Supplemental Material 1 Search strategy

1. Searcl	n strategy of PubMed:
No.	Search items
#1	((((((((Infant, Premature[MeSH Terms]) OR (Infant, Premature[Title/Abstract]))
	OR (Neonatal Prematurity[Title/Abstract])) OR (premature[Title/Abstract])) OR
	(premature baby[Title/Abstract])) OR (premature birth[Title/Abstract])) OR (premature
	infant[Title/Abstract])) OR (premature neonate[Title/Abstract])) OR (premature
	newborn[Title/Abstract])) OR (Premature Infant*[Title/Abstract])) OR
	(prematuritas[Title/Abstract])) OR (preterm baby[Title/Abstract])) OR (Preterm
	Infant*[Title/Abstract])) OR (preterm neonate[Title/Abstract])) OR (preterm
	newborn[Title/Abstract])
#2	(((((((Infant, Low Birth Weight[MeSH Terms]) OR (Infant*, Low Birth
	Weight[Title/Abstract])) OR (LBW infant[Title/Abstract])) OR (LBW
	neonate[Title/Abstract])) OR (LBW newborn[Title/Abstract])) OR (low birth
	weight[Title/Abstract])) OR (Low Birth Weight Infant*[Title/Abstract])) OR (Low Birth
	Weights[Title/Abstract])) OR (low birthweight[Title/Abstract])) OR (neonatal
	underweight[Title/Abstract])
#3	(((((Intraventricular Hemorrhage[MeSH Terms]) OR (brain bleeding[Title/Abstract]))
	OR (brain haemorrhage*[Title/Abstract])) OR (brain hemorrhage*[Title/Abstract])) OR
	(Intraventricular Haemorrhage*[Title/Abstract])) OR (Intraventricular
	Hemorrhage*[Title/Abstract])
#4	(#1 OR #2) AND #3

No.	Search items
#1	'infant, premature'/exp
#2	'infant, premature':ab,ti OR 'neonatal prematurity':ab,ti OR 'premature':ab,ti OR
	'premature baby':ab,ti OR 'premature birth':ab,ti
#3	'premature infant':ab,ti OR 'premature neonate':ab,ti OR 'premature newborn':ab,ti OR
	'premature infants':ab,ti OR 'prematuritas':ab,ti
#4	'preterm baby':ab,ti OR 'preterm infant':ab,ti OR 'preterm neonate':ab,ti OR 'preterm
	newborn':ab,ti
#5	#1 OR #2 OR #3 OR #4
#6	'infant, low birth weight'/exp
#7	'infant, low birth weight':ab,ti OR 'infants, low birth weight':ab,ti OR 'lbw infant':ab,ti OR
	'lbw neonate':ab,ti OR 'lbw newborn':ab,ti
#8	'low birth weight':ab,ti OR 'low birth weight infant':ab,ti OR 'low birth weight
	infants':ab,ti OR 'low birth weights':ab,ti OR 'low birthweight':ab,ti
#9	'neonatal underweight':ab,ti
#10	#6 OR #7 OR #8 OR #9
#11	#5 OR #10
#12	'brain hemorrhage'/exp
#13	'brain hemorrhage':ab,ti OR 'brain hemorrhages':ab,ti OR 'brain bleeding':ab,ti OR
	'brain haemorrhage':ab,ti OR 'brain haemorrhages':ab,ti

2. Search strategy of Embase:

#14	'intraventricular	haemorrhage':ab,ti	OR	'intraventricular	haemorrhages':ab,ti	OR
	'intraventricular h	nemorrhage':ab,ti OR	'intra	ventricular hemor	rhages':ab,ti	
#15	#12 OR #13 OR #	#14				
#16	#11 AND #15					

3. Search strategy of Cochrane Library

No.	Search items
#1	MeSH descriptor: [Infant, Premature] explode all trees
#2	("Infant, Premature" OR "Neonatal Prematurity" OR "premature" OR "premature baby"
	OR "premature birth"):ti,ab,kw (Word variations have been searched)
#3	("premature infant" OR "premature neonate" OR "premature newborn" OR
	"Premature Infants" OR "prematuritas"):ti,ab,kw (Word variations have been searched)
#4	("preterm baby" OR "Preterm Infant" OR "Preterm Infants" OR "preterm neonate" OR
	"preterm newborn"):ti,ab,kw (Word variations have been searched)
#5	#1 OR #2 OR #3 OR #4
#6	MeSH descriptor: [Infant, Low Birth Weight] explode all trees
#7	("Infant, Low Birth Weight" OR "Infants, Low Birth Weight" OR "LBW infant" OR
	"LBW neonate" OR "LBW newborn"):ti,ab,kw (Word variations have been searched)
#8	("low birth weight" OR "Low Birth Weight Infant" OR "Low Birth Weight Infants" OR
	"Low Birth Weights" OR "low birthweight"):ti,ab,kw (Word variations have been
	searched)
#9	(neonatal underweight):ti,ab,kw (Word variations have been searched)
#10	#6 OR #7 OR #8 OR #9
#11	#5 OR #10
#12	MeSH descriptor: [Cerebral Intraventricular Hemorrhage] explode all trees
#13	("brain bleeding" OR "brain haemorrhage" OR "brain haemorrhages" OR "brain
	hemorrhage" OR "brain hemorrhages"):ti,ab,kw (Word variations have been searched)
#14	("Intraventricular Haemorrhage" OR "Intraventricular Haemorrhages" OR
	"Intraventricular Hemorrhage" OR "Intraventricular Hemorrhages"):ti,ab,kw (Word
	variations have been searched)
#15	#12 OR #13 OR #14
#16	#11 AND #15

4. Search strategy of Web of Science

No.	Search items
#1	(((((((TS=(Infant, Premature)) OR TS=(Neonatal Prematurity)) OR
	TS=(premature)) OR TS=(premature baby)) OR TS=(premature birth)) OR
	TS=(premature infant)) OR TS=(premature neonate)) OR TS=(premature newborn)) OR
	TS=(Premature Infants)) OR TS=(prematuritas)) OR TS=(preterm baby)) OR
	TS=(Preterm Infant*)) OR TS=(preterm neonate)) OR TS=(preterm newborn)
#2	((((((TS=(Infant*, Low Birth Weight)) OR TS=(LBW infant)) OR TS=(LBW neonate))
	OR TS=(LBW newborn)) OR TS=(low birth weight)) OR TS=(Low Birth Weight
	Infant*)) OR TS=(Low Birth Weights)) OR TS=(low birthweight)) OR TS=(neonatal
	underweight)

π3	#1 OK #2
#4	((((TS=(brain bleeding)) OR TS=(brain haemorrhage*)) OR TS=(brain hemorrhage*))
	OR TS=(Intraventricular Haemorrhage*)) OR TS=(Intraventricular Hemorrhage*)
#5	#3 AND #4

Supplementary Figures

Figure S1. Sensitivity analysis for the outcome of NDI. (a) Comparison results between children with mild IVH vs. without IVH. **(b)** Comparison results between children with severe IVH vs. mild IVH.



Figure S2. Sensitivity analysis for the outcome of MDI and PDI. (a) Mean difference of MDI for children with mild IVH vs. children without IVH. **(b)** Mean difference of MDI for children with severe IVH vs. children with mild IVH. **(c)** Mean difference of PDI for children with mild IVH vs. children without IVH. **(d)** Mean difference of PDI for children with severe IVH vs. children with mild IVH. **(e)** OR for the outcome of MDI scored below 70 for mild IVH vs. without IVH. **(f)** OR for the outcome of PDI scored below 70 for mild IVH vs.



Figure S3. Sensitivity analysis for the outcome of motor scores. (a) Mean difference of motor scores for children with severe IVH vs. children with mild IVH. (b) Mean difference of motor scores for children with mild IVH vs. children without IVH. (c) OR for the outcome of motor delay comparison between mild IVH vs. without IVH. (d) OR for the outcome of motor delay comparison between severe IVH vs. mild IVH.



Figure S4. Sensitivity analysis for the outcome of cognitive score and IQ. (a) OR for the outcome of cognitive delay comparison between mild IVH vs. without IVH. **(b)** Mean difference of IQ for children with mild IVH vs. children without IVH. **(c)** OR for the outcome of IQ scored below 70 or ranked under -2SD for mild vs. without IVH. **(d)** Mean difference of IQ for children with severe IVH vs. children with mild IVH. **(e)** OR for the outcome of IQ scored below 70 or ranked under -2SD for severe vs. mild IVH.





Figure S5. Sensitivity analysis for the outcome of hearing impairment and visual impairment. (a) OR for the outcome of hearing impairment comparison between mild IVH vs. without IVH. (b) OR for the outcome of hearing impairment comparison between severe IVH vs. mild IVH. (c) OR for the outcome of visual impairment comparison between mild IVH vs. without IVH. (d) OR for the outcome of visual impairment comparison between severe IVH vs. mild IVH. (d) OR for the outcome of visual impairment comparison between severe IVH vs. mild IVH. (d) OR for the outcome of visual impairment comparison between severe IVH vs. mild IVH.



Figure S6. Sensitivity analysis for the outcome of CP and seizure events or epilepsy. (a) OR for the outcome of CP comparison between children with mild IVH vs. without IVH. **(b)** OR for the outcome of CP comparison between children with severe IVH vs. mild IVH. **(c)** OR for the outcome of seizure events or epilepsy for children with mild IVH vs. without IVH.



Figure S7. Publication bias plot of CP, visual impairment, NDI and hearing impairment. (a) CP for the comparison between children with mild IVH vs. without IVH. **(b)** CP for the comparison between children with severe IVH vs. mild IVH. **(c)** Visual impairment for the comparison between children with mild IVH vs. without IVH. **(d)** Visual impairment for the comparison between children with severe IVH vs. mild IVH. **(e)** NDI for the comparison between children with mild IVH vs. without IVH. **(f)** Hearing impairment for the comparison between children with mild IVH vs. without IVH.







Filled funnel plot with pseudo 95% confidence limits

Supplementary Tables

Table S1. PRISMA checklist

Section and	Item	Checklist item	Location where item
Торіс	#		is reported
TITLE			
Title	1	Identify the report as a systematic review.	Pg. 1
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	Pg. 3
INTRODUCTION	I		
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	Pgs. 4 and 6
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	Pg. 6
METHODS	-		
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	Pgs. 7-9
Information	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies.	Pg. 7
sources		Specify the date when each source was last searched or consulted.	
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	Pg. 7
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened	Pg. 9
		each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the	
		process.	
Data collection	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they	Pg. 10
process		worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation	
		tools used in the process.	
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain	Pgs. 8 and 9
		in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources).	Pg. 10
		Describe any assumptions made about any missing or unclear information.	
Study risk of bias	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed	Pg. 10
assessment		each study and whether they worked independently, and if applicable, details of automation tools used in the process.	
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	Pg. 10
Synthesis	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics	Pg. 10
methods		and comparing against the planned groups for each synthesis (item #5)).	
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data	Pg. 10-11

Section and	Item	Checklist item	Location where item			
Торіс	#	· · · · ·	is reported			
	10		D 10			
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	Pg. 10			
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the	Pgs. 10-11			
		model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.				
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	/			
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	Pg. 11			
Reporting bias	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	Pg. 11			
assessment						
Certainty	15Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.Pg.					
assessment	ient					
RESULTS	RESULTS					
Study selection	2. Sudy selection 16a Describe the results of the search and selection process, from the number of records identified in the search to the number of studies					
		included in the review, ideally using a flow diagram.				
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	Pg. 12			
Study	tudy 17 Cite each included study and present its characteristics.					
characteristics						
Risk of bias in	18	Present assessments of risk of bias for each included study.	Pg. 12			
studies						
Results of	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its	Pgs. 12-16			
individual studies		precision (e.g. confidence/credible interval), ideally using structured tables or plots.				
Results of	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	Pgs. 12-16			
syntheses	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision	Pgs. 12-16			
		(e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.				
	20c Present results of all investigations of possible causes of heterogeneity among study results.					
	20d Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.					
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	/			
Certainty of	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	/			
evidence						
DISCUSSION	<u>.</u>					
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	Pgs. 18-21			

Section and Topic	Item #	Checklist item	Location where item is reported
	23b	Discuss any limitations of the evidence included in the review.	Pgs. 20-21
	23c	Discuss any limitations of the review processes used.	Pgs. 22
	23d	Discuss implications of the results for practice, policy, and future research.	Pgs. 18 and 21
OTHER INFORM	IATIO	N	
Registration and	24a	Provide registration information for the review, including register name and registration number, or state that the review was not	Pg. 7
protocol		registered.	
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	Pg. 7
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	NA
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	Pg. 32
Competing	26	Declare any competing interests of review authors.	Pg. 32
interests			
Availability of	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from	Pg. 24
data, code and		included studies; data used for all analyses; analytic code; any other materials used in the review.	
other materials			

	Table S2. Baseline	characteristics	of the 37	studies for	[.] meta-analvsis
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Study	Study	country	Included infants	MAX	MAX BW	Method of IVH	IVH	Severe/Mild	Mild/none outcome	Time	to
	type		(centers)	GA	(g)	diagnosis	comparison	outcome		analyze	
							between				
							grade				
Treluyer, L.et	PC	French	3129(single)	32		cranial ultrasound	Severe/Mild;	NDI; IQ; CP;	NDI; IQ; CP;	at age 5	
al.,2023 ^[19]							Mild/none	epilepsy; visual	epilepsy; visual		
								disability; hearing	disability; hearing		
								disability	disability		
Reis, Joana	RC	Portugal	124(single)	32		cranial ultrasound	Mild/none		CP;auditory	at 24	4–36
Soares.et									deficit;blindness;	months	of
al.,2023 ^[20]									Gross motor/locomotion;	corrected	d age
									cognition		
Perisset,	RC	Switzerland	509(single)	32		cranial ultrasound	Mild/none	NDI; MDI; PDI;		at two	years
Alexandra.et								CP; visual		corrected	d age
al.,2023 ^[21]								problems; hearing			
								problems			
Yaghini, O.	PC	Iran	100(multicenter)	34	VLBW <	brain	Mild/none		cognition;	at 8 y	ears
et al., 2022					1500	ultrasonography			communication;		
[22]									receptive language;		
									expressive language;		
									fine motor and gross		
									motor		
									performance		
Wang, Y. et	PC	China	1079(single)	30		cerebral ultrasound	Severe/Mild;	CP; MDI < 70;	CP; MDI<70; deafness;	at 1	8–24
al., 2022 ^[8]							Mild/none	deafness;	blindness;	months	of
								blindness;	neurodevelopmental	corrected	d age
								neurodevelopmental	disability		
								disability			
Shah,	RC	Canada	2327(multicenter)	29		cranial ultrasound	Severe/Mild;	NDI; CP;	NDI; CP;	at 18 and	124
Vibhuti.et							Mild/none	congnition/	congnition/language/	months	of
al.,2022 ^[23]								language/	motor score	corrected	d age
								motor score			

Hwang-Bo,	RC	Korea	191(single)	32	VLBW	<	brain sonogram	Severe/Mild	seizure; cognitive		at 18 months
Seok.et al.,					1500				score;		of
2022 ^[24]									language score;		corrected age
									motor score; socio-		
									emotional score;		
									adaptive behavior		
									score		
Cha, J. H.et	RC	Korea	5734(multicenter)		VLBW	<	cranial	Mild/none		Motor/cognitive/	aged 12–42
al.,2022 ^[25]					1500		ultrasonography,			visual/hearing	months
II a 11 a baser of a s	DC	A	400(14:	20			i-1	Course AC14	CD. IO	Impairment	at 9 and and of
N L of	ĸĊ	Australia	499(Inufficenter)	20	•••		ultrasonography	Mild/none	cr, IQ, executive,	cr, IQ, executive,	
al., 2021 ^[26]							utrasonography	wind/hone	outcomes	outcomes	age
Bae, Seong	RC	Korea	240(single)		VLBW	<	brain USG	Mild/none		NDI; CP;	at a corrected
Phil.et					1500					cognitive/language/	age of 18-24
al.,2021 ^[27]										motor score	months
Shankaran,	RC	America	4216(multicenter)	26			normal cranial head	Severe/Mild;	NDI; CP;	NDI; CP;	18-22 months
Seetha.et							ultrasounds	Mild/none	cognitive/motor	cognitive/motor score;	of corrected
al.,2020 ^[28]									score; hearing/vison impairment	hearing/vison impairment	age
Scott, T.	RC	America	293(single)	32			ultrasound	Mild/none		cognitive/language/motor	between 24
E.et al.,										scores; CP; seizure;	and 42
2020 ^[29]										hearing loss	months
											chronologic
											age
Tu, Yi-	CC	China	806(multicenter)	32	VLBW	<	brain ultrasound	Severe/Mild;	epilepsy	epilepsy	5 years of age
Fang.et al., 2019 ^[10]					1500			Mild/none			
Peixoto,	CC	Portugal	172(single)	34			cranial ultrasound	Mild/none		CP; NDI; visual	at 24 months
Sara.et al., 2018 ^[30]										impairment; hearing loss	of age

Gilard, Vianney.et al., 2018 ^[31]	PC	France	122(single)	< 37		cranial ultrasound	Severe/Mild	CP; gross motor function; language development; severe visual impairment; deafness; epilepsy		24 months of corrected age
Pfahl, S.et al., 2018 ^[32]	RC	Germany	89(single)	32		cranial ultrasound	Severe/Mild; Mild/none	CP; PDI; MDI; NDI; blindness; hearing loss	CP; PDI; MDI; NDI; blindness; hearing loss	at 18–24 months of corrected age
Reubsaet, P.et al., 2017 ^[9]	CC	Netherlands	342(single)	32		cranial ultrasound	Mild/none		CP; epilepsy; NDI; visual impairment; hearing impairment; cognitive/motor score	at 2 years' corrected age
Wy, P. Ann.et al., 2015 ^[33]	PC	America	985(multicenter)	< 37	2500	cranial ultrasound	Mild/none		IQ; cognitive functioning; behavior and academic achievement	at 18 years of age
Radic, Julia A. E.et Al., 2015 ^[34]	PC	Canada	1018(multicenter)	30		cranial ultrasound	Severe/Mild; Mild/none	NDI; CP; MDI; blindness; bilateral deafness	NDI; CP; MDI; blindness; bilateral deafness	2 to 3 years of age (corrected age)
Vohr, Betty R.et al., 2014 ^[35]	PC	America	338(multicenter)	< 37	1250	cranial ultrasound	Severe/Mild; Mild/none	CP; IQ; bilateral blind or HL with amplification	CP; IQ; bilateral blind or HL with amplification	at 16 years of age
Bolisetty, Srinivas.et al., 2014 ^[14]	RC	Australia	1472(multicenter)	28		cranial ultrasound	Severe/Mild; Mild/none	CP; MDI; NDI; bilateral blindness; hearing loss	CP; MDI; NDI; bilateral blindness; hearing loss	at 2 to 3 years' corrected age

Payne, D Allison H.et al., 2013 ^[12]	PC	America	1472(multicenter)	27		cranial ultrasound	Severe/Mild; Mild/none	CP;NDI; cognitive/language score; severe visual impairment; deafness	CP;NDI; cognitive/language score; severe visual impairment; deafness	at 18 to 22 months' corrected age
Merhar, S. 1 L.et al., 2012 ^[36]	PC	America	166(multicenter)		ELBW<1000	cranial ultrasound	Severe/Mild	PDI; MDI; NDI		18–22 months
Klebermass- 1 Schrehof, K. et al.,2012 ^[37]	RC	Germany	471(single)	32		cranial ultrasound	Severe/Mild; Mild/none	CP; MDI; NDI; visual impairment; acoustic impairment	CP; MDI; NDI; visual impairment; acoustic impairment	at the age of 5.5 years
Choi, Il I Rak.et al., 2012 ^[38]	RC	Korea	49(single)	31	VLBW < 1500	cranial ultrasound	Mild/none		MDI; PDI	at a corrected age of 12 months
Broitman, Eduardo.et al.,2007 ^[39]	RC	America	2103(multicenter)		ELBW<1000	head ultrasound scanning	Severe/Mild; Mild/none	NDI; MDI; PDI; CP; blindness; deafness	NDI; MDI; PDI; CP; blindness; deafness	at 18 to 22 months corrected age
Patra, K.et al., 2006 ^[40]	PC	America	706(single)		ELBW<1000	cranial ultrasound	Mild/none		NDI; PDI; MDI; major neurologic abnormality; deafness	at 20 months' corrected age
Ancel, P. Y. 1 et al., 2006 ^[41]	PC	France	1954(multicenter)	32		cranial ultrasound	Severe/Mild; Mild/none	СР	СР	at 2 years
Sherlock, R. 1 L.et al., 2005 ^[42]	PC	Australia	298(single)	28	1000	cranial ultrasound	Severe/Mild; Mild/none	CP; IQ; major neurosensory disability	CP; IQ; major neurosensory disability	at 8 years of age
MO'Keefe. 1 et al., 2001 ^[43]	PC	Ireland	68(single)	35	2240	cranial ultrasonography	Severe/Mild	CP; visual acuity <6/60		between 12– 150 months

Doyle LW,et al., 2000 ^[44]	PC	Australia	424(single)		VLBW 1500	<	cranial ultrasound	Severe/Mild; Mild/none	СР	СР	at 5 years of age
Bendersky, M.et al., 1995 ^[45]	PC	America	105(single)	35	2000		cranial ultrasound	Severe/Mild; Mild/none	IQ; memory; language; bayley motor score	IQ; memory; language; bayley motor score	at 3 years of age
Landry, S. H.et al., 1993 ^[46]	PC	America	78(single)	34	< 1600		ultrasound or CT scan	Severe/Mild	motor score; IQ		at 6, 12,24 and 36 months of age
Vohr, B.et al., 1992 ^[47]	PC	America	112(single)	34	< 1750		cranial ultrasound	Severe/Mild; Mild/none	cognitive Index; perceptual Index; visual-motor development	cognitive Index; perceptual Index; visual-motor development	at 5 years of age
Vohr, B. R.et al., 1989 ^[48]	PC	America	112(single)	34	< 1750		cranial ultrasound	Severe/Mild; Mild/none	PDI; MDI; VER latency; Kohen-Raz subscores; Mullen scores	PDI; MDI; VER latency; Kohen-Raz subscores; Mullen scores	the first 2 years of life
Morales, W. J, 1987 ^[49]	PC	America	303(single)		VLBW 1500	<	echoencephalogram	Severe/Mild; Mild/none	PDI; MDI	PDI; MDI	at 1 years of age
Ment, L. R.et al.,1985 ^[50]	PC	America	164(single)		1250		cranial ultrasound	Severe/Mild; Mild/none	IQ; The Bayley Scales of Infant Development	IQ; The Bayley Scales of Infant Development	at 30 months' corrected age

*PC: prospective cohort study RC: retrospective cohort study CC: case-control study

		Selectio	n		Comparability		Outcome		
					(2 stars)				
Study	Representativeness of the exposed cohort	Representativeness of the non-exposed cohort	Ascertainment of exposure	Demonstration that outcome of interest was not present at start of study	Comparability of the exposed cohort and the non-exposed cohort on the basis of the design or analysis	Assessment of outcome	Was follow- up long enough for outcomes to occur	Adequacy of follow- up	Total scores
Treluyer, L.et al.,2023 ^[19]	1	1	1	1	2	1	1	0	8
Reis, Joana Soares.et al.,2023 ^[20]	1	1	1	1	2	1	1	1	9
Perisset, Alexandra.et al.,2023 ^[21]	1	1	1	1	1	1	1	1	8
Yaghini, O. et al., 2022 ^[22]	1	1	1	1	2	1	1	1	9
Wang, Y. et al., 2022 ^[8]	1	1	1	1	2	1	1	1	9
Shah, Vibhuti.et al.,2022 ^[23]	1	1	1	1	2	1	1	1	9
Hwang-Bo, Seok.et al., 2022 ^[24]	1	1	1	1	1	1	1	1	8
Cha, J. H.et al.,2022 ^[25]	1	1	1	1	2	1	1	0	8
Hollebrandse, N. L.et al., 2021 ^[26]	1	1	1	1	1	1	1	1	8
Bae, Seong Phil.et al.,2021 ^[27]	1	1	1	1	2	1	1	0	8

Table S3. Risk of bias for 34 cohort studies included as per modified Newcastle-Ottawa Scale

Shankaran, Seetha.et al.,2020 ^[28]	1	1	1	1	2	1	1	1	9
Scott, T. E.et al., 2020 ^[29]	1	1	1	1	1	1	1	1	8
Gilard, Vianney.et al., 2018 ^[31]	1	1	1	1	1	1	1	1	8
Pfahl, S.et al., 2018 ^[32]	1	1	1	1	1	1	1	1	8
Wy, P. Ann.et al., 2015 ^[33]	1	1	1	1	2	1	1	1	9
Radic, Julia A. E.et al., 2015 ^[34]	1	1	1	1	2	1	1	1	9
Vohr, Betty R.et al., 2014 ^[35]	1	1	1	1	2	1	1	1	9
Bolisetty, Srinivas.et al., 2014 ^[14]	1	1	1	1	2	1	1	0	8
Payne, Allison H.et al., 2013 ^[12]	1	1	1	1	2	1	1	1	9
Merhar, S. L.et al., 2012 ^[36]	1	1	1	1	2	1	1	1	9
Klebermass- Schrehof, K.et al.,2012 ^[37]	1	1	1	1	2	1	1	1	9
Choi, Il Rak.et al., 2012 ^[38]	1	1	1	1	1	1	1	1	8
Broitman, Eduardo.et al.,2007 ^[39]	1	1	1	1	1	1	1	1	8

Patra, K.et al., 2006 ^[40]	1	1	1	1	2	1	1	1	9
Ancel, P. Y. et al., 2006 ^[41]	1	1	1	1	1	1	1	1	8
Sherlock, R. L.et al., 2005 ^[42]	1	1	1	1	1	1	1	1	8
M O'Keefe.et al., 2001 ^[43]	1	1	1	1	1	1	1	1	8
Doyle LW,et al., 2000 ^[44]	1	1	1	1	1	1	1	1	8
Bendersky, M.et al., 1995 ^[45]	1	1	1	1	2	1	1	1	9
Landry, S. H.et al,1993 ^[46]	1	1	1	1	1	1	1	0	7
Vohr, B.et al,1992 ^[47]	1	1	1	1	1	1	1	0	7
Vohr, B. R.et al,1989 ^[48]	1	1	1	1	1	1	1	1	8
Morales, W. J, 1987 ^[489]	1	1	1	1	1	1	1	0	7
Ment, L. R.et al,1985 ^[50]	1	1	1	1	1	1	1	0	7

Item		Tu, Yi-	Peixoto,	Reubsaet,
		Fang.et	Sara.	P.et al.,
		al.,	et al.,	2017 [9]
		2019 ^[10]	2018 [30]	
Was the Case Definition				
and Diagnosis Adequate				
	A. Yes, with independent	1	1	1
	validation☆			
	B. Yes (e.g., from medical records or			
	the doctor's own records)			
	C. No description			
Representativeness of the				
Cases				
	A. Continuous cases, or the cases are	1	1	1
	representative cases 🛠			
	B. Potential for selection biases, or			
	not stated.			
Selection of Controls				
	A. Community controls☆			
	B. Hospital controls	0	0	0
	C. No description			
Definition of Controls				
	A. No history of disease(endpoint)☆	1	1	1
	B. No description of source			
Comparability (2 points)				
Comparability of Cases and				
Controls on the Basis of the				
Design or Analysis				
	A. Select and analyze controls	1	1	1
	according to the most important			
	factors☆			
	B. Select and analyze controls based	1	0	1
	on other important factors (such as			
	the second most important factor) \thickapprox			
Ascertainment of Exposure				
	A. Reliable records (such as surgical	1	1	1
	records) ☆			
	B. Blind interview (it is unknown			
	who are cases or controls) \Rightarrow			
	C. Unblinded interview			
	D. Self-documentation or medical			
	record			

	E. No description			
Same Method	of			
Ascertainment for Ca and Controls	ases			
	A. Yes ☆	1	1	1
	B. No			
No response rate				
	A. The no response rate of the two groups is the same \Rightarrow	1	1	1
	B. No description			
	C. Response rates vary but reasons			
	are not stated			
Total score		8	7	8