) | CTR (n = 69) | P-value |
|-------------------------|-------------|-------------|---------|
| ff/bb                  | 4.3% (5)    | 1.4% (1)    | 0.281   |
| ff/bB                  | 8.6% (10)   | 11.6% (8)   | 0.506   |
| ff/BB                  | 2.6% (3)    | 2.9% (2)    | 0.903   |
| fF/bb                  | 10.3% (12)  | 11.6% (8)   | 0.783   |
| fF/bB                  | 18.1% (21)  | 24.6% (17)  | 0.291   |
| fF/BB                  | 8.7% (10)   | 2.8% (2)    | 0.116   |
| FF/bb                  | 9.5% (11)   | 18.8% (13)  | 0.069   |
| FF/bB                  | 29.3% (34)  | 15.9% (11)  | 0.040   |
| FF/BB                  | 8.6% (10)   | 10.1% (7)   | 0.733   |

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In CTR group the lowest vitamin D concentrations were found in women with \( fF/bB \) haplotype, and they were similar to those found in GH pregnant women with same haplotype (Table 3).

No significant differences were found when comparing vitamin D concentrations in the nine subgroups with different \( VDR \) \( FokI/BsmI \) haplotypes between the two study cohorts.

Analysis of vitamin D status within different \( VDR \) haplotype subgroups

In order to overcome the difficulties of finding statically significant differences between the GH group and the CTR group, due to the small number of subjects included in each subgroup referred to a different \( VDR \) haplotype, pregnant women in both cohorts were stratified into three subgroups according to the set of mutant \( FokI \) \( F \) and \( BsmI \) \( B \) alleles possessed. The subgroup 1 included pregnant women bearing none/one mutant allele, hence having either \( ff/bb \), \( fF/bb \), or \( ff/bB \) haplotype; the subgroup 2 included pregnant women bearing two mutant alleles, hence having either \( fF/bB \), \( FF/bb \), or \( ff/BB \) haplotype; the subgroup 3 included pregnant women bearing three/four mutant alleles, hence having either \( FF/bB \), \( fF/BB \), or \( FF/BB \) haplotype.

Almost half of the GH pregnant women (46.5%) were grouped in the subgroup 3, while the remaining were divided with a similar frequency (about 27%) in the subgroups 1 and 2, and the remaining in the subgroup 3. Instead, 40.3% of normotensive pregnant women were included in the haplotype subgroup 2, 31.6% in the haplotype subgroup 3, and 28.1% in the haplotype subgroup 1 (Table 4). No significant difference were found with reference to the sample size of each haplotype subgroup between the two cohorts. However, a difference tending to statistical significance (\( p = 0.068 \)) was observed when comparing the number of GH pregnant women in the haplotype subgroup 3 with their normotensive counterparts.

The mean 25(OH)vitamin D3 serum levels of GH pregnant women within the three haplotype subgroups were below the normal reference range and lower than those of CTR normotensive within the same haplotype subgroups, except in the case of the haplotype subgroup 2 (Table 4). However, no statistically significant differences were found between GH women and normotensive pregnant women, likely due to the small sample size of each haplotype subgroup within each study cohort (Table 4).

Finally, GH and normotensive pregnant women within different haplotype subgroups were stratified according to their vitamin D status, namely either normal, insufficient, or deficient
vitamin D serum level, in order to better understand the relationship between VDR haplotype, vitamin D status and hypertension.

We found that about four-fifths of GH pregnant women in each haplotype subgroup had hypovitaminosis D, while the same condition was observed in less than three-fifths of normotensive pregnant women in either the haplotype subgroup 1 (ff/bb, fF/bb, ff/bB) or the haplotype subgroup 3 (FF/bB, fF/BB, FF/BB), and in about four-fifths of those in the haplotype subgroup 2 (Table 5).

No significant between-groups differences were found when comparing the relative abundance of subjects included in the haplotype subgroups stratified according to vitamin D status, except in the case of haplotype subgroup 3. In particular, a significantly lower number of GH pregnant women in the haplotype subgroup 3 was found to have normal vitamin D levels in comparison with their normotensive counterparts (Table 5). Moreover, in the haplotype subgroup 3 the frequency of GH pregnant women with deficient vitamin D levels was more than four-fold higher than that of normotensive pregnant women. However, this difference was

Table 4. Serum 25(OH)vitamin D3 concentrations in GH and normotensive pregnant women (CTR) stratified into different haplotype subgroups.

| Diagnosis | Serum 25(OH)vitamin D3 concentrations (ng/ml) |
|-----------|---------------------------------------------|
|           | Haplotype subgroup 1* (n) | Haplotype subgroup 2 ** (n) | Haplotype subgroup 3 *** (n) |
| GH (n = 99) | 20.6 ±10.0 (n = 26) | 22.1±14.3 (n = 27) | 21.0 ± 11.4 (n = 46) |
| CTR (n = 57) | 28.0 ±14.0 (n = 16) | 21.0±11.0 (n = 23) | 27.7 ± 14.6 (n = 18) |

Vitamin D3 concentrations are shown as mean ± SD.
*Haplotypes ff/bb + fF/bb + ff/bB;
**Haplotypes fF/bB + ff/BB + FF/bb;
***Haplotypes fF/BB + FF/bB + FF/BB.

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Table 5. Analysis of vitamin D status in GH and normotensive pregnant women (CTR) within different haplotype subgroups.

|                      | VDR FokI/BsmI haplotypes | Deficient vitamin D status (<10 ng/ml) | Insufficient vitamin D status (10–30 ng/ml) | Normal vitamin D status (30–80 ng/ml) |
|----------------------|--------------------------|----------------------------------------|---------------------------------------------|---------------------------------------|
| GH (n = 99)          | Haplotype subgroup 1*    | 7.7 ± 0.7                              | 19.7 ± 5.4                                  | 36.1 ± 3.3                            |
|                      | % (n = 26)               | 19.2% (5)                              | 61.6% (16)                                  | 19.2% (5)                             |
|                      | Haplotype subgroup 2**   | 8.3 ± 1.5                              | 21.0 ± 6.5                                  | 42.8 ± 11.8                           |
|                      | % (n = 27)               | 29.6% (8)                              | 48.1% (13)                                  | 22.6% (6)                             |
|                      | Haplotype subgroup 3***  | 8.4 ± 2.3                              | 20.4 ± 5.3                                  | 40.1 ± 7.4                            |
|                      | % (n = 46)               | 23.9% (11)*                            | 58.7% (27)                                  | 17.4% (8)*                            |

|                      | Haplotype subgroup 1*    | 7.2                                    | 19.9 ± 5.9                                  | 40.2 ± 10.8                           |
|                      | % (n = 16)               | 6.25% (1)                              | 50.0% (8)                                   | 43.75% (7)                            |
|                      | Haplotype subgroup 2**   | 7.4 ± 0.15                             | 18.8 ± 5.4                                  | 37.1 ± 6.1                            |
|                      | % (n = 23)               | 17.4% (4)                              | 60.9% (14)                                  | 21.7% (5)                             |
|                      | Haplotype subgroup 3***  | 6.9                                    | 18.8 ± 4.0                                  | 43.4 ± 9.6                            |
|                      | % (n = 19)               | 5.3% (1)                               | 52.6% (10)                                  | 42.1% (8)                             |

*Haplotypes ff/bb + fF/bb + ff/bB;
**Haplotypes fF/bB + ff/BB + FF/bb;
***Haplotypes fF/BB + FF/bB + FF/BB.

§ p = 0.029, significantly different frequency in comparison with normotensive pregnant women of haplotype subgroup 3 having a normal vitamin D status;
* p = 0.07, frequency value tending to statistically significant difference in comparison with normotensive pregnant women of haplotype subgroup 3 having a deficient vitamin D status.

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only tending to statistical significance, likely because of the small sample size of the two subgroups compared (Table 5). Interestingly, the risk of developing GH was decreased by about four folds in pregnant women of haplotype subgroup 3 having a normal vitamin D status (O. R = 0.26; C.I. 95% 0.079–0.87; p = 0.029).

**Discussion**

Hypovitaminosis D, classified as vitamin D insufficiency (10–30 ng/mL; 20–75 nmol/L) or vitamin D deficiency (<10 ng/mL; <20 nmol/L), has been reported worldwide among pregnant women. Women with low sun exposure and vitamin D intake as well as a poor quality diet are at greatest risk of hypovitaminosis D, that leads to pregnancy complications, including PE and GH, as well as adverse perinatal outcomes. Early pregnancy has been indicated as the likely critical window for the prevention of these adverse events [27].

Literature data from observational studies focusing on the association of vitamin D status with the onset of pregnancy hypertensive disorders are discordant, mainly because of different analytical methods employed for vitamin D level assessment, inconsistent reports on vitamin D levels, large heterogeneity of study design, and lack of adherence to standardized perinatal outcome definitions. However, the scientific community is recently moving towards the recognition of the importance of vitamin D deficiency in the development of these pathological conditions [28]. Indeed, recent meta-analyses of observational studies highlighted the existence of an inverse ratio between vitamin D levels and the development of PE as well as GH, and provided evidence for an increased risk of gestational hypertensive disorders at vitamin D concentrations <20 ng/ml (<50 nmol/L)[12, 28, 29]. Vitamin D concentrations <12 ng/mL (<30 nmol/L) in early pregnancy have been associated with a higher risk for PE in comparison to concentrations >20 ng/mL (>50 nmol/L) [30]. Finally, a 36% reduction in PE composite outcome and small-for-gestational-age birth has been reported when vitamin D concentrations were higher than 30 ng/ml (75 nmol/L) at 15 weeks’ gestation [31].

The present study confirmed the findings from the aforementioned meta-analysis, showing a significant association between hypovitaminosis D and maternal hypertensive disorder risk. Indeed, more than 80% of pregnant women with GH had vitamin D levels lower than normal reference range. Deficient vitamin D concentrations (<10 ng/mL; <25 nmol/L) or insufficient vitamin D concentrations (10–30 ng/mL; 25–75 nmol/L) were associated, respectively, with a threefold-increased and 1.5-fold increased risk for developing GH. In recent years, both evidence from clinical trials and animal experiments have been provided that vitamin D regulates blood pressure through inhibition of renin-angiotensin-aldosterone system (RAAS) activity, modulation of vascular functions and the reduction of vascular oxidative stress [17, 19, 23, 32]. In preeclamptic women the circulating levels of renin, angiotensin I (ANG I) and aldosterone are lower than their normotensive counterparts. In addition, while normotensive pregnant women demonstrate decreased vascular sensitivity to angiotensin II (ANG II), preeclamptic women exhibit increased sensitivity of the adrenal cortex and vascular system to ANG II. An increase in renin expression was demonstrated in the decidua vera of preeclamptic women in comparison with normotensive pregnant women. It has been suggested that maternal decidua acts as an additional site of RAAS activation, and that the small amount of ANG II locally produced finds its way into maternal circulation and is sufficient to down-regulate ANG II production in the kidney as seen in preeclampsia [33, 34]. Vitamin D acts as a negative endocrine regulator of RAAS, suppressing renin gene transcription or reducing autoantibodies to the ANG II type I receptor. In addition, vitamin D can influence blood pressure through the suppression of vascular smooth muscle cell proliferation [12].
Vitamin D suppresses renin gene transcription through a VDR-dependent pathway. Two VDR SNPs, *FokI* (rs2228570) and *BsmI* (rs1544410), have been reported to affect vitamin D signaling efficacy, thus increasing the risk for several pathological conditions, including hypertension, even if in this latter case conflicting results have been reported [25, 26, 35, 36]. In particular, the allele *B* of VDR*BsmI* polymorphism is strongly positively correlated with systolic and diastolic blood pressures, and the VDR *FokI* allele has been significantly and independently associated with higher plasma renin activity both in hypertensives and normotensives subjects [37, 38].

Here we aimed to investigate the VDR *FokI* and *BsmI* haplotype distribution among pregnant women with GH and normotensive pregnant women. A case-control study showed that the three common VDR SNPs (*FokI*, *ApaI*, and *BsmI*) were equally distributed in GH group compared with healthy pregnancy cohort, and none of these polymorphisms was a predisposing risk factor for PE and GH [39]. Similarly, our study reported an equal VDR *FokI* and *BsmI* polymorphism distribution between hypertensive and normotensive pregnant women, and none of the investigated polymorphisms, taken individually, were associated with maternal hypertensive disorders. However, while examining the haplotype distribution we found that *FF/bB* haplotype, accounting for slightly less than one third of GH cohort, resulted to be significantly more frequent among GH pregnant women than normotensive ones, and increase by more than two folds the risk of developing GH. This suggests an association of VDR *FF/bB* haplotype with hypertensive disorders during pregnancy. Our observations are similar with those recently reported by Rezavand and co-workers, showing that a significantly higher frequency of VDR *FokI* *F* allele is observed in preeclamptic patients (83%) than in controls (74%) and is associated with a 1.72-fold increased risk of PE. In particular, the *FokI* *FF* genotype resulted to be associated with a significantly higher blood pressure when compared to *ff* and *fF* genotypes [40].

However, the role played by the presence of *BsmI* *B* allele in heterozygosis within the *FF/bB* haplotype cannot be neglected, since it may also explain the association of this haplotype with an increased risk to develop GH. Indeed, a recent meta-analysis study highlighted a major role of *BsmI* *bB* genotype in the development of hypertension. Carriers of *bB* genotype had a higher risk of hypertension than those having *BB* or *bb* genotype (OR = 1.27, 95% CI = 1.01–1.60, *p* = 0.04) [26]. The observation that the *FF/bb* haplotype was prevalent among normotensive pregnant women recruited for this study, despite the differences in haplotype frequency between GH and control group only tended to statistical significance likely due to the small group size, suggest that the *FF/bb* haplotype may play a protective role against the onset of pregnancy hypertensive disorders. This may support the hypothesis on the important role played by *BsmI* *B* allele in GH.

Finally, we also carried out correlation analyses between genotyping data and vitamin D levels, after stratification of pregnant women in both GH and normotensive cohorts in three subgroups, including subjects carrier of haplotypes containing either none/one mutant allele (subgroup 1, haplotypes *ff/bb*, *fF/bb*, *ff/bB*), or two mutant alleles (subgroup 2, haplotypes *fF/bB*, *FF/bb*, *ff/BB*), or three/four mutant alleles (subgroup 3, haplotypes *fF/BB*, *FF/bb*, *FF/BB*) having deficient, insufficient, and normal vitamin D concentrations. We only found a statistically significant difference between hypertensive and normotensive pregnant women, when comparing subjects included in subgroup 3 having a normal vitamin D status, and a difference tending to statistical significance when comparing pregnant women in this same haplotype subgroup having deficient vitamin D levels. These findings suggest that normal vitamin D levels play a protective role against the negative effects of an unfavourable VDR haplotype. On the contrary, vitamin D deficiency plays a major role, compared with vitamin D insufficiency, in the development of GH when pregnant women are carriers of an unfavourable VDR haplotype.
The lack of other between-groups significant differences are likely due to the very small number of pregnant women included in the nine haplotype/vitamin D status subgroups in each study cohort.

Conclusions

In conclusion, our study suggests an association between GH and the VDR FF/bB haplotype, that has been shown to increase by twofold the risk for GH. Notably, almost all GH pregnant women (92%) having this haplotype also had either insufficient (65%) and deficient (27%) vitamin D levels. In general, hypovitaminosis D was largely prevalent both in GH pregnant women (81%, represented by 60% vitamin D insufficiency, 21% vitamin deficiency) and in normotensive pregnant women (69%, represented by 58% insufficiency, 11% deficiency). However, a vitamin D deficient status was more frequent among GH women. Indeed, vitamin D deficiency associated with a near three fold-increased risk of hypertension during pregnancy, while a normal vitamin D status is protective since it reduces by twofold the risk for GH development. Most importantly, a normal vitamin D status is protective against the negative effects of VDR FokI/BsmI haplotypes increasing the risk for GH, while vitamin D deficiency acts synergistically with unfavourable VDR genetic background to increase the risk of developing GH.

Despite being preliminary, these findings are promising and are worthy of replication studies in a larger population. As a whole, they suggest that vitamin D supplementation should be recommended during pregnancy, starting from the early gestation, and that genotyping of pregnant women for VDR polymorphisms may be useful for a tailored vitamin D supplementation strategy.

Supporting information
S1 File. DatabaseVDRGH, including VDR genotype and 25(OH)vitamin D3 levels of pregnant women recruited for this study. (XLSX)

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