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# BMJ Open Impact of obstetric unit closures, travel time and distance to obstetric services on maternal and neonatal outcomes in high-income countries: a systematic review

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To cite: Malouf RS. Tomlinson C. Henderson J, et al. Impact of obstetric unit closures, travel time and distance to obstetric services on maternal and neonatal outcomes in high-income countries: a systematic review. BMJ Open 2020;10:e036852. doi:10.1136/ bmjopen-2020-036852

Prepublication history and additional material for this paper is available online. To view these files, please visit the journal online (http://dx.doi.org/10. 1136/bmjopen-2020-036852).

Received 13 January 2020 Revised 07 September 2020 Accepted 23 September 2020



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#### **ABSTRACT**

Objectives To systematically review (1) The effect of obstetric unit (OU) closures on maternal and neonatal outcomes and (2) The association between travel distance/ time to an OU and maternal and neonatal outcomes.

Design Systematic review of any quantitative studies with a comparison group.

Data sources Embase, MEDLINE, PsycINFO, Applied Social Science Index and Abstracts, Cumulative Index to Nursing and Allied Health and grey literature were searched.

Methods Eligible studies explored the impact of closure of an OU or the effect of travel distance/time on prespecified maternal or neonatal outcomes. Only studies of women giving birth in high-income countries with universal health coverage of maternity services comparable to the UK were included. Identification of studies, extraction of data and risk of bias assessment were undertaken by at least two reviewers independently. The risk of bias checklist was based on the Cochrane Effective Practice and Organisation of Care criteria and the Newcastle-Ottawa scale. Heterogeneity across studies precluded meta-analysis and synthesis was narrative, with key findings tabulated.

**Results** 31 studies met the inclusion criteria. There was some evidence to suggest an increase in babies born before arrival following OU closures and/or associated with longer travel distances or time. This may be associated with an increased risk of perinatal or neonatal mortality, but this finding was not consistent across studies. Evidence on other maternal and neonatal outcomes was limited but did not suggest worse outcomes after closures or with longer travel times/ distances. Interpretation of findings for some studies was hampered by concerns around how accurately exposures were measured, and/or a lack of adjustment for confounders or temporal changes.

Conclusion It is not possible to conclude from this review whether OU closure, increased travel distances or times are associated with worse outcomes for the mother or the

PROSPERO registration number CRD42017078503.

# Strengths and limitations of this study

- ► This review is the first to synthesise systematically the current evidence on the impact of closure of obstetric units and of travel distance and travel time to obstetric units on neonatal and maternal outcomes.
- Rigorous systematic review methodology was applied including a sensitive search strategy to ensure all relevant evidence was identified.
- Heterogeneity across included studies precluded any form of meta-analysis.
- A paucity of evidence on a number of outcomes, and methodological concerns for some studies limited conclusions that could be drawn.

#### **BACKGROUND**

Closure of small obstetric units (OUs) and centralisation of obstetric services in larger units has been proposed to increase levels of consultant obstetrician cover to improve safety and limit costs. However, closure of OUs or conversion of OUs to midwifery-led units/community-based services potentially leads to an increase in travel distance or time for women in labour from their home to the nearest OU. Increases in travel time could potentially increase the risk of adverse birth outcomes.

Travel time and distance are widely used as measures to explore the geographical accessibility of health services. In a systematic review,2 the association between travelling further to healthcare facilities and having worse health outcomes was established, but the review did not include studies of maternity care. The impact of OU closure and increase in travel time/distance to the OU on perinatal and maternal outcomes have not been systematically assessed. One





review<sup>3</sup> evaluating the effects of regionalisation of perinatal services has been published. This concluded that regionalisation programmes appeared to be correlated with improvements in perinatal outcomes but that the evidence was weak. A narrative review<sup>4</sup> included 10 studies that explored travel time and distance to and between maternity services and adverse birth outcomes to inform the consultation on maternity services in Wales. The review was limited to studies reported in English and there was no clear association between travel distance or time and adverse birth outcomes

Therefore, uncertainty remains about the association between OU closure, prolonged time or distance to OUs and adverse perinatal outcomes. Specifically, there is a rise in the risk of babies born before arrival (BBA, also referred to as unplanned out of hospital births). Being BBA is more common before term and has been reported to be associated with higher perinatal mortality (PM).<sup>5</sup> Conversely, Lasswellet *et al*<sup>6</sup> found neonatal mortality (NM) was reduced when services were configured to ensure very preterm infants are born in a large maternity hospital with neonatal intensive care unit (level III NICU). In addition to mortality, Apgar scores (a standardised measure of the physical condition of a newborn infant) and neonatal admission to intensive care provide an indication of perinatal infant health.

The impact on maternal outcomes is also unclear. There are concerns that low-risk women who give birth in larger hospitals may experience more interventions, for example, increased frequency of caesarean section (CS). Along with CS, evidence on maternal mortality (MM) and maternal birth complications such as postpartum haemorrhage (PPH) and maternal blood transfusion, was also sought in this review to identify the potential impact of OU closure on maternal outcomes.

In this review, we aimed to systematically identify, critically appraise and synthesise the evidence relating to: (1) The effect of OU closures on maternal and neonatal outcomes (compared with the surrounding area or a comparable population) and (2) The association between travel distance or time to an OU and maternal and neonatal outcomes.

#### **REVIEW METHOD**

The Meta-Analyses and Systematic Reviews of Observational Studies in Epidemiology (MOOSE) reporting guideline was followed.<sup>8</sup>

# Criteria for considering studies for this review

#### Types of studies

Any quantitative study design with a comparison group was eligible for inclusion. Studies were included from 1990 onwards. The year 1990 was chosen as a cut-off date because significant advances were made in neonatal care in the early 1990s, such as surfactant therapy, assisted ventilation, prophylactic infection control and antenatal steroid therapy, which impacted on the delivery

of maternity services. <sup>9</sup> The quantitative components of mixed methods studies were also eligible. Studies were included if they:

Explored the impact of closure of an OU on maternal or neonatal outcomes either in a before-and-after comparison (same population catchment area), or a geographical comparison of different areas (comparable populations).

And/or

- ► Compared maternal and neonatal outcomes after an OU closure and retention or creation of midwifery led units to replace the OU.
- ▶ Explored the effect of travel time and/or distance on maternal and neonatal outcomes providing at least two travel times and/or distances from women's homes to the nearest OU.
- ► Explored maternal and neonatal outcomes following maternal transfer from planned or unplanned home birth to the nearest maternity centre.

We included studies of women giving birth in high-income, the Organisation for Economic Co-operation and development (OECD) countries with universal health coverage (UHC) of maternity services comparable to the UK. The list of OECD countries is shown in online supplemental appendix 1. UHC is defined as healthcare that meets everyone's right to access high quality essential health services where and when they need them without financial difficulty. <sup>10</sup>

# Types of exposures

OU closure: the closure of an OU was compared with no closure of an OU for the same or comparable geographical catchment areas prior to the closure. For a study comparing different geographical areas affected by the closure of an OU, the least affected area was used as a control group. For the purpose of this review, we used the definition of an OU used in the Birthplace Research programme in England, 11 which defined an OU as 'a clinical location in which care is provided by a team, with obstetricians taking primary professional responsibility for women at high risk of complications during labour and birth. Midwives offer care to all women in an OU, whether or not they are considered at high or low risk, and take primary responsibility for women with straightforward pregnancies during labour and birth. Diagnostic and treatment medical services including obstetric, neonatal and anaesthetic care are available on site, 24 hours a day' (P12).

Travel distance or time to the nearest OU: a shorter travel distance or time was compared with a longer travel distance or time. We used the definition of a shorter or a longer time or distance as defined by the included studies. When a study compared several different travel times or distances to the nearest OU, those with the shortest travel distance or time were used as the control group.

The following types of studies were excluded:

 Studies comparing maternal and or neonatal outcomes based on hospital size, level of NICU, type



- of hospital or model of care (eg, caseload midwifery care vs consultant care).
- ► Studies on regionalisation of neonatal care (number of centres with NICUs).
- ► Studies where a proximity rather than the actual travel time or travel distance was given (eg, rural vs urban, remote vs very remote areas).
- Studies which did not report at least one of the outcomes.

#### Review outcomes

The following outcomes were predefined in the study protocol:

#### Maternal outcomes

Maternal mortality (MM), caesarean section (CS) (overall, emergency or intrapartum), severe perineal trauma (including third and fourth degree tears), postpartum haemorrhage (PPH), maternal admission to intensive care units (ICU) and maternal blood transfusion.

#### Neonatal outcomes

Stillbirth (SB) (overall or intrapartum), neonatal mortality (NM), PM, infant mortality (IM), babies BBA, neonatal unit admission (NNU), Apgar score and hypoxic-ischaemic encephalopathy (HIE).

# **REVIEW METHODS**

A comprehensive search strategy was developed in collaboration with an information specialist (NR). We searched Embase, Medline, PsycINFO, Applied Social Science Index and Abstracts and Cumulative Index to Nursing and Allied Health databases (from 1990 to February 2019). We also searched the grey literature in the databanks of British Library EThOS, Open Grey and ProQuest Dissertations & Theses Global. National Health Service (NHS) Trusts and Health Boards in the UK were also contacted where we had been able to identify an OU closure to request information about any evaluations that were conducted. The references of eligible studies and relevant reviews were checked to identify additional studies not retrieved by the search. Searches were based on index terms and text words relating to the population/setting (eg, maternity service, pregnancy, neonatal) and exposures (eg, travel/distance or closure/regionalisation). Due to the variable nature of terms and indexing used, the strategy was kept broad by using a range of alternate terms and not limiting by outcome. No language restriction was applied. A sample search strategy for MEDLINE is shown in online supplemental appendix 2.

At least two reviewers (RSM, CT, AP, FA and JH) independently screened the references for relevance against the review eligibility criteria using Eppi-reviewer software (V.4).<sup>12</sup> Full-text study screening was also performed by at least two reviewers (RSM, CT, CO, JH and FA). Disagreements regarding study eligibility were resolved through discussion and consensus within the review

team. We contacted authors of relevant studies published as abstracts for further information. Data extraction and risk of bias assessment were undertaken by at least two reviewers (RSM, CT, CO, JH, FA and JD). The risk of bias checklist was adapted from the Effective Practice and Organisation of Care (EPOC) and the Newcastle-Ottawa scale (for case–control studies). Risk of bias assessment included selection of study groups, measurement of exposure and outcomes, missing data and appropriateness of analysis (eg, logistic regression analysis). For case–control studies, selection and comparability of cases and controls were also considered. The review team rated the quality of evidence for each domain in the tool as low, high or unclear risk of bias, or yes, no and unclear in meeting quality criteria.

Results were synthesised narratively and the key findings tabulated. The included studies varied in their study design, categories of exposure, outcomes reported, whether adjusted or unadjusted results were presented and factors adjusted for. This clinical and methodological heterogeneity across the included studies precluded any form of meta-analysis. Prespecified subgroups were risk status of woman (low vs high), parity, gestational age, UK studies compared with non-UK studies and planned versus unplanned CS; formal subgroup analyses were, however, not possible. Evidence regarding OU closure, travel distance and travel time is reported separately, and by outcome. We have highlighted where crude (unadjusted) ORs (cOR) and adjusted ORs (adjOR) have been reported.

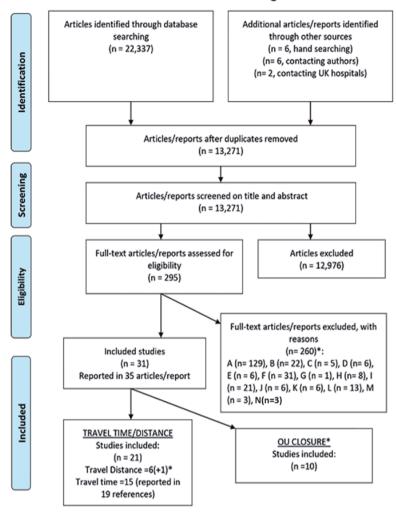
#### PATIENT AND PUBLIC INVOLVEMENT

We involved our parent, patient and public involvement (PPPI) Stakeholders Network, to explore which outcomes were important from a maternal perspective. The dissemination of findings to stakeholders will be through plain language summaries developed with members of our PPPI stakeholder network.

#### Search results

Searches of bibliographic databases and other sources from 1990 to February 2019 yielded 13 271 unique references and the steps of study selection are presented in the Preferred Reporting Items for Systematic Reviews and Meta-Analyses flow chart (figure 1). The eligibility of 295 full-text articles were assessed independently. Two hundred and sixty articles were excluded for various reasons, including: studies conducted in low-income/ middle-income countries, comparing different models or levels of maternity care, assessing women's transfer from primary to secondary maternity centres, or not providing quantifiable measures of travel time/distance (full list available from authors). Thirty-one studies, reported in 35 articles, met the review eligibility criteria (figure 1). One study included information on both OU closure and travel distance. Ten studies provided information on OU closures, 7 studies compared different travel distances

# **PRISMA Flow Diagram**



\*One study, Blondel 2011 contribute data to both travel distance and closure review questions

From: Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred Reporting items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(7): e1000097. doi:10.1371/journal.pmed1000097

For more information, visit  $\underline{www.prisma-statement.org}.$ 

#### Reason for exclusion:

- A. Different study designs
- B. Studies comparing different models of maternity care
- C. Studies comparing different hospital volumes
- D. Studies comparing different levels of care
- E. Studies comparing different hospital types (private vs. public, local vs. central)
- F. Transfer from primary to secondary birth centres
- G. Cost of care
- H. Neonatal transfer
- I. Regionalisation
- J. Not in high-income countries or in the USA
- K. Distance or travel time proximity
- L. No outcomes
- M. Modelling studies
- N. Not found

Figure 1 PRISMA flow diagram. PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses.

from women's homes to the nearest OU and 15 studies compared different travel times from women's homes to the nearest OU.

REVIEW RESULTS Evidence from OU closures

A detailed description of the ten included OU closure



studies is presented in table 1. Three studies were from the UK, with two reported as abstracts only <sup>15</sup> <sup>16</sup> and one an unpublished data series from East Lancashire Hospitals NHS Trust, UK (East Lancashire Hospitals NHS Trust, unpublished data 2017). There appeared to be overlap between populations reported in two studies (East Lancashire Hospitals NHS Trust, unpublished data 2017) and Fleming et al. <sup>15</sup> Three studies were from Scandinavia, <sup>7</sup> <sup>17</sup> <sup>18</sup> three from Canada <sup>19–21</sup> and one from France. <sup>5</sup>

Seven studies compared adverse birth outcomes before and after centralisation of services, which included closure of varying numbers of OUs. All three studies from the UK<sup>15</sup> <sup>16</sup> and (East Lancashire Hospitals NHS Trust, unpublished data 2017), examined the impact of the amalgamation of two OUs. Four studies were published after 2014<sup>15–17</sup> <sup>20</sup>; the earliest was from 1990.<sup>21</sup> Three studies included all births<sup>17</sup> <sup>18</sup> <sup>21</sup>; the other studies varied in their eligibility criteria, for example, restricting the analysis to singletons pregnancies, live births, various gestational ages and birth weight, hospital births or location. Reporting of eligibility criteria and participant characteristics across studies was inconsistent (table 1).

#### Risk of bias assessment

Risks of bias related to a lack of reporting of whether changes over time (other than closure/reconfiguration) could have influenced the findings, with only two 1720 of 10 studies reporting that temporal variation was adjusted for in the analysis (table 2). Further, 5 out of 10 studies either did not adjust results for potential confounding factors or provided insufficient information to know whether this was undertaken. Five out of 10 studies did not provide sufficient information to gauge the completeness of data. Half of the studies reported and used appropriate data analysis methods. Other potential sources of bias (eg, relating to selection, exposure and outcome) were less of a concern due to the use of routinely collected registry data before and after the closure and the objective nature of most outcomes.

#### **Findings**

A summary of maternal and neonatal outcomes is presented in table 3.

# Maternal outcomes

# Maternal mortality (MM)

In the two studies that reported MM, <sup>16</sup> <sup>20</sup> the number of deaths (<5) was too low to allow comparisons between the preclosure and postclosure groups.

# **Caesarean section (CS) (overall or intrapartum)**

Total CS rates were reported in four studies.<sup>7 15 20 21</sup> One UK study<sup>15</sup> reported a decline in CS rates following the amalgamation of two units from 26.1% to 21.5%.

A Norwegian study,<sup>7</sup> reported an increase in CS rates from 13.1% to 16.4% following OU closure, (cOR 1.31, 95% CI 1.27 to 1.35) as did two Canadian studies<sup>21</sup> (cOR

1.13, 95% CI 1.09 to 1.18) and (cOR 1.10, 95% CI 1.01 to 1.19). On adjusted results were reported.

# **Emergency CS**

Emergency CS rates were reported in one UK study, <sup>16</sup> which found no difference before/after the amalgamation of two OUs (cOR 0.95, 95% CI 0.86 to 1.05).

#### Severe perineal trauma (third or fourth degree tear)

Two studies<sup>16</sup> reported this outcome and found no statistically significant difference between the before/ after closure groups. The incidence of the outcome in both studies was low (<3%).

PPH—No studies reported this outcome.

#### Maternal admission to ICU

Two studies<sup>16</sup> <sup>20</sup> found no significant difference before/ after the amalgamation of two OUs in the number of women requiring admission to ICU (cOR 0.80, 95% CI 0.44 to 1.46).<sup>16</sup> The numbers in one study<sup>20</sup> were too small (<5) to allow a comparison.

# Maternal blood transfusion

One study<sup>20</sup> found no significant differences before/after OU closure (cOR 0.82, 95% CI 0.55 to 1.21). The incidence of the outcome was low (<1% of women).

#### **Neonatal outcomes**

#### Stillbirth (SB) (overall or intrapartum)

Three studies examined the impact of OU closure on SB. One unpublished UK study (East Lancashire Hospitals NHS Trust, unpublished data 2017) showed a statistically significant reduction in SB over the period after the amalgamation of two units (cOR 0.61, 95% CI 0.47 to 0.78). Similar findings were seen in one study from Canada<sup>22</sup> during post closure years (cOR 0.80, 95% CI 0.67 to 0.96). A third study from the UK<sup>16</sup> found no difference in SB rates after OU closure.

#### Neonatal mortality (NM)

Three studies reported this outcome. Two studies from the UK $^{16}$  and (East Lancashire Hospitals NHS Trust, unpublished data 2017) showed no statistically significant difference in the rate of NM in the years after OU closure (cOR 1.33, 95% CI 0.81 to 2.17; cOR 0.80, 95% CI 0.29 to 2.26). A study from Norway $^{17}$  also reported no difference (no OR presented).

# Perinatal mortality (PM)

Two studies reported this outcome. In a study from Norway,  $^7$  PM was significantly lower following OU closure (cOR 0.76, 95% CI 0.58 to 0.98). A Canadian study  $^{22}$  also reported a significant reduction in PM after OU closure (cOR 0.75, 95% CI 0.64 to 0.87).

# Infant mortality (IM)

One study<sup>17</sup> reported this outcome, IM rates were 'not statistically elevated' after the closure of thirteen hospitals in Norway.

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General Hospital		SB; NM; Maternal BBA mortality; Emergency CS; third and fourth degree perineal tea; Maternal transfer to ICU

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Table 1 Continued	nued								
Author, year, country Perinatal	Study design and setting	Study objectives	Study period	Eligibility criteria	Participant characteristics	Description of exposure (change over time)	Services context information	Review outcomes	
Blondel <i>et al</i> <sup>5</sup> , 2011, France*	Retrospective population-based analysis of routinely collected data, OU closure Across France	To report on BBA incidence in relation to distance from OU and the closure impact on different sociodemographic groups	Time of OU closure: 2003 and 2006 Time of analysis: 2005–2006	Included: Singleton births Excluded: Municipalities if >8% missing data, or high OOH rates. Departments excl. if >20% births already excl.	n=1 349 751 births; OCH n=5740 N Births 1349 to 751 Age (yrs) (n): <20–26 152 20–24 - 188 350 25–29 - 427 462 30–34 - 442 089 35–39 - 213 534 40+ - 52 164 Nullip (n) 774 460 SES: occupation professional (n) 217 045 intellectual 325 746 admin 266 000 retail 122 727 skilled 49 201 unnskilled 84 664 none 184 368 Ethnicity, education: NR	Closure of maternity unit Pre-change: no of OUS NR Postchange: Closure of units within 15 km radius of home, number of units closed NR	Centralisation of births in larger units due to safety concerns, financial pressure, efficiency savings, and staff shortage	BBA	
Hemminki <i>et al<sup>18</sup>,</i> 2011, Finland	Retrospective population-based analysis of routinely collected data, OU closure Across Finland and a specific district Uusimaa	To describe centralisation trend, unplanned out of hospital births, perinatal mortality (PM), health and birth outcomes in areas served by different levels hospitals different levels hospitals	Time of OU closure: 1991– 2008 Time of data analysis: Finland 1991–2008; Uusimaa district 2004–2008	Inclusion: All births Exclusion: NR	1991–2008 n=474 419 Characteristics: NR	Centralisation of births, maternity units no declined Pre-change: 49 OUs in 1991 Post-change: 34 OUs in 2008	Universal access to maternity care, minimal private care. Pre- and postnatal care decentralised, birth hospital-based service, care of high-risk pregnancies centralised. Mean no births/hospital increased from 1339 to 1733 over study period.	NR NR	

Table 1 Continued	inued								
Author, year, country	Study design and setting	Study objectives	Study period	Eligibility criteria	Participant characteristics	Description of exposure (change over time)	Services context information	Review outcomes	
Perinatal	Maternal								
Engjom <i>et al</i> <sup>7</sup> , 2014, Norway	Retrospective population-based study, 3 cohort and two cross-sectional studies, OU closure Across Norway	To assess the availability of OUs, the risk of unplanned delivery outside OU and maternal morbidity	Cohort: 1979–2009 Cross-sectional: 2000 and 2010 Time of OUs closure: 1979–2009 Time of analysis: Cohort: 1979–2009 Cross-sectional: 2000 and 2010	Included: Age 15-49 years, known place of birth, GA >22 wks and/or bthwt >5500 g Excluded: Missing maternal address, planned home birth	1979–1983 n=252 621 2004–2009 n=409 432 Characteristics: NR	Declined in no of OUs in Norway Prechange: 95 OUs in 1979 Post-change: 51 OUs in 2009	Universal access to maternity care; relatively dispersed population	M S	
Grytten et al <sup>17</sup> , 2014, Norway	Retrospective population-based analysis of routinely collected data, OU closure Across Norway	To study whether neonatal and infant mortality (IM) were I independent of the type of hospital in which the delivery was carried out	Time of closures: between 1979 and 2005 Time period for analysis: 5 years pre and postclosure for each hospital	Inclusion: All births Exclusion: NR	n=33 677 Characteristics: NR	Centralisation /OU closures No of local hospitals fell from 43 to 26 between 1979 and 2005. 17 maternity wards in local hospitals closed Prechange: 22 hospitals with neonatal department; 43 local hospitals; 30 maternity clinics Postchange: 22 hospitals with neonatal department; 26 local hospitals, 10 maternity clinics postchange:	Universal free access to maternity care. Low-risk births in local hospitals, high risk in central/ regional hospitals. Births in central/ regional hospitals increased from 65%–81% over study period.	NM; IM	
Canadian Studies									
Le Coutou <i>et al</i> e <sup>1</sup> , 1990, Canada	Retrospective population-based analysis of routinely collected data, OU closure Montreal metropolitan area	To describe the evolution of obstetric practice in Montreal metropolitan area before/ after closure of units	Time of closure: 1984–1985 Time of analysis: 1981–1985	Inclusion: All births Exclusion: NR	1981–1984 n=1 28 688 Characteristics: NR	7/13 university hospitals, 5/13 specialist hospitals, 4/13 smaller units closed Prechange: 39 units in 1981 Postchange:16	State provision of maternity care. Closures due to budgetary restrictions.	NA Q	Overall CS
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Table 1 Continued	penu								
Author, year, country	Study design and setting	Study objectives	Study period	Eligibility criteria	Participant characteristics	Description of exposure (change over time)	Services context information	Review outcomes	w w
Perinatal	Maternal								
Allen <i>et al,</i> <sup>22</sup> 2004, Canada	Retrospective population-based analysis of routinely collected data, OU closure Eastern, Northern, Western, and Central in Nova Scotia	To evaluate the effect of hospital closures on critical obstetrical interventions and perinatal outcomes in rural communities	Preclosure: 1988–1993 Post-closure: 1996–2002 Time of analysis: 1988–1993 vs 1996–2002	Inclusion: All births Exclusion: Delivery <20 weeks; bthwt <500 g; triplets+; major congenital anomaly	1988–93 n=69 213 1996–2002 n=63 510 Range % Age >34 yrs: 5.6–14.8; Nullip 39.4–46.8 Twins 1.0–1.3 Ethnicity, socioeconomic status, education: NR	1988–1993 =27 hospitals 1996–2002 =19 hospitals Reduction in maternity units from 42 to 11 between 1970 and 2002 Pre-change: 42 units in 1970 Post-change: 11 units in 2002	State provision of maternity care. Reduction in no of units and physicians due to financial constraints and difficulty maintaining clinical competence and confidence.	SB; Foetal/ neonatal mortality (NM)	E C
Hutcheon <i>et al</i> <sup>20</sup> , 2017, Canada	Retrospective population based analysis of routinely collected data, OU closure 25 communities within British Columbia, Canada	To examine the effect of obstetric service closures on intrapartum outcomes	1998–2014 Time of closures: between 2000 and 2012 Time of analysis: 1998–2014	Inclusion: All births recorded in British Columbia Perinatal Data Registry (99% of deliveries) Exclusion: Communities close to larger metropolitan areas and or uncertainty about dates of service closures.	Pre-closure n=5796 Median maternal age 27 years (IQR 23-31); Nullip 39.3% Post-closure n=6153 Median maternal age 28 years (IQR 24-32); Nullip 40.7% Ethnicity, Socioeconomic status, education: NR	Centralisation /OU closures Between 1998 and 2014 one- third of hospitals stopped providing maternity services Pre-change: 21 hospitals with obstetric services Postchange: Obstetric services closed in same 21 hospitals	State provision of maternity care. Centralisation of obstetric services, majority of hospital closures in low-volume hospitals	BBA; perinatal/ NM; NNU admission	Overall CS; Maternal mortality; third/4 <sup>th</sup> degree perineal tear, blood transfusion, maternal admission to ICU

\*Blondel et al<sup>6</sup> is also included in travel distance.
Approx, approximately; BBA, Born before arrival; bthwt, birth weight, CS, caesarean section; excl, excluded; GA, gestational age; ICU, intensive care unit; MW, midwife; NHS, National Health Services; NNU, neonatal unit; NR, not reported; Nullip, nulliparous; OU, obstetric unit; SB, stillbirth; SES, socioeconomic status; wo, without, Yr, year.

Table 2 Risk of bia	Risk of bias—obstetric unit (OU) closure studies	closure studies					
Author, year, country	Study sample selection bias	Bias in measurement of exposure	Bias in measurement of outcomes	Attrition bias	Analysis method reported and appropriate	Closure independent of other changes over time	Potential confounders adjusted for and listed
UK Studies							
Fleming/East Lancashire study	re study						
Fleming <sup>15</sup> , 2013, UK (abstract)	LOW All births in East Lancashire Maternity Services catchment	LOW All births in catchment area affected by the closure No of OUs closed reported	LOW Objective outcome (CS)	UNCLEAR Not reported	UNCLEAR None reported	UNCLEAR None reported	UNCLEAR None reported
East Lancashire Hospitals NHS Trust 2017, UK (unpublished data)	UNCLEAR Unpublished data, no details	UNCLEAR Unpublished data, no details	LOW Objective outcomes (BBA, SB, NM)	UNCLEAR Unpublished data, no details	UNCLEAR Unpublished data, no details	UNCLEAR Unpublished data, no details	UNCLEAR Unpublished data, no details
Mackie <i>et al</i> <sup>16</sup> , 2014, UK (abstract & unpublished)	LOW Data from Maternity Information System	LOW All births in catchment area affected by the closure. No of OUs closed reported	LOW Objective outcomes (SB, NM, BBA, MM, ICU admission, perineal tears)	UNCLEAR Not reported	UNCLEAR None reported	UNCLEAR None reported	UNCLEAR None reported
Other European Studies	ű						
Blondel <i>et al</i> <sup>5</sup> , 2000, France <sup>*</sup>	LOW Data from birth certificates	LOW No of OUs closed reported	LOW Objective outcome (BBA)	LOW 11% excluded due to missing data	LOW Analysis method was described and appropriate, a multi- level model analysis	UNCLEAR None reported	LOW Adjusted for maternal age, occupational category and rurality
Hemminki e <i>t al</i> <sup>18</sup> , 2011,m Finland	LOW Data from Finnish medical birth register	LOW All births in catchment area affected by the closure No of OUs closed reported	LOW Objective outcome (BBA)	LOW Births with missing information excluded (<0.05%)	LOW Analysis method was described and appropriate, a regression model with adjusted analysis	UNCLEAR None reported	LOW Adjusted for Parity, plurality, age, socioeconomic status and smoking
Engjom <i>et al</i> <sup>7</sup> , 2014, Norway	LOW Data from Medical Birth Registry of Norway	LOW All births in Norway affected by the closure No of OUs closed reported	LOW Objective outcome (BBA)	LOW All units report to Medical Birth Registry	LOW Analysis method appropriate, a logistic regression model, crude and adjusted results given	UNCLEAR None reported	LOW Adjusted for maternal age, parity, education and partner status
							Continued

Table 2 Continued							
Author, year, country	Study sample selection bias	Bias in measurement of exposure	Bias in measurement of outcomes	Attrition bias	Analysis method reported and appropriate	Closure independent of other changes over time	Potential confounders adjusted for and listed
Grytten <i>et al</i> <sup>17</sup> , 2014, Norway	LOW Data from Medical Birth Registry of Norway	LOW All births in Norway No of OUs closed reported	LOW Objective outcomes (NM, Infant mortality)	LOW All maternity units report to Medical Birth Registry	Unclear Difference-in- difference statistical method used, but reporting of findings were unclear	LOW Adjusted for trend in infant outcomes based on local hospitals that were not closed.	LOW Maternal age, immigrant status, level of education, marital status, predisposing medical factors and characteristics of the birth
Canadian Studies							
Le Coutour <i>et af</i> <sup>1</sup> , 1990, Canada	LOW Data from MED- ECHO - regional data collection system	LOW All births in catchment area No of OUs closed reported	LOW Objective outcome (CS)	UNCLEAR No information	UNCLEAR No details on data analysed method	UNCLEAR None reported	HIGH No adjustment
Allen et al, <sup>22</sup> 2004, Canada	LOW Data from Nova Scotia Atlee Perinatal Database	LOW All births in catchment area affected by the closure No of OUs closed reported	LOW Objective outcomes (SB, NM)	UNCLEAR Population based dataset but no information about missing data	LOW Analysis method appropriate and data from logistic regression models were reported	UNCLEAR None reported	LOW Maternal age, smoking and maternal diseases
Hutcheon <i>et al</i> <sup>po</sup> , 2017, LOW Canada Data Colur regist	LOW Data from British Columbia Perinatal registry	LOW All births in catchment area affect by the closure No of OUs closed reported	LOW Except for third/4 <sup>th</sup> degree tears. Objective outcomes (BBA, PM, NM, ICU admission, CS, MM, blood transfusion, Maternal admission to ICU)	>99% complete	LOW Used a within- community fixed- effects design and Poisson regression	LOW Using difference in difference analysis which separates the effect of the closure from underlying time trends of reported outcomes	HIGH No adjustment

\*Blondel et al 2011 included in travel distance and OU closure.

BBA, born before arrival; CS, caesarean section; ICU, intensive care unit; MM, maternal mortality; NHS, National Health Service; NM, neonatal mortality; PM, perinatal mortality; SB, stillbirth.

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lable 3 Outcomes—c	Outcomes—obstetric unit (OU) closure	osnre						
Outcomes	Author, Year, Country	Exposure and comparator groups	Participants (N, n, %)			Findings		
MATERNAL OUTCOMES								
Maternal mortality (MM)	Mackie <i>et al</i> <sup>16</sup> , 2014, UK	Before and after	Year	Deliveries (n=15349)	MM n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
	(unpublished data)	amalgamation of 2 OUs	Pre 2010–2011	5354	1 (0.02)	Pre 2010–2011	-	N.
			Post 2011–2013	9995	1 (0.01)	Post 2011-2013	0.54 (0.03 to 8.56)	NR
	Hutcheon <i>et al</i> <sup>20</sup> , 2017, British Columbia, Canada	Before and after OU closure (1998–2014)	Closure status	Deliveries (n=11949)	Maternal deaths n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
			Preclosure	5796	<5 (<0.09)	Preclosure	NR	NR
			Postclosure	6153	<5 (<0.08)	Postclosure	NR	NR
			No significant difference	No significant difference pre/post closure in adverse events during labour and delivery	e events during labou	ır and delivery		
Caesarean section (CS) (overall or		Before and after closure of	Closure status	Deliveries (n=NR)	CS n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
intrapartum)	(abstract)	OU in 2010	Preclosure, early 2010	NR	(NR) 26.1	Preclosure, early 2010	ш 2	W.
			Postclosure, 2012	NR	(NR) 21.5	Postclosure, 2012	NB	NR
			Proportions of CS presented with no other data.	nted with no other data.				
	Engjom et al <sup>7</sup> , 2014, Norway 2000 compared with 2009	2000 compared with 2009	Year	Deliveries (n=2,177,934)	CS n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
		during which time number of OUs declined from 47 to 41	Pre 2000	58632	7653 (13.10)	Pre 2000	-	NR
			Post 2009	61895	10154 (16.41)	Post 2009	1.31 (1.27 to 1.35)	N.
	Le Coutour <i>et al</i> <sup>21</sup> , 1990,	Before and after closure of	Year	Deliveries (n=64274)	CS n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
	Quebec, Canada	OUs between 1982 and 1983	Pre 1981	32807	5852 (17.84)	Pre 1981	<del>-</del>	NR
			Post 1984	31467	6214 (19.7)	Post 1984	1.13 (1.09 to 1.18)	NR
	Hutcheon <i>et al</i> <sup>20</sup> , 2017,	Before and after OUs, closure	Closure status	Deliveries (n=11949)	CS n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
	British Columbia, Canada	(1998–2014)	Preclosure	5796	1387 (23.93)	Preclosure	-	NR
			Postclosure	6153	1579 (25.70)	Postclosure	1.10 (1.01 to 1.19)	NR
Emergency caesarean section (CS)	Mackie <i>et al</i> <sup>16</sup> 2014, UK (abstract only)	Before and after amalgamation of two OUs	Year	Deliveries (n=15349)	Emergency CS n(%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2010–2011	5354	739 (13.80)	Pre 2010–2011	-	NR
			Post 2011-2013	9666	1322 (13.23)	Post 2011-2013	0.95 (0.86 to 1.05)	N.
Severe perineal trauma (third or	Mackie <i>et al</i> <sup>16</sup> , 2014, UK	Before and after	Year	Deliveries (n=15349)	3rd & fourth n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
rourn degree tear)	(unpublished data)	amaigamation of two UUs	Pre 2010–2011	5354	133 (2.48)	Pre 2010–2011	-	NR
			Post 2011-2013	9666	276 (2.76)	Post 2011-2013	1.11 (0.90 to 1.37)	NR
	Hutcheon <i>et al</i> <sup>20</sup> , 2017, British Columbia, Canada	Before and after OU closure (1998–2014)	Closure status	Deliveries (n=11949)	third or fourth degree tear n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
			Preclosure	9629	136 (2.40)	Preclosure	-	NR.
			Postclosure	6153	174 (2.82)	Postclosure	1.21 (0.96 to 1.52)	N.
Postpartum haemorrhage	No studies							



Table 3 Continued								
Outcomes	Author, Year, Country	Exposure and comparator groups	Participants (N, n, %)			Findings		
Maternal admission to ICU	Mackie <i>et al</i> <sup>16</sup> , 2014, UK	Before and after	Year	Deliveries (n=15349)	n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
	(abstract only)	amaigamation or two OUS	Pre 2010–2011	5354	18 (0.34)	Pre 2010–2011	-	N.
			Post 2011–2013	9885	27 (0.27)	Post 2011–2013	0.80 (0.44 to 1.46)	N.
	Hutcheon <i>et al</i> <sup>20</sup> , 2017,	Before and after OU closure	Closure status	Deliveries (n=11949)	ICU admission n	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
	British Columbia, Canada	(1998–2014)	Preclosure	9629	<5	Preclosure	N.	W.
			Postclosure	6153	<5	Postclosure	NR	N.
Maternal Blood transfusion	Hutcheon <i>et al</i> <sup>20</sup> , 2017, British Columbia, Canada	Before and after OU closure (1998–2014)	Closure status	Deliveries (n=11949)	Blood transfusion n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
			Preclosure	5796	53 (0.91)	Preclosure	-	NR
			Postclosure	6153	46 (0.75)	Postclosure	0.82 (0.55 to 1.21)	NR
NEONATAL OUTCOMES								
Stillbirth (SB)	East Lancashire Hospitals NHS Trust, 2017, UK (unpublished data)	Before and after amalgamation of two obstetric units (OUs) in 2010	Year	Deliveries (n=53870)	SB n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2009	6492	75 (1.16)	Pre 2009	-	NR
			Post 2011–2017	47.378	333 (0.70)	Post 2011–2017	0.61 (0.47 to 0.78)	NR
			Stillbirth>24 weeks					
	Allen e <i>t al,</i> <sup>22</sup> 2004, Nova Scotia, Canada	Before and after closure of OUs	Year	Deliveries (n=1 32 723)	SB n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 1988–93	69213	291 (0.42)	Pre 1988–1993	-	NR
			Post 1996-2002	63510	214 (0.34)	Post 1996-2002	0.80 (0.67 to 0.96)	NR
	Mackie <i>et al</i> <sup>16</sup> 2014, UK (unpublished data)	Before and after amalgamation of 2 OUs	Year	Deliveries (n=15552)	SB n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2010–2011	5422	29 (0.53)	Pre 2010 –2011	-	NR
			Post 2011-2013	10130	60 (0.59)	Post 2011-2013	1.11 (0.71 to 1.73)	NR
Neonatal mortality (NM)	East Lancashire Hospitals NHS Trust, 2017, UK (unpublished data)	Before and after amalgamation of two obstetric units in 2010	Year	Deliveries (n=53870)	NM n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2009	6492	4 (0.06)	Pre 2009	-	NR
			Post 2011–2017	47.378	39 (0.08)	Post 2011-2017	1.33 (0.81 to 2.17)	NR
			NM not defined					
	Mackie et al <sup>16</sup> , 2014, UK (unpublished data)	Before and after amalgamation of 2 OUs	Year	Deliveries (n=15552)	NM n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2010–2011	5422	6 (0.11)	Pre 2010–2011	1	NR
			Post 2011–2013	10130	(06:0) 6	Post 2011-2013	0.80 (0.29 to 2.26)	N.
	Grytten <i>et al</i> <sup>17</sup> , 2014, Norway	Before and after 13 hospital closures	Year	Deliveries (n=33677)	NM n (%)	Year	Crude OR (95% CI)	Adjusted OR(95% CI)
			5 years before	16297	N.	5 years before	NR	W.
			5 years after	17380	N.	5 years after	NB	W.
			No statistically significant difference					
								1000

Table 3 Continued								
Outcomes	Author, Year, Country	Exposure and comparator groups	Participants (N, n, %)			Findings		
Perinatal mortality (PM)	Engjom et al', 2014, Norway 2000 compared with 2009 during which time number OUs declined from 47 to 4	2000 compared with 2009 during which time number of OUs declined from 47 to 41	Year	Deliveries (n=2,177,934) PM n (/1000 births)	PM n (/1000 births)	Year	Crude OR (95% CI)	Adjusted OR(95% CI)
			Pre 2000	58632	124 (2.11)	Pre 2000	-	W.
			Post 2009	61 895	99 (1.60)	Post 2009	0.76 (0.58 to 0.98)	W.
			PM (Intrapartum & neonatal death<24 hours, both live & stillborn)					
	Allen <i>et af</i> <sup>22</sup> , 2004, Nova Scotia, Canada	Before and after closure of OUs	Years	Deliveries (n=132723)	PM n (%)	Years	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 1988–1993	69213	422 (0.61)	Pre 1988–93	-	W.
			Post 1996-2002	63510	278 (0.43)	Post 1966-2002	0.75 (0.64 to 0.87)	NR
			Foetal/neonatal mortality not defined					
Infant mortality	Grytten et $a^{17}$ , 2014, Norway	Before and after 13 hospital closures	Year	Deliveries (n=33677)	IM n (%)	Years	Crude OR (95% CI)	Adjusted OR (95% CI)
			5 years before	16297	N. W.	5 years before	NR	NR
			5 years after	17380	NR	5 years after	NB	NB
			No significant difference in infant mortality					

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		Exposure and comparator						
Outcomes	Author, Year, Country	groups	Participants (N, n, %)			Findings		
Born before arrival (BBA)	East Lancashire Hospitals NHS Trust, 2017, UK (unpublished data)	Before and after amalgamation of two obstetric units	Year	Deliveries (n=53870)	BBA n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2009	6492	25 (0.39)	Pre 2009	-	NB
			Post 2011–2017	47.378	341 (0.72)	Post 2011–2017	1.88 (1.25 to 2.82)	NB
	Mackie <i>et al</i> <sup>16</sup> , 2014, UK (unpublished data)	Before and after amalgamation of 2 OUs	Year	Deliveries (n=15349)	BBA n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 2010–2011	5354	11 (0.21)	Pre 2010–2011	-	NB
			Post 2011–2013	9995	26 (0.26)	Post 2011–2013	1.28 (0.63 to 2.60)	NB
	Blondel <i>et al</i> <sup>5</sup> , 2011, France	OU closure within 15km radius 2003–2006	Yrs 2003–2006	Deliveries (n=1,349,751)	BBA n (/1000 births)	Yrs 2003–2009	Crude OR (95% CI)	Adjusted OR (95% CI)
			No closure	1 001 858	4531 (4.52)	No closure	-	-
			Closure within 15km radius	347 893	1209 (3.47)	Closure within 15km radius	0.77 (0.72 to 0.82)	0.91 (0.84 to 1.00)
	Engjom et al <sup>7</sup> , 2014, Norway 1979–83 compared with 2004–09, number of emergency OUs declined from 47 to 41	v 1979–83 compared with 2004–09, number of emergency OUs declined from 47 to 41	Year	Deliveries (n=662053)	BBA n (%)	Year	Crude OR (95% CI)	Adjusted OR (95% CI)
			Pre 1979–83	252621	984 (0.39)	Pre 1979–83	-	-
			Post 2004-09	409432	2832 (0.69)	Post 2004-09	1.8 (1.6 to 1.9)	2.0 (1.9 to 2.2)
	Hemminki e <i>t al</i> <sup>18</sup> , 2011, Finland	Centralisation of hospitals over years 1991–2008	Year	Births (N)	Unplanned BBA n(/1000)	Planned or unplanned BBA n(/1000)	Year	Crude OR (95% CI)
			Pre 1991	65 632	I	68 (1.0)	Pre 1991	T
			Post 2004-2008	56873	222 (3.76)	243 (4.1)	Post 2004–2008	4.14 (3.16 to 5.41)
			Total N (1991– 2008)=122505					
	Hutcheon et al <sup>20</sup> , 2017, British Columbia, Canada	Before and after OU closure (1998–2014)	Closure status	Deliveries (n=11949)	Unplanned BBA n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
			Preclosure	5796	30 (0.5)	Preclosure	1	N.
			Postclosure	6153	109 (1.8)	Postclosure	3.47 (2.31 to 5.20)	N.
Neonatal unit admission (NNU)>2 days or transfer within	Hutcheon <i>et al</i> <sup>20</sup> , 2017, British Columbia, Canada	Before and after OU closure (1998–2014)	Closure status	Deliveries (n=11949)	NNU admission n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
24 nours of birth to ICU facility for newborn >/=2500 g			Preclosure	5796	68 (1.17)	Preclosure	-	NR
			Postclosure	6153	28 (0.46)	Postclosure	0.39 (0.25 to 0.60)	NR
Apgar score (5 min Apgar score<7)	Hutcheon <i>et al<sup>20</sup>,</i> 2017, British Columbia, Canada	Before and after OU closure (1998–2014) compared with communities unaffected by closure	Closure status	Deliveries (n=11949)	5min Apgar score<7 n (%)	Closure status	Crude OR (95% CI)	Adjusted OR (95% CI)
			Preclosure	5796	71 (1.22)	Preclosure	-	NA NA
			Postclosure	6153	85 (1.28)	Postclosure	1.13 (0.82 to 1.55)	N.
Hypoxic Ischaemic Encephalopathy (HIE)	No studies							

ICU, intensive care unit; NR, not reported.

Continued

Table 3



#### Born before arrival (BBA)

Six studies reported this outcome, with four suggesting a statistically significant increase in BBA following OU closure. Data from East Lancashire Hospitals NHS Trust (East Lancashire Hospitals NHS Trust, unpublished data 2017) showed the BBA rate almost doubled over the 10-year period (cOR 1.88, 95% CI 1.25 to 2.82). Studies from Norway<sup>7</sup> and Finland<sup>18</sup> also found that the BBA rate increased over a similar period (cOR 1.8, 95% CI 1.6 to 1.9 and cOR 4.14, 95% CI 3.16 to 5.41, respectively). A Canadian study<sup>20</sup> found that the BBA rate trebled over a 16-year period (cOR 3.47, 95% CI 2.31 to 5.20). One UK study<sup>16</sup> found no statistically significant change (cOR 1.28, 95% CI 0.63 to 2.60) and in one French study,<sup>5</sup> there was weak evidence of a small reduction in the adjusted risk of BBA in communities affected by OU closure (adjOR 0.91, 95% CI 0.84 to 1.00).

#### Neonatal unit (NNU) admission

One Canadian study<sup>20</sup> suggested a significant reduction in NNU admission following OU closure (cOR 0.39, 95% CI 0.25 to 0.60).

#### Apgar score

One Canadian study<sup>20</sup> found no statistically difference in 5 min Apgar score of less than 7 before and after OU closure (cOR 1.13, 95% CI 0.82 to 1.55).

# Hypoxic-ischaemic encephalopathy (HIE)

No studies reported this outcome.

#### Evidence from travel distance studies

#### **Description of included studies**

Seven studies described the effect of travel distance to the nearest OU on maternal and neonatal outcomes (table 4). All were published in full between 1991 and 2015. The earliest study<sup>23</sup> was conducted in the UK, three more recent studies were conducted in France,  $^{5\ 24\ 25}$  and one each in Norway,  $^{26}$  Finland  $^{27}$  and Canada.  $^{28}$  Four were retrospective population-based cohort studies, and three were case–control studies. The eligibility criteria varied across studies. Pasquier *et al*  $^{24}$  included a group with special needs in the form of babies with congenital malformations. Only singleton live births were included in two studies.  $^{5\ 28}$ 

Travel distance was estimated using geographical mapping software in all studies. However, only three studies<sup>5</sup> <sup>24</sup> <sup>27</sup> measured the actual distance from women's homes to the nearest OU. In two studies<sup>25</sup> <sup>28</sup> a central geographical point for the postal code or municipality was used to estimate distances and in one study the distance was self-reported.<sup>26</sup> Additionally, the studies differed regarding their distance categories, which ranged from 2 to 150 km (table 4).

# **Risk of bias assessment**

The main risk of bias concerns related to the measurement of exposure, as three studies <sup>25</sup> <sup>26</sup> <sup>28</sup> did not calculate the distance from the woman's home but used a central

point instead or self-reported distance (table 5). Another study<sup>23</sup> measured distance between women's homes and hospital using a straight line. Further risk of bias related to a lack of comparability between study groups in the three case–control studies,<sup>23</sup> <sup>26</sup> <sup>27</sup> a lack of adjustment for confounders in two studies and missing data in two studies.<sup>26</sup> There were no risk of bias concerns relating to sample selection in the cohort studies or outcome measurement.

#### **Findings**

# Maternal outcomes

#### Maternal mortality (MM)

In one case–control study from Finland,<sup>27</sup> no maternal deaths were reported in either group (table 6).

#### Caesarean section (CS) (overall or intrapartum)

One study from Canada $^{28}$  found no statistically significant differences in CS rates with increasing distance (<50 k, 50–150 k, >150 k) based on both crude and adjusted results.

#### Emergency CS

The same study from Canada<sup>28</sup> reported no significant difference in emergency CS rates between women living at different distances from an OU based on cORs.

#### Severe perineal trauma

No studies reported this outcome.

#### Postpartum haemorrhage (PPH)

No studies reported this outcome.

# Maternal admission to ICU

No studies reported this outcome.

#### Maternal blood transfusion

No studies reported this outcome.

# **Neonatal outcomes**

# Stillbirth (SB) (overall or intrapartum)

Two cohort studies<sup>25 28</sup> reported this outcome. A Canadian study<sup>28</sup> included births to women aged over 35 years who lived <50 km, 50-150 km and >150 km from the OU. SB rates did not change by distance category in the adjusted analysis. A French study<sup>25</sup> reported SB rates at different distances (<5 km, 5-15, 15-30, 30-44, 45+ km) from mother's municipality of residence to the closest OU. The cORs showed women living at 30–44, 15–29 and 5–14 km from an OU had a statistically significantly lower rate of SB compared with women living <5 km from an OU (5-14 km vs <5 km, cOR 0.87, 95% CI 0.85 to 0.90; 15-29 km vs < 5 km; cOR 0.85, 95% CI 0.82 to 0.88; 30-44 km vs <5 km, cOR 0.85, 95% CI 0.81 to 0.90). The findings still hold for the adjusted analysis (limited data reported). However, neither the crude nor the adjusted analysis showed a significant difference in risk of SB for individuals resident 45+ km from an OU compared with <5 km.

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Table 4 Descri	Description of included etualise — travel distance	oib loyert—soibuta	tanco						
+	iption of monded	studies—traver dis	oldilloe						
Author, year, country	Study design and / setting	Study objectives	Study period	Eligibility criteria	Participant characteristics	Description of exposures travel distance	Services context information	Review outcomes Perinatal	Maternal
UK Studies									
Bhoopalam <i>et</i> <i>al</i> <sup>23</sup> ,1991, UK	Case-control study, 2 OUs	To establish BBA prevalence and women at risk of BBA, and morbidity and mortality associated with BBA births	1983-1987	Included cases: Women and their BBA babies Included controls: Two controls for each BBA case, one random (next born in the same hospital), one matched (next born in same hospital matched by GA and BW) Excluded: BW <500g	N(BBAs)=137, 1 twins All n=398 Age yrs (n): <21 (69) 21-35 (339) >35 (27) Nullips (107) Ethnicity (n): European(191) Asian (101) Other (16) SES and education: NR	Distance (km): <2 2-7 >7	Universal state provision, 2 units six miles apart, serving rural areas of Warwickshire	BBA	Ψ <sub>Z</sub>
Other European Studies	ndies								
Pasquier <i>et al<sup>24</sup>,</i> 2007, France	Retrospective, population-based cohort, 3 Level-III maternity wards with meonatal surgical centre, Rhône-Alpes Region	To examine maternal origin, distance to the nearest maternity ward with a neonatal surgical centre, on perinatal diagnosis, elective termination of pregnancy, delivery in an adequate place and neonatal mortality (NN) for pregnancies with severe malformations	1990–1995 and 1996–2000 (two periods separated due to changes in prenatal screening)	Included: Fetuses with omphalocele, gastroschisis, diaphragmatic hernia or spina bifida that required surgical repair Excluded: Chromosomal anomalies fetuses and babies without anomalies	n=706 infants n=554 (analysed) Age: yrs (n, %) <21 (15, 2) 21–35 (550, 82)-35 (106, 16) European (393, 76) Non-Western European (124, 24) Parity, SES and education: NR	Distance (кт): <11 11–50>50	Distance to Level III maternity ward with NNU and a neonatal surgical centre, there were three in the Rhone-Alps Region	N	Ψ.
Blondel <i>et al</i> <sup>6</sup> , 2011, France*	Retrospective cohort, population based study, metropolitan France	To calculate the incidence of BBA birth in relation to distance from maternity units and the impact of recent closure on different sociodemographic groups	2005–2006	Included: Singleton live births Excluded: Municipalities with >8% missing data, unrealistically high BBA births, Departments were excl. if >20% births already excl.	n=1517599 ivebirths n=1349751(analysed) Age yrs -n<20-26 152 20-34 - 105 790 135-39 - 213 534 40+ - 52 164 Party. Nullip n -774 460 SEs n: Occupation professional- 217045 Intellectual- 325746 Admin- 266000 Retail-122,727 Skilled- 149201 Unskilled- 84 664 None- 84 368 Ethnicity nd education: NR	Distance (km): <5 5-14 15-29 30-44 45+	Centralising births in BBA; BBA by larger units parity	BBA; BBA by parity	E E
									1

Table 4   Continued	penu								
	Study design and				Participant	Description of exposures	Services context	Review outcomes	
Author, year, country	- 1	Study objectives	Study period	Eligibility criteria	characteristics	travel distance	information	Perinatal	Maternal
Pilkington et al <sup>25</sup> , 2014, France	Retrospective, population-based cohort study, French National Vital Statistics registry from mainland France	To investigate the impact of distance to closest maternity unit on perinatal outcomes	2001–2008, Stillbirth (SB) data 2002–2005	Included: All births Excluded: NR	n=3 086 128 all births n=3 085 839 (analysed) Age yrs (n) <25-4 94 689 25-34 - 2 008 320 35-39 -469,975 40+ -113 144 Singleton pregnancy n=2 988 169 Multiple pregnancy: n=97 959 Parity, ethnicity and socioeconomic status and education: NR	Distance (km): <5 5–14 15–29 30–44 ≥45	1998 to 2003, 20% of maternity units closed Mean distance to nearest maternity unit increased (6.6– 7.2km)	SB; NM	Ľ
Ovaskainen <i>et al<sup>27</sup>,</i> 2015, Finland	Case-control study, one centre, Tampere University Hospital	To establish if BBA births increased over time, to identify risk factors associated with BBAs, also if BBAs babies were more prone to neonatal morbidities compared with those delivered in hospital	1996-2011	Included cases: Planned and unplanned BBA Included Controls: 2 controls for infant and mother for each BBA case Excluded: BBA with no information whether planned or unplanned	Cases: BBAs (n=67 births): Age yrs (mean, SD) (range)- 29.0, 5.9 (15-47) Parity 1 (0-16) Controls: n=134 Plurality, ethnicity, education and SES: NR.	Distance (km): <35 ≥35	Tampere University Hospital is the catchment area for 23 municipalities, 521 700 residents 5000 births/yr	ВВА	ű Z
Fougner <i>et af</i> <sup>e</sup> , 2000, Norway	Case-control study, 14 municipalities, Oppland County	To compare the experience and care of women who delivered during transport to hospital and women who adlivered an hour after arriving to hospital	1989–1997	Included cases: Women who delivered their babies before arriving at hospital Included Controls: Women who delivered their babies with 1 hour after arriving at hospital Excluded: NR	n=202 Cases: n=115BBA Distance (km): <12.88 women Parity n ≥12.88 (%)- Nullips 15 (13%) Controls: n=87 women Parity- Nullips 18 (20%) age, ethnicity, education and SES: NR	Distance (km): <12.88 ≥12.88	Oppland county: 4 hospitals	ВВА	щ Z
Canadian Studies									
Lisonkova <i>et al</i> <sup>28</sup> , 2011, Canada	Retrospective population-based cohort study, British Columbia	To examine the association between rural residence and birth outcomes in older mothers	1999-2003	Included: Singleton mothers aged 35+ Excluded: Women with missing postcodes, babies with congenital anomaly	n=29698 women age >35 years parity n (%): Nullip 87733 (0%) Low SES (n, %) (4385 14% 22.6 vs 3615, 13.7) Ethnicity n (%) first nation 826 (2.8%) Education:	Distance (km): <50 50-150 >150	17 small maternity units (250–2500 births/yr) closed between 1999 and 2003	SB; perinatal mortality (PM); NNU admission ≥1 day	All CS; Emergency CS

\*Blondel et ale is also included in the OU closure. adjusted OR; BBA, born before arrival; BW, birth weight; CS, caesarean section; ICU, intensive care unit; NHS, National Health Services; NNU, neonatal unit; NR, not reported; OU, obstetric unit; SES, socioeconomic status.

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et a							
I. BMJ	Table 5 Risk of bias-	Risk of bias—travel distance					
or, year, country	Study sample selection bias additional criteria for case- control studies	n bias <b>additional</b> Itrol studies	Bias in measurement of exposure	Bias in measurement of outcomes	Attrition bias	Analysis method reported and appropriate	Potential confounders adj for and listed
tudies							
palam <i>et al<sup>e3</sup>,</i> UK	Case definition adequate records Representative-ness of cases Appropriate selection of controls Definition of control appropriate Comparability of cases RES BBA case hospital same hospital Occurred Comparability of cases NO Significant and controls age, parity, eth	YES From hospital records YES BBA cases from 2 hospitals over 5 years YES two controls from same hospital YES The outcome (BBA) could not have occurred s NO Significant differences in maternal age, parity, ethnicity and antenatal booking	HIGH Distance measured in straight line from home address	LOW Objective outcome (BBA)	<b>Low</b> 3/134 (2.2%) BBA cases and 10/274 (3.6%) excluded from distance analysis	HIGH Descriptive analysis only	<b>HIGH</b> No adjustments
r European Stud							
uier <i>et al<sup>24</sup>, 2007,</i> se	, <b>LOW</b> Data from France Central-East malformation registry	e Central-East	LOW GIS software used to estimate distance between maternal residence and nearest maternity ward with neonatal surgical centre	LOW Objective outcomes (NM)	<b>Low</b> 12 births (0.03%) missing survival data	LOW Method detailed and appropriate, univariate analysis & multiple logistic regression were reported	<b>LOW</b> Adjusted for parity, and other characteristics
iel <i>et al</i> <sup>5</sup> , 2011, :e*	LOW Birth certificates		LOW GIS software used to estimate distance to hospital from home	LOW Objective outcome (BBA)	LOW 11% of births excluded	LOW Method detailed and appropriate, multilevel model was reported	LOW Adjusted for matern occupation, parity and oth characteristics
gton <i>et al</i> <sup>25</sup> , 201. se	gton <i>et al</i> <sup>25</sup> , 2014, <b>LOW</b> French National Vital Statistics Registry se	Vital Statistics Registry	HIGH Distance calculated from centre of municipality not home address, using road networks provided by the French National Geography Institute	<b>Low</b> Objective outcomes (SB, NM)	<b>Low</b> 10% missing for type of pregnancy and 17% for maternal age	LOW Method detailed and appropriate, logistic regression analysis was reported	LOW Adjusted for matern plurality, unemployment re single parent households
							en a

Table 5 Continued

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or, year, country	Study sample selection bias additional criteria for case-control studies	n bias <b>additional</b> trol studies	Bias in measurement of exposure	Bias in measurement of outcomes	Attrition bias	Analysis method reported and appropriate	Potential confounders adj for and listed
ner <i>et al<sup>26</sup>,</i> 2000, nd		NO Data from a questionnaire	HIGH Data for distance was self-reported in	Low Objective outcome (BBA)	HIGH Data from women who responded to questionnaire	HIGH Descriptive analysis only	HIGH No adjustment
	Representative-ness of cases	YES Cases from three hospitals in one county over 8 years	questionnaire				
	Appropriate selection of controls	Yes Women who delivered their babies within 1 hour of arriving at hospital					
	Definition of control appropriate	<b>Yes</b> Women with no BBA					
	Comparability of cases <b>Unclear</b> and controls	3 Unclear					
kainen e <i>t al<sup>27</sup>,</i> Finland	Case definition adequate	YES Medical records	<b>LOW</b> Distance from women's home	LOW Objective outcome (BBA)	High 13 out-of-hospital deliveries LOW Method detailed and (19%) excluded as could not appropriate, logistic regres	<b>LOW</b> Method detailed and appropriate, logistic regression	LOW Adjusted for single-navity, and other characte
	Representative-ness of cases	<b>YES</b> Cases from one centre, but over 15 years	calculated using web- based route planner		ascertain whether planned or not planned	data were given	
	Appropriate selection of controls	YES Births occurring immediately preceding and following case					
	Definition of control appropriate	<b>YES</b> The outcome (BBA) could not have occurred					
	Comparability of cases No Sig. differences and controls for parity, partnersh status, smoking, antenatal visits, lab duration and distan to delivery unit	s No Sig. differences for parity, partnership status, smoking, antenatal visits, labour duration and distance to delivery unit					
dian studies							
kova <i>et al<sup>28</sup>,</i> Canada	LOW Population-based study	d study	HIGH GIS used to calculate distance from postcode central point of residence to hospital; mostly using straight line distance	<b>LOW</b> Objective outcomes (SB; PM); NNU admission)	LOW 492 (1.7%) women excluded due to missing postcodes	Low Method detailed and appropriate, multivariate regression analysis reported	Low Adjusted for parity, a mother, low income, ethni other characteristics

del *et af* 2011 included in travel distance & OU closure. born before arrival; GIS, Geographical Information System; NM, neonatal mortality; NNU, neonatal unit; PM, perinatal mortality; SB, still birth; sigs, significant.



	Author, year, country	Exposure groups	Participants (N, n, %)			Findings		
Maternal mortality	Ovaskainen <i>et al</i> <sup>27</sup> , 2015,	Travel distance (km):	Groups	N (201)	NM n (%)	No events in either group		
	Finland	<35 ≥35	<35	N.	0			
			≥35	N.	0			
Caesarean section (CS)	Lisonkova <i>et al</i> <sup>28</sup> , 2011,	Travel distance (km): <50	Groups	N (29 698)	CS n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
(overall or intrapartum)	Canada	50-150 >150	<50	27 836	9099 (32.70)	<50	-	RN
			50–150	1534	464 (30.25)	50–150	0.89 (0.80 to 1.00)	RN
			>50	328	94 (28.70)	>50	0.83 (0.65 to 1.05)	W.
Emergency CS	Lisonkova <i>et al</i> <sup>28</sup> , 2011,	Travel distance (km): <50 Groups	Groups	N (9657)	Emergency CS n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Canada	061< 061-06	<50	6606	5378 (59.11)	<50	-	RN RN
			50–150	464	258 (55.60)	50–150	0.87 (0.72 to 1.05)	RN
			>50	94	52 (55.32)	>50	0.86 (0.57 to 1.29)	N.
Severe perineal trauma	No studies							
Postpartum haemorrhage	No studies							
Admission to ICU	No studies							
Blood transfusion	No studies							
Stillbirth (SB)	Pilkington <i>et al</i> <sup>25</sup> , 2014,	Travel distance (km): <5	Groups	N (30 859) (2002-2005)	SB n (/per 1000)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	France	5-14 15-29 30-44 ≥45	<5	1 404 665	13204 (9.4)	<5	-	-
			5–14	81–1775	6657 (8.2)	5–14	0.87 (0.85 to 0.90)	Reported as RR 0.87 (NR)*
			15–29	648 495	5188 (8.0)	15–29	0.85 (0.82 to 0.88)	Reported as RR 0.85 (NR)*
			30-44	186537	1492 (8.0)	30-44	0.85 (0.81 to 0.90)	Reported as RR 0.85 (NR)*
			≥45	34367	306 (8.9)	<b>≥45</b>	0.95 (0.84 to 1.06)	Reported as RR 0.95 (NR)(NS)
	Lisonkova <i>et al</i> <sup>28</sup> , 2011,	Travel distance (km): <50 Groups	Groups	N (29 698)	NM n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Callada	001<001-00	<50	27 836	150	<50	NB	RN
			50–150	1534	N.	50–150	NB	RN
			C	4 4	<u>q</u>		0	<u> </u>

Continued

Table 6 Continued	pe							
Outcomes	Author, year, country	Exposure groups	Participants (N, n, %)			Findings		
Neonatal mortality (NM)	Pasquier <i>et al</i> <sup>24</sup> , 2007,	Travel distance (km): <11	Groups	N (554)	NM n (%)	Groups	Crude OR (95% CI)	Adjusted OR(95% CI)
	rrance	11-50 >50	<b>11</b>	239	NR	<u>+</u>	NR	-
			11–50	156	NR	11–50	NB	0.98 (0.34 to 2.88)
			>50	159	NR	>50	NB	1.37 (0.49 to 3.86)
	Pilkington et al <sup>25</sup> , 2014,	Travel distance (km): <5	Groups	N (6 202 918) (2001–2008) NM n(/per 1000)	NM n(/per 1000)	Groups	Crude OR (95% CI)	Adjusted RR (95% CI)
	France	5-14 15-29 30-44 ≥45	<5	2 808 068	7582 (2.7)	<5	-	-
			5-14	1626885	3416 (2.1)	5–14	0.78 (0.75 to 0.81)	Reported as RR 0.91 (NR) *
			15–29	1316329	2896 (2.2)	15–29	0.81 (0.78 to 0.85)	Reported as RR 0.94 (NR)(NS)
			30–44	381 288	801 (2.1)	30-44	0.78 (0.72 to 0.84)	Reported as RR 0.9 (NR)*
			≥45	69787	154 (2.2)	≥45	0.82 (0.70 to 0.96)	Reported as RR 0.96 (NR)(NS)
			NM after BBA			NM after BBA		
			Groups	N (6 202 918)(2001-2008)	NM n (/per 100,000)	Groups	Crude OR (95% CI)	Adjusted RR (95% CI)
			<5	2808068	115 (4.1)	<5	-	-
			5–14	1626885	65 (4.0)	5–14	0.98 (0.72 to 1.32)	Reported as RR 1.1 (NR) (NS)
			15–29	1316329	72 (5.5)	15–29	1.34 (0.99 to 1.79)	Reported as RR 1.58 (NR)*
			30–44	381288	23 (6.0)	30-44	1.47 (0.94 to 2.30)	Reported as RR 1.51 (NR)(NS)
			≥45	69 787	7 (10.0)	≥45	2.45 (1.14 to 5.25)	Reported as RR 3.68 (NR)*
Perinatal mortality (PM)	Lisonkova <i>et al</i> <sup>28</sup> , 2011,	Travel distance (km): <50 Groups	Groups	N (29 698)	PM n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Oallada	001<001=00	<50	27 836	221 (0.80)	<50	-	-
			50-150	1534	19 (1.24)	50–150	1.57 (0.98 to 2.51)	1.53 (1.10 to 2.12)
			>150	328	8 (2.44)	>50	3.12 (1.53 to 6.38)	3.06 (2.20 to 4.24)
Infant mortality (IM)	No studies							

Continued

Table 6 Continued	pə							
Outcomes	Author, year, country	Exposure groups	Participants (N, n, %)			Findings		
Born before arrival (BBA)	Bhoopalam <i>et al<sup>23</sup>,</i> 1991, UK	<b>Travel distance (km):</b> <2 2-7 >7	Groups	N (398) cases and controls	BBA cases n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			<2	59	4 (6.80)	<2	-	NR
			2–7	249	88 (35.34)	2-7	7.52 (2.64 to 21.43)	NB
			>7	06	42 (46.70)	>7	12.03 (4.02 to 36.01)	NR.
	Blondel <i>et al</i> <sup>5</sup> 2011, France	Travel distance (km): <5	Groups	N (1 359 756)	BBA n (rate /1000 births)	s) Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
		5-14 15-29 30-44 >45	<5	596363	1849 (3.1)	<5	-	NR.
			5–14	352 279	1395 (3.9)	5–14	1.28 (1.19 to 1.37)	NR.
			15–29	296734	1659 (5.6)	15–29	1.81 (1.69 to 1.93)	NR.
			30-44	88670	692 (7.8)	30-44	2.53 (2.32 to 2.76)	NR
			>45	15705	182 (11.)	45+	3.77 (3.23 to 4.39)	NR
							Parity 1 nd 2 n=152 426	Parity 3+ N=197325
						Groups	Adjusted OR (95% CI)	Adjusted OR (95% CI)
						<5	-	$1.73 (1.57 \text{ to } 1.90)^a$
						5–14	1.14 (1.03 to 1.27)	2.32 (2.04 to 2.63)
						15–29	1.39 (1.24 to 1.57)	3.25 (2.84 to 3.71)
						30–44	1.78 (1.55 to 2.05)	3.71 (3.13 to 4.41)
						>45	2.47 (2.02 to 3.02)	6.46 (4.92 to 8.48)
	Ovaskainen <i>et al</i> <sup>27</sup> , 2015,	Travel distance (km):	Groups	N (201)	BBA n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Finland	<35 ≥35	BBA cases	29	NR	<35	NB	-
			Controls	134	NR	≥35	NR	5.02 (1.80 to 14.04)
			<35 km	N.	N.			
			≥35 km	NB	NR			
	Fougner <i>et af</i> <sup>66</sup> , 2000, Norway	<b>Travel distance (km):</b> <12.88 ≥12.88	Groups	N (202) cases and controls	BBA n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			<12.88	06	44 (48.90)	<12.88	-	NR.
			≥12.88	112	71 (63.34)	≥12.88	1.81 (1.03 to 3.18)	NB
Neonatal unit admission		Travel distance (km): <50	Groups	N (15 325)	NNU n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
(DNN)	Canada	061< 061-06	<50	14333	648 (4.80)	<50	-	RN
			50–150	815	32 (3.92)	50–150	0.86 (0.60 to 1.24)	RN
			>150	177	12 (6.80)	>150	1.54 (0.85 to 2.77)	RN
Apgar score	No studies							
뿦	No studies							

"Significant difference.

"Significant difference.

Floating and <5 km.

Floating and <5 km.

Floating and <5 km.

Floating age: HE, hypoxic-ischemic encephalopathy; ICU, intensive care unit; NNU, neonatal unit; NR, not reported; NS, not significant; OR, odd ratio, RR, relative risk.



# Neonatal mortality (NM)

Two French cohort studies<sup>24</sup> <sup>25</sup> reported this outcome. One study<sup>24</sup> examined the distance from women's homes to the nearest OU with neonatal surgical facilities for 706 fetuses with severe malformations. Analyses adjusted for malformation type, number of malformations, amniotic fluid anomaly, previous anomaly in the family and parity showed no association between NM and distance (<11 km vs 11-50 km, adjOR=0.89, 95% CI: 0.34, 2.88;<11 km vs >50 km, adjOR=1.37, 95% CI: 0.49, 3.86). The other study<sup>25</sup> included all births and found that NM rates were significantly higher for women living <5 km compared with 5-44 km away from an OU<sup>25</sup> (5-14 km vs <5 km, cOR 0.78, 95% CI: 0.75, 0.81; 15-29 km vs <5 km cOR 0.81, 95% CI: 0.78, 0.85; 30-44 km vs <5 km, cOR 0.78, 95% CI: 0.72, 0.84; ≥45 km vs <5 km, cOR 0.82, 95% CI: 0.70, 0.96). In this latter study, the NM of babies BBA was also explored. For the BBA group, there was a statistically significant increase in the risk of NM when women had to travel 45 km or more to an OU in comparison to <5 km (≥45 km vs <5 km, cOR 2.45, 95% CI 1.14 to 5.25).

#### Perinatal mortality (NM)

A study from Canada<sup>28</sup> reported that PM risk increased with travel distance in an adjusted model (<50 km v 50-150 km adjOR 1.53, 95% CI 1.1 to 2.12; <50 km >150 km adjOR 3.06, 95% CI 2.20 to 4.24).

#### Infant mortality (IM)

No studies reported this outcome.

#### Born before arrival (BBA)

Three case–control studies, <sup>23</sup> <sup>26</sup> <sup>27</sup> and one cohort study<sup>5</sup> reported this outcome. All four studies reported a significant increase in BBA rate with longer travel distance, although only two reported adjusted analyses. <sup>5</sup> <sup>27</sup> In the UK study, <sup>23</sup> the risk of BBA increased 12-fold for women living >7 km from the OU compared with women living <2 km away (cOR 12.5, 95% CI 4.02 to 36.01). The risk of BBA increased significantly for women living >13 km from an OU in a Norwegian study <sup>26</sup> (cOR 1.81, 95% CI 1.03 to 3.18). The Finnish study <sup>27</sup> reported a fivefold increased risk of BBA for women living >35 km from the OU compared with <35 km (adjOR 5.02, 95% CI 1.80 to 14.04).

In France,<sup>5</sup> the rate of BBA significantly increased with longer distances and it tripled for all women living 45+km from the OU compared with women living <5 km away (cOR 3.77, 95% CI 3.23 to 4.39). The association persisted in an adjusted analysis which included women of parity three or higher and living 45+km from the OU, who had a sixfold increased risk of BBA compared with women living <5 km away and of parity one or two (adjOR 6.49, 95% CI 4.92 to 8.48).

# Neonatal unit (NNU) admission

A study from Canada<sup>28</sup> reported an increase in NNU admission for births to women living >150 km from an OU compared with those living <50 km away (6.8% vs 4.8%).

#### Apgar score

No studies reported this outcome.

#### Hypoxic-ischaemic encephalopathy (HIE)

No studies reported this outcome.

# Evidence from travel time studies

# **Description of included studies**

Fifteen studies explored the impact of travel time from a woman's home to an OU (see table 7). Two studies (one reported as an abstract only) were conducted in the UK, <sup>29–31</sup> three studies in France, <sup>32–34</sup> three studies (reported in five articles) in the Netherlands, <sup>35–39</sup> one study reported in two articles from Norway, <sup>40 41</sup> five studies in Canada <sup>42–46</sup> and one study in Japan. <sup>47</sup>

Eleven studies were of a retrospective cohort design, one was a prospective cohort study, <sup>39</sup> one was a before-and-after design <sup>47</sup> and two were case–control studies. <sup>33</sup> <sup>34</sup> All the studies clearly stated the eligibility criteria. Only singleton births were included in five studies. <sup>30–32</sup> <sup>35–38</sup> <sup>42</sup> One study <sup>39</sup> specifically enrolled women with postnatal haemorrhage after home birth, and one study <sup>42</sup> focused on planned home birth regardless of the actual place of birth.

The studies were heterogeneous in their travel time intervals. With the exception of one study in Canada, <sup>42</sup> longer time cut-off points were examined in studies from Norway, Japan and Canada compared with studies in other countries (all European). Travel duration was estimated using geographical mapping software in all studies. However, most studies estimated travel duration to and from central points within areas rather than actual addresses.

#### Risk of bias assessment

Risk of bias assessment and supported explanations for each of the risk of bias domains are presented in table 8. With the exception of Stolp et al, 39 sample selection and measurement of outcomes were considered to be at low risk of bias across all studies as such data were obtained from national databases and birth registries. The groups in the two case-control studies were appropriately selected and defined, however, the case and control groups were not comparable in both studies (eg, difference in antenatal care attendance and sociodemographics). Eight studies<sup>29</sup> 34-42 were considered at low risk of exposure measurement bias, as the women's actual place of residence was used to estimate travel time to nearest OU. The risk of attrition bias was low for the majority of the included studies. Similarly, analyses and adjustment for potential confounders were found to be appropriate in the majority of studies.

# Findings

Maternal outcomes

# Maternal mortality (MM):

No studies reported this outcome.

Table 7 Descr	Description of included studies - travel time	studies- travel t	time						
Author, year, country	Study designandsetting	Study objectives	Study period	Eligibility criteria	Participant characteristics	Description of exposures travel distance	Services context information	Review outcomes Perinatal	Maternal
UK Studies									
Dummere <i>t af</i> °³, 2004, UK	Retrospective population-based cohort study, Cumbria	To investigate whether geographical accessibility to hospitals affected SB rates and infant mortality	1950-1993 grouped: 1950-1959 1960-1969 1970-1979 1980-1993	Included: All births Excluded: Women with missing postcodes	n=283 668 births Other characteristics: NR	Travel time (mins): <17 18–35 >35	Universal state provision of maternity care. 1950–1993: 4 hospitals opened, 2 closed	NM; Early NM; Post NM NR	E E
Paranjothyet al <sup>20</sup> 31, 2013, 2014, UK (abstract® full paper)	Retrospective cohort study, All Wales Perinatal Survayand National Community Chilid Health Database	To study the association between travel time from home to OU on intrapartum stillbirth (SB)andNM	1995-2009	included: All registered birth >23 wks GA Excluded: Antepartum SB, lethal congenital anomalies, multiple pregnancies, invalid or missing GA, missing maternal age/postcode/ hospital of birth or baby's gender	n=466.255 singleton births Maternal age yrs %-20 90.7 20-34 76.5 34-44 1.38 454-0.1 Parity. Nullips 44.9% Social deprivation quintile %: I (least dept) 16.7 2-4 57.8 5 (most dept) 25.6 Ethnicity, education. NR	Travel time (mins): <15 15-29 30-44 >45	Universal state provision of maternity care. 50 hospitals (16 outside Wales)	Intrapartum SB; Early NM; Late NM	띺
Other European Studies	sə								
Combieret al <sup>02</sup> , 2013, France	Retrospective cohort study, based on hospital discharge summaries, Burgundy	To analyse the effect of travel time to closest OU on pregnancy outcomeandprenatal management in Burgundy	2002-2009	included: Singleton births >21 wks GA Excluded: Medical ToP, multiple pregnancy, births outside Burgundy, births in 2002 and 2008 due to closure of 3 units	n=111001 births Other characteristics=NR	Tavel time (mins): ≤15 16–30 31–45 ≥46	2000–2001; 2 private maternity units closed 2002–2009: 3 public maternity units closed. Units(n): 2000 (20) 2009 (15)	SB; PM; BBA	<u> </u>
Renesme <i>et al<sup>24</sup>,</i> 2013, France	Case-control, multicentre study, 8 units, Finistere District, Brittany	To evaluate the social-geographical factors associated with BBAs	2007-2009	Included cases: BBA of live birth Included controls: 2 controls: 2 controls for each case irrespective of delivery mode. Excluded: GA <22 weeks, BW <500g, planned home birth	n=225 Cases vs controls rare 6 vs 149 Age (median, range) vrs: 30 (16-41) vs 30 (16-41) vs 30 (16-41) parity (median, range); 2 (1-6) Maternal INSEE code n (9); 1, 2, 3 or 4=15 (23.8) vs 56 (43.4); five or 6; 20 (31.8) vs 56 (42.6); 8=28 (44.4%) vs 18(14) Ethnicity, education: NR	Travel time (mins): 15 15–29 30–44 >45	9700 births/year in Finistere in 2012 units with <300 births/yrs were closed. Universal state provision of maternity care	ВВА	N.
Nguyen <i>et al</i> <sup>23</sup> , 2016, France	Case-control study, university hospital in Caen	To estimate the incidence of BBA during the study period	2002-2009	Included cases: Unplanned BBA Included controls: Next spontraneous birth in hospital Excluded: NR	n=188Casesn=94 Mean age: 28.9 years Parity: 1. SES: 73.4 % no profession/student Control n=94 Mean age: 29.2 years Parity: 0.9 SES: 47.9% no profession/ student Ethnicity, education: NR	Travel time (mins): <20 mins >20 mins	University Hospital with neonatal care facilities. Universal state provision of maternity care	ВВА	<del>М</del>
Ravelli Study									

Table 7 Continued	nued								
						Description of exposures	Services context	Review outcomes	
Author, year, country	Study designandsetting Study objectives	Study objectives	Study period	Eligibility criteria	Participant characteristics travel distance	travel distance	information	Perinatal	Maternal
Ravelli et a <sup>(36-37</sup> , 2011, Netherlands (full papers & abstract)	Retrospective s population-based cohort study, rural and urban areas, 12 provinces	To study the effect of travel time from home to OU on mortality and other adverse outcomes in pregnant women at term in primary and secondary care	2000–2006	Included: Singleton term births Excluded: Antepartum deaths, congenital anomalies, invalid/missing postcodes or outpatient codes, or births from Wadden islands, hone deliveries, hospitals participated for 1–2 years	n=751926 singleton births Age yrs, % <20, 2 20–34, 78.3 35–39, 17.2≥40, 2.4 Party: Nullips: 49.9% Ethnicity: White 81.7% SES %: high 25.2, medium 48.2, low 26.7 Education: NR	7ravel time (mins): <20≥20	Universal state provision of maternity care. 99 OUs including tertiary perinatal centres	NM (Combined intrapartum & early & late NM up to 28 days) NM (0-27 days) NM(0-27 days); Combined (mortality and/or Apgar<4 at 5 min, and/or NNU admission)	띺
Ravelli et al <sup>38</sup> , 2012, Netherlands	Retrospective cohort study in nine regions	To investigate provincial 2000–2006 diffeences in perinatal mortality (PM) and to determine the influence of different risk factors, including tavel time from home to the OU during labour.	1 2000–2006	Included: Singleton births Excluded: Women with incorrect post codes	n=1242725 singletons Age yrs, % <20, 18-35, 19.5 Parity, % Nullips, 46.3 Ethnicity, % Non- western 16.2 SES low (10 <sup>th</sup> centile): 10% Education NR	Travel time(mins): <20>20 Universal state provision of ma care	Universal state provision of maternity care	Md	RN
Stolp et al <sup>39</sup> , 2015, Netherlands	Prospective cohort study, rural & urban areas	To assess whether the limit of 45 mins is met for ambulance transfer of women with PPH after home birth	2008-2010	Included: Women with PPH after MW supervised home birth Excluded: Cases of PPH with missing data	n=72 (54 analysed) Age median (range) yrs: 31 (23-41); Parity (n, %); Primip 27%- 50% Ethnicity, Education, SES: NR	Travel time (mins): <45 >45	Home birth for low risk women and hospital birth. Universal state provision of maternity care	æ Z	Maternal admission to intensive care; (ICU); Blood transfusion; Postpartum haemorrhage (PPH)
Egjom Study									
Engjomet a <sup>nd 41</sup> , 2017 & 2015, Norway (full paper & abstract)	Retrospective population-based cohort study, Medical Birth Registry of Norway and Statistics Norway, 19 counties	To assess peripartum mortality associated with place of birth and availability of obstetric units.	1999–2009	Included: All births in Norway with GA 222 wks or BW 2500g Excluded: Lack of address and municipality, antepartum SB, planned home births	n=646 888 960.4% singletons. Age yrs, % <20.20.4.20-35, 80.77-35, 16.9%; Multips 58.7%; Education >11 y 77.2 Ethnicity: Western 90.7%	Travel time (hrs):<1 1-2 >2	Basic obstetric care for nomal delivery; Emergency obstetric care <1500 births/ yr. Universal state provision of maternity care	ВВА	띺
Canadian Studies									
Grzybowskind Stroll Study	study								
Grzybowski <i>et al</i> <sup>rts</sup> , 2011, Canada	Retrospective cohort study, rural areas of British Columbia	To document newborn and maternal outcomes in relation to travel time to the nearest OU with CS capability	2000-2004	Included: All deliveries>20 weeks' deliveries>20 weeks' dA Excluded: Multiple birth, companital anomailes or late Top. core urban areas	n=35 426 birthsGroups:<1 hour, 1-2, 2-4,>4 hours Group N. 32 814, 1559, 747, 506 Mean maternal age yrs: 28.7, 28.67, 27.25, 27.2 Parity % primips: 42.6, 38.6, 36.7, 36.8 SES: 0.12, 0.10, 0.30, 0.33, first Nations % 0.05, 0.30, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.23, 0.22, 0	Travel time (hrs): <1 1-2 2-4 >4	Universal medical coverage for core healthcare, 13 NNUs, 42 000 births /year	PM (SB & early NND); BBA; NNU admission	8
Grzybowski <i>et af<sup>4</sup>,</i> 2013, Canada	Retrospective cohort study, rural areas of British Columbia	To compare rural maternity care by level of services	2000–2007	Included: Singleton births Excluded: Women with residential postcode of large urban centres	n=4672 births; Mean age, yrs: 27.7 Parity: primips: 39.7% SES*: 0.22%; Ethnicity: first Nations 0.3% Education: NR	Travel time (hrs):<1 >1	Universal medical coverage for core healthcare	SB; NND (late<1 month); PM; IM; BBA; NNU admission	CS; Emergency CS; PPH
									Continued

Table 7 Continued	penu								
						Description of exposures Services context	Services context	Review outcomes	
Author, year, country	Study designandsetting Study objectives	Study objectives	Study period	Eligibility criteria	Participant characteristics travel distance	travel distance	information	Perinatal	Maternal
Grzybowskier a <sup>rts</sup> , 2015, Canada	Retrospective cohort study, British Columbia (BC), Alberta, Nova Scotia (NS)	To examine the safety of rural Canadian maternity services	2003–2008	Included: Singleton deliveries Excluded: Mutiple births, infants born with congenital anomalies, planned home births, accidental BBA	Alberta, BC, NS Age yrs (n %) <18: 1618 (2.3), 1256 (2.0), 413 (2.2) >35 yrs: 5127 (7.3) 886 (14.3), 2387 (12.7) Multips n (%) 41730 (59.6), 35 089 (56.6), 10656 (56.8) Ethnicity, SES, education: NR	2-4 >4	Universal medical coverage for core healthcare, 20 small maternity closures since 2000	PM (SB & NND up to 7 days)	ઈ
Stoll <i>et al</i> <sup>46</sup> , 2014, Canada	Retrospective cohort study, rural British Columbia	To report on characteristics and perinatal outcomes of rural women with only MW involved in care	2003–2008	Included: Women residing outside core urban areas, singletons >20w & Adand care by a MW Excluded: Late ToP, congenital anomalies	<1 hour, 1–2 hours, >2 hours:n=3438, 124, 130 Mean age yrs: 29,78, 314, 30.5 Primps n (%) 1574 (45), 63 (50.8), 63 (48.5) Ethnicity, SES & Education: NR	1-2 >2 1-2 >2	Universal medical coverage for core healthcare, closure of 22 rural maternity services	PM (SB & NND up to 7 days)	8
Darling et al <sup>re</sup> , 2019, Canada	Retrospective population-based cohort study, Ontario	Whether greater diving distances to OU associated with a higher risk of adverse neonatal outcomes	2012–2015	Included: Women who planned home births regardless of actual place of births Excluded: Multiple births, Preterm <37 wks	n=11869 Age yrs, %:<25, 9.5 25-39,87.6≥40, 2.9 Primps n (%) 4208 (35.5) SES low, n(%) 2465 (20.8) Ethnicityandeducation: NR	Travel time (mins): <30 >30	Universal medical coverage for core healthcare	PM (PM); NNU admission; 5mins Apgar<7	SO
Other countries									
Aoshima <i>et al<sup>47</sup>, 2</i> 011, Japan	Before and after study design, data from perinatal care centres	Whether reducing travel time influences the neonatal mortality rate (NM)	2002–2006	Included: All births Excluded: Municipalities consisting of isolated islands	Number of births: 2002=347 284 2006=322 514 Other characteristics: NR	Travel time (hrs): ≤1 >1	Universal healthcare insurance system, 346 perinatal care centres	M	EN.

INSEE: Institute National de la Statistique et des Etudes Economiques; INSEE codes: 1: farmer, 2: craftsperson, merchant or entrepreneur; 3: businessexecutive, intellectual occupation; 4: other professionals; 5: employee; 6: worker; 8: no occupation. 1-15 occup

Table 8 Risk o	Risk of bias—travel time						
Author, year, country	Study sample selection bias <b>additional criteria for case- control studies</b>	tional criteria for case- control	Bias in measurement of exposure	Bias in measurement of outcomes	Attrition bias	Analysis method reported and appropriate	Potential confounders adjusted for and listed
UK Studies							
Dummer & Parker <sup>29</sup> , 2004, UK	LOW Cumbrian Births Database		LOW Modelled using GIS	<b>LOW</b> Objective outcome (NM)	LOW Of 3352 live births, 42 stillbirths excluded as the outcome NM	<b>LOW</b> Method detailed, results of LR were reported	LOW Adjusted for year of birth, social class, birth order, multiple births
Paranjothy <i>et aj<sup>® 31</sup>,</i> 2013 & 2014, UK	LOW National Community Child H Survey)	Low National Community Child Health Database & All Wales Perinatal Survey)	HIGH Women's address replaced by population-weighted centroid, travel time calculated using Google Maps API (v3)	LOW Objective outcomes (SB, NM)	LOW 11% excluded where information on parity was missing	LOW Analysis method described and multilevel LR data were reported	LOW Adjusted for maternal age, parity, urban/rural location, SES, and other characteristics
European Studies							
Combier <i>et al</i> <sup>92</sup> , 2013, France	LOW Burgundy perinatal network database	database	HIGH Municipality town hall not woman's home address	LOW Objective outcomes (SB, PM, BBA)	LOW All births identified included in the analysis	LOW Method described; hierarchical LR and multilevel LR reported	LOW Adjusted for maternal age, urbanisation level and other characteristics
Renesme <i>et al</i> ³4, 2013, France	Case definition	YES linked to perinatal network database	LOW Distance & travel time estimated using GIS	LOW Objective outcomes retrieved	<b>LOW</b> 5/81 (6%) BBAs missing, 3/162 (2%) controls	LOW Method described and univariate and appropriate; multivariate reported	LOW Adjusted for age, family status, INSEE
	Representativeness of cases	YES All cases in defined period		from regional and hospital databases	missing		maternal occupation, parity, and other characteristics
	Appropriate selection of controls	YES Controls chosen randomly from same databases and from births occurring at the nearest delivery date and hour to cases					
	Definition of control appropriate	YES Outcome could not have occurred					
	Comparability of cases and controls	NO Difference in antenatal care attendance					
Nguyen et a/33, 2016,	Case definition	YES Using medical records	UNCLEAR No information	Low Objective	UNCLEAR No information	HIGH No details of the analysis method HIGH No adjustment for	HIGH No adjustment for
France	Representativeness of cases	YES All cases in defined period		outcome (BBA)		and analysis was only descriptive	any potential confounders
	Appropriate selection of controls	YES Next birth, of equivalent GA					
	Definition of control appropriate	YES Outcome could not have occurred					
	Comparability of cases and controls	NO Significant differences in parity, smoking, pregnancy monitoring, profession					
Ravelli et al <sup>35–37</sup> , 2011, Netherlands (abstract & full papers)	LOW Population based study usir	LOW Population based study using the Netherlands Perinatal Registry	LOW GIS software used to measure travel time from women's postcodes	Low Objective outcomes from perinatal registry	LOW Small proportion (0.3%) of women excluded due to incorrect zip code	LOW Method reported; descriptive analysis & LR results given	LOW Analysis adjusted for age, parity, ethnicity, SES
Ravelli et al <sup>38</sup> , 2012, Netherlands	LOW Population based study usir	LOW Population based study using the Netherlands Perinatal Registry	LOW GIS software used to measure travel time from women's postcodes	Low Objective outcomes from perinatal registry	LOW Small proportion 4% of women excluded	UNCLEAR No information	LOW Adjusted for age, parity, very urban /very rural, SES
Stolp <i>et al</i> <sup>39</sup> , 2015, Netherlands	HIGH Study participants were selected by midwives	ected by midwives	LOW Ambulance interval includes total time from dispatch call to arrival at hospital	UNCLEAR Method of measuring blood loss not reported	HIGH Missing data 18/72 (25%) due to incomplete documentation	<b>НІGH</b> Data only analysed descriptively	HIGH No adjusted analysis
Engjom et at**0, 2017 and Engjom et at**1, 2015, Norway (abstract & full paper)	LOW Medical Birth Registry of Norway	veavo	LOW Travel time polygon from home address using GIS	LOW Objective outcomes from birth registry	UNCLEAR No information	LOW Analysis appropriate, details of LR, multilevel modelling were reported	LOW Adjusted for maternal age, parity, education, ethnicity
							Continued

Table 8 Continued	nued					
Author, year, country	Study sample selection bias additional criteria for case- control studies	Bias in measurement of exposure	Bias in measurement of outcomes	Attrition bias	Analysis method reported and appropriate	Potential confounders adjusted for and listed
Grzybowski et aj <sup>43</sup> , 2011, Canada	<b>LOW</b> Population based study using British Columbia Perinatal Health Programme	HIGH GIS used to create 1 hour LOW Objective travel zone for each maternity outcomes from service, but central postal code to the nearest maternity Programme care used	LOW Objective outcomes from Perinatal Health Programme	<b>LOW</b> 0.3% excluded due to incorrect zip code	LOW Analysis appropriate, descriptive analysis & hierarchical LR reported	LOW Adjusted for maternal age, parity, SES, ethnicity
Grzybowski e <i>t al<sup>44</sup>,</i> 2013, Canada	LOW Data from Perinatal Data Registry	HIGH Community central postal code used not women's home address	LOW Objective outcomes from Perinatal Data Registry	HIGH Number of women excluded due to incorrect postal address not reported	LOW Analysis appropriate descriptive analysis & LR	LOW Adjusted for maternal age, parity, lone parent status, ethnicity, SES
Grzybowski <i>et al</i> <sup>45</sup> , 2015, Canada	LOW Provincial perinatal registries	HIGH Community central point postal code used not women's home address	LOW Objective outcomes from Perinatal Data Registries	UNCLEAR No information on missing data	LOW Analysis appropriate, descriptive analysis & LR reported	<b>Low</b> Adjusted for maternal age, parity
Stoll <i>et al</i> <sup>46</sup> , 2014, Canada	LOW Based on British Columbia Perinatal Database Registry	LOW Used GIS and Google maps; travel times were adjusted for travel conditions	Low Objective outcomes (CS)	LOWNo missing data	HIGH Data were only analysed descriptively	HIGH No adjusted analysis
Darling <i>et al<sup>42</sup>,</i> 2019, Canada	LOW Data from Perinatal Registries	LOW Driving time from women's residence using online mapping tool ArcGIS	LOW Objective outcomes from Perinatal Data Registries	LOW 3.7% excluded not being able to calculate distance to nearest hospital	LOW Method reported, results of descriptive analysis & LR reported	LOW Adjusted for maternal age, parity, gestational age, season, SES
Aoshima et af <sup>r</sup> , 2011, Japan	Aoshima <i>et al</i> <sup>rr</sup> , 2011, <b>LOW</b> All Japan except for isolated islands outside road network (96.6% Japan of all Medical Service Areas)	HIGH Used central point of municipality not home address but analysis based on (larger) Medical Service Areas.	LOW Objective outcomes from Medical Service Area databases	UNCLEAR No information on missing data	LOW Method appropriate, unpaired t-test, difference-in-difference analysis	HIGH No adjusted analysis

BBA, bom before arrival; CS, caesarean section; GA, gestational age; GIS, geographicial information system; INSEE, institute national de la statistique et des etudes economiques; LR, logistic regression; NIN, neonatal mortality; SES, socio economic status; SB, still birth.



# Caesarean section (CS) (overall, or intrapartum)

Five Canadian studies <sup>42–46</sup> reported CS rates (table 9). Across three studies, <sup>43–45</sup> cORs for CS rates were higher among women who lived closer to OUs with CS rates highest for women living less than 1 hour away compared with other categories (1–2 hours, 2–4 hours and >4 hours). One study <sup>46</sup> included women who had a midwife involved in their care, and found no significant differences in CS rates for women living 1–2 hours and more than 2 hours away compared with within 1 hour of an OU (1–2 vs <1 hour, cOR 1.23, 95% CI 0.80 to 1.91 and >2 hours vs <1 hour, cOR 1.11, 95% CI 0.71 to 1.72). A further study <sup>42</sup> also showed a higher CS rate among women who planned a home birth and lived less than half an hour away from OU services (>30 min vs ≤30 min, cOR 0.74, 95% CI 0.59 to 0.92).

# **Emergency CS**

Shorter travel time to an OU was associated with a statistically significant higher emergency CS rate in one Canadian study<sup>45</sup> (>1 hour vs <1 hour, cOR 0.80, 95% CI 0.75 to 0.86).

#### Severe perineal trauma

No studies reported this outcome.

#### Postpartum haemorrhage (PPH)

One Canadian study found the risk of PPH was significantly higher for women who lived more than 1 hour away from obstetric services compared with women who lived less than 1 hour away<sup>44</sup> (>1 hour vs <1 hour, cOR 1.27, 95% CI 1.13 to 1.43).

#### Maternal admission to ICU

One study from the Netherlands<sup>39</sup> involved women who had a PPH after midwifery-supervised home births and examined adverse maternal outcomes associated with travel time longer than 45 min to hospital. No difference was found in the number of women admitted to ICU who travelled more than 45 min compared with <45 travel time to hospital, but the numbers of events were low.

# Maternal blood transfusion

One study from the Netherlands<sup>39</sup> found no significant difference in the median number of units of blood transfused to women who travelled more than 45 min to an OU compared with <45 min travel time.

#### Neonatal outcomes:

# Stillbirth (SB) (overall or intrapartum)

Three studies examined the association between increasing travel time and SB, one study each from the UK, <sup>30 31</sup> France<sup>32</sup> and Canada. <sup>44</sup>

In the UK study, <sup>30 31</sup> there was no association between travel time and SB when analysing all women (adjOR 1.13, 95% CI 0.98 to 1.30). However, subgroup analyses showed a significant increase in the risk of SB with every 15 min increase in travel time to the OU for term pregnancies (adjOR 1.36, 95% CI 1.17 to 1.59) and for nulliparous

women (adjOR 1.21, 95% CI 1.02 to 1.44). The other two studies  $^{32}$   $^{44}$  found no significant increase in the incidence of SB with increasing travel time.

# Neonatal mortality (NM)

Five studies examined the association between travel time and NM, two from the UK, <sup>29–31</sup> one from the Netherland, <sup>35–37</sup> one from Canada<sup>44</sup> and one from Japan. <sup>47</sup>

The adjusted analysis in one UK study<sup>29</sup> showed no statistically significant association between NM and travel time. The adjusted analyses in the other UK study<sup>31</sup> showed a significant increase in early and late NM, with every 15 min increase in travel time (adjOR 1.13, 95% CI 1.07 to 1.20) and (adjOR 1.15, 95% CI 1.05 to 1.26) respectively. Subgroup analysis for nulliparous women showed a statistically significant increased risk of early NM associated with every 15 min increase in travel time from home to the OU (adjOR 1.15, 95% CI 1.06 to 1.25). For term births, late (but not early) NM increased significantly with every 15 min increase travel time from home to the OU (adjOR 1.34, 95% CI 1.13 to 1.59).

In one study from the Netherlands,  $^{35}$  a travel time of 20 min or more was associated with a significant increase in the combined intrapartum, early and late NM $^{35-37}$  ( $\geq$ 20 min vs <20 min, adjOR 1.23, 95% CI 1.07 to 1.41). No NM events were reported in the study from Canada. The study from Japan $^{47}$  reported that following a median reduction in travel time from 67 min in 2002 to 39 min in 2006 that there was a decrease in NM rate from 1.67 to 1.28, however, no further analyses were presented.

# **Perinatal mortality (PM)**

Seven studies examined PM, one from France,<sup>32</sup> one from the Netherlands<sup>38</sup> and five from Canada.<sup>42-46</sup> The French study<sup>32</sup> found no significant association between increasing travel time to the nearest OU and PM based on unadjusted data. However, in the Dutch study a longer travel time (20 min or more) was significantly associated with higher PM<sup>38</sup> ( $\geq$ 20 min vs <20 min, adjOR 1.66, 95% CI 1.59 to 1.74).

The Canadian studies also reported longer travel times to OUs being associated with an elevated risk of PM. A significant increase in PM was reported in women living more than 4 hours away from OUs compared with women living less than 1 hour (>4 hours vs <1 hour adjOR 3.17, 95% CI 1.45 to 6.95). 43 However, findings from the same study suggested no significant increase for women living 1-2 hours and 2-4 hours from an OU compared with those living less than 1 hour from services. Similarly, the PM risk significantly increased in women who lived >1 hour from OUs in a further Canadian study, 44 (cOR 1.54, 95% CI 1.09 to 2.17). When this was divided into different Canadian provinces, 45 the rates of PM were highest in communities living more than 4 hours from an OU in comparison to less than 1 hour in British Colombia only (adjOR 2.84, 95% CI 2.84 to 5.10). Stoll and Kornelsen, 46 found that in women who received midwifery care only, PM was not statistically significantly different for women

Table 9 Outcomes—travel time	ravel time							
Outcomes	Study,year, country	Exposure groups	Participants (N, n, %)	n, %)		Findings		
Maternal outcomes								
Maternal mortality (MM)	No studies							
Caesarean section (CS)(overall or intrapartum)	Grzybowski et a/ <sup>43</sup> , 2011, Canada	<b>Travel time (hrs):&lt;1</b> 1-2 2-4 >4				All CS		
			Groups	N (35,429)	All CS n(%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			▽	32814	8597 (26.2)		-	NR
			1-2	1359	313 (23)	1-2	0.84 (0.74, 0.96)	NR
			2-4	747	156 (20.9)	2-4	0.74 (0.62, 0.89)	NR
			>4	909	97 (19.06)	>4	0.66 (0.53, 0.83)	NR
	Grzybowski <i>et al</i> <sup>44</sup> , 2013,	Travel time (hrs): <1 >1						All CS
	Canada		Groups	N (59 386)	n(%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			√	54 714	14882 (27.20)	₹	_	NR
			7	4672	1075 (23.01)	<u>~</u>	0.80 (0.75, 0.86)	NR.
	Grzybowski <i>et al</i> <sup>45</sup> , 2015,	Travel time (hrs): <1 1-2 2-4 >4						Alberta CS
	Canada		Groups	Alberta N (34 453)	%u	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			<1 hour	29906	NR		NR	-
			1-2	2940	NR	1-2	NR	0.86 (0.78, 0.94)
			2-4	1297	NR	2-4	NB	0.67 (0.58, 0.77)
			>4	310	NR	>4	NB	0.64 (0.48, 0.87)
								British Columbia CS
			Groups	British Columbia N (42,217)	%u	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			<1 hour	39 101	NR		NB	-
			1-2	1892	NR	1-2	NB	0.92 (0.83, 1.03)
			2-4	623	NR	2-4	NR	0.74 (0.61, 0.90)
			× 4<	601	NR	>4	NR	0.70 (0.57, 0.85)
								Nova Scotia CS
			Groups	Nova ScotiaN (17 336)	%u	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			<1 hour	15465	N.	√	N.	-
			1-2	1772	N N	1-2	N.	0.87 (0.77, 0.98)
			2-4	66	N.	2-4	N.	0.67 (0.40, 1.10)
						>4	NR	
	Stoll <i>et al</i> <sup>46</sup> , 2014, Canada	Travel time (hours): <1 1-2 >2	Groups	N (3692)	(%) u	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
				3438	633 (18.41)	.^	-	NR
			1-2	124	27 (21.80)	1-2	1.23 (0.80, 1.91)	NR
			>2	130	26 (20.0)	>2	1.11 (0.71, 1.72)	NR
	Darlinget aft, 2019, Canada Travel time (mins): ≤ 30	a <b>Travel time (mins):</b> ≤ 30 >30	Groups	Z	ın (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			≥ 30	9189	536 (5.83)	≥ 30	1	W.
			>30	2236	98 (4.44)	>30	0.74 (0.59, 0.92)	W.
								1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1

Continued

Table 9 Continued								
	Study, year, country	Exposure groups	Participants (N, n, %)	(%		Findings		
Emergency CS	Grzybowski <i>et al</i> <sup>44</sup> , 2014,	Travel time (hours): <1 >1	Groups	N (59 386)	u (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Canada		~	54,714	9247 (16.99)		1	NB
			<u>`</u>	4672	701 (15.00)	<u>~</u>	0.80 (0.75, 0.86)	NR
Severe perineal trauma (3rd or 4th degree tear)	No studies							
Postpartum haemorrhage	Stolpet al <sup>39</sup> , 2015,	Travel time (mins): <45 >45	Groups	N (54)	u (%)	Groups	Median (range) ml	
	Netherlands		<45	34	Z Z	<45	2,000 (1,100–7,000)	
			>45	20	NR	>45	2,050 (1,000-6,000) (P=0.9)	(6:
	Grzybowski <i>et al</i> <sup>44</sup> , 2013,	Travel time (hrs): <1 >1	Groups	N (59,386)	u (%)	Groups	Crude OR (95%CI)	
	Canada		√	54 714	3064 (5.6)	<u>^</u>	-	
			<u>`</u>	4672	327 (7.0)	<u>~</u>	1.27 (1.13, 1.43)	
Maternal admission to intensive	Stolpet al <sup>39</sup> , 2015,	Travel time (mins): <45 >45	Groups	N (54)	u (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
care unit	Netherlands		≤45	34	1 (2.94)	≤45	-	NB
			>45	20	1 (5.0)	>45	1.74 (0.10, 29.39)	NB
Maternal blood transfusion	Stolp <i>et a</i> <sup>(39</sup> ,2015,	Travel time (mins): <45 >45	Groups	N (54)	u (%)	Groups	Median (range) L	
	Netherlands		≤45	34	≥45	≥45	0 (0-8)	
			>45	20	>45	>45	2 (0-8)	
Neonatal outcomes								
Stillbirth (SB) (overall or intrapartum)	Paranjothy <i>et al</i> ³¹, 2014, UK	Every 15 min increase in travel time (continuous variable)	Groups	N (412,827)	SB n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			All women	412,827	135 (0.03)	All women	1.29 (1.14, 1.47)	1.13 (0.98, 1.30)
			Term births in hospital	387,429	85 (0.02)	Term births only	1.35 (1.16, 1.57)	1.36 (1.17, 1.59)
			Nullips births in hospital	185,419	69 (0.04)	Nullips only	1.33 (1.13, 1.57)	1.21 (1.02, 1.44)
	Combier <i>et al</i> <sup>32</sup> , 2013,	Travel time (mins): ≤ 15 16-30	Groups	N (111,001)	SB n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
	France	31-45 ≥46	≤ 15	70,427	333 (0.47)	≤ 15	-	-
			16-30	31,792	148 (0.47)	16-30	0.98 (0.81, 1.20)	1.16 (0.96, 1.40)
			31-45	8445	50 (0.59)	31-45	1.25 (0.93, 1.69)	1.31 (0.89, 1.93)
			≥46	337	3 (0.89)	≥46	1.89 (0.60, 5.92)	1.90 (0.70, 5.15)
	Grzybowski <i>et al<sup>44</sup>,</i> 2013,	Travel time (hrs): <1 >1	Groups	N (59,386)	SB n (Rate/1000)	Groups	Crude OR (95%CI)	Adjusted OR (95%%CI)
	Calada		√	54,714	274 (5.0)	∵	-	-
			>1	4672	28 (6.0)	>1	1.20 (0.81, 1.77)	NR

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Table 9 Continued								
Outcomes	Study,year, country	Exposure groups	Participants (N, n, %)	(%		Findings		
Neonatal mortality (NM)	Dummer et al <sup>29</sup> , 2004, UK	<b>Travel time (mins):</b> ≤ 17 17-35 > 35	Groups	N (28,7993)	Early NM (0-6 days)n (%)	Early NM (0-6 days) Crude OR (95%CI) Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			≤ 17	NR	1850 (NR)	< 17	NR	_
			17-35	NR	789 (NR)	17-35	NR	0.97 (0.89, 1.06)
			>35	NR	196 (NR)	>35	NR	0.95 (0.81,1.1)
			Groups	N (28,7993)	NM (0-27days)n (%) NM (0-27 days) Groups	NM (0-27 days) Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			< 17	NR	1854 (NR)	< 17	NR	_
			17-35	NR	946 (NR)	17-35	NR	0.96 (0.89, 10.4)
			>35	NR	239 (NR)	>35	NR	0.95 (0.83, 1.09)
			Groups	N (28,7993)	Post NM(28-1yr)n (%)	Post NM (28 days - Crude OR (95%CI) 1yr) Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			s 17	NR	961 (NR)	< 17	NR	_
			17-35	NR	400 (NR)	17-35	NR	0.97 (0.86,10.9)
			>35	NR	98 (NR)	>35	NR	0.95 (0.77, 1.17)
	Paranjothy <i>et af</i> ³¹, 2014, UK	Every 15 min increase in travel time (continuous variable)	Groups	z	Early NM n (%)	Early NM (0-6 days)Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			All women	412,827	609 (0.15)	All women	1.37 (1.31, 1.45)	1.13 (1.07, 1.20)
			Term births only	387,429	177 (0.05)	Term births only	1.02 (0.86, 1.21)	0.97 (0.80, 1.17)
			Nullips only	185,419	303 (0.16)	Nullips only	1.42 (1.33, 1.51)	1.15 (1.06, 1.25)
			Groups	z	Late NM n (%)	Late NM (7-27 days)Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			All women	412,827	251 (0.06)	All women	1.33 (1.23, 1.44)	1.15 (1.05, 1.26)
			Term births only	387,429	77 (0.02)	Term births only	1.24 (1.03, 1.50)	1.34 (1.13, 1.59)
			Nullips only	185,419	116 (0.06)	Nullips only	1.31 (1.15, 1.49)	1.11 (0.97, 1.28)

			Adjusted OR (95% CI)		1.41)	Adjusted OR (95%CI)			Adjusted OR (95%CI)		1.47)	NM (Combined intrapartum & early & late NM up to 28 days)	Adjusted OR (95%CI)		1.12)	1.36)	24 hrs	Adjusted OR (95%CI)		2.02)	l/s	Adjusted OR (95%CI)		1.67)	ays	)R(95%CI)		2.27)				Adjusted OR (95%CI)	
			Adjusted C	-	1.23 (1.07, 1.41)	Adjusted C	-	0.8 (0.4, 1.7)	Adjusted C	-	1.23 (1.04, 1.47)	NM (Combined intrapartum & earl NM up to 28 days)	Adjusted C	-	0.94 (0.79, 1.12)	1.17 (1.02, 1.36)	NM within 24 hrs	Adjusted C	-	1.51 (1.13, 2.02)	NM 0-7 days	Adjusted C	-	1.37 (1.12, 1.67)	NM 8-27 days	Adjusted OR(95%CI)	-	1.24 (0.67, 2.27)				Adjusted C	!
			Crude OR (95% CI)	-	1.23 (1.08, 1.39)	Crude OR (95%CI)	Z Z	N.	Crude OR (95%CI)	Z Z	NR		Crude OR (95%CI)	-	0.97 (0.82, 1.15)	1.22 (1.07, 1.39)		Crude OR (95%CI)	-	1.52 (1.17, 1.97)		Crude OR (95%CI)	-	1.44 (1.20, 1.72)		Crude OR (95%CI)	-	1.30 (0.74, 2.26)	nth, no events			Crude OR (95%CI)	
	Findings		Groups	< 20 mins	≥20 mins	Low risk women	< 20	>20	Low risk women became high risk during labour	<20	≥20		Groups	<15	15-19	> 20		Groups	<20	> 20		Groups	<20	> 20		Groups	<20	> 20	Late NM age <1 month, no events			Groups	
			NM n (%)	789 (0.14)	336 (0.17)	NM n(/1000)63 (0.05/1000)	NR	N.	NM n(/1000)1.9/1000	N.	RN		NM n (1125) (%)	N.	N N	336		NM within 24 hrs n (255) (%)	N N	N N		NM 0-7 dys (523) (%)	Z Z	N.		NM 8-27 dys (58) (%)	RN	NR	NM n (%)	0	0	NM n (Rate/ 1000)	
	(%)	NM (Combined intrapartum and early NM)	N (1 054 342)	558,181	193,745	N (120 896)	NR	NR	N (142,824)	NR	NR		N (751,926)	425,952	132,229	193,745		N (751,926)	558,181	193,745		N (751,926)	558,181	193,745		N (751,926)	558,181	193,745	N (59,386)	54,714	4672	Z	
	Participants (N, n, %)	NM (Combined in	Groups	< 20	≥20	Low-risk women	< 20	>20	Low risk women became high risk during labour	<20	≥20		Groups	<15	15-19	>20		Groups	<20	>20		Groups	<20	≥20		Groups	<20	≥20	Groups	<u>^</u>	<u>`</u>	Groups	
	Exposure groups	Travel time (mins): < 20 mins	≥20 mins									<b>Travel time (mins):</b> <15 15-19 ≥20																	Travel time (hrs): <1 >1			Travel time (mins): Median	
	Study,year, country	Ravelli 2011 <sup>35–37</sup> ,	Netherlands																										Grzybowskiet al <sup>44</sup> , 2013,	Canada		Aoshima et al <sup>47</sup> , 2011,	
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Table 9	Outcomes																																

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Table 9 Continued								
Outcomes	Study,year, country	Exposure groups	Participants (N, n, %)	u, %)		Findings		
Perinatal mortality (PM)	Combier <i>et al</i> <sup>32</sup> , 2013,	<b>Travel time (mins):</b> ≤15 16 -30	Groups	N (110,664)	PM n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
	France	31-45 ≥46	≤15	70,427	452 (0.64)	≤15	-	-
			16-30	31 792	195 (0.61)	16-30	0.96 (0.81, 1.13)	1.08 (0.90, 1.29)
			31-45	8445	59 (0.7.0)	31-45	1.09 (0.83, 1.43)	1.18 (0.86, 1.62)
			>46	337	4 (1.19)	≥46	1.86 (0.69, 5.01)	1.85 (0.66, 5.19)
	Ravelli <i>et al</i> <sup>38</sup> , 2012,	Travel time (mins): <20 ≥20	Groups	N (1,242,725)	PM n (Rate/1000)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
	Netherlands		<20	1,006,607	81 (0.08)	<20	-	-
			≥20	236,118	19 (0.08)	≥ 20	1.53 (1.47,1.50)	1.66 (1.59,1.74)
	Grzybowskiet al <sup>43</sup> , 2011,	<b>Travel time (hrs):</b> <1 1-2 2-4 >4	Groups	N (35,429)	PM n (Rate/1000)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
	Canada		√	32,814	197 (6.0)	<u>^</u>	-	-
			1-2	1359	8 (6.0)	1-2	0.98 (0.48, 1.99)	1.04 (0.48, 2.22)
			2-4	747	4 (5.0)	2-4	0.89 (0.33, 2.40)	0.92 (0.33, 2.53)
			>4	509	9 (18.0)	× ×	2.98 (1.52, 5.85)	3.17 (1.45, 6.95)
			PM (SB & early NM)	JM)				
	Grzybowskiet al <sup>44</sup> , 2013,	Travel time (hrs): <1 >1	Groups	N (59,386)	PM n (Rate/1000)	Groups	Crude OR (95%CI)	Adjusted (OR 95%CI)
	Canada		<u>^</u>	54 714	383 (7.0)	₹	-	NR
			<u>~</u>	4672	37 (8.0)	<u>~</u>	1.54 (1.09, 2.17)	RN
	Grzybowskiet al <sup>45</sup> , 2015,	<b>Travel time (hrs):</b> <1 1-2 2-4 >4				PM (SB & earlyNMAlberta	Nberta	
	Calada		Groups	Alberta (N=34,453)	PM n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			<1 hr	29,906	NR	<u>^</u>	-	-
			1-2	2940	RN R	1-2	N.	1.50 (1.03, 2.18)
			2-4	1297	RN	2-4	NB	1.35 (0.77, 2.38)
			× ×	310	RN	<b>*</b>	NR	1.40 (0.44, 4.39)
								PM (SB & early NM) BC
			Groups	British Columbia (N=42,317)	PM n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			<1 hr	39,101	RN	<u>^</u>	-	-
			1-2	1892	RN	1-2	NR	0.79 (0.43, 1.45)
			2-4	623	RN	2-4	NR	1.33 (0.59, 3.01)
			>4	601	RN	4×	NB	2.84 (1.58, 5.10)
								PM (SB & early NM) Nova Scotia
			Groups	Nova Scotia (N= 17,336)	PM n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			<1 hr	15,465	RN	<u>^</u>	-	-
			1-2	1772	N.	1-2	N.	0.66 (0.38, 1.14)
			2-4	66	N.	2-4	NB	NR
			<b>*</b>	0	NR	>4	NB	NR
								1

Table 9 Continued								
Outcomes	Study,year, country	Exposure groups	Participants (N, n, %)	n, %)		Findings		
	Stoll et al <sup>46</sup> 2014, Canada	<b>Travel time (hrs):</b> <1 1-2 >2	Groups	N (3,692)	PM n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
			▽	3438	15 (0.4)	▽	-	RN
			1-2	124	0	1-2	0.89 (0.05, 14.91)	NB
			>2	130	2 (1.5)	>2	3.57 (0.81, 15.76)	N.
			PM (SB and ear	PM (SB and early neonatal death up to 7 days)				
	Darling et al <sup>44</sup> , 2019 (42),	Travel time (mins): ≤30 >30	Groups	N (10 681)	PM n (%)	Groups	CrudeOR(95% CI)	Adjusted OR (95% CI)
	Canada		≥30	NR	NR	<30	NB	NR
			>30	N.	NB	>30	NR	NR as RR 2.2 (0.67, 7.43)
Infant mortality (IM)	Grzybowski et a/4, 2013,	Travel time (hrs): <1 >1						IM (age 1-12 month)
	Canada		Groups	N (59 386)	IM n (rate/1000)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
				54 714	109(2.0)	▽	-	NB.
			<u>~</u>	4672	14 (3.0)	<u></u>	1.51 (0.86, 2.63)	N.
Born before arrival (BBA)	Combier <i>et al</i> <sup>32</sup> , 2013,	Travel time (mins): <15 15-29	Groups	N (111 001)	BBA n (%)	Groups	Crude OR (95%CI)	Adjusted OR (95%CI)
	France	30-44 >45	<16	70 427	132 (0.19)	<16	-	-
			16-30	31,792	93 (0.29)	16–30	1.56 (1.20, 2.04)	1.73 (1.23, 2.46)
			31-45	8445	29 (0.34)	31–45	1.84 (1.23, 2.75)	1.64 (1.06, 2.54)
			>45	337	0	>45	ı	
	Renesme 2013 (34), France	Travel time (mins): <15 15-29	Groups	CasesN (73) (%)	ControlN (148) (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
		30-44 >45	<15	22 (30.2)	(39.9)	<15	-	-
			15-29	33 (45.2)	64 (43.2)	15–29	1.79 (0.87, 3.68)	1.92 (0.86, 4.96)
			30-44	9 (12.3)	18 (12.2)	30-44	1.68 (0.58, 4.87)	1.10 (0.35, 3.48)
			>45	9 (12.3)	7 (4.7)	>45	5.89 (1.12, 30.89)	6.18 (1.33, 8.65)
	Nguyen <i>et al</i> <sup>33</sup> , 2016,	Travel time (mins): $\leq 20 > 20$	Groups	N (188)	BBA n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	France		>20	94 controls	22 (23.4)	>20 controls	-	NR
			>20	94 cases	27 (28.7)	>20 cases	1.3 (0.7, 2.6)	NR
	Engjom <i>et al</i> <sup>41</sup> , 2017,	<b>Travel time (hrs):</b> <1 1-2 >2	Groups	N (646 898)	BBA n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Norway		-	615 896	3488 (0.60)	-	-	-
			1-2	25,494	844 (3.31)	1-2	6.01 (5.57, 6.49)	NR reported as RR* 5.3 (5.0,5.8)
			~	5508	246 (4.50)	>2	8.21 (7.19, 9.37)	NR as RR* 7.2 (6.3,8.2)
	Grzybowskiet al <sup>43</sup> , 2011,	<b>Travel time (hrs):</b> <1 1-2 2-4 >4	Groups	N (35 429)	BBA n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Canada		▽	32 814	66 (0.20)		-	-
			1-2	1359	31 (2.30)	1-2	11.58 (7.53, 17.81)	6.41(3.69,11.28)
			2-4	747	3 (0.3)	2-4	2.00 (0.63, 6.38)	0.92 (0.22, 3.88)
			>4	206	7 (1.4)	>4	6.96 (3.18, 15.25)	3.63 (1.40, 9.40)
	Grzybowski <i>et al</i> <sup>44</sup> , 2013,	Travel time (hrs): <1 >1	Groups	(98 38e) N	BBA n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
	Callada			54 714	164 (0.3)	▽	-	RN
			<u>~</u>	4672	70 (1.5)	~	5.06 (3.82, 6.70)	NA M

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Outcomes	Study,year, country	Exposure groups	Participants (N, n, %)	n, %)		Findings		
Combined mortality and or	Ravelli 2011 <sup>35–37</sup> ,	Travel time (mins): <15 15-19	Groups	N (751 926)	Event n (4543) (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
Apgar < 4 at 5 mins and or transfer to NICU	Netherlands	>20	<15	425 952	N.	<15	-	-
			15-20	132 229	RN	15–20	0.99 (091, 1.07)	1.11 (1.02, 1.21)
			>20	193 745	NR	>20	1.11 (1.04, 1.19)	1.27 (1.17, 1.38)
Neonatal Unit admission (NNU)	Grzybowski et af <sup>43</sup> , 2011,	Travel time (hrs): <1 1-2 2-4 >4	NICU level 2 ac	NICU level 2 admissions per 1000 births (2001–2004)	4)			NICU 2
	Canada		Groups	N (35 429)	NICU2 n (rate/1000) Groups	Groups	Crude OR (95% CI)	Adjusted OR (95%CI)
			▽	32 814	1082 (33.0)	▽	-	-
			1-2	1359	69 (51.0)	1-2	1.57 (1.22, 2.01)	2.20 (1.59, 3.05)
			2-4	747	8 (11.0)	2-4	0.32 (0.16, 0.64)	0.31 (0.14, 0.65)
			× 4×	909	14 (27.0)	>4	0.83 (0.49, 1.42)	1.07 (0.54, 2.12)
			NICU level 3 pe	NICU level 3 per 1000 births (2001–2004)				NICU 3
			Groups	N (34 920)	NICU3 n (rate/1000) Groups	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			▽	32 814	98 (3.0)	.^	-	NR
			1-2	1359	11 (8.0)	1-2	2.72 (1.46, 5.09)	NR
			2-4	747	4 (5.0)	2-4	1.80 (0.66, 4.90)	NR
			>4	509	2 (4.0)	>4	1.32 (0.32, 5.35)	RN
	Grzybowski et af <sup>44</sup> , 2013,	Travel time (hrs): <1 >1	NICU level 2 (2	NICU level 2 (2001–2002 nd 2006–2007) n=74 697				NICU 2
	Canada		Groups	z	NICU2 n (rate/1000) Groups	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
				54 714	1751 (32.0)	.^	-	NR
			~	4672	154 (33.0)	<u>`</u>	1.03 (0.87, 1.22)	NR
			NICU level 3 (2	NICU level 3 (2001–2002 and 2006–2007) n=74 697				NICU 3
			Groups	z	NICU3 n (rate/1000) Groups	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
				54 714	219 (4.0)	.^	-	NR
			~	4672	28 (6.0)	<u>`</u>	1.50 (1.01, 2.23)	NR
			NICU (2 and 3)					NICU admission
			Groups	N (59 386)	NICU n (rate/1000)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			▽	54 714	1970 (36.0)	<u>^</u>	-	NR
			<u>^</u>	4672	182 (39.0)	>1	1.09 (0.93, 1.27)	NR
	Darlinget alt, 2019, Canada	Darlinget aft, 2019, Canada Travel time (mins): <30 >30	Groups	N (10 687)	NICU n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			> 30	NR	NR	> 30	NR	-
			>30	W Z	NA N	>30	N.	Reported as RR 0.6 (0.44, 0.81)



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utcomes	Study,year, country	Exposure groups	Participants (N, n, %)	n, %)		Findings		
pgar <7 at 5 mins	Darling et al <sup>42</sup> , 2019,	Travel time (mins): ≤30 >30	Groups	N (10 578)	Apgar <7 n (%)	Groups	Crude RR (95% CI)	Adjusted OR (95% CI)
	Canada		≥ 30	NB	NB	≥30	NR	_
			>30	NR	NR	>30	NR	NR as RR 1.02 (0.95, 1.10)
			Nullips	N (4208)	Apgar <7 n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			≥30	3425	51 (1.5)	> 30	-	NR
			>30	621	14 (2.3)	>30	1.53 (0.84, 2.77)	NR
			Mullips	N (7661)	Apgar <7 n (%)	Groups	Crude OR (95% CI)	Adjusted OR (95% CI)
			≥ 30	5764	30 (0.5)	> 30	-	NR
			>30	1615	11 (0.7)	>30	1.31 (0.66, 2.62)	NR
<u> </u>	No studies reported							
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elative risk. sirth weight; HIE, hypoxic-ischaemic encephalopathy; NICU, neonatal intensive care unit; NR, not reported; Nulips, nulipare living more than 2 hours away from an OU compared with women living less than 1 hour from an OU based on an unadjusted analysis (cOR 3.57, 95% CI 0.81 to 15.76). In Darling *et al*, <sup>42</sup> the PM rates were not statistically significantly different for women with a planned home birth and more than 30 min drive from hospital (adjRR 2.2, 95% CI 0.67 to 7.43).

#### Infant mortality (IM)

One Canadian study<sup>43</sup> reported no significant difference in IM rates for women living less than 1-hour travel time to OU compared with more than 1-hour travel time to OU (cOR 1.51, 95% CI 0.86 to 2.63).

#### Born before arrival (BBA)

Six studies reported this outcome, four cohort studies<sup>32</sup> <sup>41</sup> <sup>43</sup> <sup>44</sup> and two case–control studies.<sup>33</sup> <sup>34</sup> Five of the six studies found some association between travel time and BBA, four based on adjusted analyses.

There were three studies conducted in France. <sup>32–34</sup> Combier *et al*, <sup>32</sup> reported that a travel time greater than 15 min was significantly associated with an increased risk of BBA (16–30 min vs <16 min, adjOR 1.73, 95% CI 1.23 to 2.46); (31–45 min vs <16 min, adjOR 1.64, 95% CI 1.06 to 2.54). <sup>32</sup> In a case–control study, <sup>34</sup> the BBA rate increased sixfold when the travel time increased to more than 45 min from home to the OU compared with women who travelled less than 15 min (>45 min vs >15 min, adjOR 6.18 95% CI 1.33 to 28.65). However, in the other case– control study the risk of BBA was not significantly increased in women who travelled for greater than 20 min. <sup>33</sup>

In a study from Norway, <sup>40 41</sup> the risk of BBA increased significantly with longer travel time to the nearest OU from home. Women who travelled more than 2 hours had an eight fold increased risk of BBA compared with women who lived within 1 hour of the nearest OU (>2 hours vs <1 hour, cOR 8.21, 95% CI 7.19 to 9.37). <sup>41</sup>

The studies from Canada<sup>43</sup> <sup>44</sup> found a significant increase in BBA in women living in communities greater than 1-hour travel time from an OU compared with those living less than 1 hour away. In Grzybowski et al, 43 women who lived 1-2 hours from an OU had the highest risk of BBA compared with less than 1 hour (adjOR 6.41, 95% CI 3.69 to 11.28) and women who lived greater than 4 hours away also had an increased risk compared with those living less than 1 hour away (adjOR 3.63, 95% CI 1.40 to 9.40); however, there was no difference between those who lived 2-4 hours from an OU and those living less than 1 hour away (adjOR 0.92, 95% CI 0.22 to 3.88). Gryzbowski et al, 44 found a five-=fold increase in BBA in women who lived more than an hour away from an OU in comparison to women who lived less than an hour away (cOR 5.06, 95% CI 3.82 to 6.70).

#### Neonatal unit (NNU) admission

Three studies from Canada reported on NNU admission.  $^{42-44}$  The two studies from British Columbia  $^{43}$   $^{44}$ 

Outcompage Appg



reported NNU depending on whether the admission was for level 2 care (high dependency) or level 3 care (intensive care). Findings from one of these studies<sup>43</sup> showed NNU level 2 admission increased significantly in babies born to women living more than 1 hour away from an OU compared with less than 1 hour (adjOR 2.20, 95% CI 1.59 to 3.05). For those living 2-4 hours away, level 2 admissions were significantly lower compared with those living less than 1 hour away (adjOR 0.31, 95% CI 0.14 to 0.65). For those living more than 4 hours away, there appeared to be no increase in NNU level 2 admission. For level 3 NNU admission, a significantly increased risk was found for the 1-2 hours category (1-2 hours vs <1 hour, cOR 2.72, 95% CI 1.46 to 5.09). For the other two categories, 2-4 and >4 hours, neither crude nor adjusted analyses showed any significant difference. The number of women in each group decreased with increasing time from an OU. In Grzybowski et al, 44 there was no increased risk of admission to NNU level 2 in babies born to women living more than 1 hour from an OU compared with less than 1 hour, however, admission to NNU level 3 was significantly higher (cOR 1.50, 95% CI 1.01 to 2.23). The third Canadian study from Ontario<sup>42</sup> showed a lower relative risk of NNU admission for planned home births with a travel time greater than 30 min when compared with less than 30 min (adjRR 0.6, 95% CI 0.44 to 0.81).

#### Apgar score

Two studies reported on Apgar Score; one from Canada and one from the Netherlands. The Canadian study, and in significant difference was found for Apgar score <7 at 5 min between women who planned home birth and lived more or less than 30 min away from an OU, either for nulliparous or multiparous subgroups (adjRR 1.02, 95% CI 0.95 to 1.10).

The study from the Netherlands<sup>37</sup> used a composite outcome of mortality and/or Apgar <4 at 5 min and/or transfer to NNU, and showed a small but significant increase in this outcome in women whose travel time to an OU exceeded 15 min (15–20 min vs <15 min, adjOR 1.11, 95% CI 1.02 to 1.21 and  $\geq 20$  min vs <15 min, adjOR 1.27, 95% CI 1.17 to 1.38).

#### Hypoxic-ischaemic encephalopathy (HIE)

No studies reported this outcome.

# DISCUSSION

This review describes studies which have explored the associations between OU closure, distance or travel time to an OU, and maternal and neonatal outcomes. The included studies were conducted in the UK, France, the Netherlands, Norway, Canada and Japan. Many studies were from parts of the world where service configuration varied and the study populations were sometimes dispersed over a large geographical area. The included studies differed in their design, geographical boundaries, outcomes measures used and included a wide

range of travel time/distance thresholds used. In addition, although many studies reported that potential confounders were adjusted for in their analyses, many of the outcomes of interest for this review were crude measures of effect without adjustment. Therefore, comparing these studies with each other was a challenge.

All of these studies were brought together to explore whether women who had to travel longer and further to their planned OU were at increased risk of adverse outcomes. There was one reasonably consistent finding which was that there appeared to be an increased risk of BBA the longer it took to reach the OU. This may have been associated with an increased risk for the baby with a suggestion of an increased risk of perinatal or NM in some studies, however, this effect was not consistent across all the studies. There was also an increase in CS rates following closure of an OU and with shorter travel distance and time, however, it is unclear if the difference was related to the exposure or unmeasured differences in CS rates.

#### Strengths and limitations of the review

This work is the first to synthesise systematically the current evidence relevant to OU closure and the impact of travel time and travel distance on maternal and neonatal outcomes. Rigorous systematic review methodology was applied, including a sensitive search strategy to identify all the relevant literature, and thorough assessment of potential risks of bias. All screening, data extraction and risk of bias assessment were performed independently by at least two reviewers.

The process of selecting studies for inclusion was challenging due to a lack of reporting of some details, for example, it was not always clear which level of maternity services the study referred to, in others, findings related to the impact of travel time and distance were not always presented despite this being described as a study objective.

# INTERPRETATION OF FINDINGS

It is difficult to conclude from this review whether reconfiguration of maternity services, with closure of OUs, resulting in increased travel distances and times for women is unequivocally associated with worse outcomes for the mother or the baby. Assessing the impact of OU closure and prolonged travel time and distance is not straightforward; to isolate the impact of the closure and travel time and distance on maternal and neonatal outcomes we need to fully understand the models of maternity care, transport services, landscape characteristics, women's satisfaction with care and places of birth available to women in that specific geographical area. Understanding how services are delivered to women is vital when assessing the impact of travel distance and time as services may be adapted to meet the challenges for women living in remote areas, for example by transferring women antenatally a few weeks before birth. Some studies found an increase in CS rates with shorter



distance/travel time. Attributing this solely to closure or reconfiguration of services is problematic as simple analytical comparisons of rates before and after changes do not account for underlying time trends. Future studies might want to consider an interrupted time series design as a more appropriate method.

There remains an urgent need to evaluate the impact of changing maternity service provision. The imperative to close and consolidate OUs into larger units is based on a belief that this will improve safety for both mother and baby. If increasing travel times and distances increases risks to mothers and babies, then the postulated benefits of larger OUs could be offset by the harms of the reconfiguration.

Waiting for closure of OUs to prospectively evaluate the impact on the surrounding maternity population will always be challenging. However, exploring the existing impact of distance and travel time from home to an OU may be a reasonable approach to explore what the impact of reconfiguration may be for a proportion of the women in the area served by the OU which would have these parameters increased by closure of one of more local OU(s). Such a study would need to be large to explore the impact of travel time and distance on substantive harms such as mortality for the baby, so will almost certainly need to use routinely collected data to obtain large numbers. Such studies will also need to include vigorous evaluation of confounders, such as maternal characteristics, socioeconomic status and maternal medical history, which are known to influence birth outcomes; controlling for these factors is vital to determine the OU closure impacts. These studies should also collect data at multiple time points after the closure and apply statistical analysis which considers time-varying relationships and the outcomes.

Measurement of travel time and distance from the woman's place of residence to an OU would also need more sophisticated approaches than previously used in many studies; for example the use of web-based route planners and adjustment for travel conditions rather than using straight line distances or relying on self-reports.

Many study designs assume that travel time and distance have a constant effect on outcomes. If local OUs are far away, it is possible that women will modify their behaviour in relation to when they set off for their OU in labour, if they know they have an hour's journey compared with a 20 min journey. The extent to which this will mitigate the effects of longer travel times would not be seen in a study looking at existing travel times and distances.

#### **CONCLUSION**

Given the substantial variation across studies we were unable to draw firm conclusions regarding the association between OU closure, travel distance or time to obstetric services and maternal and neonatal outcomes. There appears to be a consistent association with BBA with increasing distance and travel time to an OU and

a suggestion of increasing risk to the baby. However, few studies have rigorously controlled for potential confounders.

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Acknowledgements Our thanks to Pamela White for contacting NHS Trusts and obtaining papers, Nia Roberts for development of the search strategy and Mark Willett and Fiona Mackie for providingun published data.

#### Collaborators NA.

**Contributors** PB, FA, RSM and JD conceived the research. All authors developed the protocol and RSM developed the search strategy. RSM, CT, AP, FA and JH screened the search results and full papers. RSM, CT, JH, FA and CO assessed the quality of included papers, extracted the data and synthesised the results. RSM and FA drafted the manuscript and all authors agreed the final manuscript.

**Funding** This research is funded by the National Institute for Health Research (NIHR) Policy Research Programme, conducted through the Policy Research Unit in Maternal Health and Care, 108/0001.

**Disclaimer** The views expressed are those of the author(s) and not necessarily those of the NIHR or the Department of Health and Social Care.

Competing interests None declared.

Patient consent for publication Not required.

Provenance and peer review Not commissioned; externally peer reviewed.

**Data availability statement** All data relevant to the study are included in the article or uploaded as online supplemental information. All the data included in this systematic review are in the public domain.

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