Appendix

Linkage process

The linkage process was based on a set of attributes: mother's name and age, date of birth, sex and place of residence (municipality). It was not possible to use the Live Birth Certificate number as a linking attribute, as it is only available for infant deaths and has a very low completion rate.^{18,23} The relationship was performed by similarity matching, using the algorithm Center for Data and Knowledge Integration for Health-Record Linkage (CIDACS-RL),²³ an open source linkage algorithm from CIDACS that generates a similarity score based on multiple identifiers. The process was conducted in the secure environment of the CIDACS data center, in strict data protection structure following ethical and legal standards.²⁴ CIDACS-RL applies a combination of indexing and searching algorithms. Indexing searches for the most similar records in the indexed SIM for each SINASC record and submits them to a paired comparison step. Candidate linkage records are ranked according to their scores and the highest score comparison pair is retained. For this dataset, a sample of 2000 pairs, stratified into three categories of linkage scores (high score, greater than 0.95; intermediate score, values between 0.90 and 0.95; and low score, less than 0. .90) was manually reviewed to assess link quality. In this validation process, the overall sensitivity and specificity was greater than 90% in all years, with little variation (for example, 94.0% in 2015; 98.1% in 2017, and 90.4% in 2019).

Figure 1. Linkage flowchart



Year	Cutoff	Sensitivity (%)	Specificity (%)
2012	0.974	92.1	91.6
2013	0.974	93.3	94.3
2014	0.974	92.9	95.5
2015	0.973	94.0	94.0
2016	0.951	94.7	96.5
2017	0.950	95.0	98.1
2018	0.939	92.8	93.3
2019	0.956	92.6	90.4

Table 1. Exact values for sensitivity and specificity about linkage SIM X SINASC

	Manual Review		
	True	False	
≥ Cutoff	a	b	
> Cutoff	c	d	
	a+c	b+d	

Sensitivity = a / (a+c)

Specificity = d / (b+d)





Table 2. Observational Routinely-Collected health Data (RECORD) guideline

	Ite	STROBE items	Location in	RECORD items	Location in
	m		manuscript		manuscript
	No		where items		where items
			are reported		are reported
Title and abs	tract		• • • • • • • • • • • • • • • • • • •		•
	1	(a) Indicate the study's	Abstract	RECORD 1.1: The type of	Abstract
		design with a commonly		data used should be specified	
		used term in the title or		in the title or abstract. When	
		the abstract (b) Provide		possible, the name of the	
		in the abstract an		databases used should be	
		informative and		included.	
		balanced summary of			
		what was done and what		RECORD 1.2: If applicable,	
		was found		the geographic region and	
				timeframe within which the	
				study took place should be	
				reported in the title or	
				abstract.	
				RECORD 1 3. If linkage	
				hetween databases was	
				conducted for the study this	
				should be clearly stated in the	
				title or abstract	
Introduction	L		I	the of dobtidet.	
Background	2	Explain the scientific	Introduction:		Introduction:
rationale		background and	1 st to 4 th		1 st to 4 th
		rationale for the	paragraphs		paragraphs
		investigation being	1 0 1		1 0 1
		reported			
Objectives	3	State specific objectives,	Introduction:		Introduction:
-		including any	last paragraph		last
		prespecified hypotheses			paragraph
Methods					
Study	4	Present key elements of	Methods		Methods
Design		study design early in the	section: 1 st		section: 1 st
		paper	paragraph -		paragraph -
			Study design,		Study
			data source,		design, data
			population		source,
			and linkage		population
			process (page		and linkage
			0)		(page 6)
Setting	5	Describe the setting	Methods		(page 0) Methods
Setting	5	locations and relevant	section: Study		section:
		dates including periods	design data		Study
		of recruitment	source.		design data
		an reer annient,	population		source,

The RECORD statement – checklist of items, extended from the STROBE statement, that should be reported in observational studies using routinely collected health data.

		exposure, follow-up, and data collection	and linkage process (page 6) and Expousure definitions (pages 6 and 7)		population and linkage process (page 6) and Expousure definitions (pages 6 and 7)
Participants	6	 (a) Cohort study - Give the eligibility criteria, and the sources and methods of selection of participants. Describe methods of follow-up <i>Case-control study</i> - Give the eligibility criteria, and the sources and methods of case ascertainment and control selection. Give the rationale for the choice of cases and controls <i>Cross-sectional study</i> - Give the eligibility criteria, and the sources and methods of selection of participants (b) Cohort study - For matched studies, give matching criteria and number of exposed and unexposed <i>Case-control study</i> - For matched studies, give matching criteria and the number of controls per case 	Methods section: Study design, data source, population and linkage process (methods of selection of participants) (page 6), Data linkage (methods of selection of participants) (page 6)	RECORD 6.1: The methods of study population selection (such as codes or algorithms used to identify subjects) should be listed in detail. If this is not possible, an explanation should be provided. RECORD 6.2: Any validation studies of the codes or algorithms used to select the population should be referenced. If validation was conducted for this study and not published elsewhere, detailed methods and results should be provided. RECORD 6.3: If the study involved linkage of databases, consider use of a flow diagram or other graphical display to demonstrate the data linkage process, including the number of individuals with linked data at each stage.	6.1: Methods section: Study design, data source, population and linkage process (page 6), Data linkage (page 6) 6.2: Methods section: Data linkage (CIDACS- RL tool) (page 7) 6.3: Methods section: Study design and population; Figure 2 (Results section)
Variables	7	Clearly define all outcomes, exposures, predictors, potential confounders, and effect modifiers. Give diagnostic criteria, if applicable.	Methods section: pages 6-7	RECORD 7.1: A complete list of codes and algorithms used to classify exposures, outcomes, confounders, and effect modifiers should be provided. If these cannot be reported, an explanation should be provided.	All definitions can be found in the Methods section: pages 6-7
Data sources/ measuremen t	8	For each variable of interest, give sources of data and details of methods of assessment (measurement). Describe comparability of assessment methods if there is more than one group	Methods section: Study design, data source, population and linkage		Methods section: Study design, data source, population

			process (pages	and linkage
			6), Exposures	process
			definitions	(pages 6),
			(page 6 and 7),	Main
			Outcomes	exposure
			definitions	variable
			(page 7),	(page 5 and
			Statistical analysis (nage	6),
			8)	Outcomes
				definitions
				(page 7), Statistical analysis (page 8)
Bias	9	Describe any efforts to	Statistical	Statistical
		of bias	Analysis	Analysis
			(page 8)	(page 8)
Study size	10	Explain how the study	Figure 2 (Pesults	Figure 2 (Pesults
		Size was arrived at	section)	section)
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If applicable, describe which groupings were chosen, and why	Table 1 (Results section) Information regarding these variables can be found throughout the methods: (pages 6, 7 and 8)	Table 1 (Results section) Information regarding these variables can be found throughout the methods: (pages 6, 7 and 8)
Statistical	12	(a) Describe all	Methods	Methods
memous		including those used to	section:	section:
		control for confounding	Statistical	Statistical
		(b) Describe any methods used to	Analysis	Analysis
		examine subgroups and interactions (c) Explain how missing data were addressed (d) <i>Cohort study</i> - If applicable, explain how loss to follow-up was addressed <i>Case-control study</i> - If applicable, explain how	(page 8)	(page 8)

	matching of cases and			
	controls was addressed			
	Cross-sectional study -			
	If applicable, describe			
	analytical methods			
	sampling strategy			
	(e) Describe any			
	sensitivity analyses			
Data access		Methods	RECORD 12.1: Authors	RECORD
methods		section: Study	which the investigators had	12.1:
		design, data	access to the database	Methods
		source,	study population.	section:
		population		Study
		and linkage	RECORD 12.2: Authors should provide information	design, data
		process (pages	on the data cleaning methods	source,
		6), Exposures	used in the study.	population
		definitions		and linkage
		(page 6 and 7),		process
		Outcomes		(pages 6),
		definitions		Exposures
		(page 7),		definitions
		Statistical		(page 6 and
		Analysis		7),
		(page 8)		Outcomes
		Results section: Figure		definitions
		2		(page 7),
				Statistical
				Analysis
				(page 8)
				RECORD 12.2:
				Methods section
				Study
				design, data
				source,
				and linkage
				process
				(page 6)
				Results
				section: Figure 2

Linkage				RECORD 12.3: State whether the study included person-level, institutional- level, or other data linkage across two or more databases. The methods of linkage and methods of linkage quality evaluation should be provided.	Methods section: Study design, data source, population and linkage process (page 6), Statistical Analysis (page 8)
Results					
Participants	13	 (a) Report the numbers of individuals at each stage of the study (<i>e.g.</i>, numbers potentially eligible, examined for eligibility, confirmed eligible, included in the study, completing follow-up, and analysed) (b) Give reasons for non-participation at each stage. (c) Consider use of a flow diagram 	Results section: 1st paragraph; Figure 2	RECORD 13.1: Describe in detail the selection of the persons included in the study (<i>i.e.</i> , study population selection) including filtering based on data quality, data availability and linkage. The selection of included persons can be described in the text and/or by means of the study flow diagram.	Results section: 1st paragraph; Figure 2
Descriptive data	14	 (a) Give characteristics of study participants (<i>e.g.</i>, demographic, clinical, social) and information on exposures and potential confounders (b) Indicate the number of participants with missing data for each variable of interest (c) <i>Cohort study</i> - summarise follow-up time (<i>e.g.</i>, average and total amount) 	Table 1		Table 1
Outcome data	15	<i>Cohort study</i> - Report numbers of outcome events or summary measures over time <i>Case-control study</i> - Report numbers in each exposure category, or	Results section: 1 st and 2 nd paragraphs; Table 1; and 3 th to 6 th in table 2		Results section: 1 st and 2 nd paragraphs; Table 1; and 3 th to 6 th in table 2

		summary measures of exposure			
		Cross-sectional study - Report numbers of			
		summary measures			
Main results	16	(a) Give unadjusted estimates and, if applicable, confounder- adjusted estimates and their precision (e.g., 95% confidence interval). Make clear which confounders were adjusted for and why they were included (b) Report category boundaries when continuous variables were categorized (c) If relevant, consider translating estimates of relative risk into absolute risk for a	Results section: 3 th to 7 th , Tables 2 and 3		Results section: 3 th to 7 th , Tables 2 and 3
		meaningful time period			
Other analyses	17	Report other analyses done—e.g., analyses of subgroups and interactions, and sensitivity analyses	Results section (last paragraph) Supplementary Material: Tables 1, 2, 3 and 4 and figure 3		Results section (last paragraph) Supplementa ry Material: Tables 1, 2, 3 and 4 and figure 3
Discussion			0 -		0 -
Key results	18	Summarise key results with reference to study objectives	Discussion section (1st paragraph)		Discussion section (1st paragraph)
Limitations	19	Discuss limitations of the study, taking into account sources of potential bias or imprecision. Discuss both direction and magnitude of any potential bias	Discussion section (8th paragraph)	RECORD 19.1: Discuss the implications of using data that were not created or collected to answer the specific research question(s). Include discussion of misclassification bias, unmeasured confounding, missing data, and changing eligibility over time, as they pertain to the study being reported.	Discussion section (8th paragraph)
Interpretatio n	20	Give a cautious overall interpretation of results considering objectives, limitations, multiplicity of analyses, results from similar studies, and other relevant evidence	Discussion section: page 11 to 14		Discussion section: page 11 to 14

Generalisabi lity	21	Discuss the generalisability (external validity) of the study results	Discussion section: page 13 (8 th paragraph)		Discussion section: page 13 (8 th paragraph)
Other Inform	nation				
Funding	22	Give the source of funding and the role of the funders for the present study and, if applicable, for the original study on which the present article is based	Funding (pages 2)		Funding (page 2)
Accessibilit y of protocol, raw data, and programmin g code			Methods section (pages 6-8)	RECORD 22.1: Authors should provide information on how to access any supplemental information such as the study protocol, raw data, or programming code.	Methods section (pages 6-8)

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