Asthma medication prescribing before, during and after pregnancy
a study in seven European regions

Charlton, Rachel A; Pierini, Anna; Klungsøyr, Kari; Neville, Amanda J; Jordan, Susan; de Jong-van den Berg, Lolke T W; Thayer, Daniel; Bos, H Jens; Puccini, Aurora; Hansen, Anne V; Gini, Rosa; Engeland, Anders; Nybo Andersen, Anne-Marie; Dolk, Helen; Garne, Ester

Published in:
B M J Open

DOI:
10.1136/bmjopen-2015-009237

Publication date:
2016

Document version
Publisher's PDF, also known as Version of record

Document license
CC BY-NC

Citation for published version (APA):
Asthma medication prescribing before, during and after pregnancy: a study in seven European regions


OBJECTIVES: To explore utilisation patterns of asthma medication prescribing before, during and after pregnancy as recorded in seven European population-based databases.

Design: A descriptive drug utilisation study.

Setting: 7 electronic healthcare databases in Denmark, Norway, the Netherlands, Italy (Emilia Romagna and Tuscany), Wales, and the Clinical Practice Research Datalink representing the rest of the UK.

Participants: All women with a pregnancy ending in a delivery that started and ended between 2004 and 2010, who had been present in the database for the year before, throughout or during the year following pregnancy.

Main outcome measures: The percentage of deliveries where the woman received an asthma medicine prescription, based on prescriptions issued (UK) or dispensed (non-UK), during the year before, throughout or during the year following pregnancy.

Results: In total, 1 165 435 deliveries were identified. The prevalence of asthma medication prescribing during pregnancy was highest in the UK and Wales databases (9.4% (CI95 9.3% to 9.6%) and 9.4% (CI95 9.1% to 9.6%), respectively) and lowest in the Norwegian database (3.7% (CI95 3.7% to 3.8%)). In the year before pregnancy, the prevalence of asthma medication prescribing remained constant in all regions. Prescribing levels peaked during the second trimester of pregnancy and were at their lowest during the 3-month period following delivery. A decline was observed, in all regions except the UK, in the prescribing of long-acting β-2-agonists during pregnancy. During the 7-year study period, there were only small changes in prescribing patterns.

Conclusions: Differences were found in the prevalence of prescribing of asthma medications during and surrounding pregnancy in Europe. Inhaled β-2 agonists and inhaled corticosteroids were, however, the most popular therapeutic regimens in all databases.
uncertainty surrounding the effects of treatment versus the effects of the disease itself. Asthma is initially managed with short-acting β-2-agonists (SABAs) for symptom relief in the case of reasonably well-controlled disease and with a step-up approach when disease control becomes reduced, with the addition of inhaled corticosteroids (ICS) and other medications to prevent and reduce inflammation of the airways. Studies evaluating the course of asthma symptoms during pregnancy have shown that approximately one-third of women find their symptoms improve, one-third get worse and one-third remain the same. The prevalence of asthma during pregnancy in Europe has been estimated at between 4% and 8%, making it one of the most common, potentially serious, medical complications in pregnancy. Estimates of asthma prevalence during pregnancy vary by geographic location, study setting, definition of asthma and time period. The overall prevalence of prescribed antiasthma therapy during pregnancy has been reported for the UK, Sweden, Norway and the Netherlands, but for many other regions of Europe it is unknown and few studies have looked at the prevalence of prescribing of individual classes of asthma medicine. This study aimed to describe the extent and nature of asthma medication prescribing, during pregnancy and in the year before and after pregnancy, between 2004 and 2010, using data from population-based electronic healthcare databases in Europe. This study forms part of EUROMediCAT, a Seventh Framework Programme study funded by the European Commission that aims to make more systematic use of electronic healthcare databases in combination with the EUROCAT congenital anomaly registry data for reproductive safety evaluation.

METHODS

Setting

Seven population-based electronic healthcare databases, which captured data on prescriptions and pregnancies, contributed to the study; two in Italy (Tuscany/Emilia Romagna), two in the UK (the Secure Anonymised Information Linkage (SAIL) Databank in Wales and the UK-wide Clinical Practice Research Datalink (CPRD) with data from Wales excluded), and one each in Denmark, the Netherlands and Norway (see online supplementary table S1). A more detailed description of the databases can be found elsewhere. Where multiple databases were linked, such as in Denmark, where data from the Danish National Patient Register were linked to the Danish Prescription Registry, for the remainder of this paper these linked databases will be referred to as a single database.

Data extraction

A common protocol was implemented across databases. All pregnancies were identified that started and ended between 1 January 2004 and 31 December 2010 (except for Denmark and Norway where inclusion dates were 1 January 2004 to 31 December 2009 and 1 January 2005 to 31 December 2010, respectively). Pregnancies were eligible for the study if they ended in a delivery (live or stillbirth) and the woman had been contributing data to the database capturing prescription data, throughout pregnancy and for a full year before the start of pregnancy and following delivery. Women were able to contribute more than one pregnancy to the study. For each eligible delivery, a best estimate of the first day of the last menstrual period was calculated as summarised in online supplementary table S1.

All prescriptions for an asthma medication recorded in the databases during the study period were identified. In the UK databases, this included prescriptions issued and in the non-UK databases this included only prescriptions actually dispensed. None of the databases captured asthma medications given directly to the patient during a hospital stay. In Denmark, Norway and the Netherlands, all asthma medication dispensed from a pharmacy was captured, regardless of who prescribed the medication and where the prescription was made. In the UK databases, prescriptions initiated by a specialist in a hospital outpatient department and private prescriptions were rarely recorded; these numbers were likely to be small as most subsequent repeat prescribing will have been undertaken in primary care and private practice is limited. In Italy, only prescriptions reimbursed by the Italian healthcare system were captured; this included all reimbursed prescriptions regardless of whether they were prescribed by a general practitioner (GP) or a specialist who was an employee of the healthcare system. Asthma medications identified within the databases included all SABA, long-acting β-2-agonists (LABAs), ICS, combined β-2-agonists and inhaled steroids in a fixed-dose combination, leukotriene receptor antagonists, cromoglicate and related therapy, antimuscarinic bronchodilators and theophylline. These were defined as products with an Anatomical Therapeutic Chemical (ATC) classification code starting R03. Prescriptions for oral glucocorticoids (ATC code H02AB), potentially used to treat an asthma exacerbation, were also identified. Since oral glucocorticoids can be prescribed for conditions other than asthma, only those prescribed in a 3-month time period during which an asthma medication was also prescribed were eligible for inclusion.

Analyses

The percentage of pregnancies where the women received a prescription for an asthma medication in each of the databases was calculated for the year leading up to pregnancy, during pregnancy and the year following pregnancy. Prescribing patterns were described for each pregnancy trimester and for 3-month time periods during the years before and after pregnancy. For each class of asthma medication, the specific products most
frequently prescribed were compared between regions. Changes in prescribing over calendar time were also described and compared. Sensitivity analyses were carried out restricting the analyses to those deliveries where the woman received a minimum of two prescriptions for an asthma medicine during the 33-month time period of interest.

RESULTS

Within the seven databases, 1,165,435 deliveries were identified (table 1). The mean maternal age at the start of pregnancy was highest in Emilia Romagna and lowest in Wales (32.3 and 27.7 years, respectively) (table 1). The mean maternal age in all of the regions did not vary substantially between those who received and those who did not receive a prescription for an asthma medicine. The prevalence of asthma medication prescribing during pregnancy was highest in the UK and Wales databases (9.4% (CI95 9.3% to 9.6%) and 9.4% (CI95 9.1% to 9.6%), respectively) and lowest in the Norwegian database (3.7% (CI95 3.7% to 3.8%)).

During the year before pregnancy, the prevalence of asthma medication prescribing was relatively constant in all regions (figure 1A). In Italy and Norway, a reduction in prescribing was observed during the first trimester of pregnancy. During pregnancy, prescribing levels peaked during the second trimester with the largest increases being observed in the UK/Wales. Prescribing was at its lowest during the 3-month period following delivery and then gradually increased, returning towards prepregnancy levels, during the remainder of the year following delivery. Of those who received a prescription for an asthma medication in the year before, during and/or the year after pregnancy, approximately 30% in the Netherlands, Wales and the rest of the UK received only a single prescription (data not shown).

The prescribing prevalence of SABAs in each of the 3-month time periods was lowest in Tuscany and highest in the UK databases (figure 2). In the UK, approximately 90% of women who received a prescription for an asthma medication during pregnancy received a prescription for a SABA, whereas in Denmark and Norway it was approximately 75% and in Italy it was 26%. These percentages were not dissimilar to those during the year before pregnancy. In all regions, with the exception of Denmark where terbutaline was the most popular, salbutamol accounted for the majority of SABA prescriptions (figure 2). SABAs in a fixed-dose combination with an

Table 1. Percentage of deliveries between 2004 and 2010 where the woman received ≥ 1 prescription for an asthma medicine during the year before pregnancy, during pregnancy or the year following pregnancy

<table>
<thead>
<tr>
<th>Country/Region</th>
<th>Number of eligible deliveries in the entire cohort</th>
<th>Eligible deliveries in the entire cohort</th>
<th>Number of prescriptions received during pregnancy or the year following pregnancy</th>
<th>Any of the pregnancy trimesters</th>
<th>One year before pregnancy</th>
<th>One year following pregnancy</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N</td>
<td>SD*</td>
<td>N</td>
<td>Per cent (95% CI)</td>
<td>N</td>
<td>Per cent (95% CI)</td>
</tr>
<tr>
<td>Denmark</td>
<td>320,846</td>
<td>30.0</td>
<td>16,722</td>
<td>5.2 (5.1 to 5.3)</td>
<td>13,988</td>
<td>4.4 (4.3 to 4.5)</td>
</tr>
<tr>
<td>Italy—Tuscany</td>
<td>157,916</td>
<td>31.8</td>
<td>13,240</td>
<td>8.4 (8.3 to 8.5)</td>
<td>8,400</td>
<td>5.3 (5.2 to 5.4)</td>
</tr>
<tr>
<td>Italy—Emilia</td>
<td>129,220</td>
<td>32.3</td>
<td>9,317</td>
<td>12.7 (12.5 to 12.9)</td>
<td>8,904</td>
<td>8.4 (8.3 to 8.6)</td>
</tr>
<tr>
<td>Norway</td>
<td>301,820</td>
<td>29.7</td>
<td>11,205</td>
<td>3.7 (3.7 to 3.8)</td>
<td>10,627</td>
<td>3.5 (3.5 to 3.6)</td>
</tr>
<tr>
<td>The Netherlands</td>
<td>182,920</td>
<td>30.2</td>
<td>11,205</td>
<td>3.7 (3.7 to 3.8)</td>
<td>10,627</td>
<td>3.5 (3.5 to 3.6)</td>
</tr>
<tr>
<td>UK*</td>
<td>182,920</td>
<td>31.1</td>
<td>17,646</td>
<td>9.7 (9.6 to 9.8)</td>
<td>17,269</td>
<td>9.4 (9.3 to 9.5)</td>
</tr>
<tr>
<td>Wales</td>
<td>301,820</td>
<td>32.2</td>
<td>19,131</td>
<td>6.3 (6.2 to 6.4)</td>
<td>16,913</td>
<td>5.9 (5.8 to 6.0)</td>
</tr>
<tr>
<td>Total across countries</td>
<td>1,165,435</td>
<td>31.0</td>
<td>105,282</td>
<td>8.9 (8.8 to 9.0)</td>
<td>96,849</td>
<td>8.1 (8.0 to 8.2)</td>
</tr>
</tbody>
</table>

*Excluding Wales to avoid duplication of pregnancies in the Welsh Secure Anonymised Information Linkage (SAIL) Databank.

ICS were rarely prescribed in all regions, with the Italian databases having the highest prescribing prevalence. In the UK, the SABA prescribing prevalence increased during the second trimester of pregnancy and was approximately 25% higher than during the first trimester.

Norway had the lowest prescribing prevalence of ICS and the UK had the highest (figure 3). Of women who received a prescription for an asthma medication during one of the pregnancy trimesters, approximately 50% in Norway, 60% in the UK and Denmark and 89% in Emilia Romagna received a prescription for an ICS during pregnancy. Italy was the only region where prescribing of ICSs was more common than the prescribing of SABAs. In the UK/Wales and Italian databases, beclometasone was the most commonly prescribed ICS, whereas in Denmark it was budesonide and in Norway beclometasone and budesonide were equally prescribed (figure 3). In the Netherlands, the prevalence of beclometasone prescribing during pregnancy was almost double the prevalence observed before and after pregnancy, with the increase coinciding with a reduction in the prescribing of fluticasone and other ICS in a fixed-dose combination with a LABA. Norway was the only region where the prevalence of prescribing of ICS in a fixed-dose combination with a LABA was higher than the prescribing of ICS products not as part of a fixed-dose combination.

During pregnancy, evidence of a reduction in the prescribing of LABAs, both alone and as part of a fixed-dose combination, was observed in Norway, the Netherlands and Italy (figure 4). In Norway and Italy, prescribing was approximately 50% lower during the second trimester of pregnancy compared with the 3-month period prior to the start of pregnancy; in the Netherlands, it was approximately 65% lower. Of those who received a prescription for a LABA during pregnancy, between 75.2% in Tuscany and 86.5% in Norway received it in a fixed-dose combination with an ICS. Salmeterol in a fixed-dose combination was the most commonly prescribed LABA product in the UK, Italy and the Netherlands, while in Denmark it was formoterol in combination with an ICS and in Norway both formoterol and salmeterol in fixed-dose combinations were prescribed to the same degree. This was not substantially different from the therapeutic pattern observed in non-pregnant women during prepregnancy.

The prescribing of other asthma medications was low in all regions and lowest in Denmark and Norway (figure 5). Cromones were rarely prescribed, although
the number of women receiving a prescription in Norway increased to 51 during the first trimester of pregnancy compared with 17 during the 3-month period before pregnancy. The percentage of deliveries where the mother received a prescription for a leukotriene receptor antagonist during pregnancy ranged from 0.04% (CI95 0.03% to 0.04%) in Norway to 0.14% (CI95 0.11% to 0.17%) in Wales, with all regions showing a decline in prescribing during the latter trimesters. No prescriptions for an anticholinergic were dispensed during the time period of interest in Denmark and the highest level of prescribing of these during pregnancy was observed in the Netherlands, with 0.16% (CI95 0.10% to 0.23%) of women receiving a prescription. Prescribing of oral glucocorticoids was considerably higher in the UK and Italy than other regions, although in all databases the levels of prescribing declined during pregnancy and increased steadily in the year following delivery. Prednisolone was the most commonly prescribed oral corticosteroid in all regions.

**DISCUSSION**

**Main findings**

This study found variations in the prevalence of asthma medication prescribing and the specific products most commonly prescribed to women before, during and after pregnancy in different regions of Europe. Changes in prescribing were observed during pregnancy, with a peak in prescribing in the second trimester, after which prescribing declined until 4–6 months after delivery. The prescribing of asthma medications was considerably higher in the UK/Wales and Emilia Romagna than in other regions; for the UK/Wales, the differences may be accentuated because these databases captured all prescriptions issued, while the other databases captured only those actually dispensed.

**Strengths and limitations**

This study captured over 65 000 deliveries, between 2004 and 2010, where the woman received a prescription for an asthma medication during her pregnancy. Prescription information was recorded independently by the prescriber or pharmacist, removing maternal recall bias. No data, however, were available on whether the
Figure 2 Prevalence of prescribing of short-acting β₂-agonists (SABAs) in the year before, during and after pregnancy in women with a live or stillbirth during 2004–2010 for women prescribed ≥1 prescription for an asthma medication during ≥1 of the time periods of interest.
Figure 3  Prevalence of prescribing of ICS in the year before, during and after pregnancy in women with a live or stillbirth during 2004–2010 for women prescribed ≥1 prescription for an asthma medication during ≥1 of the time periods of interest (ICS, inhaled corticosteroid; LABA, long-acting β-2-agonist; SABA, short-acting β-2-agonist).
Figure 4  Prevalence of prescribing of long-acting β2-agonists (LABAs) in the year before, during and after pregnancy in women with a live or stillbirth during 2004–2010 for women prescribed ≥1 prescription for an asthma medication during ≥1 of the time periods of interest.
women took the medication and whether it was administered as instructed. The often intermittent nature of asthma symptoms and the fact that some treatments can be used on an ‘as needed’ basis makes determining the precise timing of exposure difficult in electronic health-care databases. In the UK databases, data were based on

Figure 5  Prevalence of prescribing of other asthma medications in the year before, during and after pregnancy in women with a live or stillbirth during 2004–2010 for women prescribed ≥1 prescription for an asthma medication during ≥1 of the time periods of interest.

prescriptions issued, whereas in other regions it was prescriptions dispensed. It is possible that some women who are issued prescriptions may not get them dispensed. However, even if we take a conservative estimate and assume that 25% of prescriptions issued in the UK databases were not dispensed, the levels of asthma prescribing during pregnancy would still be higher in the UK/Wales than in all other regions, with the exception of Emilia Romagna. This study did not account for prescriptions issued during one 3-month time period that could have continued to be taken during the following 3-month time period. This will have resulted in an underestimation of exposure during some time periods, especially as women may use inhalers for a long time after they are prescribed. Some women may also have several inhalers for a product issued in a single prescription and women may have multiple inhalers of the same product in use at the same time. The study period for which data were available did vary between databases and it is possible that these small differences in calendar time may have influenced the results and could explain some of the variations observed; however, for those countries contributing data for the entire study period, only small changes in prescribing patterns by calendar time were observed.

An absence of data on indication for prescribing was a limitation when investigating the use of oral glucocorticoids to treat more severe asthma and asthma exacerbation, as these products can be prescribed to treat a number of other conditions. This study attempted to reduce the impact of this by only including oral glucocorticoid prescriptions if they were issued during the same 3-month period as an asthma medication; it is likely, however, that some misclassification will still have occurred. In addition, some women will not have needed to take the oral glucocorticoids at the time of issue but will have been prescribed them in advance so that they had them available for when they experienced an exacerbation in the future. None of the databases captured prescriptions issued during a hospital stay or a visit to an accident and emergency department and this may have led to an underestimate of the use of oral glucocorticoids to treat asthma exacerbations. An absence of data on the indication for prescribing of medications was also a limitation when trying to interpret and provide explanations for the differences observed in prescribing practices. The large percentage of women receiving only a single prescription in some regions may imply that some products may be being prescribed for bronchoconstriction in association with acute bronchitis or other respiratory tract infections. Inhaled SABAs and ICS were the products most commonly prescribed to women who received only a single prescription and in these cases they are unlikely to have been for a chronic condition. It is also possible that some women are started on an inhaled SABA or ICS to see if it has any effect on the symptoms and if there is no effect, the treatment is discontinued and it is concluded that the woman was not suffering from asthma.

Comparison with the literature

The UK prescribing prevalence of asthma medications during pregnancy in this study was found to be slightly higher than that reported by a previous study looking at the prevalence of treated asthma using data from the CPRD (9.4% vs 8.3%). The previous study had required women to have evidence of an asthma diagnosis in addition to at least one prescription or a minimum of six prescriptions to be eligible for inclusion in the study, and this is likely to explain some of the difference observed. Trends and levels of overall asthma medication prescribing in the Netherlands and Norway were in line with those reported elsewhere for studies looking at a similar time period and using the same data sources (5.1% vs 4.5% during pregnancy in the Netherlands and 1.7% vs 1.7% during the first trimester in Norway). The Norwegian study, however, had not reported on the prescribing prevalence during the entire pregnancy and on the level of prescribing of specific classes of asthma medications. In Denmark, a large cohort study based on maternal self-reporting, rather than prescription data, had reported that 6.2% of women suffered from asthma during pregnancy between 1996 and 2002, which was higher than the 4.4% (CI95 4.3% to 4.4%) who had received a prescription during the slightly later time period of our study. To the best of our knowledge, this is the first study to report on the prevalence of asthma medication prescribing during pregnancy in Italy.

Interpretation

The variation observed in the levels of prescribing may to some extent reflect differences in the prevalence of asthma in the different regions. The World Health Survey conducted by the WHO has reported country-specific estimates of doctor diagnosed asthma prevalence, at any time, in individuals aged 18–44 years. This study found a considerably lower prevalence of asthma in Italy (6.0%) compared with the other regions in our study: 9.5% in Denmark, 11.1% in Norway, 15.2% in the Netherlands and 17.6% in the UK. Variations may also be explained by differences in exposure to allergens, infections, air pollution and climate, as well as the cost of individual medications and different attitudes to prescribing. Unfortunately, data on these variables were not available within the majority of databases and could not be evaluated as part of this study. Variations in prevalence and these variables may also explain the within-country regional prescribing differences observed between the two Italian databases, with the Italian National Survey data demonstrating geographic variation in asthma prevalence (5.3% vs 8.4% in Tuscany vs Emilia Romagna, respectively). Hospital admission data for asthma exacerbations in Italy support our findings, with higher hospital admissions observed in Emilia Romagna than Tuscany, as well as higher costs associated with the diagnostic label ‘bronchitis (asthmatic)/asthma’. The differences in prescribing prevalence between Tuscany and Emilia Romagna and the
representativeness of the two regions do not permit the extrapolation of the results to the rest of Italy. The findings of our study also demonstrated a peak in asthma medication prescribing during the second trimester in all regions. This may be explained by the findings of a systematic review that concluded that a worsening of asthma symptoms is most likely during the latter trimesters with a peak in the sixth month of gestation and an improvement during the last 4 weeks of pregnancy, which may also explain some of the lower levels of prescribing observed shortly after pregnancy. It is also possible that women are informed of the importance of well controlled asthma when visiting their GP during their pregnancy, and they are therefore more likely to request and use the required prescriptions during the second trimester than they were during the early stages of pregnancy. Women may also, to some extent, feel more reassured about taking their asthma medicines once the critical period of organogenesis during the first trimester of pregnancy has passed.

In all regions, the prescribing guidelines advised that treatment for asthma during pregnancy should be the same as that for other groups of patients and that the risks associated with poorly controlled asthma were greater than those posed by conventional asthma medications. There was general agreement that SABAs were not teratogenic, with salbutamol and terbutaline being the recommended first choice. In Italy, however, the possibility of increased risks of maternal and neonatal transitory hypoglycaemia, maternal and fetal tachycardia, acute heart failure, pulmonary oedema and maternal death were listed in relation to salbutamol and the guidelines stated that the manufacturer advised to avoid during pregnancy unless the potential benefit outweighs the risk. The advice from the manufacturer and the risk of a number of maternal and fetal outcomes were not reported in the guidelines of the other regions, and it is possible that this may, in part, explain the lower levels of SABA prescribing in Italy than the other regions; these risks are, however, largely thought to be associated with medication when used as an infusion and not for inhaled medications used at a normal dose.

All guidelines advised that there was no need to stop ICS during pregnancy. During the study period, budesonide was the only ICS that had the more favourable Food and Drug Administration (FDA) pregnancy category B, while all others had an FDA pregnancy category C, indicative of the fact that animal reproduction studies had shown an adverse effect on the fetus and there were no adequate and well-controlled studies in humans. Norway and Italy were the only countries, however, where the guidelines recommended budesonide as the ICS of choice and Denmark was the only region where budesonide was found to be the ICS most commonly prescribed. In Italy, despite the guidelines recommending budesonide, beclometasone was by far the most commonly prescribed ICS. The Italian guidelines on beclometasone also stated that the manufacturer recommends avoiding its use during the first trimester; although a small reduction was observed during this time period, it was still quite commonly prescribed. It is possible that other factors, including the cost of different products, may influence prescribing practices in the different regions, but this study was not able to investigate this.

The UK guidelines were the only ones to specify that LABAs should be used with an ICS and ideally as part of a combination product. It was stated that although little information is available on the safety of combination products, there is no reason, in this instance, to think that the products would be more harmful in combination than when given separately. In the Danish guidelines, no concerns were raised about the use of LABAs and the UK guidelines referenced evidence from prescription event monitoring to suggest that salmeterol was safe in pregnancy. In Norway, it was advised that the older and better known LABAs should be used before the newer ones and this study found salmeterol and formoterol to be the most popular and prescribed products in equal measures. In the Netherlands, the guidelines on LABAs differed from other regions, with the emphasis being put on the lack of information relating to their safety and guidance from the Dutch College of General Practitioners advising against the use of these medications in pregnancy. The differences in prescribing guidelines relating to LABAs are likely to explain some of the variations observed between regions. Our study observed a sharp decline in the prescribing of LABAs during pregnancy in the Netherlands, which is in line with the guidelines and has also been observed in a Dutch study by Zetstra-van der Woude et al. The decline in the prescribing of LABAs during pregnancy, observed in all regions except the UK, indicates that clinicians and pregnant women may worry about using these rather new inhaled medications during pregnancy, as there are currently few published studies reporting on their safety.

The limited information on the safety of leukotriene receptor antagonists was acknowledged within all guidelines, with a general consensus that they should not be started during pregnancy but could be continued in women who were using them to successfully control their asthma before pregnancy; all regions in our study saw some decline in the level of their prescribing during pregnancy. For oral glucocorticoids, several of the guidelines mentioned the possibility of an increased risk of cleft lip/palate if used during the first trimester of pregnancy, and our study observed a decline in their use during pregnancy in all regions. In Denmark, the UK and the Netherlands, it was stated that any well indicated treatment should be initiated and not withheld during pregnancy as the advantages of well controlled asthma would outweigh any associated risk to the fetus. In Denmark and the Netherlands, it was recommended that these were prescribed at the lowest dose and for the shortest time period. The low prescribing levels of oral
asthma medications in general in all regions and the decline observed during pregnancy are reassuring, as treatment guidelines focus on inhaled medications and recommend the addition of oral products only for severe asthma.

CONCLUSION
This study identified differences in the percentage of women who received a prescription for an asthma medication during and surrounding pregnancy in different regions of Europe. Differences were also observed in relation to the specific products that were most commonly prescribed; however, no major differences were observed in the treatment in general with inhaled β2 agonists and ICS being prescribed to the majority of women.

Author affiliations
1 Department of Pharmacy and Pharmacology, University of Bath, Bath, UK
2 Institute of Clinical Psychology, National Research Council (IFC-CNPR), Pisa, Italy
3 Medical Birth Registry of Norway, The Norwegian Institute of Public Health, Bergen, Norway
4 Department of Global Public Health and Primary Care, University of Bergen, Bergen, Norway
5 Emilia Romagna Birth Registry, Centre for Clinical and Epidemiological Research, University of Ferrara, Ferrara, Italy
6 Department of Nursing, College of Human and Health Sciences, Swansea University, Swansea, UK
7 Pharmacoepidemiology and Pharmacoeconomics Unit, University of Groningen, Groningen, The Netherlands
8 Centre for Health Information, Research and Evaluation, Swansea University, Swansea, UK
9 Drug Policy Service, Emilia Romagna Region Health Authority, Bologna, Italy
10 Paediatric Department, Hospital Lillebaelt, Copenhagen, Denmark
11 The Regional Agency for Public Health of Tuscany, Florence, Italy
12 Department of Pharmacoepidemiology, The Norwegian Institute of Public Health, Bergen, Norway
13 Department of Public Health, University of Copenhagen, Copenhagen, Denmark
14 Institute of Nursing, University of Ulster, Newtownabbey, UK

Acknowledgements The authors wish to thank members of the EURoMediCAT Steering Group and, most particularly, Professor Corinne de Vries for her invaluable contribution to the design of the study. They would also like to thank Professor Vibeke Backer from the Advisory Board for her comments on the draft manuscript. They would like to acknowledge Dr Sandra Baldacci for her comments on the Italian data and all the data providers who make anonymised data available for research. The work presented in this paper describes anonymised data held in the Secure Anonymised Information Linkage (SAIL) Databank, which is part of the national e-health records research infrastructure for Wales. They thank Karen Tingay and David Tucker for their contribution to the work carried out with SAIL data. This study also describes data from the Full Feature Clinical Practice Research Datalink obtained under licence from the UK Medicines and Healthcare products Regulatory Agency. The Tuscany Registry of Birth Defects is funded by the “Direzione Generale Diritti di cittadinanza e Coesione sociale—Regione Toscana.” The Emilia Romagna Registry of Birth Defects is funded by the Emilia Romagna Region Health authority grant number Delibera 56412/2010.

Contributors LTWdJ-vdB and HD contributed to the conception of the study, RAC, SJ, API, EG, AJN, KK and LTWdJ-vdB contributed to the design of the work. Data acquisition and analysis was carried out by AVH, EG and A-MNA, KK and AE, RG and API, APu and AJN, HJB and LTWdJ-vdB, Karen Tingay, DT, SJ and RAC. RAC compiled the results for all regions. All the authors were involved in the interpretation of the study results as well as the drafting and revision of the manuscript and all approved the final version to be published.

Funding This study was part of the EURoMediCAT research project (http://www.euromedicat.eu) which has been supported by the European Commission under the Seventh Framework Programme (grant agreement number 260598).

Disclaimer The interpretation and conclusions contained in this report are those of the authors alone.

Competing interests All authors, with the exception of A-MNA, RG, APu, AJN and AE, had financial support from the European Union for the submitted work under the Seventh Framework Programme (grant agreement HEALTH-F5-2011-260598). RAC owns shares in GlaxoSmithKline.

Ethics approval Ethical and data access approvals were obtained for each database from the relevant governance infrastructures.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

REFERENCES


Asthma medication prescribing before, during and after pregnancy: a study in seven European regions


BMJ Open 2016 6:
doi: 10.1136/bmjopen-2015-009237

Updated information and services can be found at:
http://bmjopen.bmj.com/content/6/1/e009237

These include:

Supplementary Material
Supplementary material can be found at:
http://bmjopen.bmj.com/content/suppl/2016/01/19/bmjopen-2015-009237.DC1

References
This article cites 32 articles, 3 of which you can access for free at:
http://bmjopen.bmj.com/content/6/1/e009237#BIBL

Open Access
This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections
Articles on similar topics can be found in the following collections

Epidemiology (2210)
Respiratory medicine (355)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/