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Caesarean Section in Uninsured Women in the United States: Systematic Review and Meta-analysis

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Caesarean Section in Uninsured Women in the United States: Systematic Review and Meta-analysis

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variation, health services, financial incentives, underuse, underserved

Word count

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Abstract

Objective: The aim of this study is to assess the odds of caesarean section for uninsured women in the United States and understand the underlying mechanisms as well as consequences of lower use.

Study design: Systematic review and meta-analysis.

Data sources: MEDLINE, Embase, The Cochrane Library and CINAHL from the first year of records through April 2018.

Eligibility criteria: We included studies that reported data to allow the calculation of odds ratios of caesarean section of uninsured as compared to insured women.

Outcomes: The pre-specified primary outcome was the adjusted odds ratio of deliveries by caesarean section of uninured women as compared with privately or publicly insured women. The pre-specified secondary outcome was the crude odds ratio of deliveries by caesarean section of uninsured women as compared with insured women.

Results: Twelve articles describing sixteen separate studies involving more than 8.8 million women were included in this study. We found: 0.70 times lower odds of caesarean section in uninsured as compared to privately insured women (95%CI 0.63 to 0.78), with no relevant heterogeneity between studies ($\tau 2=0.01$); and, 0.92 times lower odds for caesarean section in uninsured as compared to publicly insured women (95%CI 0.80, 1.07), with no relevant heterogeneity between studies ($\tau 2=0.02$). The lower odds were noticed in subgroup analyses as well as in crude analysis. We found 0.70 times lower odds in uninsured as compared to publicly insured women (95%CI 0.69, 0.72).

Conclusions: Caesarean sections are less likely to be performed in uninsured women as compared with insured women. In many regions the rates for uninsured women are close or below the benchmarks for appropriate caesarean section rates, therefore efforts to assess the

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Strengths and limitations of this study

- Extensive literature search, screening and data extraction performed in duplicate, review and analysis of study characteristics as well as thorough quality assessment of included studies.
- ✓ All studies are from one country, i.e. the United States, and this limits the effect of contextual factors.
- A major limitation of our study is the variation across studies, in terms of the study populations characteristics, type of data used, types of caesarean section analyzed and adjusting variables used in statistical analyses.
- ✓ Despite similar features, the uninsured are a diverse group of United States citizens.
- ✓ While a population level caesarean section rate of less than 15 or 19 percent suggests underuse, we cannot determine the mix of under, over, and appropriate use in a specific population.

Introduction

Introduction of clinical procedures in medical practice have saved and improved the lives of many people worldwide. But with time, these clinical procedures become subject to overuse or underuse; i.e. some people get them without really needing them while others do not get them although in need of them.¹ As a result, overuse of procedures may result in unnecessary harm due to the side effects of the procedures or, in case of underuse, not receiving the care they need.¹⁻³ These adverse effects occur due to differing health systems and other contextual factors.³⁴ These factors include financial and non-financial barriers in accessing healthcare even in the most advanced economies in the world, such as the United States (US). Consequently, specific segments of the population may be underserved as healthcare systems are unable to address structural problems that leave patients without the care they need.¹ Globally, Caesarean section (CS) is an example of overuse and underuse of clinical procedures. Once introduced into clinical practice, it greatly improved maternal and newborn outcomes.⁵ Presently, many countries have long exceeded the 15 or 19 percent benchmarks for CS out of total deliveries, argued to be the ideal rate of CS in terms of improving the health of women and newborns.⁶⁻⁸ CS rates average as high as 40.5 percent among countries in Latin America and the Caribbean region,⁹ 32.3 percent in Northern America⁹ (32.2 percent in US),¹⁰ while on the other extreme, it is as low as 7.3 percent in Africa⁹ and known to range even lower in specific countries: 1 percent in Nepal and Cambodia to 0.6 percent in Ethiopia and Niger.¹¹ Variations are also observed within countries,⁵¹¹ for instance, in the US⁵ a recent study reported a range between 4 to 65 percent across health markets.¹²

Insurance coverage is one health system factors known to influence the use of medical procedures,^{13 14} including CS.¹⁵⁻¹⁸ While private insurance, for example, seems to increase the odds of having a CS delivery,¹⁵ the lack of insurance appears to decrease it.¹⁸⁻²⁰ Millions of people worldwide, as well as in the US, are not covered by any insurance scheme and are

exposed to the hazard of being underserved with clinical procedures,²¹⁻²⁵ including perinatal services.²⁶ The US has a mixed health insurance system dominated by private insurance.²² The Federal Medicare program, covers people over 64 years old and/or disabled, which accounts for about 16.7 percent of the population.²² State Medicaid programs cover children and parents from low income families as well as partially caring for Medicare beneficiaries with low incomes and, in total, accounts for about 19.4 percent of the US population.²⁷ Over half of US population is covered with voluntary employer based private insurance.²⁷ The remaining population is uninsured and can range from 2.5 (Massachusetts) to 16.6 (Texas) percent according to 2016 estimates.²⁷ For decades, in the US, there has been an ongoing debate for and against universal health coverage and related topics with limited but substantial progress towards more coverage through the Affordable Care Act (ACA).^{21 28-35} Nonetheless, millions of Americans remain uninsured for various reasons and are not able to access the healthcare they need.^{21 29 36} The aim of this study is to assess the odds of CS for uninsured women in the US and understand the underlying mechanisms as well as consequences of lower use in the US context.³⁷

Materials and methods

Search strategy and data sources

Search words referring to CS, such as 'caesarean section', 'caesarean delivery', 'caesarean', were combined with words referring to factors contributing to variation and increase of CS rates, such as 'insurance', 'social class', 'socioeconomic', and words referring to study design, such as 'geographic variation', 'medical practice variation' (Appendix 1). No publication date or language restrictions were applied. We searched MEDLINE, Embase, the Cochrane Library and CINAHL from the beginning of records to the end of April 2018, when we last updated our search. A manual search was applied on the reference lists of included studies and previous systematic reviews.

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Study selection and outcomes

To be included in the analysis, studies had to report odds ratio (OR) of CS comparing uninsured against privately and/or publicly insured women. Adjusted OR of deliveries by CS of uninsured women in comparison to insured women was the pre-specified primary outcome. Crude OR of deliveries by CS of uninsured women in comparison to insured women was the pre-specified secondary outcome.

Data extraction 🥒

Papers screening and independent data extraction was done by two researchers (IH and MB). Differences were resolved based on consensus. We extracted data on study population, study design, data sources, setting, type of CS analyzed, statistical analysis, and (primary and secondary) outcome measures.

Quality assessment

We used Quality In Prognostic Studies (QUIPS) to assess the risk of bias across six study domains.³⁸ Each study was evaluated independently by two researchers (IH and MB) and any differences among evaluators were discussed and resolved. A single rating was assigned for all studies. As specified in the QUIPS tool, a "high", "moderate", or "low" rating was applied for individual domains and overall rating of a study.³⁸ If a study was rated with a low risk of bias across all the six domains, it would receive an overall rating of low risk of bias.¹⁵ If one or more domains of a study were rated with a moderate risk of bias, it would receive an overall moderate risk of bias, it would receive an overall high risk of bias.¹⁵

Main analysis

Standard inverse-variance random effects meta-analysis was used to combine the overall ORs. An OR lower than one implies a lower frequency of CS in uninsured than in insured women. We calculated τ^2 to measure heterogeneity between studies.³⁹ Pre-specified cutoffs of

 τ 2 of 0.04, 0.16 and 0.36 were used to represent low, moderate, and high heterogeneity between studies.⁴⁰ Subgroup analysis by study design, period of data collection, state, type of CS analyzed, parity, inclusion of women with previous CS, pregnancy risk of included women and level of (QUIPS) risk of bias was performed to examine between-study heterogeneity and chi-square test was used to calculate p-values for interaction among subgroups. Test for linear trend was performed in case of more than two ordered strata. All pvalues were two-sided. STATA, release 13, was used for analyses (Stata-Corp, College Station, Texas).

Additional analysis

We calculated CS rates among different insurance subgroups for the studies included in the analysis.

Patient involvement

No patients were involved in this study. We used data from published papers only.

Results

We identified a total of 1837 records: 1123 from Medline; 556 from Embase; 39 from the Cochrane Library, 119 from CINAHL and 28 from manual search (Figure 1). We removed 240 duplicates. 1597 records were screened for eligibility. We performed full text examination on 177 records. We excluded 139 that did not report insurance status of women and 26 that were otherwise irrelevant. Finally, 12 records describing 16 separate studies¹⁸⁻²⁰ ⁴¹⁻⁴⁹ including more than 8.8 million women were included in review and meta-analysis.

Characteristics of studies are presented in Table 1 and Appendices 2, 3, 4 and 5. All studies were from the US. Thirteen studies were cross-sectional and three were retrospective cohort studies. Population size of studies ranged from 9,017 to 6,717,486 cases. Studies used data from years 1986 to 2011 and most studies used hospital records data (Appendix 2). Case

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exclusion criteria varied considerably (Appendix 3) as well as variables studies used for statistical adjustment (Appendix 4). Appendix 5 reports evaluation of studies using QUIPS risk of bias tool. Four studies were classified with low risk of bias, two studies with moderate risk, and ten studies with high risk of bias (Appendix 5).

Figure 2 presents meta-analyses for primary outcome measure, i.e. adjusted ORs of CS in uninsured women as compared to privately or publicly insured. Since there was a positive interaction between uninsured vs privately insured group and uninsured vs publicly insured group (p=0.016), we performed meta-analyses for each group separately. In the meta-analysis comparing uninsured with privately insured women, including seven studies in 556,454 women, we found that the odds of CS were 0.70 times lower in uninsured as compared to privately insured women (95%CI 0.63 to 0.78), with no relevant heterogeneity between studies in 510,010 women, we found that the odds of CS were 0.92 times lower in uninsured as compared to publicly insured women (95%CI 0.80, 1.07), with no relevant heterogeneity between studies (τ 2=0.02). An additional study in 6,717,486 women, which did not distinguish between privately and publicly insured women (95%CI 0.69, 0.72).

Figure 3 presents results of subgroup analyses of adjusted odds ratios in uninsured vs privately insured women (upper panel) and in uninsured vs publicly insured women (lower panel). In the analysis of uninsured vs privately insured women, estimates varied for subgroups state (p for interaction<0.001), type of CS (p for interaction<0.001), parity (p for interaction=0.07), and pregnancy risk (p for interaction<0.001). There was no positive trend in the period of data collection subgroup. In the lower panel, which presents subgroup analyses of adjusted odds ratios in uninsured vs publicly insured women, estimates varied for

subgroups period of data collection (p for interaction=0.03), state (p for interaction=0.004), type of CS (p for interaction=0.03), parity (p for interaction=0.03) and QUIPS risk of bias (p for interaction=0.03).

In Figure 4 we present meta-analyses for crude ORs of CS in uninsured as compared to privately or publicly insured women as secondary outcome. In the meta-analysis comparing uninsured with privately insured women, including eleven studies in 2,010,483 women, we found that the odds of CS were 0.71 times lower in uninsured as compared to privately insured women (95%CI 0.66 to 0.76), with no relevant heterogeneity between studies ($\tau 2=0.018$). In the meta-analysis comparing uninsured with publicly insured women, including eleven studies in 2,010,483 women, we found that the odds of CS were 0.93 times lower in uninsured as compared to publicly insured women, including eleven studies in 2,010,483 women, we found that the odds of CS were 0.93 times lower in uninsured as compared to publicly insured women (95%CI 0.85, 1.01), with no relevant heterogeneity between studies ($\tau 2=0.017$).

Appendix 6 presents rates of CS among groups with different insurance status for individual studies. Two studies found CS rates for uninsured women below the 19 percent benchmark, another five studies found CS rates below the 15 percent benchmark. The rates of other studies range from 19.3 percent to 23.0 percent, close to 19 percent benchmark.

Discussion

Our systematic review and meta-analyses estimated that the overall odds of receiving a caesarean section are on average 0.70 times lower for uninsured women as compared with privately insured women, 0.92 times lower for uninsured women as compared with publicly insured women and 0.70 times lower for uninsured women as compared to privately and publicly insured women. The lower odds were noticed across all subgroups of studies in subgroup analyses as well as in crude analyses.

Context

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To our knowledge, this is the first meta-analysis that examines CS rates of uninsured women compared to insured women. Two recently published meta-analyses by our group reported the association of CS with for profit status of hospitals and type of insurance.^{15 50} Investigating the association of for-profit vs non-profit status of hospital with the odds of CS, we found that the odds of CS were 1.41 higher in for-profit hospitals as compared with non-profit hospitals (95% CI 1.24 to 1.60).⁵⁰ The findings were consistent in subgroup analyses.⁵⁰ Investigating the association of CS with private insurance, we found that the odds of CS were 1.13 times higher for privately insured women compared with women covered with public insurance.¹⁵ Again, the increased risk was observed across all subgroups.¹⁵

Strengths and limitations

The major strengths of our meta-analysis include an extensive literature search, screening and data extraction performed in duplicate, review and analysis of study characteristics as well as thorough quality assessment of included studies. In addition, all studies are from one country, i.e. the US, and this limits the effect of contextual factors. A major limitation is the variation across studies, in terms of the study populations characteristics (i.e. parity, inclusion of women with previous CS, risk for CS), type of data used, types of CS analyzed and adjusting variables used in statistical analyses. It should also be taken into consideration, that despite similar features, the uninsured are a diverse group of US citizens.^{24 25} Finally, while a population level CS rate of less than 15 or 19 percent suggests underuse, we cannot determine the mix of under, over, and appropriate use in a specific population.

Mechanism

There are several possible explanations why uninsured women have lower odds of CS when compared to insured women. One likely factor is that financial incentives are stronger with private insurance than in the publicly insured or uninsured.^{15 16} These incentives result from higher payment for CS by private insurers through reimbursement arrangements that

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encourage more expensive procedures as means to increase profits, as well as providers' (hospitals and individual physicians) responses to these incentives.^{15 50 51} The responses to incentives by hospitals exist in the form of patient scheduling policies that direct privately insured patients to profit inclined physicians.^{18 50} It is also a known association that physicians who have a higher share of privately insured patients will tend to overuse CS.^{19 20} ⁵⁰ They do so as they perceive patients to have a higher social class, i.e. able to pay higher fees, or fear malpractice liability.^{16 41 50 52}

Additional reasons are likely reflected in the comparison between uninsured and publicly insured women. A first set of reasons are related to deliberate or forced decisions of uninsured women to keep out-of-pocket payments low.¹⁶ The uninsured patients are more likely to seek less expensive care when they face the need for healthcare services.¹⁶ In the case of giving birth, this would lead to a greater preference for vaginal delivery. A second set of reasons may be discrimination of providers towards uninsured women. Providers have a preference for profitable, i.e. privately insured patients, a preference commonly referred to as "cream skimming".^{19 20 50 53}

Implications for uninsured women

Most studies included in our meta-analysis, including the most recent studies from California⁴⁷ and Florida,⁴⁸ show that rates for CS among uninsured women are below or close to the 15 and 19 percent benchmarks previously reported.⁶⁻⁸ Even in instances where the average state rates are slightly above the 19 percent benchmark, some hospitals service areas are likely to have CS rates lower than 19 percent or even 15 percent for uninsured women because of the well-established within state variation in CS rates.^{5 52} Uninsured women in these areas are highly likely to be underserved with caesarean section during delivery. Uninsured patients generally have higher unmet needs than insured patients due to access barriers.^{21 22 24 26 54-58} Such barriers encourage inappropriate health seeking behaviors among

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uninsured.^{21-23 30 56 59-61} Consequently, uninsured populations face higher health risks and have worse healthcare outcomes.^{21-24 30 54-57 59 60 62 63}

The uninsured also face financial burdens which result from out-of-pocket payments that are more severe/extensive than co-payments or premiums that are paid by people that are publicly or privately insured. The uninsured are known to pay higher prices for services as compared to other payers for the same care,^{25 64} spend a high portion of income to cover medical expenses²² (although they spend less for their health compared to patients who have insurance),²⁴ are frequently charged for full price for healthcare services,^{22 64} often do not benefit from discounts from providers,^{22 25} and face severe financial difficulties.^{21 22} Uninsured manage to pay only part of the costs for their care.²⁴ The remaining costs are uncompensated costs^{21 24 65 66} and most of such costs are covered by the local, state or federal government,^{24 65} eventually resulting in tax increases.²⁴

Implications for research and policy making

Future studies should examine the association of a lack of insurance in pregnant women across health care markets with varying CS rates and assess if delivery outcomes were correspondingly worse, in the effort to investigate the presence of underuse of CS.

In parallel, policy options that could lead to improvements of insurance coverage for delivering women should be assessed in terms of their ability to address healthcare outcomes while keeping overall costs at minimum. In the past, states have adopted different strategies for covering uninsured people.^{22 23 37 67} While there are many known benefits to insurance coverage,^{21 22 30-33 35 57 60 66 68-70} other important policy aspects should be considered. At a time of rising healthcare costs ^{22 33 70 71} regulation of financial incentives is crucial. A revision of payment policies should be pursued ^{15 16 22 50} to align financial incentives with proper health outcomes.^{15 22 50} Reimbursement policies that would pay the same amount for CS and vaginal delivery is one option.^{50 72}

Conclusion

Caesarean sections are less likely to be performed in uninsured women as compared with insured women. The lower odds are consistent in all subgroups and in crude analyses. In many regions the rates for uninsured women are close or below the benchmarks for appropriate CS rates, therefore efforts to assess the delivery outcomes as well as policy options that could improve insurance coverage for women giving birth are important.

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Contributorship Statement

IH, DG, PJ conceived and designed the study. IH, MB performed the data extraction and preparation. IH, LS analyzed the data. IH, MB, LS drafted the paper, which was critically reviewed and approved by all authors.

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Competing interests statement

All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coi disclosure.pdf and declare: no support from any organisation for the

submitted work; no financial relationships with any organisations that might have an interest in the submitted work in the previous three years; no other relationships or activities that could appear to have influenced the submitted work.

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No funding was received to perform this study. All authors, had full access to all of the data (including statistical reports and tables) in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

Data sharing statement

from the study. No additional unpublished data are available from the study.

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Table 1. Characteristics of included studies

| Author | Year | State | Study design | Number | Number | Year of | Population | Sampling | Type of |
|---------------|------|---------------|---------------|----------|----------|------------|-----------------------|-------------|----------|
| | | | | of cases | of | data | | | CS |
| | | | 0, | | hospital | collection | | | analyzed |
| | | | | 0 | units | | | | |
| Stafford | 1990 | California | Cross | 461066 | Not | 1986 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | | any risk | | |
| Haas et al. A | 1993 | Massachusetts | Cross | 57257 | Not | 1984 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | 0 | any risk | | |
| Haas et al. B | 1993 | Massachusetts | Cross | 64346 | Not | 1987 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | | any risk | | |
| Braveman et | 1995 | California | Retrospective | 213761 | Unclear | 1991 | Primipara; no | Consecutive | Any |
| al. | | | cohort | | | | previous CS; any risk | | |
| Burns et al. | 1995 | Arizona | Cross | 33233 | 36 | 1989 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | | | any risk | | |

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| Onion et al. | 1999 | Maine | Cross | 41177 | Not | 1990- | Primipara; no | Consecutive | Any |
|--------------|------|------------|---------------|--------|----------|-------|-----------------------|-------------|---------|
| А | | | sectional | | reported | 1992 | previous CS; any risk | | |
| Onion et al. | 1999 | New | Cross | 41401 | Not | 1990- | Primipara; no | Consecutive | Any |
| В | | Hampshire | sectional | | reported | 1992 | previous CS; any risk | | |
| Onion et al. | 1999 | Vermont | Cross | 19077 | Not | 1990- | Primipara; no | Consecutive | Any |
| С | | | sectional | | reported | 1992 | previous CS; any risk | | |
| Aron et al. | 2000 | Ohio | Retrospective | 25697 | 21 | 1993- | Primipara; no | Consecutive | Any |
| | | | cohort | | 6 | 1995 | previous CS; any risk | | |
| Grant A | 2005 | All states | Cross | 9017 | Not | 1988 | Primi- and multipara; | Random | Any |
| | | | sectional | | reported | 0 | any risk | | |
| Grant B | 2005 | Florida | Cross | 147821 | Not | 1992 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | | any risk | | |
| Coonrod et | 2008 | Arizona | Cross | 28863 | 40 | 2005 | Primipara; low risk | Consecutive | Any |
| al. | | | sectional | | | | | | |
| Huesch | 2011 | New Jersey | Cross | 182108 | Not | 2004- | Primi- and multipara; | Consecutive | Planned |
| | | | sectional | | reported | 2007 | no previous CS; low | | |

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| Kozhimannil | 2013 | All states | Cross | 6717486 | Over | 2002- | Primi- and multipara; | Random | Any |
| et al. | | | sectional | | 1000 | 2009 | any risk | | |
| Huesch et al. | 2014 | California | Cross | 408355 | 254 | 2010 | Primi- and multipara; | Consecutive | Planned |
| | | | sectional | | | | no previous CS; any | | |
| | | | í D | 0 | | | risk | | |
| Sebastião et | 2016 | Florida | Retrospective | 412192 | 122 | 2004- | Primipara; no | Consecutive | Emergency |
| al. | | | cohort | | 10 | 2011 | previous CS; low risk | | |
| CS = caesarean section | | | | | | | | | |
| | | | | | | | | | |
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Figure legends

- Figure 1. The flow diagram of review
- Figure 2. Adjusted odds ratios of caesarean section
- Figure 3. Subgroup analyses for adjusted estimates/Legend: *P for trend
- Figure 4. Crude odds ratios of caesarean section

Supporting information

Appendix 1. Search Strategy

- Appendix 2. Type of data used
- Appendix 3. Reported exclusion criteria

Appendix 4. Covariates used for statistical adjustment

Appendix 5. QUIPS risk of bias

Appendix 6. Caesarean section rates among groups with different insurance status







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Figure 3. Subgroup analyses for adjusted estimates

- For peer review only http://bmjopen.bmj.com/site/about/guidelines.xhtml









Figure 4. Crude odds ratios of caesarean section

Uninsured vs privately insured





Figure 4. Crude odds ratios of caesarean section

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Research Checklist

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According to MOOSE statement for meta-analyses of observational studies

| Reporting of background should include | Where to find in manuscript |
|---|---------------------------------|
| Problem definition | Manuscript (page 5, 6) |
| Hypothesis statement | Manuscript (page 5, 6) |
| Description of study outcome(s) | Manuscript (page 6) |
| Type of exposure or intervention used | Manuscript (page 6) |
| Type of study designs used | Manuscript (page 6, 7) |
| Study population | Manuscript (page 6, 7) Table 1, |
| | Appendix 1 |
| Reporting of search strategy should include | |
| Qualifications of searchers (eg, librarians and investigators) | Manuscript (page 1) |
| Search strategy, including time period included in the synthesis and | Manuscript (page 6), Appendix 1 |
| keywords | |
| Effort to include all available studies, including contact with authors | Manuscript (page 6) |
| Databases and registries searched | Manuscript (page 6) |
| Search software used, name and version, including special features | Manuscript (page 6) |
| used (eg, explosion) | |
| Use of hand searching (eg, reference lists of obtained articles) | Manuscript (page 6) |
| List of citations located and those excluded, including justification | Figure 1 |
| Method of addressing articles published in languages other than | n/a |
| English | |
| Method of handling abstracts and unpublished studies | Manuscript (page 6, 7) |
| Description of any contact with authors | No contact made |
| Reporting of methods should include | |
| Description of relevance or appropriateness of studies assembled for | Manuscript (page 6, 7) |
| assessing the hypothesis to be tested | |
| For peer review only - http://bmjopen.bmj.com/site/ał | oout/guidelines.xhtml |

Manuscript (page 6, 7)

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Documentation of how data were classified and coded (eg, multiple Manuscript (pages 6, 7) raters, blinding, and interrater reliability) Assessment of confounding (eg, comparability of cases and controls Manuscript (page 6-7), Appendix 2, 3, in studies where appropriate) 4 Assessment of study quality, including blinding of quality assessors; Manuscript (page 7), Figure 2, stratification or regression on possible predictors of study results Appendix 5, Assessment of heterogeneity Manuscript (page 7) Description of statistical methods (eg, complete description of fixed Manuscript (page 7) or random effects models, justification of whether the chosen models account for predictors of study results, dose-response models, or cumulative meta-analysis) in sufficient detail to be ê. Ç. replicated Provision of appropriate tables and graphics Table 1, Figure 1-3 and Appendixes 1-7 **Reporting of results should include** Figure 2, Appendix 6 Graphic summarizing individual study estimates and overall estimate Table giving descriptive information for each study included Table 1 Results of sensitivity testing (eg, subgroup analysis) Figure 3 Indication of statistical uncertainty of findings Manuscript, Figure 2-4 **Reporting of discussion should include** Quantitative assessment of bias (eg, publication bias) Manuscript (page 8) Justification for exclusion (eg, exclusion of non—English-language Manuscript (page 8) citations) Assessment of quality of included studies Manuscript (page 8)

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| Reporting of conclusions should include | |
|---|-------------------------|
| Consideration of alternative explanations for observed results | Manuscript (pages 9-13) |
| Generalization of the conclusions (ie, appropriate for the data | Manuscript (page 13) |
| presented and within the domain of the literature review) | |
| Guidelines for future research | Manuscript (page 13) |
| Disclosure of funding source | Manuscript (page 13) |
| | |

PRISMA checklist

| IIILE | | | |
|---------------------------|---|---|--|
| Title | 1 | Identify the report as a systematic review, meta-analysis, or both. | Manuscript (page 1) |
| ABSTRACT | | | |
| Structured summary | 2 | Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number. | Manuscript (page 3,4) |
| INTRODUCTION | • | | |
| Rationale | 3 | Describe the rationale for the review in the context of what is already known. | Manuscript (page 5,6) |
| Objectives | 4 | Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS). | Manuscript (page 6) |
| METHODS | • | (Q) | |
| Protocol and registration | 5 | Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number. | No published protocol or registration |
| Eligibility criteria | 6 | Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale. | Manuscript (page 6) |
| Information sources | 7 | Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched. | Manuscript (page 6) |
| | | Present full electronic second strategy for at least one database, including any limits used such that it could be | Appendix 1 |

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| Study selection | 9 | State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis). | Manuscript (page 6), Figure 1 |
|------------------------------------|----|--|---|
| Data collection process | 10 | Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators. | Manuscript (page 6,7) |
| Data items | 11 | List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made. | Manuscript (page 6,7) |
| Risk of bias in individual studies | 12 | Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis. | Manuscript (page 7) |
| Summary measures | 13 | State the principal summary measures (e.g., risk ratio, difference in means). | Manuscript (page 6) |
| Synthesis of results | 14 | Describe the methods of handling data and combining results of studies, if done, including measures of consistency $(e.g., I^2)$ for each meta-analysis. | Manuscript (page 6, 7) |
| Risk of bias across studies | 15 | Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies). | Manuscript (page 6) |
| Additional analyses | 16 | Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified. | Manuscript (page 7) |
| RESULTS | | | |
| Study selection | 17 | Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram. | Manuscript (page 8), Figure 1 |
| Study characteristics | 18 | For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations. | Manuscript (page 8) Table 1, Appendix 2, 3, 4 |
| Risk of bias within studies | 19 | Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12). | Manuscript |

| Results of individual studies 20 For all outcomintervention gr Synthesis of results 21 Present results Risk of bias across studies 22 Present results Additional analysis 23 Give results of DISCUSSION 24 Summarize the key groups (e.: List is is in the second state of the seco | hes considered (benefits or harms), present, for each study: (a) simple summary data for each roup (b) effect estimates and confidence intervals, ideally with a forest plot. s of each meta-analysis done, including confidence intervals and measures of consistency. s of any assessment of risk of bias across studies (see Item 15). f additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]). | Figure 2, Appendix 6 Manuscript (page 8,9) Figure 2, Appendix 6 Manuscript (page 8), Appendix 5 Manuscript (page 9), Figure 3, Appendix 7 |
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| Risk of bias across studies 22 Present results Additional analysis 23 Give results of DISCUSSION 24 Summarize the key groups (e.g) Line in the interview 25 Discussion | s of any assessment of risk of bias across studies (see Item 15). f additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]). | Manuscrip (page 8), Appendix 5 Manuscrip (page 9), Figure 3, Appendix 7 |
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| Summary of evidence 24 Summarize the key groups (e.g. Linitation 25 Dimension | e main findings including the strength of evidence for each main outcome: consider their relevance to | |
| | g., healthcare providers, users, and policy makers). | Manuscript (page 9, 10 |
| Limitations 25 Discuss limitati identified research | tions at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of arch, reporting bias). | Manuscript (page 10) |
| Conclusions 26 Provide a gene | eral interpretation of the results in the context of other evidence, and implications for future research. | Manuscript (page 10- 13) |
| FUNDING | | |
| Funding 27 Describe source systematic rev systematic rev | ces of funding for the systematic review and other support (e.g., supply of data); role of funders for the iew. | Manuscript (page 13) |

Appendix 1. Search Strategy

1. For Medline (PubMed)

((((((causes OR determinants OR statistics OR rates OR factors OR decision* OR physician* OR socioeconomic OR state medicine OR evidence-based OR hospital OR hospitals OR hospitalization OR hospitalized OR uncertain* OR educational status OR social class OR obstetric* OR gynecolog* OR supply OR distribut* OR utilization OR insurance OR choice OR attitude OR patient OR economics OR maternal OR accessib* OR health service* OR rural population OR urban population[Title/Abstract])) NOT medline[sb])) OR ("Decision Making"[Mesh] OR "Physician's Practice Patterns" [Mesh] OR "Socioeconomic Factors" [Mesh] OR "State Medicine" [Mesh] OR "Evidence-Based Medicine" [Mesh] OR "Hospitals" [Mesh] OR "Uncertainty" [Mesh] OR "Educational Status" [Mesh] OR "Hospital Costs" [Mesh] OR "Physician Incentive Plans" [Mesh] OR "Social Class" [Mesh] OR "Obstetrics and Gynecology Department, Hospital" [Mesh] OR "supply and distribution"[Subheading] OR "utilization"[Subheading] OR "Insurance"[Mesh] OR "Choice Behavior"[Mesh] OR "Attitude to Health"[Mesh] OR "Patient Participation"[Mesh] OR "Physician-Patient Relations" [Mesh] OR "Economics, Hospital" [Mesh] OR "Maternal Health Services" [Mesh] OR "Health Services Accessibility" [Mesh] OR "Health Services Research" [Mesh] OR "Rural Population"[Mesh] OR "Urban Population"[Mesh]))) OR factors OR rates OR statistics OR causes OR determinants AND (((((operative delivery OR caesarean section OR cesarean section OR c section OR caesarean OR caesarean delivery OR cesarean delivery OR caesarean rates OR cesarean rates)))) OR cesarean section [MeSH Terms])) AND ((((("Catchment Area (Health)"[Mesh] OR "Small-Area Analysis"[Mesh]))) OR ((((small area analysis OR small area analyses OR medical practice variation OR regions OR geographic variation OR variation)))))

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| Appendix | 2. Type | of dat | a used | 1 | | |
|--------------------|---------|--------|------------------|--------------------------------|-------------|--------|
| Author | Year | Survey | Hospital records | Birth certificates/registry | Census data | |
| Stafford | 1990 | | + | | | |
| Haas et al. A | 1993 | | + | + | | |
| Haas et al. B | 1993 | | + | + | | |
| Braveman et al. | 1995 | | | + | + | |
| Burns et al. | 1995 | | + | + | | |
| Onion et al. A | 1999 | | + | | | |
| Onion et al. B | 1999 | | + | | | |
| Onion et al. C | 1999 | | + | | | |
| Aron et al. | 2000 | | + | | | |
| Grant A | 2005 | + | | | | |
| Grant B | 2005 | | + | | | |
| Coonrod et al. | 2008 | | | + | | |
| Huesch | 2011 | | + | | | |
| Kozhimannil et al. | 2013 | | + | | | |
| Huesch et al. | 2014 | | + | | | |
| Sebastião et al. | 2016 | | + | + | | |
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| Authors Stafford Haas et al. A | Year | Source population | ≤14 | al or ethnic orities | para | us caesarean a | risk factors for ean section | th | e delivery r more) | n weighting | presentation | entation | delivery (less weeks) | sk factors for n section | bor | th missing da | characteristi | |
|---|--------------|---|-----|-------------------------|-------|-------------------|---------------------------------|----------|-----------------------|-------------------|--------------|------------------|--------------------------|-----------------------------|-----------|---------------|---------------|---|
| Stafford Haas et al. A | 1990 | ····· | Age | Raci mino | Multi | Previc | Other | Stillbir | Multipl (twin or | Newbor <500 gr | Breach | Other malpres | Preterm than 37 | Other ris caesarea | Not in la | Cases wi | Provider | |
| Haas et al. A | 1770 | All deliveries in California, United States | | | | | | | | | | | | | | | + | |
| | 1993 | All deliveries in Massachusetts, United States | | | | | | + | + | + | | | | | | + | | |
| Haas et al. B | 1993 | All deliveries in Massachusetts, United States | | | | | | + | + | + | | | | | | + | | |
| Braveman et al. | 1995 | All deliveries in California, United States | | | + | + | | + | + | | | | + | | | + | | Γ |
| Burns et al. | 1995 | All deliveries in Arizona, United States | | | | | | | | | | | | | | + | + | |
| Onion et al. A | 1999 | All deliveries in Maine, United States | | | + | + | | | + | | | | | | | | | |
| Onion et al. B | 1999 | All deliveries in New Hampshire, United States | | | + | + | | | + | | | | | | | | | T |
| Onion et al. C | 1999 | All deliveries in Vermont, United States | | | + | + | | | + | | | | | | | | | T |
| Aron et al. | 2000 | All deliveries in Cleveland, Ohio, United States | | | | + | | | | +* | | | | | | + | + | T |
| Grant A | 2005 | All deliveries, United States | | | | | | | | | | | | | | + | | T |
| Grant B | 2005 | All deliveries in Florida, United States | | | | | | | | | | | | | | + | + | Γ |
| Coonrod et al. | 2008 | All deliveries in Arizona, United States | | + | + | | | + | + | | + | + | + | | | | + | Γ |
| Huesch | 2011 | All deliveries in New Jersey, United States | | | | + | + | | + | | + | + | + | + | + | | + | Γ |
| Kozhimannil et al. | 2013 | All deliveries in 44 states, United States | | | | | | | | | | | | | | | + | T |
| Huesch et al. | 2014 | All deliveries in California, United States | + | | | + | | | | | | | | + | | + | | |
| Sebastião et al. | 2016 | All deliveries in Florida, United States | | | + | + | | + | + | | + | + | + | | + | + | + | T |
| Huesch et al. Sebastião et al. *500 or less grams | 2014 2016 | All deliveries in California, United States All deliveries in Florida, United States | + | | + | + + | | ÷ | + | 2 | + | + | + | + | + | ++ | | ł |

| Appendix 4. | Covariates | used for | statistical | adiustment |
|-------------|------------|----------|-------------|------------|
| | 00.00 | | | |

| | | | | I | Aaterna | al preco | oncepti | on statı | 15 | | | Mat | ernal cl | inical s | tatus | cha | Fetus racteris | stics | | | | | |
|--------------------|------|----------------|-------------------|----------------|-----------------|-------------------|--------------|----------|--------|-----------------|-----|--------|-------------------------------|---|--|-----------------|-------------------|-----------------------|---------------|--------------------------|--------------------------|-----------------|----------------------------|
| Author | Year | Ethnicity/Race | Educational level | Marital status | Economic status | Insurances status | Urban status | Weight | Height | Body mass index | Age | Parity | Previous caesarean section | Pre-existing (before pregnancy) conditions | Conditions developed during pregnancy | Gestational age | Birth weight | Other characteristics | Prenatal care | Delivery characteristics | Provider characteristics | Other variables | Total number of covariates |
| Stafford* | 1990 | | • | | | | | | | | | | | | | | | | | | | | ſ |
| Haas et al. A* | 1993 | | | | | | | | | | | | | | | | | | | | | | |
| Haas et al. B* | 1993 | | | | | | | | | | | | | | | | | | | | | | |
| Braveman et al. | 1995 | + | + | + | + | + | | | | | + | | | - | ł | | + | + | + | + | ++ | + | 1 |
| Burns et al. | 1995 | + | + | | | | C | | | | + | + | + | | ++ | + | + | ++ | + | | ++ | ++ | 3 |
| Onion et al. A | 1999 | | | | | | | | | | + | | | | | | | | | | | | |
| Onion et al. B | 1999 | | | | | | | | | | + | | | | | | | | | | | | |
| Onion et al. C | 1999 | | | | | | | | | | + | | | | | | | | | | | | |
| Aron et al. | 2000 | | | | | | | | | | + | + | | ++ | ++ | ++ | + | ++ | | | | | 3 |
| Grant A* | 2005 | | | | | | | | | | | | | | | | | | | | | | |
| Grant B* | 2005 | | | | | | | | | | | | | | | | | | | | | | |
| Coonrod et al.* | 2008 | | | | | | | | | | | | | | | | | | | | | | |
| Huesch | 2011 | + | | + | | | + | | | | + | | | | | | | | | | + | ++ | |
| Kozhimannil et al. | 2013 | + | | | | | | | | | + | + | + | ++ | ++ | + | | ++ | | | ++ | | 1 |
| Huesch et al. | 2014 | + | | | + | | | | | | + | | | ++ | ++ | + | | ++ | + | ++ | ++ | ++ | 12 |
| Sebastião et al. * | 2016 | | | | | | | | | | | | | | | | | | | | | | |

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| Appendix 5. | QUIPS risk of bias |
|-------------|--------------------|
|-------------|--------------------|

| | | | | Prognostic | | | Statistical | |
|-------------------------|------|---------------|-----------------|-------------|-------------|-------------|--------------|----------------|
| | | Study | | Factor | Outcome | Study | Analysis and | |
| Author | Year | Participation | Study Attrition | Measurement | Measurement | Confounding | Reporting | Overall rating |
| Stafford 1990 | 1990 | low | low | low | low | high | moderate | high |
| Haas et al. 1993 A | 1993 | low | low | low | low | high | moderate | high |
| Haas et al. 1993 B | 1993 | low | low | low | low | high | moderate | high |
| Braveman et al. 1995 | 1995 | low | low | low | low | moderate | low | moderate |
| Burns et al. 1995 | 1995 | low | low | low | low | moderate | low | moderate |
| Onion et al. 1999 A | 1999 | low | low | low | low | high | low | high |
| Onion et al. 1999 B | 1999 | low | low | low | low | high | low | high |
| Onion et al. 1999 C | 1999 | low | low | low | low | high | low | high |
| Aron et al. 2000 | 2000 | low | low | low | low | low | low | low |
| Grant 2005 A | 2005 | moderate | high | low | low | high | low | high |
| Grant 2005 B | 2005 | low | low | low | low | high | low | high |
| Coonrod et al. 2008 | 2008 | low | low | low | low | high | low | high |
| Huesch 2011 | 2011 | low | low | low | low | low | low | low |
| Kozhimannil et al. 2013 | 2012 | low | low | low | low | low | low | low |
| Huesch et al. 2014 | 2013 | low | low | low | low | low | low | low |
| Sebastião et al. 2016 | 2014 | low | low | low | low | high | low | high |

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Appendix 6. Caesarean section rates among groups with different insurance status

| afford 19901990California198626.822.119.3aas et al. 1993 A1993Massachusetts198423.019.417.2aas et al. 1993 B1993Massachusetts198725.920.822.4aveman et al. 19951995California199127.121.223.0urns et al. 19951995Arizona1989n/an/an/anion et al. 1999 A*1999Maine1990-199215.914.913.4nion et al. 1999 B*1999New Hampshire1990-199216.113.213.0nion et al. 1999 C*1999Vermont1990-199214.513.59.4on et al. 20002000Ohio1993-199517.014.210.7ant 2005 A2005All states198827.023.717.1ant 2005 B2005Florida199230.021.620.7onrod et al. 20082008Arizona200526.019.020.0uesch 20112011New Jersey2004-200726.722.520.3ozhimannil et al. 20132012All states2002-2009n/an/an/auesch et al. 20142013California201013.910.713.0bastião et al. 20162014Florida2004-201125.222.819.7 |
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| The rates are adjusted as compared to the rates from other studies which are crude rates. Strates bellow 15% benchmark Strates bellow 19% benchmark |

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Caesarean Section in Uninsured Women in the United States: Systematic Review and Meta-analysis

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| Keywords: | caesarean section, health insurance, uninsured, self-pay, medical practice variation, underuse |
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Caesarean Section in Uninsured Women in the United States: Systematic Review and Meta-analysis

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Keywords

caesarean section, health insurance, uninsured, self-pay, access to care, medical practice

variation, health services, financial incentives, underuse, underserved

Word count

Manuscript: 3147 words excluding title page, abstract, references, figures and tables.

Abstract

Objective: The aim of this study is to assess the odds of caesarean section for uninsured women in the United States and understand the underlying mechanisms as well as consequences of lower use.

Study design: Systematic review and meta-analysis.

Data sources: PubMed, Embase, The Cochrane Library and CINAHL from the first year of records through April 2018.

Eligibility criteria: We included studies that reported data to allow the calculation of odds ratios of caesarean section of uninsured as compared to insured women.

Outcomes: The pre-specified primary outcome was the adjusted odds ratio of deliveries by caesarean section of uninured women as compared with privately or publicly insured women. The pre-specified secondary outcome was the crude odds ratio of deliveries by caesarean section of uninsured women as compared with insured women.

Results: Twelve articles describing sixteen separate studies involving more than 8.8 million women were included in this study. We found: 0.70 times lower odds of caesarean section in uninsured as compared to privately insured women (95%CI 0.63 to 0.78), with no relevant heterogeneity between studies ($\tau 2=0.01$); and, 0.92 times lower odds for caesarean section in uninsured as compared to publicly insured women (95%CI 0.80, 1.07), with no relevant heterogeneity between studies ($\tau 2=0.02$). We found 0.70 times lower odds in uninsured as compared to publicly insured women (95%CI 0.69, 0.72).

Conclusions: Caesarean sections are less likely to be performed in uninsured women as compared with insured women. While the higher rates for CS among privately insured women can be explained with financial incentives associated with private insurance, the lower odds among uninsured women draw attention at barriers to access for delivery care. In

many regions the rates for uninsured women are above, close or below the benchmarks for appropriate caesarean section rates and could imply both, underuse and overuse.

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Strengths and limitations of this study

- Extensive literature search, screening and data extraction performed in duplicate, review and analysis of study characteristics as well as thorough quality assessment of included studies.
- ✓ All studies are from one country, i.e. the United States, and this limits the effect of contextual factors.
- A major limitation of our study is the variation across studies, in terms of the study populations characteristics, type of data used, types of caesarean section analyzed and adjusting variables used in statistical analyses.
- ✓ The results of this study are driven by the largest study which contains over two thirds of the population included in this review. Only five out of 16 studies included in the review report data after year 2000.
- While a population level caesarean section rate of less than 9, 10 or 19 percent suggests underuse, we cannot determine the mix of under, over, and appropriate use in a specific population.

Introduction

Introduction of clinical procedures in medical practice have saved and improved the lives of many people worldwide. But with time, these clinical procedures become subject to overuse or underuse; i.e. some people get them without really needing them while others do not get them although in need of them.¹ As a result, overuse of procedures may result in unnecessary harm due to the side effects of the procedures or, in case of underuse, not receiving the care they need.¹⁻³ These adverse effects occur due to differing health systems and other contextual factors.³⁴ These factors include financial and non-financial barriers in accessing healthcare, present even in the most advanced economies of the world, such as the United States (US). Consequently, specific segments of the population may be underserved as healthcare systems are unable to address structural problems that leave patients without the care they need.¹ Globally, Caesarean section (CS) is an example of overuse and underuse of clinical procedures. Once introduced into clinical practice, it greatly improved maternal and newborn outcomes.⁵ Presently, many countries have long exceeded the 9 to 16 percent or 10 to 15 percent thresholds or 19 percent benchmark for CS out of total deliveries, argued to be the ideal rates of CS in terms of improving the health of women and newborns.⁶⁻⁹ CS rates average as high as 40.5 percent among countries in Latin America and the Caribbean region,¹⁰ 32.3 percent in Northern America¹⁰ (32.2 percent in US),¹¹ while on the other extreme, it is as low as 7.3 percent in Africa¹⁰ and known to range even lower in specific countries: 1 percent in Nepal and Cambodia to 0.6 percent in Ethiopia and Niger.¹² Variations are also observed within countries,^{5 12 13} for instance, in the US a recent study reported a range between 4 to 65 percent across health markets.¹⁴

Insurance coverage is one health system factors known to influence the use of medical procedures,^{15 16} including CS.¹⁷⁻²⁰ While private insurance, for example, seems to increase the odds of having a CS delivery,¹⁷ the lack of insurance appears to decrease it.²⁰⁻²² Millions of

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people worldwide, as well as in the US, are not covered by any insurance scheme and are exposed to the hazard of being underserved with clinical procedures.²³⁻²⁷ including perinatal services.²⁸ The US has a mixed health insurance system dominated by private insurance.²⁴ The Federal Medicare program, covers people over 64 years old and/or disabled, which accounts for about 16.7 percent of the population.²⁴ State Medicaid programs cover children and parents from low income families as well as partially caring for Medicare beneficiaries with low incomes and, in total, accounts for about 19.4 percent of the US population.²⁹ Over half of US population is covered with voluntary employer based private insurance.²⁹ The remaining population is uninsured and can range from 2.5 (Massachusetts) to 16.6 (Texas) percent according to 2016 estimates.²⁹ For decades, in the US, there has been an ongoing debate for and against universal health coverage and related topics with limited but substantial progress towards more coverage through the Affordable Care Act (ACA).^{23 30-37} Nonetheless, millions of Americans remain uninsured for various reasons and are not able to access the healthcare they need.^{23 31 38} The aim of this study is to assess the odds of CS for uninsured women in the US and understand the underlying mechanisms as well as consequences of lower use in the US context.³⁹

Materials and methods

Search strategy and data sources

Search words referring to CS, such as 'caesarean section', 'caesarean delivery', 'caesarean', were combined with words referring to factors contributing to variation and increase of CS rates, such as 'insurance', 'social class', 'socioeconomic', and words referring to study design, such as 'geographic variation', 'medical practice variation' (Appendix 1). No publication date or language restrictions were applied. We searched PubMed, Embase, the Cochrane Library and CINAHL from the beginning of records to the end of April 2018, when

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we last updated our search. A manual search was applied on the reference lists of included studies and previous systematic reviews.

Study selection and outcomes

To be included in the analysis, studies had to report odds ratio (OR) or data that enabled the calculation of OR of CS comparing uninsured against privately and/or publicly insured women. More specifically, we didn't exclude studies based on any population characteristic. Studies had to report normal (vaginal) and CS deliveries with uninsured and privately and/or publicly insured comparisons. In an ideal situation, studies would report adjusted OR of uninsured as compared to privately and/or publicly insured women, but in cases ORs were not calculated by the authors, we would extract data (rates and regression coefficients) and perform calculations that would allow for the derivation of OR. We didn't exclude studies by type of study design, variables used for adjustment or any other study characteristic. Adjusted OR of deliveries by CS of uninsured women in comparison to insured women i

Data extraction

Papers screening and independent data extraction was done by two researchers (IH and MB). Differences were resolved based on consensus. We extracted data on study population, study design, data sources, setting, type of CS analyzed, statistical analysis, and (primary and secondary) outcome measures. (Appendix 2)

Quality assessment

We used Quality In Prognostic Studies (QUIPS) to assess the risk of bias across six study domains.⁴⁰ Each study was evaluated independently by two researchers (IH and MB) and any differences among evaluators were discussed and resolved. A single rating was assigned for all studies. As specified in the QUIPS tool, a "high", "moderate", or "low" rating was applied

for individual domains and overall rating of a study.⁴⁰ If a study was rated with a low risk of bias across all the six domains, it would receive an overall rating of low risk of bias.¹⁷ If one or more domains of a study were rated with a moderate risk of bias, it would receive an overall moderate risk of bias.¹⁷ If one or more domains of a study were rated with a high risk of bias, it would receive an overall high risk of bias.¹⁷

Main analysis

Standard inverse-variance random effects meta-analysis was used to combine the overall ORs. An OR lower than one implies a lower frequency of CS in uninsured than in insured women. We calculated τ^2 to measure heterogeneity between studies.⁴¹ Pre-specified cutoffs of τ^2 of 0.04, 0.16 and 0.36 were used to represent low, moderate, and high heterogeneity between studies.⁴² Subgroup analysis by study design, period of data collection, state, type of CS analyzed, parity, inclusion of women with previous CS, pregnancy risk of included women and level of (QUIPS) risk of bias was performed to examine between-study heterogeneity and chi-square test was used to calculate p-values for interaction among subgroups. Test for linear trend was performed in case of more than two ordered strata. All p-values were two-sided. STATA, release 13, was used for analyses (Stata-Corp, College Station, Texas).

Additional analysis

We calculated CS rates among different insurance subgroups for the studies included in the analysis.

Patient involvement

No patients were involved in this study. We used data from published papers only.

Results

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We identified a total of 1837 records: 1123 from PubMed; 556 from Embase; 39 from the Cochrane Library, 119 from CINAHL and 28 from manual search (Figure 1). We removed 240 duplicates. 1597 records were screened for eligibility. We performed full text examination on 177 records. We excluded 139 that did not report insurance status of women⁴³⁻¹⁸¹ and 26 that were otherwise irrelevant.¹⁸²⁻²⁰⁷ (Appendix 3) Finally, 12 records describing 16 separate studies^{20-22 62 208-215} including more than 8.8 million women were included in review and meta-analysis.

Characteristics of studies are presented in Table 1 and Appendices 4, 5, 6 and 7. All studies were from the US. Thirteen studies were cross-sectional and three were retrospective cohort studies. Population size of studies ranged from 9,017 to 6,717,486 cases. Studies used data from years 1986 to 2011 and most studies used hospital records data (Appendix 4). Case exclusion criteria varied considerably (Appendix 5) as well as variables studies used for statistical adjustment (Appendix 6). Appendix 7 reports evaluation of studies using QUIPS risk of bias tool. Four studies were classified with low risk of bias, two studies with moderate risk, and ten studies with high risk of bias (Appendix 7).

Figure 2 presents meta-analyses for primary outcome measure, i.e. adjusted ORs of CS in uninsured women as compared to privately or publicly insured. Since there was a positive interaction between uninsured vs privately insured group and uninsured vs publicly insured group (p=0.016), we performed meta-analyses for each group separately. In the meta-analysis comparing uninsured with privately insured women, including seven studies in 556,454 women, we found that the odds of CS were 0.70 times lower in uninsured as compared to privately insured women (95%CI 0.63 to 0.78), with no relevant heterogeneity between studies (τ 2=0.01). In meta-analysis comparing uninsured with publicly insured women, including four studies in 510,010 women, we found that the odds of CS were 0.20 times lower in uninsured women, including four studies in 510,010 women, we found that the odds of CS were 0.95%CI 0.80 to 1.07), with no

relevant heterogeneity between studies ($\tau 2=0.02$). An additional study in 6,717,486 women, which did not distinguish between privately and publicly insured women,²¹⁵ reported that the odds of CS were 0.70 times lower in uninsured as compared to insured women (95%CI 0.69 to 0.72).

Figure 3 presents results of subgroup analyses of adjusted odds ratios in uninsured vs privately insured women (upper panel) and in uninsured vs publicly insured women (lower panel). In the analysis of uninsured vs privately insured women, estimates varied for subgroups state (p for interaction<0.001), type of CS (p for interaction<0.001), parity (p for interaction=0.07), and pregnancy risk (p for interaction<0.001). There was no positive trend in the period of data collection subgroup. In the lower panel, which presents subgroup analyses of adjusted odds ratios in uninsured vs publicly insured women, estimates varied for subgroups period of data collection (p for interaction=0.03), state (p for interaction=0.004), type of CS (p for interaction=0.03), parity (p for interaction=0.03) and QUIPS risk of bias (p for interaction=0.03).

In Figure 4 we present meta-analyses for crude ORs of CS in uninsured as compared to privately or publicly insured women as secondary outcome. In the meta-analysis comparing uninsured with privately insured women, including eleven studies in 2,010,483 women, we found that the odds of CS were 0.71 times lower in uninsured as compared to privately insured women (95%CI 0.66 to 0.76), with no relevant heterogeneity between studies ($\tau 2=0.018$). In the meta-analysis comparing uninsured with publicly insured women, including eleven studies in 2,010,483 women, we found that the odds of CS were 0.93 times lower in uninsured as compared to publicly insured women, including eleven studies in 2,010,483 women, we found that the odds of CS were 0.93 times lower in uninsured as compared to publicly insured women (95%CI 0.85 to 1.01), with no relevant heterogeneity between studies ($\tau 2=0.017$).

Table 2 presents rates of CS among groups with different insurance status for individual studies. Six studies found CS rates for uninsured women below the 19 percent benchmark.

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One study found CS rates below the 10 percent benchmark. The rates of other studies range from 19.3 percent to 23.0 percent, close to 19 percent benchmark.

Discussion

Our systematic review and meta-analyses estimated that the overall odds of receiving a caesarean section are on average 0.70 times lower for uninsured women as compared with privately insured women (95%CI 0.63 to 0.78), 0.92 times lower for uninsured women as compared with publicly insured women (95%CI 0.80 to 1.07) and 0.70 times lower for uninsured women as compared to privately and publicly insured women (95%CI 0.69 to 0.72). The lower odds were noticed across all subgroups of studies in subgroup analyses as well as in crude analyses.

Context

To our knowledge, this is the first meta-analysis that examines CS rates of uninsured women compared to insured women. Two recently published meta-analyses by our group reported the association of CS with for profit status of hospitals and type of insurance.^{17 216} Investigating the association of for-profit vs non-profit status of hospital with the odds of CS, we found that the odds of CS were 1.41 higher in for-profit hospitals as compared with non-profit hospitals (95% CI 1.24 to 1.60).²¹⁶ The findings were consistent in subgroup analyses.²¹⁶ Investigating the association of CS with private insurance, we found that the odds of CS were 1.13 times higher for privately insured women compared with women covered with public insurance (95% CI 1.07 to 1.18).¹⁷ Again, the increased risk was observed across all subgroups.¹⁷

Strengths and limitations

The major strengths of our meta-analysis include an extensive literature search, screening and data extraction performed in duplicate, review and analysis of study characteristics as well as

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thorough quality assessment of included studies. In addition, all studies are from one country, i.e. the US, and this limits the effect of contextual factors. A major limitation is the variation across studies, in terms of the study populations characteristics (i.e. parity, inclusion of women with previous CS, risk for CS), type of data used, types of CS analyzed and adjusting variables used in statistical analyses. The results of this study are driven by the largest study which contains over two thirds of the population included in this review. Only five out of 16 studies included in the review report data after year 2000. It should also be taken into consideration, that despite similar features, the uninsured are a diverse group of US citizens.²⁶ ²⁷ We considered but could not make use of the Robson criteria to classify studies and analyze CS rates among the studies reviewed. Only two out of sixteen studies could be classified using the Robson criteria.^{62,214} While a population level CS rate of less than 9, 10 or 19 percent suggests underuse, we cannot determine the mix of under, over, and appropriate use in a specific population.

Mechanism

There are several possible explanations why uninsured women have lower odds of CS when compared to insured women. One likely factor is that financial incentives are stronger with private insurance than in the publicly insured or uninsured.^{17,18} These incentives result from higher payment for CS by private insurers through reimbursement arrangements that encourage more expensive procedures as means to increase profits, as well as providers' (hospitals and individual physicians) responses to these incentives.^{17,70,216} The responses to increase by hospitals exist in the form of patient scheduling policies that direct privately insured patients to profit inclined physicians.^{20,216} It is also a known association that physicians who have a higher share of privately insured patients will tend to overuse CS.^{21,22} ²¹⁶ They do so as they perceive patients to have a higher social class, i.e. able to pay higher fees, or fear malpractice liability.^{18,111,208,216}

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Additional reasons are likely reflected in the comparison between uninsured and publicly insured women. A first set of reasons are related to deliberate or forced decisions of uninsured women to keep out-of-pocket payments low.¹⁸ The uninsured patients are more likely to seek less expensive care when they face the need for healthcare services.¹⁸ In the case of giving birth, this would lead to a greater preference for vaginal delivery. A second set of reasons may be discrimination of providers towards uninsured women. Providers have a preference for profitable, i.e. privately insured patients, a preference commonly referred to as "cream skimming".^{21 22 216 217}

Implications for uninsured women

Most studies included in our meta-analysis, including the most recent studies from California²¹³ and Florida,²¹⁴ show that rates for CS among uninsured women are below or close to the 10 and 19 percent benchmarks previously reported.⁶⁻⁸ Even in instances where the average state rates are slightly above the 19 percent benchmark, some hospitals service areas are likely to have CS rates lower than 19 percent or even 9 percent for uninsured women because of the well-established within state variation in CS rates.^{5 111} Uninsured women in these areas are highly likely to be underserved with caesarean section during delivery. Uninsured patients generally have higher unmet needs than insured patients due to access barriers.^{23 24 26 28 218-222} Such barriers encourage inappropriate health seeking behaviors among uninsured.^{23-25 32 220 223-225} Consequently, uninsured populations face higher health risks and have worse healthcare outcomes.^{23-26 32 218-221 223 224 226 227}

The uninsured also face financial burdens which result from out-of-pocket payments that are more severe/extensive than co-payments or premiums that are paid by people that are publicly or privately insured. The uninsured are known to pay higher prices for services as compared to other payers for the same care,^{27 228} spend a high portion of income to cover medical expenses²⁴ (although they spend less for their health compared to patients who have

insurance),²⁶ are frequently charged for full price for healthcare services,^{24 228} often do not benefit from discounts from providers,^{24 27} and face severe financial difficulties.^{23 24} Uninsured manage to pay only part of the costs for their care.²⁶ The remaining costs are uncompensated costs^{23 26 229 230} and most of such costs are covered by the local, state or federal government,^{26 229} eventually resulting in tax increases.²⁶

Implications for research and policy making

Future studies should examine the association of a lack of insurance in pregnant women across health care markets with varying CS rates and assess if delivery outcomes were correspondingly worse, in the effort to investigate the presence of underuse of CS.

In parallel, policy options that could lead to improvements of insurance coverage for delivering women should be assessed in terms of their ability to address healthcare outcomes while keeping overall costs at minimum. In the past, states have adopted different strategies for covering uninsured people.^{24 25 39 231} While there are many known benefits to insurance coverage,^{23 24 32-35 37 221 224 230 232-234} other important policy aspects should be considered. At a time of rising healthcare costs ^{24 35 234 235} regulation of financial incentives is crucial. A revision of payment policies should be pursued ^{17 18 24 216} to align financial incentives with proper health outcomes.^{17 24 216} Reimbursement policies that would pay the same amount for CS and vaginal delivery is one option.^{216 236}

Conclusion

Caesarean sections are less likely to be performed in uninsured women as compared with insured women. The lower odds are consistent in all subgroups and in crude analyses. While the higher rates for CS among privately insured women can be explained with financial incentives associated with private insurance, the lower odds among uninsured women draw attention at barriers to access for delivery care. In many regions, the rates for uninsured women are above, close or below the benchmarks for appropriate CS rates and imply both,

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underuse and overuse. Therefore, efforts to assess the delivery outcomes as well as policy options that could improve insurance coverage for women giving birth are important.

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Contributorship Statement

IH, DG, PJ conceived and designed the study. IH, MB performed the data extraction and preparation. IH, LS analyzed the data. IH, MB, LS drafted the paper, which was critically reviewed and approved by all authors.

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Competing interests statement

All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coi_disclosure.pdf and declare: no support from any organisation for the submitted work; no financial relationships with any organisations that might have an interest in the submitted work in the previous three years; no other relationships or activities that could appear to have influenced the submitted work.

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No funding was received to perform this study. All authors, had full access to all of the data (including statistical reports and tables) in the study and take responsibility for the integrity of the data and the accuracy of the data analysis.

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Data sharing statement

No additional unpublished data are available from the study.

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Table 1. Characteristics of included studies

| Author | Year | State | Study design | Number | Number | Year of | Population | Sampling | Type of |
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| | | | | of cases | of | data | | | CS |
| | | | 0 | | hospital | collection | | | analyzed |
| | | | Ó, | | units | | | | |
| Stafford | 1990 | California | Cross | 461066 | Not | 1986 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | | any risk | | |
| Haas et al. A | 1993 | Massachusetts | Cross | 57257 | Not | 1984 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | e4, | any risk | | |
| Haas et al. B | 1993 | Massachusetts | Cross | 64346 | Not | 1987 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | | any risk | | |
| Braveman et | 1995 | California | Retrospective | 213761 | Unclear | 1991 | Primipara; no | Consecutive | Any |
| al. | | | cohort | | | | previous CS; any risk | | |
| Burns et al. | 1995 | Arizona | Cross | 33233 | 36 | 1989 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | | | any risk | | |

| Onion et al. | 1999 | Maine | Cross | 41177 | Not | 1990- | Primipara; no | Consecutive | Any |
|--------------|------|------------|---------------|--------|----------|-------|-----------------------|-------------|-----|
| Α | | | sectional | | reported | 1992 | previous CS; any risk | | |
| Onion et al. | 1999 | New | Cross | 41401 | Not | 1990- | Primipara; no | Consecutive | Any |
| В | | Hampshire | sectional | | reported | 1992 | previous CS; any risk | | |
| Onion et al. | 1999 | Vermont | Cross | 19077 | Not | 1990- | Primipara; no | Consecutive | Any |
| C | | | sectional | | reported | 1992 | previous CS; any risk | | |
| Aron et al. | 2000 | Ohio | Retrospective | 25697 | 21 | 1993- | Primipara; no | Consecutive | Any |
| | | | cohort | | 10. | 1995 | previous CS; any risk | | |
| Grant A | 2005 | All states | Cross | 9017 | Not | 1988 | Primi- and multipara; | Random | Any |
| | | | sectional | | reported | en, | any risk | | |
| Grant B | 2005 | Florida | Cross | 147821 | Not | 1992 | Primi- and multipara; | Consecutive | Any |
| | | | sectional | | reported | | any risk | | |
| Coonrod et | 2008 | Arizona | Cross | 28863 | 40 | 2005 | Primipara; low risk | Consecutive | Any |
| al. | | | sectional | | | | | | |

| Huesch | 2011 | New Jersey | Cross | 182108 | Not | 2004- | Primi- and multipara; | Consecutive | Planned |
|----------------|---------|------------|---------------|---------|----------|-------|-----------------------|-------------|-----------|
| | | | sectional | | reported | 2007 | no previous CS; low | | |
| | | | | | | | risk | | |
| Kozhimannil | 2013 | All states | Cross | 6717486 | Over | 2002- | Primi- and multipara; | Random | Any |
| et al. | | | sectional | | 1000 | 2009 | any risk | | |
| Huesch et al. | 2014 | California | Cross | 408355 | 254 | 2010 | Primi- and multipara; | Consecutive | Planned |
| | | | sectional | 20- | | | no previous CS; any | | |
| | | | | | 10 | | risk | | |
| Sebastião et | 2016 | Florida | Retrospective | 412192 | 122 | 2004- | Primipara; no | Consecutive | Emergency |
| al. | | | cohort | | | 2011 | previous CS; low risk | | |
| CS = caesarean | section | 1 | | 1 | | | 06. | I | |
| | | | | | | | | | |
| | | | | | | | | | |

| | | | | CS rate | CS rate | |
|-----------------|------|---------------|------------|-----------|----------|-----------|
| | | | | of | of | CS rate |
| | | | Year of | privately | publicly | of |
| | | | data | insured | insured | uninsured |
| Author | Year | State | collection | (%) | (%) | (%) |
| Stafford | 1990 | California | 1986 | 26.8 | 22.1 | 19.3 |
| Haas et al. A | 1993 | Massachusetts | 1984 | 23.0 | 19.4 | 17.2 |
| Haas et al. B | 1993 | Massachusetts | 1987 | 25.9 | 20.8 | 22.4 |
| Braveman et al. | 1995 | California | 1991 | 27.1 | 21.2 | 23.0 |
| Burns et al. | 1995 | Arizona | 1989 | n/a | n/a | n/a |
| | | | 1990- | | | |
| Onion et al. A | 1999 | Maine | 1992 | 15.9 | 14.9 | 13.4 |
| | | New | 1990- | | | |
| Onion et al. B | 1999 | Hampshire | 1992 | 16.1 | 13.2 | 13.0 |
| | | | 1990- | | | |
| Onion et al. C | 1999 | Vermont | 1992 | 14.5 | 13.5 | 9.4 |
| | | | 1993- | | | |
| Aron et al. | 2000 | Ohio | 1995 | 17.0 | 14.2 | 10.7 |
| Grant A | 2005 | All states | 1988 | 27.0 | 23.7 | 17.1 |
| Grant B | 2005 | Florida | 1992 | 30.0 | 21.6 | 20.7 |
| Coonrod et al. | 2008 | Arizona | 2005 | 26.0 | 19.0 | 20.0 |
| | | | 2004- | | | |
| Huesch | 2011 | New Jersey | 2007 | 26.7 | 22.5 | 20.3 |

Table 2. Caesarean section rates among groups with different insurance status

| | | | 2002- | | | |
|--------------------|------|------------|-------|------|------|------|
| Kozhimannil et al. | 2013 | All states | 2009 | n/a | n/a | n/a |
| Huesch et al. | 2014 | California | 2010 | 13.9 | 10.7 | 13.0 |
| | | | 2004- | | | |
| Sebastião et al. | 2016 | Florida | 2011 | 25.2 | 22.8 | 19.7 |

*The rates are adjusted as compared to the rates from other studies which are

crude rates.

CS rates bellow 10% benchmark

CS rates bellow 19% benchmark

Figure legends

Figure 1. The flow diagram of review

Figure 2. Adjusted odds ratios of caesarean section

Figure 3. Subgroup analyses for adjusted estimates/Legend: *P for trend

Figure 4. Crude odds ratios of caesarean section

Supporting information

Appendix 1. Search Strategy

Appendix 2. List of the extracted variables

Appendix 3. List of excluded articles

Appendix 4. Type of data used

Appendix 5. Reported exclusion criteria

Appendix 6. Covariates used for statistical adjustment

Appendix 7. QUIPS risk of bias

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Uninsured vs privately insured



Uninsured vs publicly insured



Lower rate of caesarean section if insured

Uninsured vs publicly or privately insured





529x791mm (96 x 96 DPI)

- ,

Uninsured vs privately insured



Uninsured vs publicly insured





529x784mm (96 x 96 DPI)

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Uninsured vs privately insured



Uninsured vs publicly insured



Figure 4. Crude odds ratios of caesarean section

862x1308mm (96 x 96 DPI)

Appendix 1. Search Strategy

1. For Medline (PubMed)

((((((causes OR determinants OR statistics OR rates OR factors OR decision* OR physician* OR socioeconomic OR state medicine OR evidence-based OR hospital OR hospitals OR hospitalization OR hospitalized OR uncertain* OR educational status OR social class OR obstetric* OR gynecolog* OR supply OR distribut* OR utilization OR insurance OR choice OR attitude OR patient OR economics OR maternal OR accessib* OR health service* OR rural population OR urban population[Title/Abstract])) NOT medline[sb])) OR ("Decision Making"[Mesh] OR "Physician's Practice Patterns" [Mesh] OR "Socioeconomic Factors" [Mesh] OR "State Medicine" [Mesh] OR "Evidence-Based Medicine" [Mesh] OR "Hospitals" [Mesh] OR "Uncertainty" [Mesh] OR "Educational Status" [Mesh] OR "Hospital Costs" [Mesh] OR "Physician Incentive Plans" [Mesh] OR "Social Class" [Mesh] OR "Obstetrics and Gynecology Department, Hospital" [Mesh] OR "supply and distribution"[Subheading] OR "utilization"[Subheading] OR "Insurance"[Mesh] OR "Choice Behavior"[Mesh] OR "Attitude to Health"[Mesh] OR "Patient Participation"[Mesh] OR "Physician-Patient Relations" [Mesh] OR "Economics, Hospital" [Mesh] OR "Maternal Health Services" [Mesh] OR "Health Services Accessibility" [Mesh] OR "Health Services Research" [Mesh] OR "Rural Population"[Mesh] OR "Urban Population"[Mesh]))) OR factors OR rates OR statistics OR causes OR determinants AND (((((operative delivery OR caesarean section OR cesarean section OR c-section OR c section OR caesarean OR caesarean delivery OR cesarean delivery OR caesarean rates OR cesarean rates)))) OR cesarean section [MeSH Terms])) AND ((((("Catchment Area (Health)"[Mesh] OR "Small-Area Analysis"[Mesh]))) OR ((((small area analysis OR small area analyses OR medical practice variation OR regions OR geographic variation OR variation)))))

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| Appendix | 2. List of | the extracted variables |
|------------|------------|--|
| General in | nformatio | n |
| | | Author |
| | | Year |
| | | State |
| | | Study design |
| | | Number of cases |
| | | Number of hospital units |
| | | Year of data collection |
| | | Population |
| | | Sampling |
| | | Type of CS analyzed |
| | | 6 |
| Type of da | ata used | |
| | | Type of data used |
| | | |
| Reported | exclusion | criteria |
| | | Source population |
| | Maternal | Age ≤14 |
| | character | Racial or ethnic minorities |
| | istics | Multipara 💦 |
| | | Previous caesarean section |
| | | Other risk factors for caesarean section |
| | Fetus | Stillbirth |
| | character | Multiple delivery (twin or more) |
| | istics | Newborn weighting <500 gr |
| | | Breach presentation |
| | | Other malpresentation |
| | | Preterm delivery (less than 37 weeks) |
| | | Other risk factors for caesarean section |
|] | Not in lab | or |
| (| Cases with | h missing data |
|] | Provider c | characteristics |
| (| Other fact | ors |
| | | |
| Covariate | s used for | statistical adjustment |
|] | Maternal | Ethnicity/Race |
|] | preconce | Educational level |
|] | ption | Marital status |
| | status | Economic status |
| | | Insurances status |
| | | Urban status |

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| | | Weight | | | | | |
|--------|--|--|--|--|--|--|--|
| | | Height | | | | | |
| | | Body mass index | | | | | |
| | | Age Parity | | | | | |
| | Maternal | | | | | | |
| | clinical | Previous caesarean section | | | | | |
| | status | Pre-existing (before pregnancy) conditions | | | | | |
| | | Conditions developed during pregnancy | | | | | |
| | Fetus | Gestational age | | | | | |
| | character | Birth weight | | | | | |
| | istics | Other characteristics | | | | | |
| | Prenatal c | are | | | | | |
| | Delivery of | characteristics | | | | | |
| | Provider of | characteristics | | | | | |
| | Other vari | ables | | | | | |
| | Total num | iber of covariates | | | | | |
| | | | | | | | |
| QUIPS | 5 risk of bias | | | | | | |
| | Study Par | ticipation | | | | | |
| | Study Att | rition | | | | | |
| | Prognosti | c Factor Measurement | | | | | |
| | Outcome | Measurement | | | | | |
| | Study Con | nfounding | | | | | |
| | Statistical | Analysis and Reporting | | | | | |
| | Overall ra | ting | | | | | |
| | | 4 | | | | | |
| Caesar | rean section | rates among groups with different insurance status | | | | | |
| | State | | | | | | |
| | Year of da | ata collection | | | | | |
| | CS rate of | Eprivately insured (%) | | | | | |
| L | CS rate of | Epublicly insured (%) | | | | | |
| | | $F_{\text{uningurad}}(0/)$ | | | | | |
| | CS rate of | | | | | | |
| Effect | CS rate of | | | | | | |
| Effect | CS rate of estimate Determina | ant being compared | | | | | |
| Effect | CS rate of estimate Determina Comparat | ant being compared or (reference) | | | | | |
| Effect | CS rate of estimate Determina Comparat Unadjuste | ant being compared or (reference) ed outcome measure | | | | | |
| Effect | CS rate of estimate Determina Comparat Unadjuste Effect size | ant being compared or (reference) ed outcome measure | | | | | |
| Effect | CS rate of estimate Determina Comparat Unadjuste Effect size Lower CI | ant being compared or (reference) ed outcome measure e 95% | | | | | |
| Effect | CS rate of estimate Determina Comparat Unadjuste Effect size Lower CI Upper CI | ant being compared or (reference) ed outcome measure e 95% 95% | | | | | |
| Effect | CS rate of estimate Determina Comparat Unadjuste Effect size Lower CI Upper CI SE | ant being compared or (reference) ed outcome measure e 95% | | | | | |
| Effect | CS rate of estimate Determina Comparat Unadjuste Effect size Lower CI Upper CI SE Determina | ant being compared or (reference) ed outcome measure e 95% 95% 05% | | | | | |

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| | Adjusted outcome measure | | | | | |
|------|---------------------------|--|--|--|--|--|
| | Effect size | | | | | |
| | Lower CI95% | | | | | |
| | Upper CI95% | | | | | |
| | SE | | | | | |
| | | | | | | |
| Numb | per of cases by groups | | | | | |
| | Total births (all groups) | | | | | |
| | Total No-CS (all groups) | | | | | |
| | Total CS (all groups) | | | | | |
| | Total births in group 1 | | | | | |
| | No-CS in group 1 | | | | | |
| | CS in group 1 | | | | | |
| | Total births in group 2 | | | | | |
| | No-CS in group 2 | | | | | |
| | CS in group 2 | | | | | |
| | Total births in group 3 | | | | | |
| | No-CS in group 3 | | | | | |
| | CS in group 3 | | | | | |

Appendix 3. List of excluded articles

| 4 | Author, Year | Reason for exclusion |
|----------|-----------------------------------|--|
| 5 | 1 Adhikari Dahal et al. 2017 | Didn't report on insurance status of women |
| 0 7 | 2 American en et el 2016 | Didn't report on insurance status of women |
| 8 | 2 Armstrong, et al., 2016 | Didn t report on insurance status of women |
| 9 | 3 Bailit, et al., 2006 | Didn't report on insurance status of women |
| 10 | 4 Bannister-Tyrrell, et al., 2015 | Didn't report on insurance status of women |
| 11 | 5 Blais, 1993 | Didn't report on insurance status of women |
| 12 | 6 Brown, 2007 | Didn't report on insurance status of women |
| 14 | 7 Brown, et al., 2013 | Other non-relevant studies |
| 15 | 8 Butcher, et al., 1997 | Didn't report on insurance status of women |
| 16 | 9 Caceres, et al., 2013 | Didn't report on insurance status of women |
| 17 | 10 Carayol, et al., 2007 | Didn't report on insurance status of women |
| 19 | 11 Carayol, et al., 2007 | Didn't report on insurance status of women |
| 20 | 12 Carayol, et al., 2008 | Didn't report on insurance status of women |
| 21 | 13 Carlisle, et al., 1996 | Didn't report on insurance status of women |
| 22 | 14 Chauhan, et al., 2008 | Other non-relevant studies |
| 24 | 15 Chen, et al., 2003 | Other non-relevant studies |
| 25 | 16 Chen, et al., 2014 | Didn't report on insurance status of women |
| 26 27 | 17 Chen. et al., 2016 | Didn't report on insurance status of women |
| 28 | 18 Cheng. et al., 2015 | Other non-relevant studies |
| 29 | 19 Cisse, et al., 1998 | Other non-relevant studies |
| 30 | 20 Clark, et al., 2007 | Didn't report on insurance status of women |
| 32 | 21 Clark et al. 2014 | Didn't report on insurance status of women |
| 33 | 22 Clarke et al 1995 | Didn't report on insurance status of women |
| 34 | 23 Clarke et al 1996 | Didn't report on insurance status of women |
| 35 | 24 Clayton et al. 2013 | Didn't report on insurance status of women |
| 37 | 24 Clayton, et al., 2013 | Didn't report on insurance status of women |
| 38 | 25 Coolind, et al., 2008 | Didn't report on insurance status of women |
| 39 | 26 Coulm, et al., 2012 | Didn't report on insurance status of women |
| 40 41 | 27 Cressie, 1993 | Other non-relevant studies |
| 42 | 28 Da Silva Campi, et al., 2014 | Didn't report on insurance status of women |
| 43 | 29 da Silva, et al., 2003 | Other non-relevant studies |
| 44 | 30 Danishevski, et al., 2008 | Other non-relevant studies |
| 45 | 31 Daw, et al., 2018 | Didn't report on insurance status of women |
| 40 | 32 de Regt, et al., 1986 | Didn't report on insurance status of women |
| 48 | 33 Di Mario, et al., 2013 | Didn't report on insurance status of women |
| 49 | 34 Dimitrov, 1998 | Other non-relevant studies |
| 50 51 | 35 Eckerlund, et al., 1998 | Didn't report on insurance status of women |
| 52 | 36 Edmonds, et al., 2015 | Other non-relevant studies |
| 53 | 37 Edmonds, et al., 2017 | Didn't report on insurance status of women |
| 54 | 38 Emmett, et al., 2010 | Other non-relevant studies |
| 55 56 | 39 Epstein, et al., 2009 | Didn't report on insurance status of women |
| 57 | 40 Franca, et al., 2016 | Didn't report on insurance status of women |
| 58 | · · · · | 1 |
41 Gama, et al., 2009 42 Garcia, et al., 2001 43 Gates, 1995 44 Gittelsohn, et al., 1995 45 Gomes, et al., 1999 46 Gonzalez-Perez, et al., 2001 47 Goyert, et al., 1989 48 Gregory, et al., 2001 49 Gross, et al., 2015 50 Grytten, et al., 2011 51 Grytten, et al., 2012 52 Gumede, et al., 2017 53 Hanley, et al., 2010 54 Haraldsdottir, et al., 2015 55 Haupt, 1982 56 Heffner, et al., 2003 57 Helfand, et al., 1997 58 Henke, et al., 2014 59 Hofmeyr, et al., 2015 60 Hopkins, et al., 2014 61 Hsu, et al., 2008 62 Hueston, et al., 2001 63 Jessee, et al., 1982 64 Johnson, et al., 1995 65 Joyce, et al., 2002 66 Kennare, 2003 67 Keskimaki, et al., 1994 68 Khan, et al., 2017 69 Kim, et al., 2012 70 Kim, et al., 2016 71 Kimsey, et al., 2017 72 Klassen, 1975 73 Klemetti, et al., 2010 74 Koroukian, et al., 2001 75 Korst, et al., 2005 76 Kozhimannil, et al., 2014 77 Krivenko, et al., 1994 78 Kyu-Tae, et al., 2017 79 Lee, et al., 2007 80 Lee, et al., 2014 81 Leung, et al., 2001 82 Li, et al., 2017 83 Librero, et al., 2000

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Other non-relevant studies Didn't report on insurance status of women Other non-relevant studies Didn't report on insurance status of women

Didn't report on insurance status of women Didn't report on insurance status of women Other non-relevant studies

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| 2 | 84 | Lidegaard, et al., 1994 | Didn't report on insurance status of women |
|----------|-----|-----------------------------|--|
| 3 | 85 | Little, et al., 2015 | Didn't report on insurance status of women |
| 4 | 86 | Liu. et al., 2007 | Didn't report on insurance status of women |
| 5 | 87 | Localio, et al., 1993 | Didn't report on insurance status of women |
| 7 | 88 | Lundsberg, et al., 2017 | Didn't report on insurance status of women |
| 8 | 89 | Lutomski, et al. | Didn't report on insurance status of women |
| 9 10 | 90 | Lutomski et al. 2014 | Didn't report on insurance status of women |
| 10 | 91 | Maeda et al 2018 | Didn't report on insurance status of women |
| 12 | 92 | Marquez-Calderon et al 2011 | Didn't report on insurance status of women |
| 13 | 93 | McKenzie et al 1993 | Didn't report on insurance status of women |
| 14 15 | 0/ | Menard 1000 | Didn't report on insurance status of women |
| 16 | 05 | Mondlovia et al 2017 | Didn't report on insurance status of women |
| 17 | 95 | Mestartan et al. 2017 | Didn't report on insurance status of women |
| 18 | 90 | Milaslaisanda et al. 2017 | Didn't report on insurance status of women |
| 19 20 | 97 | Mikolajczyk, et al., 2013 | Didn't report on insurance status of women |
| 21 | 98 | Mindell, et al., 1982 | Didn't report on insurance status of women |
| 22 | 99 | Misra, 2008 | Didn't report on insurance status of women |
| 23 | 100 | Mitler, et al., 2000 | Didn't report on insurance status of women |
| 24 25 | 101 | Mossialos, et al., 2005 | Didn't report on insurance status of women |
| 26 | 102 | Movsas, et al., 2012 | Didn't report on insurance status of women |
| 27 | 103 | Murray, 2000 | Didn't report on insurance status of women |
| 28 | 104 | Murray, et al., 1997 | Didn't report on insurance status of women |
| 29 30 | 105 | Naiditch, et al., 1997 | Didn't report on insurance status of women |
| 31 | 106 | Newton, et al., 1989 | Didn't report on insurance status of women |
| 32 | 107 | Nicholson, et al., 2009 | Didn't report on insurance status of women |
| 33 | 108 | Nigam, 2011 | Didn't report on insurance status of women |
| 34 35 | 109 | Nilsen, et al., 2014 | Didn't report on insurance status of women |
| 36 | 110 | Nirupam, et al., 1995 | Other non-relevant studies |
| 37 | 111 | Oleske, et al., 1991 | Didn't report on insurance status of women |
| 38 | 112 | Ono, et al., 2016 | Didn't report on insurance status of women |
| 39 40 | 113 | Paranjothy, et al., 2005 | Didn't report on insurance status of women |
| 41 | 114 | Parazzini, et al., 2015 | Didn't report on insurance status of women |
| 42 | 115 | Pel et al 1995 | Other non-relevant studies |
| 43 44 | 116 | Phinps et al 2014 | Other non-relevant studies |
| 44 | 117 | Placek et al 1980 | Didn't report on insurance status of women |
| 46 | 118 | Rabilloud et al. 1998 | Didn't report on insurance status of women |
| 47 | 110 | Radifioud, et al., 1998 | Didn't report on insurance status of women |
| 48 49 | 120 | Ramman, et al., 2014 | Didn't report on insurance status of women |
| 50 | 120 | Ratifier, 1990 | Other non-relevant studies |
| 51 | 121 | Ravindran, 2003 | Diller non-relevant studies |
| 52 | 122 | Ravindran, 2008 | Didn't report on insurance status of women |
| 53 54 | 123 | Renzi, et al., 2012 | Didn't report on insurance status of women |
| 55 | 124 | Ribeiro, et al., 2007 | Didn't report on insurance status of women |
| 56 | 125 | Riddell, et al., 2017 | Didn't report on insurance status of women |
| 57 | 126 | Rohrer, 1993 | Other non-relevant studies |
| 58 59 | | | |

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127 Roohan, et al., 2001 128 Rossignol, et al., 2013 129 Rowe, et al., 2014 130 Sandall, et al., 2013 131 Sarria Santamera, et al., 1994 132 Schemann, et al., 2015 133 Schemann, et al., 2016 134 Sentell, et al., 2016 135 Shiono, et al., 1987 136 Shorten, et al., 2007 137 Signorelli, et al., 1991 138 Singata, et al., 2013 139 Snyder, et al., 2011 140 Souza, et al., 2016 141 Sufang, et al., 2007 142 Tang, et al., 2006 143 Tang, et al., 2006 144 Tracy, et al., 2006 145 Tucker, et al., 146 Tussing, et al., 1994 147 Vadnais, et al., 2017 148 Vankan, et al., 2017 149 Vayda, et al., 1984 150 Vecino-Ortiz, et al., 151 Wang, et al., 2017 152 Ward, et al., 2010 153 Weber, 1990 154 Wei, et al., 2013 155 Woolbright, 1996 156 Xing Lin, et al., 2012 157 Xirasagar, et al., 2004 158 Xirasagar, et al., 2006 159 Xirasagar, et al., 2007 160 Yang, et al., 2014 161 Yi-Chen, et al., 2012 162 Zdeb, et al., 1980 163 Zere, et al., 2010 164 Zhang, et al., 2013 165 Zwecker, et al., 2011

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Appendix 5. Reported exclusion criteria

| | | | N | laterna | l chara | cteristi | cs | | | Fetus o | charact | eristics | | | | а | s | |
|--------------------|------|--|---------|--------------------------------|-----------|-------------------------------|---|------------|-------------------------------------|------------------------------|---------------------|-----------------------|--|---|--------------|------------------------|-------------------------|---------------|
| Authors | Year | Source population | Age ≤14 | Racial or ethnic minorities | Multipara | Previous caesarean section | Other risk factors for caesarean section | Stillbirth | Multiple delivery (twin or more) | Newborn weighting <500 gr | Breach presentation | Other malpresentation | Preterm delivery (less than 37 weeks) | Other risk factors for caesarean section | Not in labor | Cases with missing dat | Provider characteristic | Other factors |
| Stafford | 1990 | All deliveries in California, United States | | | | | | | | | | | | | | | + | |
| Haas et al. A | 1993 | All deliveries in Massachusetts, United States | | | | | | + | + | + | | | | | | + | | |
| Haas et al. B | 1993 | All deliveries in Massachusetts, United States | | | | | | + | + | + | | | | | | + | | |
| Braveman et al. | 1995 | All deliveries in California, United States | | | + | + | | + | + | | | | + | | | + | | |
| Burns et al. | 1995 | All deliveries in Arizona, United States | | | | | | | | | | | | | | + | + | |
| Onion et al. A | 1999 | All deliveries in Maine, United States | | | + | + | | | + | | | | | | | | | + |
| Onion et al. B | 1999 | All deliveries in New Hampshire, United States | | | + | + | | | + | | | | | | | | | + |
| Onion et al. C | 1999 | All deliveries in Vermont, United States | | | + | + | | | + | | | | | | | | | + |
| Aron et al. | 2000 | All deliveries in Cleveland, Ohio, United States | | | | + | | | | +* | | | | | | + | + | + |
| Grant A | 2005 | All deliveries, United States | | | | | | | | | | | | | | + | | |
| Grant B | 2005 | All deliveries in Florida, United States | | | | | | | | | | | | | | + | + | + |
| Coonrod et al. | 2008 | All deliveries in Arizona, United States | | + | + | | | + | + | | + | + | + | | | | + | |
| Huesch | 2011 | All deliveries in New Jersey, United States | | | | + | + | | + | | + | + | + | + | + | | + | |
| Kozhimannil et al. | 2013 | All deliveries in 44 states, United States | | | | | | | | | | | | | | | + | |
| Huesch et al. | 2014 | All deliveries in California, United States | + | | | + | | | | | | | | + | | + | | |
| Sebastião et al. | 2016 | All deliveries in Florida, United States | | | + | + | | + | + | | + | + | + | | + | + | + | + |

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| App | endix 6. | Covariates | used for | statistical | adjustment |
|-----|----------|------------|----------|-------------|------------|
|-----|----------|------------|----------|-------------|------------|

| | | | | I | Materna | al preco | onceptio | on statu | IS | | | Mate | ernal c | linical s | tatus | cha | Fetus racteris | stics | | | | | |
|-------------------|------|----------------|-------------------|----------------|-----------------|-------------------|--------------|----------|--------|-----------------|-----|--------|----------------------------|---|--|-----------------|-------------------|-----------------------|---------------|--------------------------|--------------------------|-----------------|----------------------------|
| uthor | Year | Ethnicity/Race | Educational level | Marital status | Economic status | insurances status | Urban status | Weight | Height | Body mass index | Age | Parity | Previous caesarean section | Pre-existing (before pregnancy) conditions | Conditions developed Juring pregnancy | Gestational age | Birth weight | Other characteristics | Prenatal care | Delivery characteristics | Provider characteristics | Other variables | Total number of coverintee |
| tafford* | 1990 | + | | | | _ | | | | | 1 | - | | | | | _ | • | | | | | Ť |
| aas et al. A* | 1993 | | | | | | | | | | | | | | | | | | | | | | 1 |
| aas et al. B* | 1993 | | | 4 | | 6 | | | | | | | | | | | | | | | | | 1 |
| raveman et al. | 1995 | + | + | + | + | + | | | | | + | | | | + | | + | + | + | + | ++ | + | |
| urns et al. | 1995 | + | + | | | | | | | | + | + | + | | ++ | + | + | ++ | + | | ++ | ++ | |
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| Appendix 7. | OUIPS | risk | of bias |
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| representation / · | QUID | 1191 | or blub |

| Author Year Participation Study Attrition Measurement Goalounding Reporting Overall rating Stafford 1990 low low low low low high moderate high Hass et al. 1993 low low low low low moderate high Braveman et al. 1995 low low low low moderate low | VathorYearParticipationStudy AttritionMeasurementConfoundingReportingOverall ratinisafford1990lowlowlowlowlowhighmoderatehighasa et al.1993lowlowlowlowlowhighmoderatehighasa et al.1993lowlowlowlowlowmoderatelowmoderatearrena et al.1995lowlowlowlowmoderatelowmoderatearrena et al.1995lowlowlowlowmoderatelowmoderatearrena et al.1995lowlowlowlowmoderatelowmoderatearrena et al.1995lowlowlowlowlowmoderatelowmoderatearrena et al.1999lowlowlowlowlowmoderatelowmoderatearrena et al.1999lowlowlowlowlowlowlowlowlowarrena et al.1999lowlowlowlowlowlowlowlowlowlowarrena et al.2000lowlowlowlowlowlowlowlowlowlowarrena et al.2005moderatehighlowlowlowlowlowlowlowarrena et al.2005lowlowlowlow <td< th=""><th>Author Vear Participation Study Attrition Industry and the summer of the summer</th><th></th><th></th><th>Study</th><th></th><th>Prognostic Factor</th><th>Outcome</th><th>Study</th><th>Analysis and</th><th></th></td<> | Author Vear Participation Study Attrition Industry and the summer of the summer | | | Study | | Prognostic Factor | Outcome | Study | Analysis and | |
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Research Checklist

According to MOOSE statement for meta-analyses of observational studies

| Reporting of background should include | Where to find in manuscript |
|---|---------------------------------|
| Problem definition | Manuscript (page 5, 6) |
| Hypothesis statement | Manuscript (page 5, 6) |
| Description of study outcome(s) | Manuscript (page 6) |
| 5 Type of exposure or intervention used | Manuscript (page 6) |
| Type of study designs used | Manuscript (page 6, 7) |
| Study population | Manuscript (page 6, 7) Table 1, |
| | Appendix 1 |
| Reporting of search strategy should include | |
| Qualifications of searchers (eg, librarians and investigators) | Manuscript (page 1) |
| Search strategy, including time period included in the synthesis and | Manuscript (page 6), Appendix 1 |
| keywords | |
| Effort to include all available studies, including contact with authors | Manuscript (page 6) |
| Databases and registries searched | Manuscript (page 6) |
| Search software used, name and version, including special features | Manuscript (page 6) |
| used (eg, explosion) | |
| Use of hand searching (eg, reference lists of obtained articles) | Manuscript (page 6) |
| List of citations located and those excluded, including justification | Figure 1 |
| Method of addressing articles published in languages other than | n/a |
| English | |
| Method of handling abstracts and unpublished studies | Manuscript (page 6, 7) |
| Description of any contact with authors | No contact made |
| Reporting of methods should include | |
| Description of relevance or appropriateness of studies assembled for | Manuscript (page 6, 7) |
| assessing the hypothesis to be tested | |

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Rationale for the selection and coding of data (eg, sound clinical Manuscript (page 6, 7) principles or convenience) Documentation of how data were classified and coded (eg, multiple Manuscript (pages 6, 7) raters, blinding, and interrater reliability) Assessment of confounding (eg, comparability of cases and controls Manuscript (page 6-7), Appendix 2, 3, in studies where appropriate) 4 Assessment of study quality, including blinding of quality assessors; Manuscript (page 7), Figure 2, stratification or regression on possible predictors of study results Appendix 5, Assessment of heterogeneity Manuscript (page 7) Description of statistical methods (eg, complete description of fixed Manuscript (page 7) or random effects models, justification of whether the chosen models account for predictors of study results, dose-response models, or cumulative meta-analysis) in sufficient detail to be replicated Provision of appropriate tables and graphics Table 1, Figure 1-3 and Appendixes 1-7 **Reporting of results should include** Graphic summarizing individual study estimates and overall Figure 2, Appendix 6 estimate Table giving descriptive information for each study included Table 1 Results of sensitivity testing (eg, subgroup analysis) Figure 3 Indication of statistical uncertainty of findings Manuscript, Figure 2-4 **Reporting of discussion should include** Quantitative assessment of bias (eg, publication bias) Manuscript (page 8) Justification for exclusion (eg, exclusion of non-English-language Manuscript (page 8) citations) Assessment of quality of included studies Manuscript (page 8)

Reporting of conclusions should include

| Consideration of alternative explanations for observed results | Manuscript (pages 9-13) |
|---|-------------------------|
| Generalization of the conclusions (ie, appropriate for the data | Manuscript (page 13) |
| presented and within the domain of the literature review) | |
| Guidelines for future research | Manuscript (page 13) |
| Disclosure of funding source | Manuscript (page 13) |
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