Research Article

Incidence and Risk Factors of Severe Post-Partum Haemorrhage: A Nationwide Population-Based Study from a Hospital Database

Karine Goueslard[#]

Biostatistics and Bioinformatics (DIM), University Hospital, Bourgogne Franche-Comté University, Dijon, France

Mathilde Revert[#]

Versailles Saint Quentin University, France

Silvia Iacobelli

CHU La Réunion, Centre d'Études Périnatales de l'Océan Indien, Réanimation Néonatale et Pédiatrique, Néonatologie, Saint Pierre, CHU La Réunion

Jonathan Cottenet

Biostatistics and Bioinformatics (DIM), University Hospital, Bourgogne Franche-Comté University, Dijon, France

Adrien Roussot

Biostatistics and Bioinformatics (DIM), University Hospital, Bourgogne Franche-Comté University, Dijon, France

Evelyne Combier

Centre d'épidémiologie des populations, Univ de Bourgogne, Dijon, France

Catherine Quantin

INSERM, Dijon University Hospital, Clinical Investigation Center, clinical epidemiology/ clinical trials unit, Dijon, France

*Both the authors contributed equally to this work

ABSTRACT

Objective: The aim of this study was to ascertain the risk factors of major postpartum haemorrhage (PPH), as substantiated from diagnoses and procedures recorded in the French hospital administrative database.

Materials and methods: This nationwide populationbased study looked at 723 905 deliveries in mainland France in 2011, from the hospital database. The outcome of interest was major PPH identified by the International Classification of DiseasesICD-10 code for PPH in association with i) coding procedures for advanced measures for haemostasis or ii) need for intensive care or iii) death. Determinants studied included maternal and pregnancy characteristics, birth environmental factors and home-to-hospital distance. Adjusted odds ratios (aOR) for the outcome were calculated using a multilevel random-intercept model of logistic regression.

Results: Major PPH was associated with age \geq 35 years (aOR 1.41; 95% confidence interval, 95% CI [1.25–1.59]), multiple pregnancy (aOR 3.40 [2.85–4.05], pre-eclampsia (aOR 2.80 [2.32–3.38]), chorioamnionitis (aOR 2.57 [1.64–4.03]), caesarean section (aOR 4.80, [4.27-5.39]) and delivery in a level III maternity unit. Home-to-hospital distance was not a significant risk factor for PPH. The AUC was 0.8 [0.83-0.85].

Conclusion: The model used gave accurate predictions of PPH occurrence according to risk factors identifiable from a French hospital database.

Keywords: Post-partum haemorrhage; Incidence; Risk factors; Hospital data; Accessibility

What do we know?

Post-Partum Haemorrhage is recognized as a major public health problem. Several epidemiological studies have focused on maternal, obstetric and therapeutic predictors. Only one of these studies, in Norway, considered home-to-hospital distances and reported increased maternal morbidity with increased distance to the healthcare establishment. In France, it is an important issue as 23% of deliveries occur >30 min from the woman's home.

What does this paper add?

Home-to-hospital distance was not a significant risk factor for PPH even though we found a positive odds ratio gradient for suburban and rural women.

The major strength of this nationwide population-based study lies in the fact that we analysed several important maternal variables for all deliveries in a whole country and drew the same conclusion as those from studies based on medical records.

Contrary to models in previous studies, our model has a very strong predictive ability and revealed important implications for the screening of women at risk of severe PPH.

How this fits in with quality in primary care?

The major finding of this paper was the need for the exhaustive collection of the maternal characteristics in order to predict PPH requiring advanced interventional procedures. We then highlight the role of all primary care professionals in the pregnancy follow-up as their contribution to the collection of this information is essential. Coordination between outpatient and inpatient care should be strongly promoted in order to provide hospital practitioners with indicators of severe PPH in time.

Introduction

Post-Partum Haemorrhage (PPH) is recognized as a major public health problem by both clinicians and researchers, because it is the leading cause of maternal mortality around the world [1] and also contributes to substantial maternal morbidity [2].

When looking at data sources on the incidence of PPH in highresource countries, it appears that research in this area needs to be improved. One review recommended an improvement in data collection, in particular with regard to measurements of severity [3]. This work also reported that overall the incidence of PPH was increasing, especially because of severe PPH from uterine atony.

Atonic PPH is the leading cause of PPH, whatever the mode of delivery and the severity [4] and its severe forms are often unexpected and may occur in the absence of recognized risk factors [5]. Several epidemiological studies have focused on maternal, obstetric and therapeutic predictors [6]. None of these studies considered home-to-hospital distances. However, 23% of deliveries occur >30 min from the woman's home in France [7] and a Norwegian study reported increased maternal morbidity, including haemorrhage >1500 ml or blood transfusion, with increased distance to the healthcare establishment [8].

The French Health Authority (HAS) has stated that a decrease in mortality could be achieved by paying particular attention to severe forms, namely those requiring advanced interventional procedures. Recently, the HAS underlined that organizational aspects may contribute to maternal morbidity/mortality due to PPH and that they should be studied in order to make possible changes. As a result, all hospitals are now obliged to collect quality indicators for PPH, including those based on French hospital data.

Currently, each healthcare facility in France is legally obliged to produce a discharge abstract which codes all of the diagnoses made and procedures carried out. All of these abstracts are included in a French national administrative database, which provides a huge amount of epidemiological information concerning hospitalized patients in France [9-11]. The high proportion of in-hospital births in France makes this database particularly useful [10]. In the field of severe maternal morbidity, the validity of hospital data was assessed by comparing them with medical records in three teaching hospitals. Procedures for PPH reported in medical records showed a high degree of sensitivity and specificity in identifying severe PPH [12].

The aim of the present investigation was to ascertain the incidence and risk factors of severe atonic PPH in a nationwide, population-based study using diagnoses and procedures recorded in the hospital database.

Methods

Study design

The principle of this population-based retrospective cohort study was to examine hospital data for all deliveries in mainland France in 2011. This study was based on the hospital database, which gathers administrative and medical information from diagnoses coded according to the International Classification of Diseases (ICD-10) and from procedures coded according to the French Common Classification of Medical Procedures (CCMP), collected in a standardized fashion. To ensure the quality of data collection, quality control procedures are carried out *a posteriori* on sample data sets by medical inspectors and territorial medical advisors.

Setting

All deliveries in mainland France, recorded in the database with the codes Z37 of the ICD-10, noted in the women's abstract data, were examined. For the French agency responsible for collecting, hosting and analysing medical-economic data of French health establishments, Z37 codes are considered the most reliable and exhaustive to investigate deliveries. All deliveries were included, except for those that occurred in Paris, Lyon and Marseille: the "Assistance Publique-Hôpitaux de Paris (APHP)", "Hospices Civils de Lyon (HCL)" and "Assistance Publique-Marseille (APM)" all of which comprise many establishments scattered over several geographical sites. These three institutions each use a unique legal address identified by their specific legal number (FINESS). As we used this number to locate each individual birthplace, the above establishments were excluded from the analysis.

Parameters

The outcome of interest was cases of atonic severe PPH, which reflects major maternal morbidity, as substantiated by the need for advanced interventional procedures. In clinical practice, at the beginning of the haemorrhagic process, the "first-line" therapy includes manual removal of the placenta, uterine check, the infusion of crystalloid solutions and cardiopulmonary monitoring. If the haemorrhagic process continues and haemodynamic instability sets in, the second-line therapy is put in place. This consists of invasive techniques for haemostasis, blood transfusion or transfer to an intensive care unit.

In the present study, severe PPH was defined for one ICD-10 code (O72.0) linked to at least one of the following CCMP procedures, which correspond to second-line therapy and thus reflect the severity of the PPH: arterial embolization, uterine or hypogastric artery ligation, haemostasis hysterectomy and for the ICD-10 codes associated with transfer to an intensive care unit, or death [12]. Blood transfusion was not taken into account, as we showed in a recent report that the rate of transfusion is seriously underestimated in both pregnancy-related and birth-related abstracts in the French medico-administrative database [13].

As only atonic PPH was targeted in this study, we excluded all cases of severe PPH associated with one of the following major diagnoses: placenta praevia (codes O44), retroplacental haematoma (code O45), uterine rupture (codes O71.0 and O71.1).

Variables

The variables retained concerned the characteristics of the women and the healthcare establishments.

At the individual level, the following variables were considered: maternal age (<19, 20 to 34 and >35 years), parity, term, obesity, multiple pregnancy, preterm premature rupture of membranes, possible premature delivery, chronic or gestational high blood pressure, gestational diabetes, previous caesarean, prolonged labour, history of uterine scarring, preeclampsia, chorioamnionitis, mode of delivery, home-to-hospital distance.

The home-to-hospital distance was defined as 'the distance by road' in kilometres. For each delivery, the point of departure for the journey corresponded to the geographical coordinates (latitude, longitude) of the centroid of the district's main town derived from the geographic code recorded in the hospital data. The point of arrival corresponded to the geographical coordinates of the legal address of the establishment identified in hospital data by the legal number. The home-to-hospital distance was calculated using "Google Maps", which is able to calculate distance and journey time depending on the terrain, in standard traffic situations. In France, road network models are appropriate because almost all journeys take place by road. Some studies concluded that "Google Maps" is a useful tool for epidemiologists [14]: it proved to be an acceptable alternative to geocoded addresses [15] and to be reliable not only to calculate distances between postcode coordinates and care units [16], but also to predict with acceptable accuracy ambulance time of arrival at the emergency unit [17]. SAS software made it possible to automatically record requests to Google Maps.

For women domiciled in the same city as the maternity unit, we were unable to calculate home-to-hospital distance from hospital databases. For these, a distance of zero kilometres was attributed. Thus, the terms 'urban area' included all women who lived in an urban area according to the international standards and 'suburban area' included all women who lived in urban areas surrounding a town with an obstetric unit.-

Variables retained for the health establishment were those that may affect the management and have an impact on the severity of the PPH [5,18]. The type of maternity unit (I, II or III level of care), the annual number of deliveries, the number of beds and of delivery rooms, the number of staff (number of staff and full-time equivalents) were studied. To identify these elements, we used a compulsory exhaustive administrative survey that collected statements from public and private establishments.

Statistical analysis

Baseline characteristics of the participants are presented as means or proportions. The reference group included women without severe PPH reflecting major maternal morbidity. A mixed model, which took into account the hierarchical structure of the data, was used to assess associations with potential risk factors. As some women delivering in a given establishment had the same birth environment, the independence of the observations could not be confirmed. Hierarchical logistic regression was performed using the individual maternal variables as level 1 data and the establishment as level 2 data. Discrimination, which is the ability of the model to correctly predict severe PPH, was assessed by plotting the receiver operating characteristic (ROC) curve and calculating the area under the curve (AUC) [19]. An AUC value of 0.5 indicates no ability to discriminate, and larger values indicate increasing ability. A value of 0.8 is considered good.

Other than type of maternity unit, correlations between covariates of the birth environment were searched for. A French Decree established standard parameters based on the annual number of deliveries. As these parameters correlated with each other, the annual number of deliveries was retained for the analysis, because this information was more frequently available.

As distances in rural or urban areas could be very different, a sensitivity analysis was done. Descriptive analyses and multilevel analyses with the GLIMMIX procedure were performed using SAS 9.3. This study was approved by the National Committee for data protection (Commission Nationale de l'Informatique et des Libertés, registration number 1576793) and was conducted in accordance with French legislation. Written consent was not needed for this study. The PMSI database was transmitted by the national agency for the management of hospitalization data (ATIH number 2015-111111-47-33).

Results

The PMSI recorded 723,905 deliveries in 2011 in mainland France,

without APHP, HCL, APM. Among these, 1,393 severe PPH which reflects major maternal morbidity were identified (Incidence 0.19%), and 1,219 of these women (87.51%) underwent an invasive technique for haemostasis. Nearly 38% of them had arterial embolization, 32.4% ligature of the uterine or hypogastric arteries and 17.2% haemostasis hysterectomy.

The characteristics of the study population are presented in Table 1. There were more women aged over 35 years in the severe PPH group. Women presented a similar term of pregnancy in both groups. Many factors were significantly different between the two groups (Table 1). Twin pregnancy (12.06% vs. 1.57%), high blood pressure (13.5% vs. 3.1%), pre-eclampsia (10.7% vs. 1.5%) and previous caesarean (15.3% vs. 5.7%) were more frequent in the severe PPH group. The proportion of caesarean sections was also significantly higher in women with severe PPH (60.1% vs. 20.0%).

Table 1 presents the comparison between women with severe PPH and those without severe PPH with regard to the health establishment's characteristics. The frequency of severe atonic PPH was significantly lower in type I maternity units (14.5% vs. 30.4%) and type II units (37.3% vs. 48.4%), and much higher in type III units (47.8% vs. 21.1%).

The results of the multilevel logistic regression analysis are shown in Table 2. Age, multiple pregnancy, pre-eclampsia, chorioamnionitis, caesarean section and type III maternity unit were significantly associated with a higher risk of severe PPH reflecting major maternal morbidity, after adjusting for the other variables. Although the hometo-hospital distances were significantly different between the two groups in the bivariate analysis (p<0.001), the risk of severe PPH was not significantly affected by distance after adjustment for maternal variables and birth environment.

The results of the sensitivity analysis of home-to-hospital distance for women who lived in an urban, suburban or rural area are presented in Table 3. For suburban and rural women, when the home-to-hospital distance increased, we found a positive odds ratio gradient which nonetheless did not reach significance.

The AUC of the model in predicting severe PPH was 0.84, 95% Confidence Interval (0.83-0.85) (Figure 1). In fact, among the 1,393



Figure 1: ROC curve for the hierarchical model, severe postpartum haemorrhage in France in 2011.

ROC CURVE FOR THE FINAL MODEL

Catherine Quantin

Table 1: Pregnancy characteristics, adverse outcomes and environment of birth in France in 2011.								
	No sever	∙e PPH*	Severe PPH* (n=1,393)					
	(n=722	2,512)						
Term (Gestational Age), mean SD	39		38					
	Ν	%	Ν	%				
Age (years)								
≤19	16,941	2.34	28	2.01				
20-34	571,851	79.15	964	69.20				
≥ 35	133,72	18.51	401	28.79				
Area of residence								
Urban	411,63	61.57¥	879	68.46				
Rural	256,887	38.43¥	405	31.54				
Not documented	53,995	7.47	109	7.82				
Home to hospital distance [¥]								
<15 km	412,049	57.03	792	56.86				
15-29 km	173,475	24.01	285	20.46				
≥ 30 km	134,315	18.59	306	21.97				
Obesity	20,063***	2.81¥	85****	6.22				
Primiparous women	315,145	43.62	595	42.71				
Pregnancy								
Singleton pregnancy	710,985	98.41¥	1,219	87.51				
Multiple pregnancy	11,514	1.59¥	174	12.49				
Possible premature delivery	24,709	3.42 [¥]	97	6.96				
Haemorrhage during pregnancy	3,865	0.53¥	113	8.11				
High blood pressure	22,422	3.10 [¥]	188	13.5				
Gestational Diabetes	39,35	5.45¥	112	8.04				
Pre-eclampsia	10,953***	1.53 [¥]	147****	10.75				
Previous caesarean**	23,060***	5.73¥	120****	15.36				
Prolonged labour	80***	0.01	1****	0.07				
Mode of delivery								
Not documented	837	0.12¥	3	0.22				
Vaginal delivery	577,138	79.88 [¥]	552	39.62				
Caesarean	14,4537	20.00¥	838	60.16				
Elective caesarean section	24,066***	<i>3.37</i> [¥]	220****	16.09				
Emergency caesarean section	65,305***	<i>9.14</i> [¥]	371****	27.14				
Chorioamnionitis	1,851***	0.26^{F}	21****	1.54				
Type of establishment [¥]								
Туре 1	217,247ª	(30.36)	201 ^b	(14.47)				
Type 2	347,169ª	(48.42)	524 ^b	(37.32)				
Type 3	151,121ª	(21.12)	664 ^b	(47.80)				
Annual number of deliveries [¥]								
<500	40,664	(5.63)	40	(2.87)				
501-1000	113,298	(15.68)	109	(7.82)				
1001-2000	239,328	(33.12)	326	(23.40)				
2001-3000	177,866	(24.62)	436	(31.30)				
3001-4000	110,027	(15.23)	324	(23.26)				
4001-5000	30,411	(4.21)	128	(9.19)				
>5000	10,918	(1.51)	30	(2.15)				
* Post-Partum Haemorrhage	** Only multiparous women							
*** N=714,278 **** N=1,367	^a N=715,537 ^b N=1,389							
[¥] p<0.0001 versus comparative group with PPH								

58

	Seve	_	
	(n=	p-value	
	aOR	(95% CI)	
Age			< 0.0001
≤19	1.09	(0.74-1.62)	
20-34	1		
≥35	1.41	(1.25-1.59)	
Parity			0.41
Primiparous women	1		
Multiparous women	1.05	(0.94-1.17)	
Pregnancy			< 0.0001
Singleton	1		
Multiple	3.40	(2.85-4.05)	
Obesity	1.21	(0.96-1.52)	0.11
Pre-eclampsia	2.80	(2.32-3.38)	< 0.0001
Chorioamnionitis	2.57	(1.64-4.03)	< 0.0001
Mode of delivery			< 0.0001
Not documented	2.36	(0.58-9.54)	
Caesarean	4.80	(4.27-5.39)	
Vaginal delivery	1		
Home-to-hospital distance**			0.68
<15 km	0.96	(0,83-1.11)	
15-29 km	0.92	(0.78-1.10)	
>=30 km	1	(0.91-1.19)	
Annual number of deliveries			0.48
≤ 500	0.69	(0.36-1.31)	
501-1000	0.69	(0.48-0.99)	
1001-2000	0.83	(0.63-1.10)	
2001-3000	1		
3001-4000	0.84	(0.59-1.18)	
>4000	0.94	(0.54-1.62)	
Type of establishment			< 0.0001
1	0.27	(0.19-0.40)	
2	0.36	(0.26-0.49)	
3	1		
* Post-Partum Haemorrhage			
** Distance up to 160 km and 120 min			

Table 2: Association between severe PPH and maternal characteristics, adverse outcomes and birth environment in France in 2011.

women with severe PPH, only 191 (13.7%) had no risk factors identified by the model.

Discussion

This study used the national French hospital database, which includes the whole French population. It estimated an overall incidence rate for severe PPH due to uterine atony of 0.19%. This study also indicated that maternal age, multiple pregnancy, pre-eclampsia, chorioamnionitis, caesarean section and a type III maternity unit were associated with an increased risk of severe atonic PPH. We had very complete information on these factors. For example, only 0.11% of data regarding the mode of delivery were not documented. We could have excluded these women as the corresponding bias could be considered negligible. However, we decided to retain these missing data as a specific category. In the literature, data on the incidence of PPH due to uterine atony are heterogeneous, but the results of other studies that used procedures to define PPH are consistent with ours. In a nationwide inpatient sample in the USA, Kramer et al. reported an incidence rate of 0.3%, when severe haemorrhage was defined by PPH plus the receipt of a blood transfusion, hysterectomy and/or surgical repair of the uterus [20]. In 2008, using data from a computerized maternity database, Mousa et al. found an incidence closer to ours of 0.2%, for «major primary PPH» defined as «haemorrhage not responding to first-line treatment» [21].

59

Overall, the risk factors of severe PPH which reflects major maternal morbidity showed by our study correspond to the same risk factors found in previous studies carried out on medical records or hospital data. **Table 3:** Association between severe PPH and maternal characteristics, adverse outcomes and birth environment based on housing environment in France in 2011.

Outcome	Rural areas (N=390)		Severe PPH* Suburban areas (N=323)		Urban areas (N=827)	
	aOR	(95% CI)	aOR	(95% CI)	aOR	(95% CI)
Age (years)						
≤ 1 9	1.05	(0.49-2.24)	0.94	(0.30-2.95)	1.12	(0.69-1.79)
20-34	1		1		1	
≥35	1.51	(1.21 - 1.90)	1.31	(1.02-1.68)	1.33	(1.14-1.56)
Parity						
Primiparous women	1		1		1	
Multiparous women	0.89	(0.73 - 1.09)	1.23	(0,98-1.54)	1.14	(0.99-1.31)
Pregnancy						
Singleton	1		1		1	
Multiple	3.75	(2.78-5.04)	3.28	(2.32-4.64)	3.46	(2.77-4.33)
Obesity	1.09	(0.72 - 1.64)	1.48	(0.97 - 2.27)	1.27	(0.95 - 1.69)
Pre-eclampsia	2.68	(1.92-3.74)	3.30	(2.33-4.69)	2.77	(2.18-3.51)
Chorioamnionitis	1.69	(0.68-4.22)	2.34	(1.01-5.43)	2.89	(1.70-4.92)
Mode of delivery						
Vaginal delivery	1		1		1	
Caesarean	4.83	(3.90-5.99)	4.24	(3.35-5.37)	4.81	(4.15-5.56)
Not documented	4.82	(0.67-34.72)	3.89	(0.52-28.91)	1.74	(0.24-12.62)
Home-to-hospital distance						
<15 km	0.81	(0.60 - 1.08)	0.91	(0.67 - 1.23)	1.13	(0.77 - 1.26)
15-29 km	0.87	(0.69-1.09)	0.95	(0.71-1.28)	1.02	(0.76-1.37)
≥ 30 km	1		1		1	
Annual number of deliveries						
<500	0.96	(0.42 - 2.17)	0.92	(0.21-4.08)	1.19	(0.44 - 3.24)
500-1000	0.88	(0.53-1.45)	0.88	(0.48-1.63)	0.86	(0.55-1.35)
1001-2000	0.90	(0.61-1.33)	1.18	(0.80-1.75)	0.98	(0.72 - 1.34)
2001-3000	1		1		1	
3001-4000	0.67	(0.43-1.05)	0.97	(0.64-1.47)	0.87	(0.60-1.25)
>4000	0.86	(0.48-1.54)	1.04	(0.59-1.81)	0.92	(0.52 - 1.60)
Type of establishment						
1	0.17	(0.10-0.30)	0.33	(0.19-0.56)	0.27	(0.17-0.41)
2	0.24	(0.16-0.35)	0.42	(0.29-0.61)	0.38	(0.27-0.52)
3	1		1		1	
* Post-Partum Haemorrhage						
** Distance up to 160 km and	120 min					

Indeed, all of the papers retrieved from the literature showed an association between PPH or severe PPH and maternal age [6,20,22-24]. Kramer et al. reported an adjusted odds ratio (aOR) of 1.2 (1.2-1.3)

before 19 years and of 1.5 (1.5-1.6) beyond 35 years [20].

Because multiple pregnancies are associated with an enlarged uterus, they are classically considered a risk factor for PPH. In our study, multiple pregnancies presented an aOR of 3.40 versus singleton pregnancies. This is consistent with all of the studies that investigated PPH or severe PPH, as they reported OR from 2.8 to 3.4 for multiple versus singleton pregnancies [20,23].

Both pre-eclampsia and high blood pressure have been strongly associated with PPH in several reports, with OR varying between 1.5 and 5 [20,23]. We found an OR of 2.80 for pre-eclampsia. The blood loss that defines PPH in the case of vaginal birth is different from that for caesarean birth, and previous studies have shown that caesarean section increases the risk of PPH. Davis et al. estimated an OR equal to 3.59 [24], while Al-Zirqi et al. separated elective caesarean (OR=2.2) from emergency caesarean (OR=3.4) [6]. Similarly, Bateman et al. analysed caesarean without labour (OR=1.3) and caesarean with labour (OR=1.7) [25]. Our results revealed an OR of 4.80 for caesarean sections (60.0%) versus vaginal deliveries (20.0%). It is important to point out that data from the medico-administrative database showed not only the same risk factors described by previous studies based on medical records, but also that the proportions were the same, as illustrated by comparing the OR values found in several other studies with ours.

Our results showed that severe atonic PPH occurred less frequently in type I and type II maternity units than in type III units. We speculate that this finding may be the proof of the efficacy of the national maternity network in France, since it suggests that type I and II maternity units transfer the most severe cases and at-risk women to an establishment with an appropriate level of care. However, other interpretations are possible. Davis et al. [24] reported that the planned place of birth did not influence the risk of blood loss greater than 1,000 ml when adjusted for active management of labour compared with physiological management. This suggests that higher rates of severe haemorrhage could be explained by the active management of third stage of labor, usually more frequent in type III establishments.

The major result is the strong predictive ability of our model, which takes into account maternal characteristics and birth environment. This finding is new. Usually, the onset of PPH is considered unpredictable because of the lack of known predictors. However, our study showed that certain combinations of known risk factors accurately predicted the onset of PPH that required invasive procedures.

The major strength of this study lies in the fact that we analysed several important maternal variables for all deliveries in France. Z37 codes are able to identify deliveries in mainland France with a difference of 0.6% compared with INSEE data and of 0.3% compared with the civil registry, which records all births in our country [10].

Furthermore, French hospital data for the perinatal period are checked against various sources: medical records, national perinatal surveys or the civil registry. For the year 2010, hospital data, when compared with the national perinatal survey based on a representative sample of the French population, were able to properly recognize maternal age, multiple pregnancies, modes of delivery and birth weights [10]. Finally, since 2011, major changes to the data collection process have improved the quality of collection for other maternal morbidities, including high blood pressure and diabetes, especially in cases of delivery haemorrhage.

Even though the influence of medical practices was globally taken into account by level 2 of statistical model, our data did not allow us to adjust for each potential confounder as some of them were not recorded in our database – i.e., third-stage management of labour (active versus physiological), augmentation of labour (yes or no), place of birth (planned versus actual), duration of labour stages and oxytocin doses. As a result, we cannot draw any firm conclusions about the mechanisms underlying the association between severe PPH and obstetrical practices. This is one limitation. Other limitations concerned the absence from our database of data regarding socio-economic status.

Conclusion

The major strength of this nationwide population-based study lies in the fact that we analysed several important maternal variables for all deliveries in a whole country and drew the same conclusion as those from studies based on medical records. Contrary to models used in previous studies, our model has a very strong predictive ability and revealed important implications for the screening of women at risk of severe PPH.

In drawing the same conclusion as those from studies based on medical records, this study emphasizes the need for the exhaustive collection of the maternal characteristics in order to predict PPH requiring advanced interventional procedures. We then highlight the role of all primary care professionals in the pregnancy follow-up as their contribution to the collection of this information is essential. Coordination between outpatient and inpatient care should be strongly promoted in order to provide hospital practitioners with indicators of severe PPH in time.

Acknowledgement

This work was supported by the Fondation pour la Recherche Médicale, the Direction de la Recherche, des Etudes, de l'Evaluation et des Statistiques and Agence Nationale de la Recherche. We acknowledge E. Benzenine for his participation searching the "SAE" database and P. Bastable for his thorough proofreading of this article.

REFERENCES

 Say L, Chou D, Gemmill A, Tuncalp O, Moller AB, et al. Global causes of maternal death: A WHO systematic analysis. Lancet Glob Health 2014; 2: e323-e333.

- Rocha Filho EA, Costa ML, Cecatti JG, Parpinelli MA, Haddad SM, et al. Contribution of antepartum and intrapartum hemorrhage to the burden of maternal near miss and death in a national surveillance study. Acta Obstet Gynecol Scand 2015; 94: 50-58.
- Knight M, Callaghan WM, Berg C, Alexander S, Bouvier-Colle MH, et al. Trends in postpartum hemorrhage in high resource countries: A review and recommendations from the International Postpartum Hemorrhage Collaborative Group. BMC Pregnancy Childbirth 2009; 9: 55.
- Bouvier-Colle MH D-TC, Saucedo M, Comité National d'Experts sur la Mortalité Maternelle. Enquête Nationale confidentielle sur les morts maternelles France, 2007-2009. Les Morts Maternelles en France Mieux comprendre pour mieux prévenir. Inserm, Paris 2013.
- Mathai M, Gulmezoglu AM, Hill S. Saving womens lives: Evidence-based recommendations for the prevention of postpartum haemorrhage. Bull World Health Organ 2007; 85: 322-323.
- Al-Zirqi I, Vangen S, Forsen L, Stray-Pedersen B. Prevalence and risk factors of severe obstetric haemorrhage. BJOG 2008; 115: 1265-1272.
- 7. Baillot A EF. Les maternités: Un temps d'accès stable malgré les fermetures. Etudes et résultats DREES 2012
- Engjom HM, Morken NH, Norheim OF, Klungsoyr K. Availability and access in modern obstetric care: A retrospective population-based study. BJOG 2014; 121: 290-299.
- Lorgis L, Cottenet J, Molins G, Benzenine E, Zeller M, et al. Outcomes after acute myocardial infarction in HIV-infected patients: Analysis of data from a French nationwide hospital medical information database. Circulation 2013; 127: 1767-1774.
- Quantin C, Cottenet J, Vuagnat A, Prunet C, Mouquet MC, et al. Quality of perinatal statistics from hospital discharge data: comparison with civil registration and the 2010 National Perinatal Survey. J Gynecol Obstet Biol Reprod 2014; 43: 680-690.
- Gusmano M, Rodwin V, Weisz D, Cottenet J, Quantin C. Comparison of rehospitalization rates in France and the United States. J Health Serv Res Policy 2015; 20: 18-25.
- 12. Chantry AA, Deneux-Tharaux C, Cans C, Ego A, Quantin C, et al. Hospital discharge data can be used for monitoring procedures and intensive care related to severe maternal morbidity. J Clin Epidemiol 2011; 64: 1014-1022.
- 13. Quantin C, Benzenine E, Ferdynus C, Sediki M, Auverlot B, et al. Advantages and limitations of using national administrative data on obstetric blood transfusions to estimate the frequency of obstetric hemorrhages. J Public Health 2013; 35: 147-156.
- Berrazeg M, Diene S, Medjahed L, Parola P, Drissi M, et al. New Delhi Metallo-beta-lactamase around the world: An eReview using Google Maps. Euro Surveill 2014; 19.

62 Catherine Quantin

- 15. Dasgupta S, Vaughan AS, Kramer MR, Sanchez TH, Sullivan PS. Use of a google map tool embedded in an internet survey instrument: Is it a valid and reliable alternative to geocoded address data? JMIR Res Protoc 2014; 3: e24.
- 16. Raknes G, Hunskaar S. Method paper-Distance and travel time to casualty clinics in Norway based on crowdsourced postcode coordinates: A comparison with other methods. PLoS ONE 2014; 9: e89287.
- Fleischman RJ, Lundquist M, Jui J, Newgard CD, Warden C. Predicting ambulance time of arrival to the emergency department using global positioning system and Google maps. Prehosp Emerg Care 2013; 17: 458-465.
- 18. Messarat-Haddouche Z, Leleu H, Nitenberg G, Couralet M, Minvielle E, et al. Development and validation of indicators relating to the quality of prevention and early management of post-partum haemorrhage (COMPAQ-HPST research project). J Gynecol Obstet Biol Reprod 2012; 41: 271-278.
- Hanley JA, McNeil BJ. The meaning and use of the area under a receiver operating characteristic (ROC) curve. Radiology 1982; 14: 29-36.
- 20. Kramer MS, Berg C, Abenhaim H, Dahhou M, Rouleau J, et

al. Incidence, risk factors and temporal trends in severe postpartum hemorrhage. Am J Obstet Gynecol 2013; 209: e1-7.

- 21. Mousa HA, Cording V, Alfirevic Z. Risk factors and interventions associated with major primary postpartum hemorrhage unresponsive to first-line conventional therapy. Acta Obstet Gynecol Scand 2008; 87: 652-661.
- 22. Mehrabadi A, Hutcheon JA, Lee L, Kramer MS, Liston RM, et al. Epidemiological investigation of a temporal increase in atonic postpartum haemorrhage: A population-based retrospective cohort study. BJOG 2013; 120: 853-862.
- Combs CA, Murphy EL, Laros RK. Factors associated with post-partum hemorrhage with vaginal birth. Obstet Gynecol 1991; 77: 69-76.
- 24. Davis D, Baddock S, Pairman S, Hunter M, Benn C, et al. Risk of severe postpartum hemorrhage in low-risk childbearing women in New Zealand: Exploring the effect of place of birth and comparing third stage management of labor. Birth 2012; 39: 98-105.
- 25. Bateman BT, Berman MF, Riley LE, Leffert LR. The epidemiology of postpartum hemorrhage in a large, nationwide sample of deliveries. Anesth Analg 2010; 110: 1368-1373.

ADDRESS FOR CORRESPONDENCE:

Professeur Catherine Quantin, Service de Biostatistique et Informatique Médicale, Centre Hospitalier Universitaire, BP 77908, 21079 DIJON CEDEX, France, Tel: 33 3 80 29 36 29; Fax: 33 3 80 29 39 73; E-mail: catherine.quantin@chu-dijon.fr

Submitted: April 11, 2016; Accepted: April 20, 2017; Published: April 27, 2017