

Indications leading to termination of pregnancy between 22+0 and 31+6 weeks of gestational age in France: A population-based cohort study

Isabelle Monier, Nathalie Lelong, Pierre-Yves Ancel, Alexandra Benachi, Babak Khoshnood, Jennifer Zeitlin, Béatrice Blondel

▶ To cite this version:

Isabelle Monier, Nathalie Lelong, Pierre-Yves Ancel, Alexandra Benachi, Babak Khoshnood, et al.. Indications leading to termination of pregnancy between 22+0 and 31+6 weeks of gestational age in France: A population-based cohort study. European Journal of Obstetrics & Gynecology and Reproductive Biology, 2019, 233, pp.12-18. 10.1016/j.ejogrb.2018.11.021. hal-04511560

HAL Id: hal-04511560 https://hal.science/hal-04511560v1

Submitted on 26 Mar 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

- 1 Indications leading to termination of pregnancy between 22⁺⁰ and 31⁺⁶ weeks of
- 2 gestational age in France: a population-based cohort study

3

- 4 Isabelle Monier, a, b Nathalie Lelong, Pierre-Yves Ancel, Alexandra Benachi, Babak
- 5 Khoshnood, a Jennifer Zeitlin, Béatrice Blondel

6

- 7 Affiliations
- 8 a Inserm UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research
- 9 Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, DHU
- 10 Risks in pregnancy, Paris Descartes University, Paris, France.
- 11 b Antoine Beclere Maternity Unit, Department of Obstetrics and Gynaecology,
- 12 University Paris Sud, AP-HP, Paris, France

13

14

- 15 Corresponding Author
- 16 Isabelle Monier
- 17 Inserm UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team
- 18 (Epopé)
- 19 Port Royal Maternity Unit, 53 Avenue de l'Observatoire, 75014 Paris, France
- 20 Tel: +33 (0)1 42 34 55 86
- 21 Fax: +33 (0)1 43 26 89 79
- 22 E-mail: isabelle.monier@inserm.fr

23

24 Word count: Abstract: 258 Main text: 2533

Abstract

27

50

26

Indications leading to termination of pregnancy between 22⁺⁰ and 31⁺⁶ weeks of 28 gestational age in France: a population-based cohort study 29 Isabelle Monier, Nathalie Lelong, Pierre-Yves Ancel, Alexandra Benachi, Babak 30 Khoshnood, Jennifer Zeitlin, Béatrice Blondel 31 32 **Objective:** To estimate the prevalence and indications of terminations of pregnancy 33 (TOP) between 22⁺⁰ and 31⁺⁶ weeks of gestational age in France and to examine the 34 35 characteristics of women by indication of TOP. Study design: From the EPIPAGE 2 population-based cohort study of preterm births 36 in France in 2011, we selected 5009 singleton live births, stillbirths and TOP that 37 38 occurred between 22 and 31 weeks. We estimated the proportion of TOP by gestational age. We then classified terminations by indications into 4 categories: fetal 39 anomalies (TOPFA), preterm premature rupture of the membranes (PPROM), 40 maternal conditions and fetal growth restriction (FGR). We also classified TOPFA by 41 type of anomaly. Maternal characteristics were compared between TOPFA and TOP 42 43 for maternal or fetal conditions without congenital anomaly. **Results:** 23.1% of all births and 54.3% of stillbirths were terminations. The proportion 44 of terminations was 36.9% of all births at 22 weeks, 50.2% at 24 weeks and <10% at 45 30-31 weeks. 85.8% of terminations were for fetal anomaly, 4.4% for PPROM, 6.1% 46 for maternal complications and 3.7% for severe FGR. Compared to women with a 47 TOPFA, those with a termination for maternal or fetal conditions were more often 48 nulliparous, single, African, obese, smokers and covered by non-standard insurance 49

for women in socially deprived circumstances.

Conclusion: In France, there is a high proportion of TOP of which 14% are for indications other than congenital anomalies. Because rates of terminations have an impact on very preterm birth and perinatal mortality rates, studies on pregnancy outcome should report all terminations, not only those for congenital anomalies. **Key-words:** terminations of pregnancy, preterm births, stillbirths, EPIPAGE 2

Introduction

76

77

78

79

80

81

82

83

84

85

86

87

88

89

90

91

Terminations of pregnancy (TOP) are pregnancies that end to fetal deaths due to multiple conditions, most often congenital anomalies if the child is expected to have a risk of serious handicap, or pregnancy complications if there is substantial risk for the mother's life.(1) Most TOP are carried out at early gestational ages.(2,3) In some countries, terminations are also possible at later gestational age, after 21 completed weeks of gestational age (GA). These differences in policies and practices relating to TOP can affect estimates of perinatal mortality rates, if TOP are included in perinatal mortality statistics, and can influence evaluations of medical practices and survival of very preterm births.(4-7) A study on births between 22 and 31 weeks GA showed that more than 20% of stillbirths were TOP in many European regions and this proportion reached more than 40% in regions from Italy and France.(6) Therefore. information on the prevalence of TOP by gestational age and indication is important for international comparisons of perinatal health indicators. However, national-level data on TOP are scarce and most existing studies have focused on TOP for congenital anomalies, (8–13) whereas TOP are also carried out for other pregnancy complications.(14-16)

93

94

95

96

97

98

99

100

92

In a recent study, France reported the highest proportion of terminations among stillbirths at or after 22 weeks of gestation in Europe.(5) This study used aggregate data and was not able to describe the medical decision-making processes leading to the high number of TOP in France. Therefore, we aimed to estimate the prevalence of TOP among very preterm births, describe the indications of terminations and study the maternal characteristics according to the indication for termination using a population-based study of all births from 22 to 34 weeks GA in France in 2011.

Methods

Study design and population

The EPIPAGE 2 study is a national population-based prospective cohort of infants born between 22⁺⁰ and 34⁺⁶ weeks GA in 25 French regions in 2011 (N=7804).(17) All live births, stillbirths and TOP were included during three different periods according to gestational age at birth: 8 months for births between 22-26 weeks GA, 6 months for those between 27-31 weeks GA and 5 weeks for those between 32-34 weeks GA. The study population included 5009 singleton births between 22⁺⁰ and 31⁺⁶ weeks GA in 415 maternity units. Multiple pregnancies were excluded because the indications for TOP can differ from singleton pregnancies. We also excluded births between 32 and 34 weeks GA because of the short recruitment period for these births.

Variables

Data were abstracted from medical charts using a standardized questionnaire. TOP were classified by medical indication into 4 exclusive and hierarchical classes: 1) fetal anomaly (TOPFA), 2) preterm premature rupture of the membranes (PPROM) defined as the rupture of membranes at least 12 hours before the beginning of labor, or anomaly of amniotic fluid, 3) maternal conditions including pregnancy complications related to diseases such as hypertensive and psychiatric disorders and 4) severe and isolated fetal growth restriction (FGR) with no hypertensive disorder and no apparent congenital anomaly.

We classified TOPFA according to the classification of multiple congenital anomaly (MCA) algorithm defined in the European Surveillance of Congenital Anomalies (EUROCAT) guidelines.(18) Congenital anomalies were classified into 8 exclusive groups following this order: 1) chromosomal anomalies, 2) genetic syndromes, 3) associated infections during pregnancy with fetal abnormalities (mainly toxoplasmosis and cytomegalovirus infections), 4) isolated neural tube defects, 5) isolated heart cardiac defects, 6) isolated renal disorders, 7) isolated other anomalies (eye, ear, face and neck anomalies, limb reduction defects, oro-facial clefts, digestive system, genital anomalies) and 8) potential multiple anomalies defined as two or more anomalies.(19) We created a ninth group for fetal anomalies with no information on the type of anomaly (or insufficient information on the type of anomaly to permit classification).

138

139

140

141

142

143

126

127

128

129

130

131

132

133

134

135

136

137

Maternal characteristics included maternal age, parity, family situation (cohabiting or single), country of birth, medical insurance, body mass index (BMI) before pregnancy and smoking during pregnancy. We used the type of medical insurance as an indicator of underprivileged situation: women have a non-standard insurance, if they are undocumented or have very low income.

144

145

146

147

148

149

150

Prenatal care and management of terminations in France

In France, an ultrasound is recommended at each trimester of pregnancy for women with no complication including an ultrasound screening for congenital anomalies between 22 and 24 weeks GA.(20) Terminations are allowed at any gestational age for congenital anomalies, medical conditions or social reasons if the continuation of the pregnancy would lead to risk for women's health or if the child is expected to

have a "serious illness, recognized as incurable at the time of diagnosis".(1)

Decisions for pregnancy terminations must be approved by a multidisciplinary

committee for prenatal diagnosis.

Statistical analysis

We estimated the proportion of TOP with 95% confidence interval by gestational age, as a percentage of all births and as a percentage of stillbirths. For these calculations, all births included live births, TOP and spontaneous fetal deaths, and stillbirths included TOP and spontaneous fetal deaths. We also reported the proportion of feticides by gestational age. Then, we compared maternal characteristics between TOP for congenital anomalies and TOP for maternal conditions or FGR, using Chi square tests. Finally, we estimated the percentage of all TOPFA by type of fetal anomaly. In order to compare our results with other studies, we also reported the percentage of TOPFA between 22 and 26 weeks and between 27 and 31 weeks GA.

Percentage were weighted to account for the 9-week longer recruitment period for births at 22-26 weeks GA. Analyses were performed using STATA 13.0 software (StataCorpLP, College Station, TX, USA).

Results

Among the 5009 births between 22⁺⁰ and 31⁺⁶ weeks GA, there were 1266 terminations (23.1%). (Table 1) The percentage of TOP was 36.9% (95% CI: 32.6-41.3) at 22 weeks; it increased to 50.2% (95% CI: 45.9-54.4) at 24 weeks GA and then decreased to less than 10% at 30-31 weeks GA. The percentage of TOP among stillbirths was 54.3% and it varied by gestational age from 35.9% to 70.4%. A feticide

was recorded in medical charts for 78.2% of terminations: 47.9% (95% CI: 40.5-55.5) at 22 weeks, 58.6% (95% CI: 51.7-65.1) at 23 weeks and more than 90% after 26 weeks. For instance, at 22 weeks GA, no mention of feticide was found for 48.8% of TOP for congenital anomalies, 60.9% of TOP for PPROM, 47% of TOP for maternal indications and 100% of TOP for severe FGR (n=4).

Table 2 presents the percentage of TOP by medical indication and gestational age. Overall, 85.8% of TOP were carried out for congenital anomaly, 4.4% for PPROM or anomaly of amniotic fluid, 6.1% for maternal complications and 3.7% for severe FGR. At 22-23 weeks GA, 78.4% of terminations were for congenital anomalies and this percentage increased with gestational age to 97.5% at 30-31 weeks GA. At these GA, the percentages of terminations for maternal conditions or FGR were 9.3% and 1.3% respectively. At 22 and 23 weeks of gestation, the 9 fetuses born after TOP carried out for isolated FGR had severe FGR with birthweight ranging between 110 and 350 grams.

Table 3 compared terminations for congenital anomaly to those for maternal indications or FGR. Women with terminations for maternal indications or FGR were often nulliparous (52.7% vs 40.8%, p=0.003), single (11.8% vs 4.7%, p<0.001), African (18.3% vs 7.6%, p<0.001), obese (16.6% vs 9.8%, p=0.012), smokers (22.9% vs 15.2%, p=0.011) and covered by non-standard medical insurance for pregnant women in socially deprived circumstances (12.1% vs 7.5%, p=0.049). No difference between the groups was found for maternal age.

Among TOP for fetal anomaly between 22 and 31 weeks, 22.6% (95% CI: 20.2-25.2) were for chromosomal abnormalities, 21.9% (95% CI: 19.5-24.5) for isolated neural tube defects, 13.8% (95% CI: 11.8-16.0) for isolated cardiac defects and 11.5% (95% CI: 9.7-13.6) for multiple anomalies. (Table 4) Among chromosomal anomalies, 44.2% were for Down's syndrome (Trisomy 21), 25.8% for Edward syndrome (Trisomy 18), 8.4% for Patau syndrome (Trisomy 13) and 21.6% for other chromosomal anomalies. Less than 10% of TOP for congenital anomaly were for genetic syndromes, infections or other isolated anomalies. These distributions were similar for TOPFA between 22 and 26 weeks and between 27 and 31 weeks GA.

Comment

Main findings

In France in 2011, terminations represented almost one quarter of all singleton births and half of stillbirths between 22 and 31 weeks GA. Most terminations were carried out for fetal anomaly, mainly for chromosomal abnormalities and isolated neural tube defects. About 15% of terminations were for maternal and fetal pregnancy complications unrelated to congenital anomalies but these indications were rare after 28 weeks GA. Women who had a termination for maternal or fetal pregnancy complications were more often nulliparous, obese, African, smokers and covered by non-standard medical insurance compared to those who had a termination for congenital anomaly.

Strengths and limitations

The principal strength of our study is its population-based prospective design. We included all TOP between 22 and 31 weeks GA in 25 out of 26 French regions. The inclusion of all eligible births was ascertained by members of the regional coordinating committee in all maternity units.(17) Further, we included terminations for other reasons than congenital anomalies and we studied maternal characteristics related to these terminations, whereas most previous studies have focused only on terminations for congenital anomalies.(8–13)

Our study also has some limitations. We did not include terminations at 32-34 weeks GA because of the shorter recruitment period and consequently the small number of cases (N=15) and we did not have any information on terminations after 34 weeks. Additionally, there may be some classification errors: for instance spontaneous intrauterine deaths classified as TOP if scheduled TOP ended in fetal death before termination, (21) or conversely, some terminations being coded as stillbirths.

Interpretation

We calculated that 23.1% of very preterm singleton births and 54.3% of stillbirths were terminations. These high proportions corroborate results from previous studies. For instance in 2003, in one French region, these proportions were 21.5% and 52.5% respectively whereas less than 6% of very preterm births were terminations in two regions in UK.(6) More generally, there is a high proportion of TOP regardless of gestational age in France, when compared to other European countries,(2) and this is especially true at low gestational ages.(5) Several factors contribute to this situation. France has an active policy of screening for congenital anomalies resulting in a high proportion of TOP among fetuses with congenital anomaly. For instance,

between 2002 and 2004, 96% and 98% of pregnancies diagnosed with a Down's syndrome and neural tube defects respectively ended in a termination.(7) In addition, terminations are allowed at any gestational age and this makes possible to wait for results of additional investigations to provide the best prognostic information to parents for their decision.(1) Finally, the second-trimester screening ultrasound is planned at 22-24 weeks, as opposed to before 20 weeks in many other countries, leading to a shift on the timing of terminations in France.

In our study, a feticide was not mentioned in medical charts for 48% of terminations at 22 weeks and 40% at 23 weeks of gestation. Some terminations without feticide may be explained because of absence of beating of fetal heart before the onset of labor or because some parents do not want a feticide. The information on the feticide may also have been forgotten in some medical files. In addition, there is no recommendation for feticide in France. This difference in medical guidelines may explain that no feticide was reported for 18% of terminations between 22 and 26 weeks in the EPICure study compared to 27.5% in our study.(21) Nevertheless, it seems that a feticide is routinely offered for terminations after 22 weeks GA in France.(22)

We found that women who had a termination for maternal or fetal compromise were different to those who had a termination for congenital anomaly. They were more often nulliparous, obese, African, with non-standard medical insurance and smokers. These characteristics are well-known risk factors for vascular diseases including preeclampsia and FGR.(23–25) We did not find any association with older maternal age which is known to be a major risk factor for hypertensive disorders during

pregnancy.(26) This result can be explained by the use of terminations for congenital anomaly as a comparison group, as older women also have an increased risk of having a fetus with a congenital anomaly.(27)

Chromosomal anomalies were the most frequent indication for terminations among all TOPFA (22.6%). Most of these anomalies were trisomies in our study as in UK(21) and more generally in Europe,(8) although there is an active policy of antenatal screening for Down's syndrome resulting in terminations before 22 weeks. We also found that isolated neural tube defects were a frequent indication. Between 22 and 26 weeks, 22.1% of terminations were carried out for isolated neural tube defects in our study and 31.2% in the EPICure study.(21) Several randomized controlled trials have demonstrated the efficacy of folic acid supplementation before pregnancy for the reduction of neural tube defects.(28) However, a population-based study showed that the prevalence of neural tube defects has not decreased in Europe during the last two decades.(29) This may be explained by difficulties in the implementation of folic acid supplementation, despite national recommendations issued in each country. In France, only 15% of pregnant women had folic acid before pregnancy in 2010.(30) In contrast, there was a decrease in the live birth prevalence of neural tube defects, as a result of prenatal diagnosis and terminations.(29)

In our study, we estimated that 15% of terminations were for maternal or fetal conditions. Maternal reasons were responsible of 6% of terminations. In 2003, 5% of terminations among very preterm births were for maternal reasons in one French region.(6) Two studies focusing on terminations for maternal conditions showed that maternal hypertensive disorders were the most frequent reason for these

terminations.(14,15) We also estimated that 3.7% of terminations were for severe isolated FGR. A study using the EPIPAGE 2 data showed that 11.6% of pregnancies suspected with early-onset FGR without a fetal anomaly resulted in terminations between 22 and 31 weeks and this proportion was higher when FGR was detected before 26 weeks of gestation.(31)

A high proportion of late terminations leads to increased preterm birth and stillbirth rates.(5,6,32) Less is known about the impact of late terminations on indicators of newborn and infant health, but some studies have shown an effect of policies and practices of terminations on mortality rates. Studies in Canada suggested that the increase in terminations for congenital anomaly between 2000 and 2010, especially at 20-21 weeks, led to a decrease in neonatal mortality rates and the prevalence of live-born infants with congenital anomalies.(4,12) Studies from countries where terminations are restricted also report higher rates of infant deaths associated with congenital anomalies.(6,33) These effects underscore the importance of taking into account regulations and medical practice regarding late terminations when comparing stillbirth and neonatal mortality rates between countries.

Conclusion

Terminations of pregnancy between 22 and 31 weeks GA represent a high proportion of singleton very preterm births and stillbirths in France. The inclusion of all terminations in studies, not only those for congenital anomalies, is needed to understand medical practices, variations in preterm birth and stillbirth rates as well as infant health indicators in the perinatal period.

Acknowledgements

324

325 We thank the members of the EPIPAGE 2 Study Group and all the regional teams for their participation in the study and for their contribution to data acquisition. We 326 acknowledge the collaborators of the EPIPAGE 2 Obstetric Writing Group including 327 the following: Pierre-Yves Ancel, MD, PhD (Inserm UMR 1153, Obstetrical, Perinatal 328 and Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and 329 Statistics Sorbonne Paris Cité, DHU Risks in pregnancy, Paris Descartes University, 330 Recherche Clinique – Centre d'Investigations Cliniques P1419, 331 Département Hospitalo-Universitaire Risks in Pregnancy, Cochin Hotel-Dieu 332 Hospital, Assistance Publique-Hôpitaux de Paris, Paris F-75014, France), Catherine 333 Arnaud, MD, PhD (Research Unit on Perinatal Epidemiology, Childhood Disabilities 334 and Adolescent Health, INSERM UMR 1027, Paul Sabatier University, Toulouse, 335 336 France), Julie Blanc, MD (Department of Obstetrics and Gynecology, Aix Marseille University, Marseille, France), Pascal Boileau, MD, PhD (Department of Neonatal 337 338 Pediatrics, Poissy Saint Germain Hospital, France, EA7285 Versailles Saint Quentin 339 en Yvelines University, France), Thierry Debillon, MD, PhD (Department of Neonatal Pediatrics, University Hospital, Grenoble, France), Pierre Delorme, MD, MSc (Inserm 340 UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team 341 (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département 342 Hospitalo-Universitaire Risks in pregnancy, Paris Descartes University, Department 343 of Obstetrics and Gynecology, Cochin, Broca, Hôtel Dieu Hospital, AP-HP, Paris, 344 France), Claude D'Ercole, MD (Department of Obstetrics and Gynecology, Nord 345 Hospital, Assistance Publique des Hôpitaux de Marseille (AP-HM), Aix Marseille 346 347 Université, AMU, Marseille, France), Thomas Desplanches, RM, MSc (Inserm UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team (Epopé), 348

349 Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-350 Universitaire Risks in pregnancy, Paris Descartes University), Caroline Diguisto, MD, MSc (Inserm UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research 351 352 Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in pregnancy, Paris Descartes University, 353 Maternité Olympe de Gouges, University Francois Rabelais, Tours, France), 354 Laurence Foix-L'Hélias, MD, PhD (Inserm UMR 1153, Obstetrical, Perinatal and 355 356 Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in 357 pregnancy, Paris Descartes University, Sorbonne Universités, University Pierre and 358 Marie CURIE Paris 06, Institut de Formation Doctorale, 4 Place Jussieu, 75252 359 PARIS cedex 05, Paris, France, Department of Neonatal Pediatrics, Trousseau 360 361 Hospital, AP-HP, Paris, France), Aurélie Garbi, MD (Department of Neonatology, Assistance Publique Hopitaux de Marseille, Marseille, France), Géraldine Gascoin, 362 363 MD, PhD (Department of Neonatal Medicine, Angers University Hospital, Angers, 364 France), Adrien Gaudineau, MD (Department of Obstetrics and Gynecology, Hautepierre Hospital, Strasbourg, France), Catherine Gire, MD (Department of 365 Neonatology, North Hospital, Marseille, France), François Goffinet, MD, PhD (Inserm 366 367 UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team 368 (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in pregnancy, Paris Descartes University, Department 369 of Obstetrics and Gynecology, Cochin, Broca, Hôtel Dieu Hospital, AP-HP, Paris, 370 France), Gilles Kayem, MD, PhD (Inserm UMR 1153, Obstetrical, Perinatal and 371 372 Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in 373

pregnancy, Paris Descartes University, Sorbonne Universités, University Pierre and 374 Marie CURIE Paris 06, Institut de Formation Doctorale, 4 Place Jussieu, 75252 375 PARIS cedex 05, Paris, France, Department of Obstetrics and Gynecology, 376 Trousseau Hospital, AP-HP, Paris, France), Bruno Langer, MD (Department of 377 Obstetrics and Gynecology, Hautepierre Hospital, Strasbourg, France), Mathilde 378 Letouzey, MD, MSc (Department of Neonatal Pediatrics, Poissy Saint Germain 379 Hospital, France), Elsa Lorthe, RM, PhD (Inserm UMR 1153, Obstetrical, Perinatal 380 381 and Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in 382 383 pregnancy, Paris Descartes University), Emeline Maisonneuve, MD, MSc (Department of Obstetrics and Gynecology, Trousseau Hospital, APHP, Paris, 384 France), Stéphane Marret, MD, PhD (Department of Neonatal Medicine, Rouen 385 386 University Hospital and Région-INSERM (ERI 28), Normandy University, Rouen, 387 France), Isabelle Monier, RM, PhD (Inserm UMR 1153, Obstetrical, Perinatal and 388 Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and 389 Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in pregnancy, Paris Descartes University), Andrei Morgan, MD, PhD (Inserm UMR 390 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team (Epopé), 391 392 Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-393 Universitaire Risks in pregnancy, Paris Descartes University), Jean-Christophe Rozé, 394 MD, PhD (Department of Neonatal Medicine, Nantes University Hospital, Nantes, 395 France, Epidémiologie Clinique, Centre d'Investigation Clinique (CIC004), Nantes 396 University Hospital, Nantes, France), Thomas Schmitz, MD, PhD (Inserm UMR 1153, 397 Obstetrical, Perinatal and Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-398

Universitaire Risks in pregnancy, Paris Descartes University, Department of Obstetrics and Gynecology, Robert Debré Hospital, Assistance Publique-Hôpitaux de Paris, Paris, France), Loïc Sentilhes, MD, PhD (Department of Obstetrics and Gynecology, Bordeaux University Hospital, Bordeaux, France), Damien Subtil, MD, PhD (Department of Obstetrics and Gynecology, Jeanne de Flandre Hospital, Lille, France). Héloïse Torchin, MD, MSc (Inserm UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in pregnancy, Paris Descartes University, Neonatal Medicine and Resuscitation Service in Port-Royal, Cochin Hospital, Assistance Publique-Hôpitaux de Paris, Paris, France), Barthélémy Tosello, MD (Department of Neonatology, Assistance Publique Hopitaux de Marseille, Marseille, France), Christophe Vayssière, MD, PhD (Department of Obstetrics and Gynecology, University Hospital, Toulouse, France, Research Unit on Perinatal Epidemiology, Childhood Disabilities and Adolescent Health, INSERM UMR 1027, Paul Sabatier University, Toulouse, France), Norbert Winer, MD, PhD (Department of Obstetrics and Gynecology, University Hospital, INRA, UMR 1280 Physiologie des adaptations nutritionnelles, Nantes, France), Jennifer Zeitlin (Inserm UMR 1153, Obstetrical, Perinatal and Pediatric Epidemiology Research Team (Epopé), Center for Epidemiology and Statistics Sorbonne Paris Cité, Département Hospitalo-Universitaire Risks in pregnancy, Paris Descartes University).

420

421

422

423

399

400

401

402

403

404

405

406

407

408

409

410

411

412

413

414

415

416

417

418

419

All the collaborators of the EPIPAGE 2 Obstetric writing group have no conflict of interest or compensation in relation with this article to disclose. All of them consented to such acknowledgment.

424 **Disclosure of interests** 425 The authors report no conflict of interest 426 427 **Authorship** 428 429 IM, JZ and BB analysed the data and drafted the manuscript. NL, AB, BK and PYA 430 participated in the interpretation of the results and made suggestions for revisions. 431 PYA contributed to the design of the study. 432 433 **Details of ethics approval** Ethics approval was received from the National Data Protection Authority (CNIL no. 434 911009) and Consultative Committee on the Treatment of Information on Personal 435 436 Health for Research Purposes (approval granted 18 November 2010; reference number 10.626) and Committee for the Protection of People Participating in 437 438 Biomedical Research (approval granted 18 March 2011, reference CPP SC-2873). 439 **Funding** 440 441 The EPIPAGE 2 study was funded and supported by: (1) the French Institute of Public Health Research/Institute of Public Health and its partners: the French Health 442 443 Ministry, the National Institute of Health and Medical Research (INSERM), the National Institute of Cancer, and the National Solidarity Fund for Autonomy (CNSA); 444 445 (2) the French EQUIPEX programme of investments in the future; and (3) the PREMUP Foundation. 446 447

References

- 459 1. French Public Health Code. Article L162-12 modify by the Law n°94-654 July 460 art. 13 JORF July 30th 1994. Available 461 https://www.legifrance.gouv.fr. [Last Accessed April 2017].
- 462 2. Garne E, Loane M, Dolk H, De Vigan C, Scarano G, Tucker D, et al. Prenatal diagnosis of severe structural congenital malformations in Europe. Ultrasound 463 Obstet Gynecol 2005 Jan;25(1):6-11. 464
- EUROCAT. EUROCAT Special Report: Prenatal Screening Policies in Europe 465 466 2010, EUROCAT Central Registry, University of Ulster. 2010. Available at: http://www.eurocat-network.eu/content/Special-Report-Prenatal-Screening-467 Policies.pdf. 468
- 469 Joseph KS, Kinniburgh B, Hutcheon JA, Mehrabadi A, Basso M, Davies C, et al. 470 Determinants of increases in stillbirth rates from 2000 to 2010. CMAJ 2013 May 14;185(8):E345-51. 471
- 472 Blondel B, Cuttini M, Hindori-Mohangoo AD, Gissler M, Loghi M, Prunet C, et al. How do late terminations of pregnancy affect comparisons of stillbirth rates in 473 474 Europe? Analyses of aggregated routine data from the Euro-Peristat Project. BJOG 2018 Jan;125(2):226-34. 475
- 476 Papiernik E, Zeitlin J, Delmas D, Draper ES, Gadzinowski J, Kunzel W, et al. Termination of pregnancy among very preterm births and its impact on very 477 preterm mortality: results from ten European population-based cohorts in the 478 MOSAIC study. BJOG 2008 Feb;115(3):361-8. 479
- Boyd PA, Devigan C, Khoshnood B, Loane M, Garne E, Dolk H, et al. Survey of 480 prenatal screening policies in Europe for structural malformations and 481 chromosome anomalies, and their impact on detection and termination rates for 482 neural tube defects and Down's syndrome. BJOG 2008 May;115(6):689-96. 483

- 484 8. Garne E, Khoshnood B, Loane M, Boyd P, Dolk H, Group EW. Termination of pregnancy for fetal anomaly after 23 weeks of gestation: a European register-based study. BJOG 2010 May;117(6):660–6.
- 9. Rauch ER, Smulian JC, DePrince K, Ananth CV, Marcella SW, New Jersey Fetal Abnormalities R. Pregnancy interruption after second trimester diagnosis of fetal structural anomalies: the New Jersey Fetal Abnormalities Registry. Am J Obstet Gynecol 2005 Oct;193(4):1492–7.
- 491 10. Khoshnood B, De Vigan C, Vodovar V, Goujard J, Goffinet F. A population-492 based evaluation of the impact of antenatal screening for Down's syndrome in 493 France, 1981-2000. BJOG 2004 May;111(5):485–90.
- 11. Tararbit K, Bui TT, Lelong N, Thieulin AC, Goffinet F, Khoshnood B. Clinical and socioeconomic predictors of pregnancy termination for fetuses with congenital heart defects: a population-based evaluation. Prenat Diagn 2013 Feb;33(2):179–86.
- 498 12. Liu S, Joseph KS, Kramer MS, Allen AC, Sauve R, Rusen ID, et al. Relationship 499 of prenatal diagnosis and pregnancy termination to overall infant mortality in 500 Canada. JAMA 2002 Mar 27;287(12):1561–7.
- 501 13. Vaknin Z, Ben-Ami I, Reish O, Herman A, Maymon R. Fetal abnormalities 502 leading to termination of singleton pregnancy: the 7-year experience of a single 503 medical center. Prenat Diagn 2006 Oct;26(10):938–43.
- 14. Van Eerden L, Van Oostwaard MF, Zeeman GG, Page-Christiaens GC, Pajkrt E,
 Duvekot JJ, et al. Terminating pregnancy for severe hypertension when the fetus
 is considered non-viable: a retrospective cohort study. Eur J Obstet Gynecol
 Reprod Biol 2016 Nov;206:22–6.
- 508 15. Van Eerden L, Zeeman GG, Page-Christiaens GC, Vandenbussche F, Oei SG, Scheepers HC, et al. Termination of pregnancy for maternal indications at the limits of fetal viability: a retrospective cohort study in the Dutch tertiary care centres. BMJ Open 2014 Jun 17;4(6):e005145.
- 512 16. Piel B, Azria E, Oury JF, Carbillon L, Mandelbrot L. [Terminations of pregnancy for maternal indications in the Paris area: a retrospective multicenter study in the period between the 2001 French law on termination of pregnancy and the new bioethics law]. J Gynecol Obstet Biol Reprod (Paris) 2013 Jun;42(4):342–50.
- 17. Ancel P-Y, Goffinet F, EPIPAGE 2 Writing Group. EPIPAGE 2: a preterm birth cohort in France in 2011. BMC Pediatr 2014 Apr 9;14:97.
- 18. EUROCAT Guide 1.3 and reference documents Instructions for the Registration and Surveillance of Congenital Anomalies. Available at: http://www.eurocat-network.eu/content/EUROCAT-Guide-1.3.pdf.
- 19. Calzolari E, Barisic I, Loane M, Morris J, Wellesley D, Dolk H, et al. Epidemiology of multiple congenital anomalies in Europe: a EUROCAT population-based registry study. Birth Defects Res A Clin Mol Teratol 2014 Apr;100(4):270–6.

- 525 20. Haute Autorité de Santé. Suivi et orientation des femmes enceintes en fonction des situations à risque identifiées. Recommandations professionnelles. 2007. 526
- 527 Available at: www.has-sante.fr [Last accessed February 2018].
- 528 21. Draper ES, Alfirevic Z, Stacey F, Hennessy E, Costeloe K, Group EPicS. An
- 529 investigation into the reporting and management of late terminations of
- pregnancy (between 22 +0 and 26 +6 weeks of gestation) within NHS Hospitals 530
- 531 in England in 2006: the EPICure preterm cohort study. BJOG 2012
- May;119(6):710-5. 532
- 22. Jouannic JM, Favre R, Benachi A, Laurichesse H, Office of the French 533
- 534 Federation of Multidisciplinary Centers For Prenatal D. [Answer to "Termination
- of pregnancy without feticide: a French national survey"]. J Gynecol Obstet Biol 535
- Reprod (Paris) 2016 Jan;45(1):97-8. 536
- 537 23. Kramer MS. Determinants of low birth weight: methodological assessment and
- 538 meta-analysis. Bull World Health Organ 1987;65(5):663–737.
- 539 24. Conde-Agudelo A, Belizan JM. Risk factors for pre-eclampsia in a large cohort of
- 540 Latin American and Caribbean women. BJOG 2000 Jan;107(1):75–83.
- 541 25. Samadi AR, Mayberry RM, Zaidi AA, Pleasant JC, McGhee N Jr, Rice RJ.
- 542 Maternal hypertension and associated pregnancy complications among African-
- 543 American and other women in the United States. Obstet Gynecol 1996
- 544 Apr;87(4):557–63.
- 26. Bartsch E, Medcalf KE, Park AL, Ray JG, High Risk of Pre-eclampsia 545
- Identification G. Clinical risk factors for pre-eclampsia determined in early 546
- 547 pregnancy: systematic review and meta-analysis of large cohort studies. BMJ
- 548 2016 Apr 19;353:i1753.
- 27. Hollier LM, Leveno KJ, Kelly MA, DD MCi, Cunningham FG. Maternal age and 549
- 550 malformations in singleton births. Obstet Gynecol 2000 Nov;96(5 Pt 1):701–6.
- 28. De-Regil LM, Pena-Rosas JP, Fernandez-Gaxiola AC, Rayco-Solon P. Effects 551
- and safety of periconceptional oral folate supplementation for preventing birth 552
- 553 defects. Cochrane Database Syst Rev 2015 Dec 14;(12):CD007950.
- 29. Khoshnood B, Loane M, de Walle H, Arriola L, Addor MC, Barisic I, et al. Long 554
- term trends in prevalence of neural tube defects in Europe: population based 555
- study. BMJ 2015 Nov 24;351:h5949. 556
- 557 30. Tort J, Lelong N, Prunet C, Khoshnood B, Blondel B. Maternal and health care
- 558 determinants of preconceptional use of folic acid supplementation in France:
- results from the 2010 National Perinatal Survey. BJOG 2013 Dec;120(13):1661-559
- 560 7.
- 31. Monier I, Ancel P-Y, Ego A, Guellec I, Jarreau P-H, Kaminski M, et al. 561
- 562 Gestational age at diagnosis of early-onset fetal growth restriction and impact on
- management and survival: a population-based cohort study. BJOG 2017 563
- 564 Nov;124(12):1899–906.

- 565 32. Delnord M, Hindori-Mohangoo AD, Smith LK, Szamotulska K, Richards JL, Deb-566 Rinker P, et al. Variations in very preterm birth rates in 30 high-income 567 countries: are valid international comparisons possible using routine data? 568 BJOG 2017 Apr;124(5):785–94.
- 569 33. Gatt M, England K, Grech V, Calleja N. Contribution of Congenital Anomalies to 570 Neonatal Mortality Rates in Malta. Paediatr Perinat Epidemiol 2015 571 Sep;29(5):401–6.

572

573