

Meta Analysis



Prevalence of preterm birth in Scandinavian countries: a systematic review and meta-analysis

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Abstract

Objectives: As welfare societies, Scandinavian countries share characteristics of equality related to healthcare access, gender, and social services. However, cultural and lifestyle variations create country-specific health differences. This meta-analysis assessed the prevalence of preterm birth (PTB) and its categories in Scandinavian countries.

Methods: A systematic search in key databases of literature published between 1990 and 2021 identified studies of the prevalence of PTB and its categories. Following the use of the Freeman—Tukey double arcsine transformation, a meta-analysis of weighted data was performed using the random-effects model and meta-prop method.

Results: We identified 109 observational studies that involved 86,420,188 live births. The overall pooled prevalence (PP) of PTB was 5.3% (PP = 5.3%, 95% confidence interval [CI] 5.1%, 5.5%). The highest prevalence was in Norway (PP = 6.2%, 95% CI 5.3%, 7.0%), followed by Sweden

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(PP = 5.3%, 95% CI 5.1%, 5.4%), Denmark (PP = 5.2%, 95% CI 4.9%, 5.3%), and Iceland (PP = 5.0%, 95% CI 4.4%, 5.7%). Finland had the lowest PTB rate (PP = 4.9%, 95% CI 4.7%, 5.1%). **Conclusions:** The overall PP of PTB was 5.3%, with small variations among countries (4.9%–6.2%). The highest and lowest PPs of PTB were in Norway and Finland, respectively.

Keywords

Preterm birth, prevalence, Scandinavian country, systematic review, moderate/late preterm birth, very preterm birth, extremely preterm birth

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Introduction

Preterm birth (PTB) is defined by the World Health Organization as live births before 37 completed weeks of gestation. PTB is a major risk factor for neonatal mortality and morbidity and is one of the most important health problems worldwide. Approximately one-third of neonatal deaths are related to PTB. In 2015, Delnord et al. reported that approximately 75% of all neonatal deaths and 60% of all infant deaths in Europe were directly attributed to PTB.^{3,4}

PTB is the main hindrance to the achievement of the United Nations 2015 Millennium Development Goals, which aim to reduce child mortality by two-thirds. Surviving preterm-born infants are at an increased risk of long-term neurodevelopmental and cognitive impairment, chronic diseases, and mortality later in life. 5-7 Additionally, PTB potentially influences health, welfare, and development during adulthood. 8

The rate of PTB varies from 5% to 18% worldwide according to geographic region and income level. 8–13 During the last 20 years, the prevalence of PTB has increased worldwide and is expected to continue rising with the growth in other risk factors including advanced maternal age and the popularity of assisted reproduction among women. 9

Given their high human development indices and developed healthcare systems, Scandinavian countries have some of the lowest rates of PTB worldwide.¹⁴ However, a similar growth trend in the aforementioned risk factors has been reported.^{15,16} Variations exist in the PTB rate between these studies; however, little is known about the rates of PTB categories in these countries. Importantly, the development of effective preventive measures to reduce the incidence of PTB requires a thorough understanding of the epidemiology of this condition.

Therefore, this systemic review and metaanalysis aimed to present the best available peer-reviewed published evidence of the prevalence of PTB and its categories in Scandinavian countries including Norway, Sweden, Denmark, Finland, and Iceland.

Materials and methods

A review protocol was predesigned based on Cochrane's method for this systematic review and meta-analysis, which followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guideline. The study protocol includes a detailed description of the study and has been registered on the PROSPERO database (CRD42023396628). The review objectives were as follows:

 To study the overall pooled prevalence (PP) of PTB and its categories (including extremely, very, and moderate/late PTB) among the general population of pregnant women in Scandinavian countries, which consisted of Norway, Sweden, Denmark, Finland, and Iceland.

 To study the PP of PTB and its categories (including extremely, very, and moderate/late PTB) among the general population of pregnant women in each Scandinavian country.

The following Population, Intervention or Exposure, Comparison, Outcome elements were applied as inclusion criteria in this review:

Population: general population of pregnant women; Intervention: None; Comparator: pregnant women with PTB; Outcome: PTB rate; Study design: Observational.

Eligibility criteria

The inclusion/exclusion criteria were established before the literature search. Studies had to fulfill the following criteria for eligibility: peer-reviewed articles that were published in the English language and that included the general populations in each Scandinavian country (Norway, Sweden, Denmark, Finland, and Iceland); having a population-based study design; presenting the total population size and the number, percentage, incidence, or prevalence of PTBs; and accessible in full-text.

Gray literature and non-original studies including reviews, commentaries, editorials, letters, meeting abstracts, case reports, conference proceedings, governmental or organizational reports, dissertations, theses, books, unpublished data, presentations, or any studies that did not provide accurate and clear data were excluded. Moreover, studies were excluded if their definitions of the outcome differed from the pre-specified definition in our review.

Search strategy

A comprehensive computerized literature search was independently performed in the

PubMed (including MEDLINE), Scopus, and Web of Science databases by two reviewers (SB-G and MV) and covered the period between 1 January 1990 and 1 February 2022. To maximize the sensitivity of the search, free-text and medical subject heading terms were pilot-tested and were used to search the electronic databases. Furthermore, a manual search of the reference lists of selected studies and other relevant reviews was conducted to maximize the identification of eligible studies. The following keywords were used alone or in combination during the search process: (premature birth OR preterm birth OR premature labor OR preterm labor OR premature labor preterm labor OR adverse pregnancy outcome OR adverse neonatal outcome OR pregnancy complication) AND (prevalence OR incidence OR epidemiology OR rate) AND (Scandinavia OR Nordic OR Denmark OR OR Sweden OR Finland OR Iceland OR Finnish OR Danish OR Nordic OR Swedish). The search string that was used in the PubMed search is presented as an example in Supplementary Table 1.

Study selection and data extraction

Data were independently extracted by two review authors (SB-G and RB-Y) based on the eligibility criteria. When a study separately provided data for the various categories of PTB, the relevant datasets were used for subgroup analyses. Any disagreement between the two reviewers who were responsible for data extraction was resolved through discussion. In the case of missing data or ambiguities in the study design or trial conduct, the study authors were contacted via email and were asked for additional information; in case of no response, the study was excluded from the review. To ensure data collection accuracy before the meta-analysis and prevent bias in data extraction and entry, the data extraction process was double-checked by the two reviewers.

Outcome measures

The primary outcomes of interest were the PP of PTB overall and for the individual countries. The secondary outcome was the pooled risk by PTB category, which included extremely PTB, very PTB, and moderate/late PTB. PTB was defined as any live birth occurring between 22 to <37 weeks of gestation. PTB was further subcategorized into extremely PTB (22 weeks of gestation or birth weight below 500 g to <28 weeks), very PTB (28 to <32 weeks), and moderate/late PTB (32 to <37 weeks).^{2,18}

Quality appraisal

The quality of the included studies was critically appraised for methodological structure and the presentation of results. The Newcastle-Ottawa scale was used to assess the quality of each study. This scale is suitable for the evaluation of non-randomized trials in meta-analyses and evaluates studies based on three criteria: participant selection, comparability of study groups, and assessment of outcome or exposure. A study can be awarded a maximum of four stars for the selection category, a maximum of two stars for the comparability category, and a maximum of three stars for the outcome/exposure category. 19 Studies with scores above 6, 4 to 6, and less than 4 are considered of high, moderate, and low quality, respectively. In addition, the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach was used to downgrade the certainty of evidence by considering study limitations, inconsistency, imprecision, publication bias, and indirectness. The GRADE approach categorizes the quality of evidence into four levels: high, moderate, low, and very low. In the GRADE rating system, the initial quality of evidence for observational studies is low. The quality of evidence can be downgraded from a higher to a lower level based on the risk of bias, inconsistency, indirectness, imprecision, and publication bias. Conversely, the quality of evidence can be upgraded from a lower to a higher level based on a large magnitude of effect, the evidence of a dose–response, and the assessment of all plausible residual confounding.²⁰

Statistical analysis

Heterogeneity was evaluated using the chisquare test, and a p-value >0.05 was interpreted as homogeneity. Heterogeneous and non-heterogeneous results were analyzed using the fixed effects and random-effects inverse variance models to calculate the pooled effect. Publication bias was assessed using the Begg's and Egger's tests. A p-value >0.05 was interpreted as indicating no significant publication bias in the meta-analysis. Funnel plots for each outcome were drawn to illustrate publication bias; an asymmetric plot was interpreted as indicating the presence of publication bias.

The trim and fill method was used for correction in case of significant publication results. The meta-prop method was used for the pooled estimation of PTB prevalence after the Freeman-Tukey Double Arcsine Transformation was used to stabilize variances. Forest plots for each outcome and subgroup of the country of origin were also performed. Sensitivity analyses were conducted to identify studies that influenced the estimation of the overall meta-analysis summary. A graph was drawn of the results of an influence analysis in which the metaanalysis was re-estimated by omitting one study at a time. A p-value <0.05 was considered significant, and all statistical analyses were performed using STATA software (version 14; StataCorp Inc., College Station, TX, USA).

Results

Identification of studies and quality appraisal

The literature search yielded 1136 studies, of which 329 studies were further evaluated by retrieving their full texts; 193 studies were subsequently excluded. Ultimately, 109 population-based eligible studies were included in the review. Together, the studies offered extractable data that encompassed 86,420,188 live births for the meta-analysis. A flow diagram of this process is presented in Figure 1.

The characteristics and prevalences of PTB for each study are provided in Table 1. The studies were published between 1991 and 2021.

Sweden had the highest number of eligible studies (44 studies), ^{14,21-63} followed by Denmark (29 studies), ^{15,33,52,64-89} Norway (26 studies), ^{16,33,52,54,59,74,84,90-108} Finland (19 studies), ^{16,33,52,71,108-122} and Iceland (four studies). ^{33,123-125} Given that some studies presented information on more than one country, ^{16,33,52,54,71,74,84,108} the overall sum was greater than 109.Details of the quality assessment of the included studies are presented in Supplementary

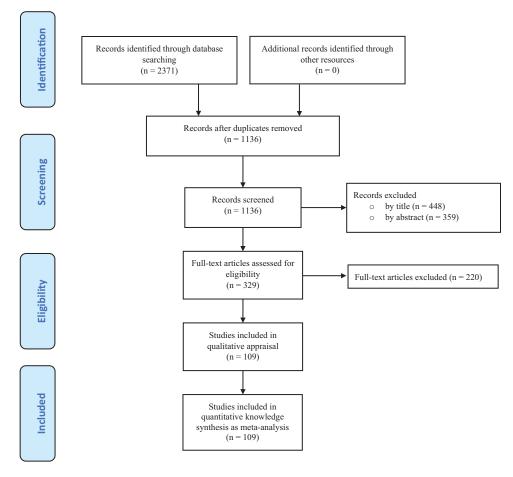


Figure 1. Flow diagram of the search strategy and study selection.

 Table I. Summary of studies that assessed the prevalence of preterm births.

				Prevalence of preterm births (%)	of preterm	h births (%)	
			Sample	Extremely	Very	Moderate/	Total
Author, year	Data source	Research year	Size (n)	PTB	PTB	late PTB	PTB
Norway							
Bjellmo et al., 2020	MBRN	1996–2015	294,598	ΣΖ	ΣΖ	ΣΖ	4.04
Deb-Rinker et al., 2015	MBRN	1995–2005	642,139	0.31	0.70	5.59	9.60
Ebbing et al., 2020	MBRN	1999–2014	918,933	ΣΖ	Σ	ΣΖ	5.89
Ebbing et al., 2013	MBRN	1999–2009	623,478	ΣΖ	Σ	4.60	90.9
Heggland et al., 2011	MBRN	1967–2006	1,130,251	ΣΖ	Σ	ΣΖ	13.19
Heitmann et al., 2013	MBRN and MoBa * study	1999–2008	68,522	ΣΖ	Σ	ΣΖ	5.16
Henriksen et al., 2014	MBRN	1999–2008	76,870	ΣΖ	Σ	ΣΖ	4.71
Huitfeldt et al., 2020	MBRN and MoBa study	1999–2008	82,038	ΣΖ	Σ	ΣΖ	4.29
Kvalvik et al., 2020	MBRN	1999–2015	302,192	ΣΖ	Σ	ΣΖ	5.95
Oftedal et al., 2016	MBRN	1999–2009	494,073	ΣΖ	Σ	ΣΖ	4.24
Panasevich et al., 2016	MoBa study	1999–2008	17,533	ΣΖ	Σ	ΣΖ	4.67
Poulsen et al., 2019	MoBa study	ΣΖ	78,267	ΣΖ	Σ	ΣΖ	4.90
Richards et al., 2016	MBRN	2006–2015	468,954	ΣΖ	ΣΖ	ΣΖ	5.29
Skorpen et al., 2018	MBRN	2006–2015	498,849	ΣΖ	ΣΖ	ΣΖ	5.43
Swamy et al., 2008	MBRN	1967–1988	1,167,506	0.39	0.81	3.97	5.17
Thompson et al., 2006	MBRN	1980–1998	943,193	ΣΖ	Σ	ΣΖ	4.81
Trønnes et al., 2014	MBRN	1967–2003	1,974,226	ΣΖ	Σ	4.91	5.92
Vandraas et al., 2013	MBRN	1967–2009	2,270,363	ΣΖ	Σ	ΣΖ	6.41
Vik et al., 2020	MBRN	1990–2016	1,083,099	ΣΖ	ΣΖ	4.43	5.36
Waldenström et al., 2014	MBRN	1990–2010	311,620	ΣΖ	ΣΖ	5.32	6.50
Zeitlin, et al. 2013	MBRN	1996, 2000, 2004, 2008	60,744	ΣΖ	ΣΖ	ΣΖ	6.70
Skulstad et al. 2017	MBRN	1967–2010	1,629	ΣΖ	ΣΖ	6.20	7.98
Naimy et al. 2013	MBRN	1986–2005	1,062,744	ΣΖ	Σ	ΣΖ	12.70
Villadsen et al. 2010	MBRN	1990–2005	675,387	ΣΖ	ΣΖ	ΣΖ	6.90
Halldorsson et al. 2021	MoBa study	1999–2008	66,302	ΣΖ	ΣΖ	ΣΖ	4.92
Tingleff et al. 2021	MBRN	1999–2014	910,752	0.38	1.20	3.76.	5.33

(continued)

Table I. Continued.

				Prevalence of preterm births (%)	f preterm	births (%)	
Author, year	Data source	Research year	Sample Size (n)	Extremely PTB	Very PTB	Moderate/ late PTB	Total PTB
Sweden							
Bodén et al. 2012	SMBR	2005–2009	332,137	ΣΖ	ΣΖ	ΣΖ	4.77
Brander et al. 2016	SMBR	1973–1996	2,421,284	ΣΖ	ΣΖ	4.38	2.00
Buchmayer et al. 2004	SMBR	1987–2000	601,883	0.24	09.0	5.42	6.26
Calling et al. 2013	SHDR and SCDR	1984–2006	2,297,134	ΣΖ	ΣΖ	ΣΖ	5.91
Cnattingius et al. 1992	SMBR	1983–1987	173,715	ΣΖ	ΣΖ	5.40	ΣΖ
Cnattingius et al. 2013	SMBR	1973–2007	1,857,822	0.23	0.47	4.36	5.03
Crump et al. 2019	SMBR	1973–2014	4,186,615	0.19	ΣΖ	ΣΖ	4.95
Crump et al. 2019	SMBR	1973–1995	2,235,012	60.0	ΣΖ	ΣΖ	7.76
Crump et al., 2019	SMBR	1973–2014	4,186,615	0.19	ΣΖ	ΣΖ	4.95
Crump et al., 2011	SMBR	1973–1979	908'629	ΣΖ	ΣΖ	1.4	4.44
Crump et al., 2010	SMBR	1973–1979	635,933	0.04	0.53	3.96	4.53
Crump et al., 2011	SMBR	1973–1979	636,552	0.02	0.49	3.9	4.45
Dahlin et al., 2016	SMBR	1999–2012	1,371,247	0.23	0.45	4.26	4.90
Deb-Rinker et al., 2015	SMBR	1995–2005	1,029,849	0.29	0.64	5.35	6.28
Derraik et al., 2016	SMBR	1991–2009	192,432	ΣΖ	ΣΖ	ΣΖ	9.00
Fransson et al., 2011	SMBR	1999–2000	2,904	ΣΖ	ΣΖ	ΣΖ	5.27
Granfors et al., 2019	Stockholm-Gotland study	2008–2014	74,087	ΣΖ	ΣΖ	ΣΖ	5.98
Gudmundsson et al., 1997	Malmo Hospital database	1990–1992	7,056	ΣΖ	ΣΖ	ΣΖ	7.03
Hesselman et al., 2019	SPR	2013–2017	218,030	0.25	0.43	4.15	4.83
Holmgren et al., 2001	SMBR	9661-1661	66,646	0.30	.059	5.24	6.13
Höglund et al., 2012	SMBR	1999–2007	340,950	0.50	ΣΖ	ΣΖ	9.10
Kornfeld et al., 1997	SMBR	1991–1992	239,773	ΣΖ	ΣΖ	4.30	5.34
Koupilová et al., 1998	SMBR	1661-6861	239,773	ΣΖ	ΣΖ	ΣΖ	5.40
Kyrklund-Blomberg et al., 1998	SMBR	1991–1993	311,977	ΣΖ	ΣΖ	4.19	5.17
Li et al., 2010	National research	1990–2004	816,743	ΣΖ	ΣΖ	ΣΖ	5.38
	database, WomMed II						
	database						

(continued)

Table I. Continued.

				Prevalence of preterm births (%)	of preterm	births (%)	
Author, year	Data source	Research year	Sample Size (n)	Extremely PTB	Very PTB	Moderate/ late PTB	Total PTB
Li et al., 2013	National research database, WomMed II	1982–2006	1,874,567	ΣΖ	ΣΖ	ΣΖ	4.90
-	garacasa	7	717	2	2	2	0
Liu et al., 2019	SPK SPK	7107-4-017	254,973	ΣΖ	ΣΖ	ΣΖ	4.90
Liu et al., 2018	SPR	1973–1977	175,821	ΣΖ	Σ	ΣΖ	3.55
Morken et al., 2005	SPR	1991–2001	1,796,769	ΣΖ	ΣΖ	ΣΖ	5.05
Murray et al., 2019	SPR	1998–2013	1,335,802	ΣΖ	ΣΖ	ΣΖ	4.02
Nager et al., 2008	SPR	1975–2003	1,133,368	ΣΖ	ΣΖ	ΣΖ	92.9
Persson et al., 2014	SPR	1998–2007	1/98/16	ΣΖ	ΣΖ	4.33	5.07
Rasmussen et al., 1995	SPR	1978–1990	1,270,407	ΣΖ	ΣΖ	ΣΖ	5.50
Richards et al., 2016	SPR	2006–2012	737,754	ΣΖ	ΣΖ	ΣΖ	4.79
Stokkeland et al., 2017	SPR	2001–2011	1,093,969	ΣΖ	ΣΖ	4.22	4.90
Waldenström et al., 2014	SPR	1990–2010	644,184	ΣΖ	ΣΖ	5.30	6.27
Wikström et al., 2010	SPR	1999–2006	569,951	ΣΖ	ΣΖ	4.15	4.75
Essén et al., 2000	SMBR	1990–1995	10,784	ΣΖ	ΣΖ	ΣΖ	5.76
Juárez et al., 2019	SPR	1997–2012	872,049	ΣΖ	ΣΖ	ΣΖ	4.24
Khanolkar et al., 2015	SPR	1982–2002	1,435,286	ΣΖ	ΣΖ	4.33	4.88
Villadsen et al., 2010	SPR	1990–2005	1,344,237	ΣΖ	ΣΖ	ΣΖ	6.40
Crump et al., 2019	SPR	1973–1997	2,566,699	0.21	1.29	ΣΖ	5.81
Hesselman et al., 2020	SPR	2007–2014	377,381	ΣΖ	ΣΖ	4.93	5.86
Mantel et al., 2020	SPR	2003–2014	1,232,863	ΣΖ	ΣΖ	ΣΖ	4.81
Finland							
Deb-Rinker et al., 2015	FMBR	1995–2005	635,910	0.15	0.55	4.55	5.26
Goisis et al., 2017	FMBR	1987–2000	124,098	ΣΖ	ΣΖ	ΣΖ	3.70
Hirvonen et al., 2017	FMBR	1991–2008	1,033,349	ΣΖ	ΣΖ	ΣΖ	3.35
Kc et al., 2017	FMBR	1996–2013	419,879	0.32	ΣΖ	ΣΖ	4.65
Klemetti et al., 2014-1	FMBR	1661	24,765	ΣΖ	ΣΖ	ΣΖ	5.80
Klemetti et al., 2014-1	FMBR	2008	23,511	ΣΖ	ΣΖ	ΣΖ	9.10

(continued)

Table I. Continued.

				Prevalence of preterm births (%)	of preterm	births (%)	
Author, year	Data source	Research year	Sample Size (n)	Extremely PTB	Very PTB	Moderate/ late PTB	Total PTB
Mortensen et al., 2020	FMBR	1980–2008	38,0291	ΣΖ	ΣΖ	ΣΖ	3.52
Olsën et al., 1995	NFBC	1985–1986	8,888	ΣΖ	ΣΖ	ΣΖ	4.78
Richards et al., 2016	FMBR	2006–2015	571,937	ΣΖ	Σ	ΣΖ	4.35
Räisänen et al., 2013	FMBR	1987–2010	1,390,742	032	0.45	3.90	4.66
Räisänen et al., 2014	FMBR	2005–2010	345,952	ΣΖ	Σ	ΣΖ	4.51
Räisänen et al., 2013	FMBR	2006–2010	291,004	ΣΖ	Σ	ΣΖ	Σ
Tingleff et al., 2021	FMBR	1999–2004	91,0752	0.38	1.20	3.76	5.33
Zeitlin et al., 2013	FMBR	1996, 2000, 2004, 2008	59,486	ΣΖ	Σ	ΣΖ	5.50
Jakobsson et al., 2008	FMBR	1987–2005	1,137,515	ΣΖ	Σ	ΣΖ	5.19
Ronkainen et al., 2019	NFBC	9961	18,953	ΣΖ	Σ	ΣΖ	5.29
Ronkainen et al., 2019	NFBC	9861	350,548	ΣΖ	Σ	ΣΖ	4.48
Malin et al., 2009	FMBR	1999–2001	158,469	ΣΖ	ΣΖ	ΣΖ	4.80
Rumrich, et al., 2020	FMBR	1991–2016	1,376,778	ΣΖ	Σ	ΣΖ	4.27
Strang-Karlsson, et al., 2021	FMBR	1987–1990	228,616	0.23	0.43	4.25	16.4
Toijonen, et al., 2020	FMBR	2004–2014	737,788	ΣΖ	0.62	3.64	4.54
Denmark							
Andreasen, et al., 2021	DNPR	2012–2015	66,059	91.0	0.32	3.59	4.07
Auger, et al., 2013	DMBR	1981–2008	1,674,308	ΣΖ	ΣΖ	ΣΖ	4.72
Bilsteen, et al., 2018	DMBR	2000–2009	649,905	0.28	0.72	5.63	6.64
Bilsteen et al., 2018	DMBR	1982–1986	22,8030	0.05	0.64	3.36	4.06
Deb-Rinker et al., 2015	DMBR	1995–2005	728,651	0.27	89.0	5.39	6.33
Garioud et al., 2020	DMBR	1992–2007	1,043,411	91.0	0.63	5.27	6.07
Hedermann et al., 2021	DNSB	2015–2020	31,180	0.19	0.57	4.27	5.02
Jepsen et al., 2018	DMBR	2013–2016	10,436	ΣΖ	Σ	ΣΖ	6.12
Kristensen et al., 1995	DMBR	1982–1987	13,967	ΣΖ	ΣΖ	4.12	4.95
Micali et al., 2016	DMBR	1996–2002	77,552	ΣΖ	ΣΖ	ΣΖ	4.71
Mortensen et al., 2020	DMBR	1980–2008	625,179	ΣΖ	Σ	ΣΖ	4.44
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				Prevalence of preterm births (%)	of preterm	births (%)	
Author, year	Data source	Research year	Sample Size (n)	Extremely PTB	Very PTB	Moderate/ late PTB	Total PTB
Pasternak et al., 2012	DMBR	2009–2010	23,262	ΣΖ	ΣΖ	ΣΖ	4.91
Pedersen et al., 2012	DMBR	1978–2007	1,557,944	ΣΖ	Σ	3.70	4.50
Poulsen et al., 2019	DMBR	ΣΖ	77,020	ΣΖ	Σ	ΣΖ	4.80
Ratanajamit et al., 2003	DMBR	1991–2001	65,977	ΣΖ	Σ	ΣΖ	4.43
Rebordosa et al., 2009	DNSB	1996–2003	98,140	ΣΖ	Σ	ΣΖ	4.26
Richards et al., 2016	DMBR	2006–2010	305,947	ΣΖ	Σ	ΣΖ	16.4
Wüst et al., 2015	DMBR	1998–2003	366,367	ΣΖ	Σ	ΣΖ	4.90
Albertsen et al., 2004	DNBC	1997–2000	40,892	ΣΖ	Σ	ΣΖ	4.60
Juhl et al., 2008	DNBC	1996–2002	87,232	0.38	0.50	4.02	16.4
Morgen et al., 2008	DNBC	1996–2002	75,890	0.39	0.52	4.10	2.00
Langhoff-Roos et al., 2006	DMBR	1995	67,840	0.22	0.55	4.40	5.17
Langhoff-Roos et al., 2006	DMBR	9661	65,635	0.27	19.0	4.85	5.73
Langhoff-Roos et al., 2006	DMBR	1997	65,783	0.21	0.58	4.39	5.18
Langhoff-Roos et al., 2006	DMBR	8661	64,578	0.20	19.0	4.57	5.37
Langhoff-Roos et al., 2006	DMBR	6661	64,578	0.22	19.0	4.54	5.38
Langhoff-Roos et al., 2006	DMBR	2000	65,541	0.21	19.0	4.76	5.58
Langhoff-Roos et al., 2006	DMBR	2001	63,814	0.26	0.59	5.09	5.94
Langhoff-Roos et al., 2006	DMBR	2002	62,515	0.24	0.63	5.18	90.9
Langhoff-Roos et al., 2006	DMBR	2003	63,038	0.31	99.0	5.26	6.23
Langhoff-Roos et al., 2006	DMBR	2004	62,814	0.31	0.67	5.36	6.33
Runge et al., 2013	DNBC	1996–2002	62,803	0.18	99.0	3.73	4.57
Yu et al., 2019	DMBR	1981–2015	1,994,618	ΣΖ	Σ	ΣΖ	4.26
Halldorsson et al., 2021	DNBC	1996–2002	60,141	ΣΖ	ΣΖ	ΣΖ	4.60
Koch et al., 2021	DMBR	1997–2010	918,280	ΣΖ	Σ	5.45	6.47
Matthiesen et al., 2021	DMBR	1997–2013	1,092,740	ΣΖ	Σ	5.26	92.9
Mørk et al., 2021	DMBR	1997–2016	1,199,610	ΣΖ	ΣΖ	4.38	5.26
Rasmussen et al., 2020	DMBR	1997–2015	1,202,870	ΣΖ	Σ	3.94	4.68

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				Prevalence of preterm births (%)	of preterm	births (%)	
Author, year	Data source	Research year	Sample Size (n)	Extremely PTB	Very PTB	Moderate/ late PTB	Total PTB
Iceland							
Deb-Rinker et al., 2015	Hospital registry	1995–2005	46,309	0.28	0.57	4.53	5.38
Eiríksdóttir et al., 2013	IMBR	2006–2009	16,271	ΣΖ	ΣΖ	ΣΖ	4.46
Grétarsdóttir et al., 2020	IMBR	1997–2016	87,076	0.27	ΣΖ	ΣΖ	5.73
Haraldsdottir et al., 2015	IMBR	2000–2009	40,982	ΣΖ	ΣΖ	ΣΖ	4.46

PTB: preterm birth, MoBa: Norwegian Mother and Child Cohort, MBRN: Medical Birth Registry of Norway, SMBR: Swedish Medical Birth Register, SHDR: Swedish Hospital Discharge Register, SCDR: Swedish Cause of Death Register, SPR: Swedish Pregnancy Register, FMBR: Finnish Medical Birth Register, NFBC: Northern Finland Birth Cohorts, DNPR: Danish National Patient Registry, DMBR: Danish Medical Birth Registry, DNSB: Danish National Birth Cohort, DNSB: Danish Neonatal Screening Biobank, IMBR: Birth Registry, DNBC: Danish National Birth Cohort, NM: not mentioned. celandic Medical

Tables 2 and 3. Given that narrow inclusion criteria were used in this review, all studies were of moderate or high quality (Supplementary Tables 2 and 3). The overall confidence in the body of evidence in this review was graded as 'very low' (Supplementary Table 4).

Meta-analysis

The PP, heterogeneity, and publication bias results are presented in Supplementary Table 5. The overall PP of PTB in the Scandinavian countries regardless of PTB category was 5.3% (95% confidence interval [CI] 5.1%, 5.5%; 4,737,106 events in 86,420,188 participants). However, in various sub-categories, the prevalences of PTB were 4.5% (95% CI 4.4%, 4.7%; 1,537,502 events in 34,517,389 participants) for moderate/late PTB, 0.6% (95% CI 0.5%, 0.7%; 140,806 events in 19,305,652 participants) for very PTB, and 0.2% (95% CI 0.2%, 0.3%; 67,509 events in 30,024,011 participants) for extremely PTB (Supplementary Table 5).

The results of the PP of PTB and its categories in each Scandinavian country are presented in Figures 2–6. The highest prevalence of PTB was observed in Norway (overall PP=6.2%, 95% CI 5.3%, 7.0%), followed by Sweden (overall PP=5.3%, 95% CI 5.1%, 5.4%), Denmark (overall PP=5.2%, 95% CI 4.9%, 5.3%), and Iceland (overall PP=5.0%, 95% CI 4.4%, 5.7%). The lowest prevalence was observed in Finland (overall PP=4.9%, 95% CI 4.7%, 5.1%; Figures 2 and 3).

A subgroup analysis was performed by PTB category across the Scandinavian countries (Figures 4–6). The same approximate trend was observed in the prevalence of the PTB categories in these countries. The pooled estimates of moderate/late PTB were 4.9% (95% CI 4.6%, 5.2%) in Norway, 4.6% (95% CI 4.4%, 4.9%) in Sweden, 4.6% (95% CI 4.2%, 4.9%)

Scandinavian countries (109 studies, 86,420,188 participants):

Total PTB prevalence: 5.3% (5.1%, 5.5%) Moderate/late PTB prevalence: 4.5% (4.4%, 4.7%) Very PTB prevalence: 0.6% (0.5%, 0.7%) Extremely PTB prevalence: 0.2% (0.2%, 0.3%)

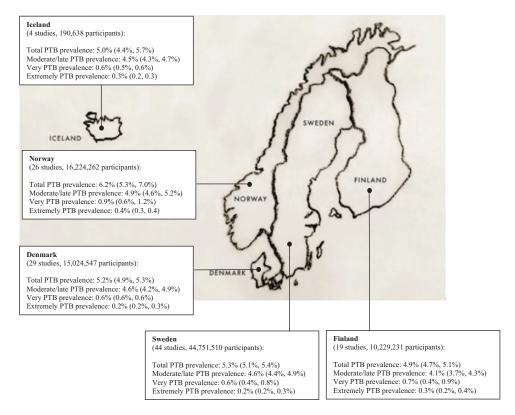


Figure 2. The prevalence of preterm birth (PTB) and its categories in Scandinavian countries.

in Denmark, 4.5% (95% CI 4.3%, 4.7%) in Iceland, and 4.1% (95% CI 3.7%, 4.3%) in Finland (Figure 4). The highest very PTB (0.9%, 95% CI 0.6%, 1.2%) and extremely PTB (0.4%, 95% CI 0.3%, 0.4%) prevalences were reported in Norway (Figures 5 and 6).

The publication bias results are presented in Figure 7 and Supplementary Table 5. The results of Begg's test indicated no substantial publication bias among various outcomes (all P > 0.05). The funnel plot showed some scatter for the moderate/late PTB outcome that may have been

caused by outliers; however, the results remained unchanged after correction using the trim and fill method (4.2%, 95% CI 4.0, 4.4). The scatter observed in the other funnel plot may have been caused by the observed heterogeneity of the included studies, fitting with the assumptions of the random-effects model. Therefore, no evidence suggested that the analysis was affected by publication bias.

The heterogeneity results indicated substantial heterogeneity between populations; most of the heterogeneity may be attributable to maternal age, body mass index,

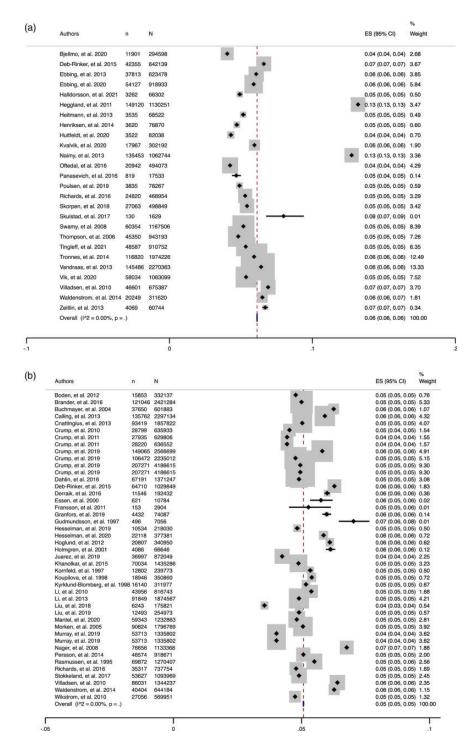


Figure 3. Forest plot of pooled prevalence of preterm birth in each Scandinavian country (a. Norway, b. Sweden, c. Finland, d. Denmark, and e. Iceland).

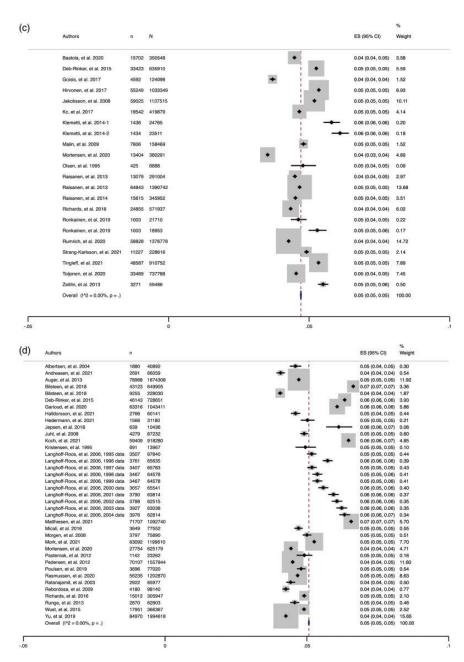


Figure 3. Continued.

or ethnicity. Nevertheless, sensitivity analysis showed that no single study fundamentally changed the PP of any outcome (Supplementary Figure 1).

Discussion

This systematic review and meta-analysis of population-based studies provided data on the prevalence of PTB and its categories in

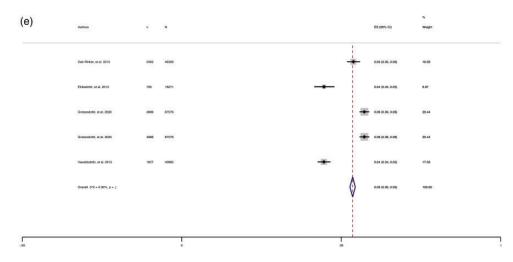


Figure 3. Continued.

Scandinavian countries. The overall PP of PTB in these countries was 5.3%, with small variations between countries that ranged from 4.9% to 6.2%. The highest and lowest PPs of PTB were observed in Norway and Finland, respectively.

Scandinavian countries have a strong tradition of collaboration and share features of policy and welfare systems and social security including free education at all levels, a social safety network for people with reduced health and work capacities, and a universal and predominantly publicly financed healthcare system. 126 In general, these countries are highly ranked on a range of sociodemographic and health-relevant measures such as the United Nations Human Development Indices and indicators, and gender equality, educational attainment, and labor force participation measures. 126

The issue of PTB is one of the major growing global health concerns. The United Nations Sustainable Development Goals aim to end all preventable deaths of newborns and children aged under 5 years by 2030. 127

The overall PTB rate was lower in the Scandinavian countries than in the global population, with the lowest rate in the

Finnish population. Although the worldwide prevalence of PTB is unknown due to the lack of data in many countries, the estimated prevalence of PTB in 184 lower-income countries based on 2010 data indicated a PTB of 11%. ¹²⁸ Additionally, major variations in PTB rates are observed by geographic region and income level. ¹⁰ Nearly 90% of all PTBs are reported to occur in low- and middle-income countries, particularly in sub-Saharan Africa. ¹

Nevertheless, Scandinavian countries provide access to quality healthcare—particularly prenatal care—which leads to the improvement of maternal and neonatal outcomes and the prevention of premature mortality. Moreover, it has been well documented that maternal omega-3 consumption during pregnancy, which is common in these countries, can significantly decrease the risk of PTB. 130

Although the general prevalence of PTB in Scandinavian countries is low, some variations exist across countries. For example, according to our review, Norway and Finland have the highest and lowest prevalence rates of PTB. Even though the Scandinavians share common characteristics that build on fundamental ideas of

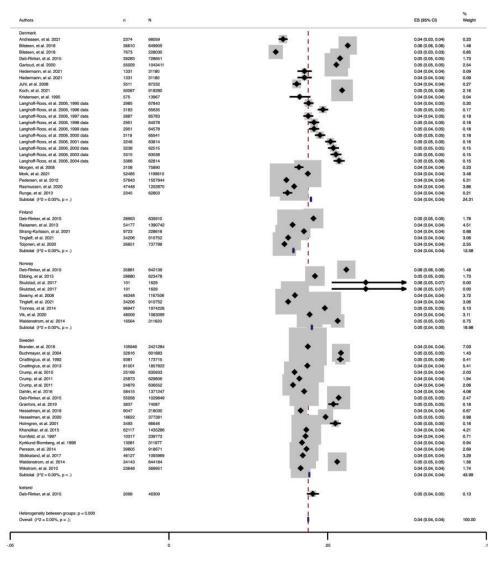


Figure 4. Forest plot of pooled prevalence of moderate/late preterm birth in each Scandinavian country.

equality related to health care access, gender, and social services, notable differences exist between the countries concerning history, societal development, immigration population, culture, and lifestyle, which may affect the risk of PTB. 8,126,131 These differences may also be responsible for variations in the rate of PTB among the countries.

This meta-analysis further provided details of PTB categories based on the gestational age subgroups. These details allow the identification of high-risk populations and the development and implementation of prevention strategies. In addition, PTB classification facilitates the surveillance of PTBs and allows standardized local and international data comparisons. According to our

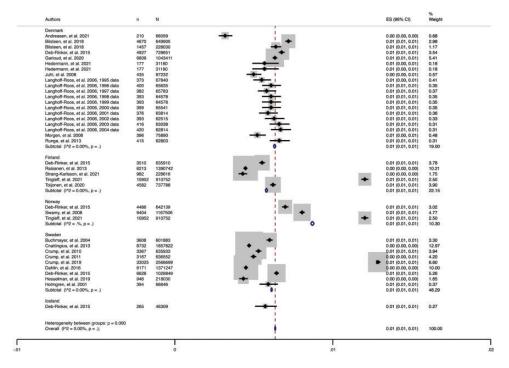


Figure 5. Forest plot of pooled prevalence of very preterm birth in each Scandinavian country.

review, among all live births, the rate of moderate/late preterm births—the common form of PTB in Scandinavian countries—ranged from 4.1% to 4.9%. Only roughly 0.2% fall into the extremely PTB category and the other 0.6% were born at 28 to < 32 weeks of gestation. Our findings are supported by the common notion of PTB found in the international literature. ¹³²

Notably, the etiopathogenesis of PTB is multifactorial and characterized by various clinical presentations. Women with singleton pregnancies without a history of a previous PTB have a low risk of PTB. However, previous PTB, a short cervix, or multiple gestation increase the risk of PTB. A recent study defined a generalized method for developing an evidence-based holistic risk assessment for PTB based on relevant and essential factors including maternal history and cervical length

ultrasound measurement by the use of a machine learning statistical approach for use in clinical practice.¹³³ This screening approach can contribute to the worldwide reduction of the prevalence of PTB.

In summary, this study expands on previous single studies in Scandinavian countries by including the prevalence of PTB and its categories. Our review results are based on population-based data that were extracted from high-quality data registries, thus ensuring the coverage of all available literature. The standardized study selection data extraction processes employed narrow inclusion criteria led to the inclusion of high-quality papers, ensured the validity and reliability of the review results and meta-analysis, and added valuable information to the current international knowledge of PTB.

The main limitation of this review may be the lack of data about potential risk

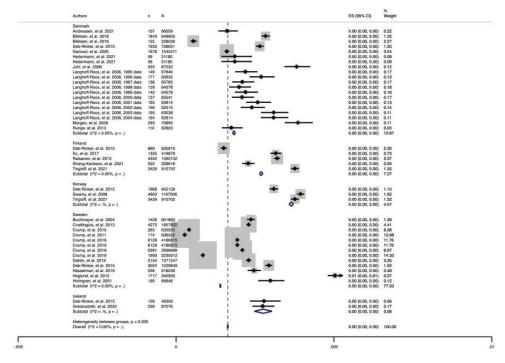


Figure 6. Forest plot of pooled prevalence of extremely preterm birth in each Scandinavian country.

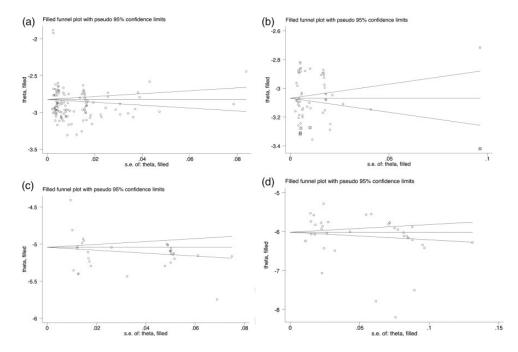


Figure 7. Funnel plot for the assessment of publication bias: a. Total preterm birth, b. Moderate/late preterm birth, c. Very preterm birth, and d. Extremely preterm birth.

factors in the included studies, which are needed to improve our knowledge of PTB. We were unable to perform subgroup analyses based on the core outcomes of PTB because of the lack of data in the included studies. 134 In addition, the studies generally did not exclude twin or multiple pregnancies in their reports; therefore, the population prevalence of late PTB may have been impacted by the multiple birth rates.³ However, given that multiple pregnancies constitute approximately 3% of births, 135,136 multiple births may not have confounded our review results. Although gray literature may be an important source of data on PTBs, this type of literature was excluded from the review because it was not peer-reviewed. The lack of methodological details and accurate data on research variables in the gray literature hinders the quality appraisal of this literature.

Conclusion

The overall PP of PTB in Scandinavian countries was 5.3%, with small variations between countries ranging from 4.9% to 6.2%. The highest and lowest PP rates of PTB in this group of countries were observed in Norway and Finland, respectively. Moderate/late PTB, which occurred between 32 to <37 weeks of gestation, was the most common category of PTB and represented 4.5% of all live births. The review findings reflect the burden of diseases on certain populations and can support policymakers who are responsible for the allocation of resources in identifying healthcare priorities and developing preventive strategies.

Data availability statement

Access to the raw data is possible upon reasonable request to the corresponding author.

Declaration of conflicting interests

The authors declare that they have no known competing financial interests or personal

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