



Menstrual and obstetrical bleeding in women with inherited platelet receptor defects—A systematic review

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Abstract

Introduction: Women with inherited platelet receptor defects (IPRD) may have an increased risk of heavy menstrual bleeding (HMB) and postpartum haemorrhage (PPH).

Aim: To present a systematic overview of the literature on the prevalence and management of menstrual and obstetrical bleeding in women with IPRD.

Methods: Electronic databases were searched for original patient data on the prevalence and management of HMB and PPH in women with known IPRD or who were being investigated for IPRD.

Results: Sixty-nine papers (61 case reports/series and 8 cohort studies) were included. Overall, studies were rated as 'poor quality'. The included cohort studies reported HMB in 25% (13/52) of women with Bernard-Soulier syndrome and in 22.1% (34/154) of women with Glanzmann thrombasthenia. In total, 164 deliveries in women with IPRD were described. Excessive bleeding occurred in 16.9% (11/65) of deliveries described in the largest cohort. PPH occurred in 63.2% (55/87) of deliveries described in case reports/series. PPH occurred in 73.7% (14/19) of deliveries that were not covered by prophylaxis compared with 54.2% (32/59) of deliveries that were (OR = 2.36, 95% CI 0.75-7.40). Neonatal bleeding complications were reported in 10.0% (8/80) of deliveries. In all (6/6) deliveries with neonatal bleeding complications wherein the presence of alloantibodies was investigated, either antiplatelet or anti-HLA antibodies were detected.

Discussion/Conclusion: Menstrual and particularly obstetrical bleeding problems frequently occur in women with IPRD, based on small case reports and series of poor quality. International collaboration, preferably on prospective studies, is needed to improve clinical management of women-specific bleeding in IPRD.

KEYWORDS

Bernard-Soulier syndrome, Glanzmann thrombasthenia, gynaecological bleeding, heavy menstrual bleeding, inherited platelet receptor defects, obstetrical bleeding, postpartum haemorrhage

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1 | INTRODUCTION

Inherited platelet receptor defects (IPRD) are a subgroup of inherited platelet disorders (IPD) characterized by qualitative or quantitative deficiencies in platelet membrane receptors.¹ The group comprises six disorders: Glanzmann thrombasthenia (GT), Bernard-Soulier syndrome (BSS), platelet-type von Willebrand disease (PT-VWD), ADP receptor (P2Y12) defects, thromboxane A2 receptor (TxA2) defects and glycoprotein VI (GPVI) deficiency. These receptor defects cause impaired primary haemostasis at the site of a blood vessel injury, often leading to a bleeding diathesis.¹ In Western countries, the population prevalence of GT and BSS is approximately 1 in 1 000 000.² Prevalence of the other IPRD is unknown, but less than a hundred affected individuals with TxA2 and P2Y12 deficiency worldwide have been reported in the literature, unlikely reflecting the true prevalence.³ PT-VWD and TxA2 defects are inherited in an autosomal dominant pattern, whereas GT, BSS, P2Y12 defects and GPVI deficiency are all autosomal recessive disorders. GT and BSS are more common in populations with a high incidence of consanguineous marriages, such as some communities in Iran.³

Clinical manifestations are diverse and may vary from recurrent mucocutaneous bleeding such as epistaxis, gingival bleeding, petechiae and abnormal bruising to less common but potentially fatal manifestations such as intracranial and postpartum haemorrhage (PPH).³ Moreover, affected women often suffer from heavy menstrual bleeding (HMB).⁴⁻⁶ HMB in women with IPRD can be challenging to manage and has been associated with a reduced quality of life.^{7,8} Furthermore, women appear to have an increased risk of develop PPH. Previous reviews on pregnancy outcomes in women with GT and BSS report a PPH prevalence of 34% and 33%, respectively.^{9,10} These numbers are generally higher than the population prevalence of PPH, which is approximately 19%.^{11,12} Neonates are at increased risk of bleeding, such as intracranial haemorrhage and subsequent death, due to placental transfer of maternal antiplatelet antibodies directed against foetal thrombocytes.³ Thus, pregnancy in women with IPRD can cause severe maternal and neonatal morbidity and even mortality.

Accurate data on the prevalence of HMB and PPH in women with IPRD are lacking, and clinical guidelines on the management are currently based on empirical evidence, as the separate IPRD are rare and randomized controlled trials have not been performed.¹³ In 2006, the UKHCDO formulated guidelines for pregnancy in women with severe platelet disorders, including GT and BSS.³ Most of these guidelines were based on expert opinion and case reports. In a review paper by Grainger *et al* (2018), these guidelines were re-evaluated, yet no consistent approach to the prevention or treatment of bleeding complications is provided.²

The aim of this paper is to present an overview of the literature on the prevalence and management of menstrual and obstetrical bleeding in women with IPRD to provide a complete literature background aiding clinical decision-making.

2 | METHODS

2.1 | Protocol and registration

The protocol details for this systematic review can be found in the International Prospective Register of Systematic Reviews (registration number CRD42018115116).¹⁴ This review was conducted in accordance with the Preferred Reporting Items for Systematic Review and Meta-Analyses (PRISMA) statement and Cochrane methods.^{15,16} The database search covered all IPD, but due to the amount of data, the authors decided to focus on IPRD in this review.

2.2 | Review questions

This systematic review addresses the following research questions: (a) What has been published on the prevalence of HMB in women with IPRD? (b) What has been published on the prevalence of pregnancy-related bleeding in women with IPRD reported in literature? (c) Which management strategies have been published on women with IPRD and HMB? and (d) Which management strategies have been published on women with IPRD and pregnancy-related bleeding?

2.3 | Inclusion and exclusion criteria

Studies were considered eligible for inclusion if they had an observational or interventional study design and included women already diagnosed with IPRD or women with HMB or PPH who were investigated for platelet receptor defects. Only studies in English or Dutch, concerning the prevalence and management of HMB and pregnancy-related bleeding and containing original patient data, were included. Abstracts, posters and articles without full-text access as well as articles that merely mentioned the prevalence of IPRD in a group of women with HMB were excluded.

2.4 | Data sources and search strategy

The electronic bibliographic databases PubMed, The Cochrane Library, Embase and CINAHL were searched up to the 16th of January 2019. A combination of search terms and MeSH/Emtree terms related to IPD, HMB and pregnancy was used (Appendix S1). No search limits were applied.

2.5 | Study selection

Search results were merged using Mendeley Reference Management Software, and duplicate references were removed using the Mendeley Deduplicate Tool and by hand. Two reviewers (PS and

MP) independently screened the titles and abstracts to identify potentially relevant articles. Subsequently, full-text papers were retrieved and assessed on eligibility by two independent reviewers (PS and MP). Whenever the full text of an article was unobtainable, the corresponding author was contacted once. Any difference of opinion between the reviewers concerning study selection was resolved by consulting a third reviewer (KG). Cross-referencing was conducted in the included studies and relevant reviews.

2.6 | Data extraction

One reviewer (PS) extracted data from the included articles using a standardized data collection form (Appendix S2). A second reviewer (MP) double-checked all articles for accuracy of data extraction. HMB was defined as a Pictorial Blood Loss Assessment Chart (PBAC) score > 100, reflecting at least one menstrual cycle.¹⁷ In the absence of PBAC scores, the author's definition of HMB was used. Primary PPH was defined as estimated blood loss \geq 500 mL occurring within 24 hours of delivery, and severe PPH was defined as estimated blood loss \geq 1000 mL.¹⁸ Secondary PPH was defined as excessive bleeding requiring medical attention between 24 hours and three months after delivery.¹⁸ Treatment of HMB or PPH could consist of desmopressin (DDAVP), iron supplements, hormonal treatment, antifibrinolytics, uterotonic agents, blood products, surgery or other interventions (eg fibrinogen concentrates, local compression devices and crystalloids). Prophylactic treatment for delivery and the postpartum period included measures taken to prevent PPH and could consist of blood

products (erythrocytes, platelets, plasma, coagulation factors, plasmapheresis), recombinant Factor VIIa (rFVIIa), DDAVP or other preventive measures (eg steroids, uterotonic agents, antifibrinolytics, fibrinogen concentrate and operative and invasive procedures such as preventive hysterectomy and embolization). Study design was based upon the following criteria: a cohort study in case all eligible patients during a certain time period were included in the study, and a case series if patient selection was not described.¹⁹ Any doubt regarding the extraction of data was resolved by consulting a third reviewer (KG).

2.7 | Quality assessment

One reviewer (PS) assessed the quality of each included study through an adjusted Chambers scale, and each assessment was checked by a second reviewer (MP) (Appendix S3).²⁰ This adjusted version of the Chambers scale including only criteria relevant to the included study designs was used to provide an illustrative reflection of the quality of small case series. Studies were rated as 'good', 'satisfactory' or 'poor'. Any doubt regarding quality assessment was resolved by consulting a third reviewer (KG).

2.8 | Data synthesis

The quantitative results were described separately for case reports and series and cohort studies. A descriptive overview was provided for prevalence and (prophylactic) treatment of HMB in women with

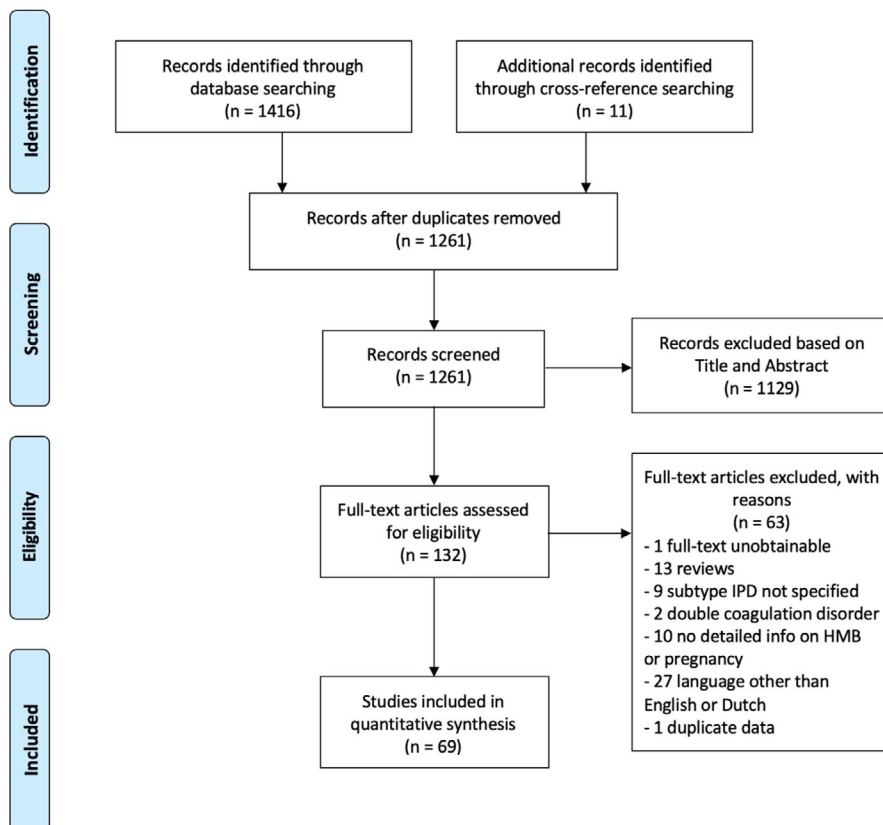


FIGURE 1 PRISMA flow diagram of study selection [Colour figure can be viewed at wileyonlinelibrary.com]

IPRD and for deliveries and the postpartum period in women with IPRD. Due to the limited amount of data on the different subtypes of IPRD, it was not possible to perform any subgroup analyses. A narrative synthesis on menstruation includes patient and menstruation characteristics and treatment of HMB. A narrative synthesis on pregnancy includes patient and pregnancy characteristics, prophylaxis for delivery, PPH and neonatal outcome.

3 | RESULTS

A total of 1261 articles were screened after duplicate removal and addition of eleven articles identified through cross-reference searching (Figure 1). One hundred and thirty-two articles were examined in full text, of which 69 were eventually included in this systematic review. These comprised 61 case reports and series^{13,21-80} and eight cohort studies.^{4,81-87} The exact number of women with HMB is uncertain due to incomplete reporting in one article.⁸⁷ In total, 164 deliveries in women with IPRD were described.

3.1 | Quality assessment

A summary of the quality assessment of the included studies according to the study population size is provided in Table 1. A complete overview of the quality assessment of each included study is provided in Appendix S4. None of the included studies were considered of high quality. Three studies (4.3%) were rated as 'satisfactory' and the remaining 66 studies (95.7%) were rated as 'poor'. Main limitations were seen across the entire quality assessment including the lack of diagnostic criteria, inadequate reporting of the outcome measures and the absence of reporting on prognostic/confounding factors.

3.2 | Narrative synthesis of results on menstruation

3.2.1 | Data from case reports and series

A total of 32 women with at least one episode of HMB were described in 18 case reports and series. These included 18 women with GT (56.3%) and 14 women with BSS (43.8%). No data were available on PT-VWD, TxA2, P2Y12 and GPVI deficiency. Mean age (\pm SD) was 19 years (7.67) in the 21 cases with available information. Median age at diagnosis of IPRD was 15 years (range 14 months – 50 years) in the 17 women for whom this was documented. HMB had been present since menarche in 80% (16/20) of cases in which the timing of the first manifestation of HMB was reported.

None of the individual cases used PBAC scores to diagnose HMB. HMB was defined as blood loss > 80 mL per menstrual cycle in 9.4% (3/32) of individual cases. Menstrual bleeding was described as 'mild' or 'moderate' in 9.4% (3/32) of women, described as 'prolonged', 'severe', 'excessive' or 'unrestrainable' in 65.6% (21/32) of

TABLE 1 Risk of bias assessment of small, medium and large studies using the adjusted Chambers scale

Study population size ^a	Percentage of studies in which criterion is fulfilled (% yes)						
	1. Were diagnostic criteria adequately reported?	2. Was the selected population representative of that seen in normal practice?	3. Was the outcome measure adequately reported? ^b	4. Was loss to follow-up reported or explained?	5. Were those included at baseline sufficiently long followed up? ^c	6. Were patients recruited prospectively?	7. Did the study report relevant prognostic/confounding factors?
Small (n = 1-10)	1.8	1.8	36.8	0.0	0.0	0.0	33.3
Medium (n = 11-50)	66.7	50.0	33.3	0.0	83.3	50.0	16.7
Large (n = 51-382)	83.3	100.0	33.3	0.0	100.0	50.0	33.3

Note: Overall quality rating: good (if the answer is 'yes' to all criteria); satisfactory (if the answer is 'yes' to 5-6 criteria); poor (if the answer is 'yes' to less than 5 criteria). In this review, overall quality of included studies is 'poor'.

References: small studies,^{20-37,39-61,63-77,79} medium studies,^{13,38,62,78,83,86} large studies.^{4,80-82,84,85}

^aRefers to all patients included in a study.

^bMeaning blood loss in mL for postpartum hemorrhage (PPH) and PBAC scores for heavy menstrual bleeding (HMB).

^cAt least three months for PPH and two consecutive menstrual cycles for HMB.

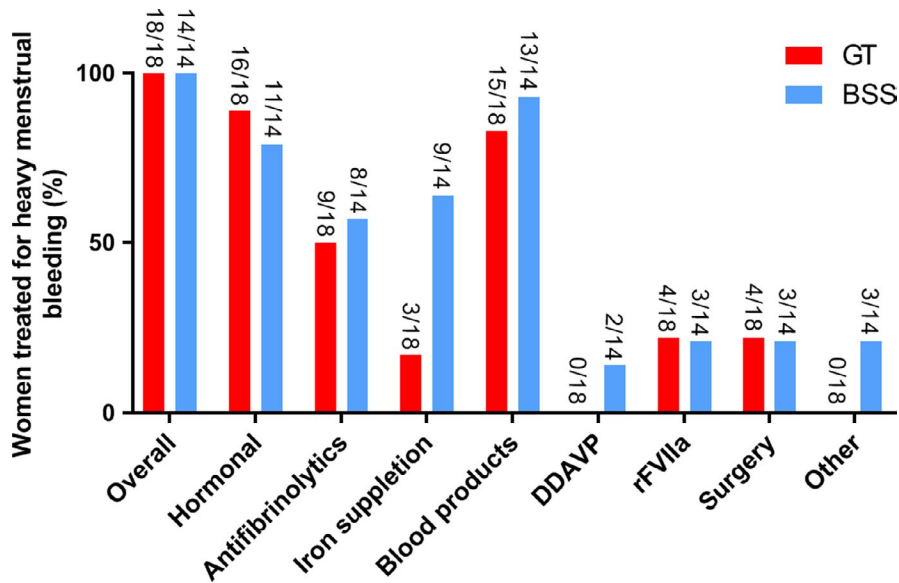


FIGURE 2 Treatment of heavy menstrual bleeding according to the subtype of inherited platelet receptor defect. GT, Glanzmann thrombasthenia; BSS, Bernard-Soulier syndrome; rFVIIa, recombinant Factor VIIa; DDAVP, desmopressin; Other = steroids and immunoglobulins [Colour figure can be viewed at wileyonlinelibrary.com]

women. In the remaining 15.6% (5/32) of cases, no other description than 'meno- or metrorrhagia' was given.

All included women received treatment for HMB (Figure 2). Most frequently prescribed treatments were blood products, hormonal therapy and antifibrinolytics, in 87.5% (28/32), 84.4% (27/32) and 53.1% (17/32) of women, respectively. Blood products consisted of red blood cell transfusions in 22 women, platelet transfusions in twenty women, whole blood transfusions in five women, plasma transfusions in four women and for one woman, the type of transfusion was unspecified. Two women were treated with DDAVP. They did not experience any adverse events such as hyponatremia due to this treatment. Surgery for HMB was performed in 21.9% (7/32) of women and mainly included thermal endometrial ablation and examination under anaesthesia (EUA) with uterine tamponade. Three (9.4%) women also received other forms of therapy including uterotonic agents, steroids and IVIG. No deaths due to HMB were reported.

3.2.2 | Data from cohort studies

Seven cohort studies with information about menstruation described a total of 189 women with GT and 58 women with BSS (Table 2).

Four cohort studies investigated the prevalence of HMB in women with IPRD.^{4,82,86,87} Overall, 25% (13/52) of women with BSS experienced at least one episode of HMB^{82,86} and 22.1% (34/154) of women with GT experienced HMB,^{4,86} but the exact number was uncertain due to incomplete reporting in one article.⁸⁷ Awidi *et al* and Vijapurkar *et al* report the following blood product use: 48% (12/25) of women together received 140 units of blood and 80 units of platelets for treatment of HMB during a total follow-up of 145 patient years and 11.8% (2/17) of women required more than a hundred units of platelets and whole blood transfusions over the course of more than twenty years, respectively.^{86,87}

Three cohort studies investigated the prevalence of IPRD in women with HMB.⁸³⁻⁸⁵ Out of 104 women who presented at the

emergency department in India with complaints of menorrhagia not caused by local pelvic pathology or hormonal disorders, 6.7% (7/104) was diagnosed with GT and 1.0% (1/104) with BSS (Table 2).

3.3 | Narrative synthesis of results on pregnancy

3.3.1 | Narrative synthesis of data from case reports and series

A total of 90 deliveries were described in 45 case reports and series in 66 women with IPRD. In 37.9% (25/66) of women, diagnosis had been made before pregnancy, in 6.1% (4/66) during pregnancy and in 6.1% (4/66) after pregnancy. For the remaining 50.0% (33/66) of women, the timing of diagnosis in relation to pregnancy was not stated. 10.6% (7/66) of women were known to be born to consanguineous parents. Bleeding history before pregnancy was documented in 84.8% (56/66) of women, and a bleeding tendency was reported in 96.4% (54/56) of women. The most common bleeding symptoms were epistaxis (48.1%), menorrhagia (46.3%), easy bruising (22.2%) and gingival bleeding (13.0%). Antenatal bleeding was reported in 41.2% (21/30) of pregnancies where the antepartum period was described. The mean maternal age at delivery was 28 years \pm 5.08 SD. Information on alloantibody formation (against HLA and/or the lacking glycoprotein) was available in 43.3% (39/90) of deliveries, and they were present in 66.7% (26/39). The mode of delivery was reported in 92.2% (83/90) of deliveries and comprised 39.8% (33/83) normal vaginal deliveries, 7.2% (6/83) assisted vaginal deliveries (3 forceps deliveries, 2 vacuum extractions and 1 unspecified instrumental delivery) and 53.0% (44/83) caesarean sections.

Peri- and postpartum prophylactic treatment

Information on prophylactic therapy administration was available in 88.9% (80/90) of individual cases. Overall, 75.0% (60/80) of deliveries were covered by some kind of prophylaxis and 25.0% (20/80) of

TABLE 2 Cohort studies on heavy menstrual bleeding in women with inherited platelet receptor defects

Author, year	Study design, study population, HMB definition	Results	HMB treatment
HMB in women with inherited platelet receptor defects			
Awidi (1992) ⁸⁷	Prospective cohort study 25 women with GT HMB definition unknown	HMB incidence unclear	48% (12/25) of women together received 140 units of blood and 80 units of platelets for treatment of HMB during a total follow-up of 145 years. In 57.1% (4/7) of women that used OCPs, HMB was successfully managed, whereas in 42.9% (3/7), the need for transfusions was unchanged
Vijapurkar (2009) ⁸⁶	Prevalence of HMB among Indian women with rare bleeding disorders HMB defined as > 80 mL blood loss per menstruation lasting more than seven days	HMB was reported in 66.7% (2/3) of women with BSS and in 77.3% (17/22) of women with GT. In two women, HMB led to iron deficiency anaemia despite the use of OCPs	Management mainly included tranexamic acid and OCPs. Both women with BSS received platelet and blood transfusions in combination with OCPs. 11.8% (2/17) of women required more than a hundred units of platelets and whole blood transfusions over the course of more than twenty years, one of whom was later efficaciously treated with endometrial ablation.
Toogeh (2010) ⁸²	Retrospective cohort study 49 Iranian women with BSS HMB definition unknown	22.4% (11/49) of women had had at least one episode of HMB.	Not described
Toogeh (2004) ⁴	Retrospective cohort study 132 Iranian women of reproductive age diagnosed with GT HMB definition unknown	12.9% (17/132) of women had had at least one episode of HMB	Not described
Inherited platelet receptor defects in women with HMB			
Kushwaha (2017) ⁸³	Prospective study 104 women who presented at the emergency department with complaints of menorrhagia in whom local pelvic pathology and hormonal disorders had been eliminated as a cause HMB definition not provided	Identified seven women with GT (6.7%) and one woman with BSS (1.0%)	Not described
Caki Kılıç (2013) ⁸⁵	Sixty women of reproductive age women who presented at the gynaecology and haematology outpatient and emergency care units HMB defined as > 80 mL blood loss per menstrual cycle	Identified two women with GT (3.3%) and three women with BSS (5.0%)	Detailed information on HMB treatment was not available, except that all included women received oral tranexamic acid and none of them underwent endometrial ablation or hysterectomy
Hutspardol (2010) ⁸⁴	28 adolescents with a history of menorrhagia HMB defined as > 80 mL blood loss per menstrual cycle	One woman with GT (3.6%) and two women with BSS (7.1%)	HMB was successfully treated using a combination of OCPs and tranexamic acid

Abbreviations: GT, Glanzmann thrombasthenia; BSS, Bernard-Soulier syndrome; HMB, heavy menstrual bleeding; OCP, oral contraceptive pills.

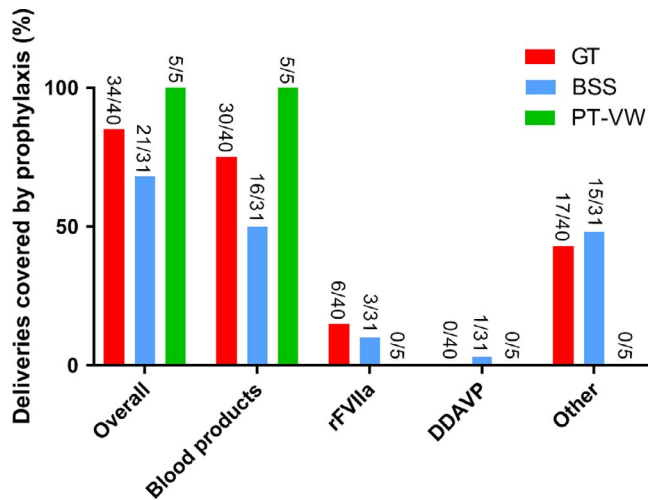


FIGURE 3 Prophylaxis administration according to the subtype of inherited platelet receptor defect. GT, Glanzmann thrombasthenia; BSS, Bernard-Soulier syndrome; PT-VWD, platelet-type von Willebrand disease; TxA2, thromboxane A2 receptor defects; P2Y12, ADP receptor defects. Prophylactic treatment for delivery is defined as measures taken to prevent PPH and could consist of blood products (erythrocytes, platelets, plasma, coagulation factors, plasmapheresis), recombinant Factor VIIa (rFVIIa), desmopressin (DDAVP), or other measures. Other = preventive hysterectomy, antifibrinolytics, uterotonic agents, fibrinogen concentrate and steroids [Colour figure can be viewed at wileyonlinelibrary.com]

deliveries were not (Figure 3). The included papers did not describe the reason for prophylaxis usage and, therefore, we could not distillate any differences between the group which received prophylactic treatment and the group which did not. Blood products were most often used as prophylaxis (85.0%, 51/60), followed by rFVIIa (15.0%, 9/60) and DDAVP (1.7%, 1/60). Blood products consisted of platelet transfusions in 47 cases, platelet plus red blood cell transfusions in two cases and platelet-rich plasma in two other cases. Other prophylactic therapies were given in 53.3% (32/60) of deliveries and included intravenous immunoglobulin (IVIG), steroids, oxytocin, antifibrinolytics and in one case a caesarean hysterectomy. In the latter case, this preventive measure was taken because the patient had anti-glycoprotein IB/IX antibodies, but there were no suitable platelets available if life-threatening bleeding would occur.⁶⁷

Postpartum haemorrhage

Information on the prevalence of PPH was available in 96.7% (87/90) of individual cases (Figure 4). In the 27.8% (25/90) of deliveries in which estimated blood loss during delivery was mentioned, median blood loss was 866 mL with a range of 50–2000 mL.^{23,25,27,30,32,34–42,44,48,49,54,56,76} In all twenty of these articles, the method of blood loss estimation (eg weighted or visual estimation) was not specified. Other studies defined PPH as blood loss \geq 500 mL without further specifying the exact amount of blood loss or elaboration on the definition of PPH.^{13,22,24,26,28,29,31,33,43,45,46,50–53,55,57,58,65,77–80}

Overall, PPH occurred in 63.2% (55/87) of deliveries. Severe PPH (\geq 1000 mL) occurred in 11.5% (10/87) of deliveries. Primary

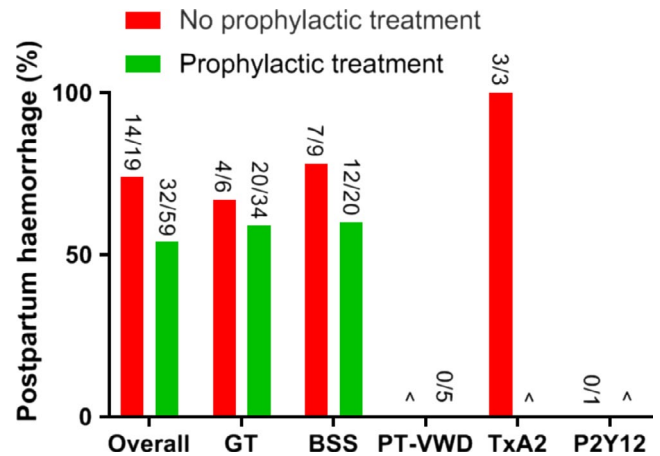


FIGURE 4 Prevalence of PPH according to the subtype of inherited platelet receptor defect and use of prophylactic treatment. ^No data available. GT, Glanzmann thrombasthenia; BSS, Bernard-Soulier syndrome; PT-VWD, platelet-type von Willebrand disease; TxA2, thromboxane A2 receptor defects; P2Y12, ADP receptor defects. Prophylactic treatment for delivery is defined as measures taken to prevent PPH and could consist of blood products (erythrocytes, platelets, plasma, coagulation factors, plasmapheresis), recombinant Factor VIIa (rFVIIa), desmopressin (DDAVP), or other measures (preventive hysterectomy, antifibrinolytics, uterotonic agents, fibrinogen concentrate and steroids). [Colour figure can be viewed at wileyonlinelibrary.com]

PPH was reported in 52.7% (29/55), secondary PPH in 47.3% (26/55), and in fifteen deliveries (27.3%), the timing of PPH was not stated. Both primary and secondary PPH were reported in 27.3% (15/55) of deliveries.

PPH occurred in 73.7% (14/19) of deliveries that were not covered by prophylaxis compared with 54.2% (32/59) of deliveries that were (odds ratio (OR) = 2.36, 95% confidence interval (CI): 0.75–7.40, $P = .14$). For caesarean sections and vaginal deliveries, PPH incidence was 64.7% (22/34) and 71.9% (23/32), respectively (OR = 0.72, 95% CI: 0.25–2.04, $P = .53$).

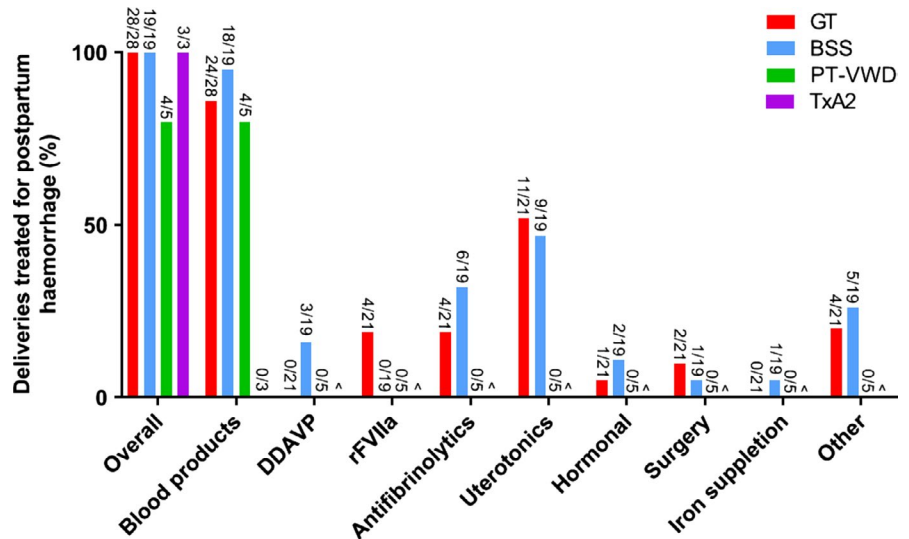
Treatment of PPH was described in 94.8% (55/58) of deliveries (Figure 5). The most common therapies included blood products (83.9%, 47/55), uterotonics (45.7%, 21/55) and antifibrinolytics (23.9%, 11/55). No thrombotic complications or maternal deaths due to PPH were reported in any of the individual cases.

Neonatal outcome

Neonatal bleeding complications were reported in 10.0% (8/80; maternal diagnosis being GT $n = 42$, BSS $n = 29$, PT-VWD $n = 5$, TxA2 $n = 3$ and P2Y12 $n = 1$) of deliveries with data on neonatal outcome. These included one foetal death due to severe third trimester gastrointestinal bleeding (maternal BSS),⁷⁹ one in utero foetal death due to intracranial haemorrhage (maternal GT),³⁸ one neonatal death six hours after delivery due to antepartum intracranial haemorrhage (maternal BSS),²¹ one neonatal death 48 hours after delivery due to haemolytic disease of the newborn (maternal GT),⁴⁷ one neonate suffering from bruising

FIGURE 5 Treatment of postpartum haemorrhage according to the subtype of inherited platelet receptor defect.

^No data available. GT, Glanzmann thrombasthenia; BSS, Bernard-Soulier syndrome; PT-VWD, platelet-type von Willebrand disease; TxA2, thromboxane A2 receptor defects; rFVIIa, recombinant Factor VIIa; DDAVP, desmopressin. Other = fibrinogen concentrates, local compression devices and crystalloids [Colour figure can be viewed at wileyonlinelibrary.com]



after vaginal delivery (maternal GT)⁴¹ and three neonates that developed petechiae and ecchymoses shortly after caesarean sections (maternal BSS in all three cases).²⁶ In 6/8 deliveries with neonatal bleeding complications, the presence of alloantibodies was investigated, and in all six, either antiplatelet or anti-HLA antibodies were present. None of these neonatal bleeding complications occurred during an instrumental delivery. Screening for postpartum neonatal thrombocytopenia was reported in 27 deliveries, of which the results were 'normal' or 'no thrombocytopenia' in fifteen neonates. A platelet count (range 0-307,000/ μ L, seven neonates with a platelet count of < 100,000/ μ L) was provided in twelve neonates.

3.3.2 | Narrative synthesis of data from cohort studies

Three cohort studies with information on postpartum blood loss described a total of 74 deliveries.^{4,81,82} These included 41 women with BSS and two women with PT-VWD. The exact number of women with GT is uncertain because of incomplete reporting, yet seven deliveries in women with GT were included.⁴

Noris *et al* (2014) conducted a retrospective, multicentre study evaluating 339 pregnancies in 181 women with different forms of inherited thrombocytopenia.⁸¹ They adapted two definitions of excessive bleeding at delivery: 'excessive bleedings requiring blood transfusion' (EBBT), which included transfusion of platelets and/or red blood cells, and 'all excessive bleedings' (AEB), which included all blood loss deemed larger than normal by the treating physician. Out of 21 deliveries in thirteen women with biallelic BSS (bBSS), EBBT was reported in 4.7% (1/21, 95% CI 0.1-23.8) and AEB in 14.2% (3/21, 95% CI 3-36.3). In 24 women with monoallelic BSS (mBSS), EBBT and AEB occurred in 2.5% (1/39, 95% CI 0.1-13.4) and 12.8% (5/39, 95% CI 4.2-27.4) of deliveries, respectively. Out of five deliveries in two women with PT-VWD, 20% (1/5, 95% CI 0.5-71.6) was complicated by EBBT and 60% (3/5, 95% CI 14.6-94.7) by AEB. 38.2% (21/55) of neonates

were affected by either bBSS, mBSS or PT-VWD; one of them (with mBSS) had postpartum bleeding symptoms in the form of petechiae.

A large single-centre retrospective study by Toogeh *et al* (2004) reported nine pregnancies in 131 investigated Iranian women of reproductive age with GT.⁴ Two out of nine pregnancies ended in spontaneous abortion, and one out of six caesarean sections was complicated by bleeding. No bleeding complications were reported in the one vaginal delivery. No information on prophylactic treatment or neonatal outcome was provided.

Another retrospective study by Toogeh *et al* (2010) described four pregnancies in 49 investigated Iranian women with BSS.⁸² Two (100%) vaginal deliveries were complicated by bleeding requiring a hysterectomy, while the remaining two pregnancies ended in spontaneous abortion.

4 | DISCUSSION

In this paper, we reviewed all published literature on the prevalence and management of menstrual and obstetrical bleeding in women with IPRD. This review provides sufficient data to assume an increased incidence of HMB in these women and highlights a markedly increased risk of peripartum maternal and neonatal bleeding.

The included cohort studies reported HMB in 25% (13/52) of women with BSS and in 22.1% (34/154) of women with GT. These numbers are higher than previously reported, yet previous HMB definitions are unclear. It is difficult to determine whether HMB can solely be attributed to the existing receptor defect, or whether gynaecological risk factors (eg uterine fibroids or ovulatory dysfunction) play a role too.⁸⁸ Nonetheless, the high rates of HMB observed in our study vastly exceed the estimated population prevalence of 8%-10% and call for increased awareness of HMB in women with IPRD.⁸⁹ More importantly, the high rate of transfusion of blood products to treat HMB, greatly burdens women with IPRD and the use of blood products is associated with high costs and risks, and should

therefore be minimized.^{86,87} The significant use of blood products may be explained by the lengthy period in which the included studies were published. Still, we believe greater effort should be made to move away from the use of blood products, where possible, to decrease the chance of alloantibody formation. For example, to prevent severe anemia due to HMB or PPH by timely administering hormonal treatment, antifibrinolytics, iron supplementation, rFVIIa and uterotinics. Currently, the optimal clinical management in this respect remains to be elucidated. In addition, HMB often required specifically platelets, with resultant high rates of alloimmunization, which in turn increases the risk of foetal complications in future pregnancies. Unfortunately, the majority of included cohort studies did not describe HMB treatment in detail, such as the use of rFVIIa, hormonal treatment and antifibrinolytics. Hence, we cannot draw any firm conclusions regarding the optimal management of HMB in women with IPRD.

Overall, PPH occurred in 63.2% (55/87) of deliveries in individual cases, which is vastly higher than previously reported in systematic reviews of pregnancy in GT and BSS.^{9,10} On the contrary, excessive bleeding occurred in only 16.9% (11/65) of deliveries described in the largest included cohort ($n = 39$ women).⁸¹ This difference may be explained by the effect of reporting and selection bias in case reports and series and the definition used by the authors of the mentioned cohort study (for example, abnormal bleeding based on the obstetrician's judgement). Visual estimation of blood loss has been proven to underestimate blood loss.⁹⁰ Known risk factors for PPH such as uterus atony, a prolonged second stage of labour, previous caesarean section and retained placenta, are potential confounders that may have affected our results, yet have been infrequently described.¹² Our data show the ongoing risk of PPH for women receiving prophylactic treatment of 54.2% (32/59), compared with a PPH incidence of 73.7% (14/19) in the non-prophylactic group, with both incidences being much higher than currently seen in the general population.⁹¹ Although the number of included deliveries is small, these data indicate that more women might benefit from prophylactic treatment and that current prophylaxis might be insufficient to prevent PPH. Furthermore, we observed a high rate of secondary PPH (47.3%, 26/55). These findings are in accordance with previously published results and highlight the need for optimizing peripartum management strategies.^{9,10} The difference in management strategies for vaginal deliveries and caesarean sections, as recommended by the UK guidelines, is not evident from the included literature in this review.³ The heterogeneity of the included study population, the few deliveries per type of treatment strategy and unknown obstetric confounders, makes subpopulation analysis and subsequent recommendations unreliable. For example, rFVIIa was mentioned as prophylaxis for PPH in only a few cases. The results of the current review are further limited given that the data were either relatively outdated, based on anecdotal evidence^{9,10} or lack objective uniform definitions of PPH.^{13,81}

This systematic review utilizes a thorough literature search in various online databases without publication date restrictions. The main limitations of this review relate to the low quality of evidence

and incomplete reporting. The majority of included studies were small case reports and series of low quality, susceptible to reporting and selection bias, more commonly seen in rare diseases. The included cohort studies were of higher quality, but generally less comprehensive in describing HMB and PPH management than the case reports and series. Moreover, most studies did not use objective or identical outcome measurements of HMB and PPH, such as PBAC scores and total blood loss in millilitres. Lastly, in some cases it was hard to distinguish whether a certain measure was used to prevent or treat PPH.

Our findings support current clinical guidelines that women with IPRD should be managed by a multidisciplinary team.³ Considering the remarkably high percentage (66.7%, 26/39 deliveries) of maternal alloimmunization in our study, and its strong association with neonatal bleeding complications, we underline the importance of compatible HLA-matched platelets and monitoring of alloantibody titres during pregnancy to prevent platelet refractoriness and foetal alloimmune thrombocytopenia. Furthermore, we would like to highlight the importance of optimization of prophylactic treatment, for example through continued administration of tranexamic acid and closer observation during the postpartum period. More women who currently do not receive prophylactic treatment might benefit from it.

We call for international collaboration to systematically gather data on the management of menstrual and obstetrical bleeding in women with IPRD. Large prospective studies using uniform objective definitions are needed to evaluate the true safety and efficacy of different management strategies to lay the foundation for evidence-based clinical guidelines.

5 | CONCLUSION

Literature on the management of menstrual and obstetrical bleeding in women with IPRD mainly comprises small case reports and series of poor quality. This review provides sufficient data to assume an increased incidence of HMB associated with an undesirable high consumption of blood products. There is a markedly increased risk of PPH, which is lowered with prophylactic haemostatic therapy, but still high. Neonates appear to be at risk of bleeding complications if antiplatelet or anti-HLA antibodies are present. Large prospective studies are required to lay the foundation for evidence-based guidelines on the management of HMB and delivery to reduce morbidity and mortality among women with IPRD and their newborns.

ACKNOWLEDGEMENTS

MC Punt designed and performed the research, analysed and interpreted the data and wrote the manuscript. PCE Schuitema performed the research, analysed and interpreted the data and wrote the manuscript. KPM van Galen designed the research, analysed and interpreted the data and critically reviewed the manuscript. KWM Bloemenkamp and ICL Kremer Hovinga interpreted the data and

critically reviewed the manuscript. All authors gave their consent to the final version of the manuscript.

DISCLOSURES

The authors stated that they had no interests which might be perceived as posing a conflict or bias.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.

How to cite this article: Punt MC, Schuitema PCE, Bloemenkamp KWM, Kremer Hovinga ICL, van Galen KPM. Menstrual and obstetrical bleeding in women with inherited platelet receptor defects—A systematic review. *Haemophilia*. 2020;26:216-227. <https://doi.org/10.1111/hae.13927>