Supplementary Online Content

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eFigure 1. Funnel plots of effect sizes and standard errors used for the trim and fill analyses for assessing possible publication biases

(A) No indications of a publication bias were observed in the Visual Following (Neonates) meta-analysis. (B) Assessment of the funnel plot using the trim-and-fill method indicated a possible minor publication bias in the Visual Following...
meta-analysis, which led to the imputation of one additional study. The effect of this adjustment was negligible and slightly increased the total effect size, with an adjusted pooled estimate of -0.94 (95% CI, -1.43 to -0.45, z = -3.77, p < 0.001), suggesting that a publication bias could not better explain the result. (C) No indications of a publication bias were observed in the Latency to Fixate meta-analysis. (D) No indications of a publication bias were observed in the Habituation meta-analysis. (E) Assessment of the funnel plot using the trim-and-fill method indicated a possible minor publication bias in the Novelty Preference meta-analysis, which led to the imputation of two additional studies. The effect of this adjustment was negligible and slightly attenuated the total effect size, with an adjusted pooled estimate of -0.17 (95% CI, -0.29 to -0.06, z = -2.92, p = 0.003), suggesting that a publication bias could not better explain the result. (F) No indications of a publication bias were observed in the Focused Attention meta-analysis.
eFigure 2. A flow diagram of studies’ selection process

1841 Articles identified through PubMed

624 Articles identified through PsycINFO

157 Additional articles identified through other sources

2176 Articles screened after duplicates removed

1993 Articles excluded

183 Full-text articles assessed for eligibility

130 Articles excluded for the following reasons:

39 Other attention outcomes

28 Insufficient data

26 Overlapping study sample

20 Non-attention outcomes

17 Miscellaneous

53 Articles included in the meta-analyses
eFigure 3. Forest and funnel plots for the meta-analysis on the differences in visual following (animate stimuli) between preterm and full-term born neonates.

### eFigure 3A. A Forest Plot for the Differences in Visual Following (Animate Stimuli) Between Preterm and Full-term Born Neonates

| Study                        | Cohen’s d (95% CI) | Favors control | Favors preterm | Weight, % |
|------------------------------|--------------------|----------------|----------------|-----------|
| Leijon et al. 1982           | -0.06 (-0.56 to 0.44) |                |                | 30.0      |
| Paludetto et al. 1982        | -0.37 (-0.98 to 0.25) |                |                | 24.3      |
| Ferrari et al. 1983          | -1.65 (-1.71 to -0.39) |                |                | 22.5      |
| Stjernqvist & Svenningsen 1990 | -0.45 (-1.10 to 0.19) |                |                | 23.2      |
| Total                        | -0.45 (-0.86 to -0.04) |                |                | 100       |

- $Q_3 = 5.56, p = 0.14, I^2 = 46\%$

### eFigure 3B. A funnel plot of effect sizes and standard errors; no indication of a publication bias was found.

eFigure 3. (A) An additional analysis was conducted to address the four studies in neonates that also presented a measure of following an animate stimulus (i.e., the examiner’s face). The analysis indicated that in regard to socially-charged objects full-term neonates are more likely to present superior visual following, with a pooled effect size of -0.45 (95% CI, -0.86 to -0.04; z = -2.15, $p = 0.03$; $k = 4$, $N = 218$; $Q_3 = 5.56, p = 0.14; I^2 = 46\%$). This implies that even though preterm-born neonates are more likely to show better reactivity to neutral stimuli at term, their visual system is less primed to track human figures. However, this finding should be interpreted with caution due to the small number of studies and moderate evidence of heterogeneity. (B) A funnel plot of effect sizes and standard errors; no indication of a publication bias was found.
## eTable 1. Characteristics of the studies included in the meta-analyses

| Study | Included attention measure/s | Age in months | GA<sub>RT</sub> (weeks) | BW<sub>RT</sub> (g) | Cohort | Setting | Exclusion criteria |
|-------|-----------------------------|---------------|-------------------------|------------------|--------|---------|------------------|
| Bonin et al,¹ 1998 | Novelty preference<sup>PD</sup>  
Ratio of looking at the novel stimulus to the looking time at the familiar and novel stimuli; stimulation: abstract patterns and naturalistic faces.  
Habituation<sup>PD</sup>  
Total looking time to reach criterion (infant-controlled); criterion: two consecutive fixations lasting 50% or less than the mean duration of the two longest fixations among the first three looks. | 3.95<sup>–</sup>  
(T1: 2.0;  
T2: 3.92;  
T3: 5.94) | 32.1  
(2.4; 27–36) | 1712  
(392) | Montreal, Canada | Lab | CNS disorder; congenital malformation syndromes; BPD; IVH > 1; 5' Apgar < 5; assisted ventilation > 28 d. |
| Butcher et al,² 2002 | Latency to fixate<sup>PD</sup>  
Latency to fixate (RT) and correct frequency on refixations on a peripheral stimulus; fixation stimuli: either shifting abstract shapes or a phase-reversing schematic face appearing on either the left or the right side of a monitor; peripheral target was a flashing shape; off-line scoring via video recordings; only noncompetition trials considered. | 4<sup>–</sup>  
(once a month from 6 to 26 weeks) | 29.5  
(1.0; < 32) | 1183  
(234) | Groningen, Netherlands 1995–1999 | Lab | GA at birth > 32 weeks; serious neonatal medical complication according to the cutoff point of the NBRS; ROP; IVH > 1. |
| Cherkas-Julkowski,³ 1998 | Focused attention | 14<sup>–</sup>  
(T1: 13;  
T2: 15) | 36.3 (0.7) | 1878  
(848) | USA | Home | GA at birth > 37 weeks; BW > 2300 g; congenital disorder; ROP; NICU hospitalization after GA of 42 weeks; neurological impairment group was excluded. |
| de Jong et al,³ 2015 | Focused attention  
"On-Task Persistence" (i.e., consistent focus on activity: 5-point scale) during 5 min of free play with toys and 10 min of structured tasks (reading a book and making a puzzle with the mother). | 17.3 | 34.7  
(1.3; 32–36) | 2585  
(517) | Utrecht, Netherlands 2010–2011 | Lab | Dysmaturity; multiple births; admission to a tertiary NICU; severe congenital malformations; maternal antenatal alcohol or drug use; chronic maternal antenatal use of psychiatric drugs. |
| De Schuymer et al,⁴ 2012 | Latency to fixate | 5.02<sup>–</sup>  
(T1: 4.02;  
T2: 6.01) | 30.9  
(1.4; 28–34) | 1551  
(406) | Ghent, Belgium | Home | IVH > 2; PVL; ROP > 2; abnormal Algo test. |
| DiPietro et al,⁵ 1992 | Focused attention  
"Examining" (based on Ruff et al,⁶ 1984); free play during 3 trials (2 min each). | 7.67 | 30.2  
(2.1; 26–33) | 1409  
(353) | USA | Lab | Congenital anomalies; maternal intravenous drug use; assisted ventilation > 10 d. |
| Ferrari et al,⁶ 1983 | Visual following | Neonates | 31.0  
(2.4; 28–33) | 1520  
(250) | Modena, Italy 1978–1980 | Hospital | 1' Apgar < 5; 5' Apgar < 5; medical problems; abnormal somatic development; fetal distress; infection or sepsis; metabolic problem; RDS; CNS disorder; abnormal EEG functioning. |
| Foreman et al,⁷ 1991 | Latency to fixate<sup>PD</sup>  
Latency (RT) to turn toward a peripheral stimulus (random check patterns); eight trials; coding by a single experienced observer (no videotape). | 5.10  
(range: neonates–12 m) | (25–34) | 1590  
(462) | Leicester, UK | Hospital | IVH; mild hydrocephalus; respiratory difficulties. |
| Forslund & Bjerre,¹¹ 1983 | Visual following  
Following the movement of a red ball (4-point scale); FT group assessed at a mean age of 3.6 days; PT group assessed at a mean GA of 40.2 weeks. | Neonates | ~32.0 (< 35) | | | Hospital | |
**eTable 1. Characteristics of the studies included in the meta-analyses**

| Study                  | Included attention measure/s                                                                 | Age in months | GA<sub>ref</sub> (weeks) M (SD; Range) | BW<sub>ref</sub> (g) M (SD) | Cohort                       | Setting                       | Exclusion criteria                        |
|------------------------|-----------------------------------------------------------------------------------------------|---------------|----------------------------------------|-----------------------------|-------------------------------|--------------------------------------------|---------------------------------------------|
| Friedman et al,<sup>12</sup> 1981 | **Habituation**<br>Only visual condition; **stimuli:** green/red lighted box; two measures were considered: "Time to Response Decrement" and "Amount of Response decrement" indicating the time to reach decrement criterion (infant-controlled), and the difference in visual attention between the first trials and the last trial; both FT and PT groups were assessed at a mean GA of 40 weeks.<br>**Latency to fixate**<br>"Quickness of response" in the visual condition; latency (RT) to turn to the stimulus presented above as reflected on the pupils. | Neonates      | 33.7 (2.2)                           | 1819 (388)                  | Washington, USA           | Hospital                     | Abnormal BW for GA; poor medical history; inadequate rearing conditions. |
| Harel et al,<sup>13</sup> 2011  | **Novelty preference**<br>Ratio of looking at the novel stimulus to the total looking time; one trial; **stimuli:** abstract patterns presented on a computer screen. **Habituation**<br>Total looking time to reach criterion (fixed fixation time); criterion: accumulated 10 sec of looking at the stimulus. | 3.29          | 32.2 (1.4; < 34)                        | 1752 (364)                   | Ramat Gan, Israel          | Home                           | High medical risk; chromosomal anomalies; IVH > 2. |
| Hodel et al,<sup>14</sup> 2017  | **Novelty preference**<br>Difference in cumulative looking-times for the novel and familiar stimuli; **stimuli:** naturalistic faces presented on a monitor (same in habituation test); video coding of gaze direction (same in habituation and latency to fixate tests). **Focused attention**<br>Duration of visual attention toward a toy, including focused face and/or examining behaviors (based on Ruff et al,<sup>15</sup> 1990); one trial (2 min) of free play with a toy. **Habituation**<br>Trials to reach criterion (infant-controlled); criterion: mean duration of looking for three consecutive fixations < 50% than the mean duration for the first three trials. **Latency to fixate**<br>Latency (RT) to shift attention from a central to a peripheral stimulus; **stimuli:** colorful line drawings presented on either the left or right edges of a monitor. | 9.13          | 35.0 (1.7; 30–36)                       | 2467 (542)                   | MN, USA                      | Lab                           | Neurological insult or disease; intrauterine growth restriction; congenital heart disease; serious medical illness. |
| Hunnius et al,<sup>16</sup> 2008  | **Latency to fixate**<br>Latency to fixate (RT) and frequency of correct refixations on a peripheral stimulus; **stimuli:** either an abstract or a naturalistic face video appearing on either the left or the right side of a monitor; off-line scoring via a video recording; only noncompetition trials considered. | 4<sup>th</sup> (once a month from 6 to 26 weeks) | 29.6 (1.8; 27–32)                 | 1267 (488)                   | Groningen, Netherlands 2000–2002 | Lab                           | GA at birth > 32 weeks; prolonged ventilation; hemorrhagic and ischemic brain lesions; serious infections; ROP > 1. |
| Kooiker et al,<sup>17</sup> 2019  | **Latency to fixate**<br>Reaction time to fixation (RT) and correct detection of salient targets; five visual stimuli (i.e., cartoon, form, color, contrast and motion); gaze recorded via a computerized eye-tracking system. | 12.1          | 27.6 (1.8; 24–32)                       | 1038 (293)                   | Rotterdam, Netherlands 2010–2014 | Lab                           | GA at birth > 33 weeks; visual acuity < 0.15; ROP > 3. |
### eTable 1. Characteristics of the studies included in the meta-analyses

| Study                  | Included attention measure/s                                                                 | Age in months | GA_{PT} (weeks) M (SD; Range) | BW_{PT} (g) M (SD) | Cohort     | Setting        | Exclusion criteria                                                                 |
|------------------------|-----------------------------------------------------------------------------------------------|---------------|--------------------------------|-------------------|------------|----------------|------------------------------------------------------------------------------------|
| Landry & Chapieski,18  | **Focused attention**<br>Mean length of time an infant spent looking at and/or physically examining a toy (based on Ruff et al,19 1984; one trial (2 min) of free play with a toy. | 6             | 30.3 (1.6)                     | 1205 (235)        | Texas, USA | Lab            | Low-risk group: moderate to severe degrees or clear precursors of cerebral palsy; sensory handicaps; non-IVH-related forms of hydrocephalus; IVH > 2; BPD. |
| Landry et al,19 1985   | **Novelty preference**<br>Difference in the amount of time spent looking at the familiar stimulus (trial 7) vs. the novel stimulus (trial 8); stimuli: abstract patterns slides projected on a screen. | 7             | 30.3 (2.5; 27–34)              | 1296 (260)        | Texas, USA | Lab            | No IVH group: sensory handicaps; diagnosis of cerebral palsy; non-IVH-related forms of hydrocephalus; IVH. |
| Landry et al,20 1996   | **Focused attention**<br>Frequency of toy manipulation events during a 5 min toy-centered play with their mother. | 6             | 31.0 (1.7; < 34)               | 1250 (248)        | Texas, USA | Lab            | Low-risk group: severe respiratory complications; IVH > 2.                          |
| Lawson et al,21 1984   | **Novelty Preference**<br>Difference in time looking at the novel vs. the familiar stimulus on the visual-only test trial (simultaneous presentation); stimuli: objects moving on a track; gaze direction measured online using an event recorder. | 2.93          | 31.7 (1.6)                     | 1200 (201)        | New York, USA | Lab            |                                                                                   |
| Leijon,22 1982         | **Visual following**<br>SNBAS: orienting to an inanimate visual stimulus (9-point scale); FT group assessed at the fifth PND; for the PT group only the assessment at term age (i.e., GA of 39–41 weeks) was considered. | Neonates      | 30.3 (1.6; < 35)               | 1550 (323)        | Linköping, Sweden | Hospital | Abnormal BW for GA; serious obstetrical problems; respiratory symptoms 48 h after birth; severe infections or hyperbilirubinemia; nutritional problems; inadequate postnatal growth; neurological symptoms; abnormal neurological function up to 18 m of age. |
| Lejeune et al,23 2019  | **Focused attention**<br>“Sustained attention”: reflecting the intensity of facial interest, duration of looking and duration of manipulation in a 3 min free play with decorated cubes (every 10 sec were coded on a 4-point scale, and then averaged for a composite score). | 14.4          | 29 (2.2; < 32)                 | 1207 (274)        | Geneva, Switzerland 2013-2015 | Hospital | Prematurity control group: Major brain lesions detected on MRI (e.g., IVH > 2) |
| Lobo et al,24 2015     | **Focused attention**<br>“Looking while acting”: infant’s eyes directed toward the object while exhibiting another behavior (frequency); seven trials of free play with 7 separate objects (each trial up to 30 sec). | 8.6^7 (nine times between birth and 2 years) | 26.6 (0.4; 22–30) | 881 (57) | Delaware, USA | Home            | PT no BI group: PVL; IVH > 2                                                     |
| Loi et al,25 2017      | **Focused attention**<br>“Sustained attention with objects”: the degree to which the child engaged in focused and elaborative play during a 15 min play session with toys in the presence of their caregiver (7-point scale). | 22.2          | 29.7 (1.9; 25–33)              | 1263 (308)        | CA, USA    | Lab            | Bilingual environment; medical conditions that might preclude participation in the study. |
| Mash et al,26 1998     | **Novelty preference**<br>Ratio of looking at the novel stimulus to the looking time at the familiar and novel stimuli (only Experiment 1 considered); four trials; stimuli: black press-on elements on white cards (above vs. bellow patterns); simultaneous presentation of familiar and novel stimuli; fixation times | 3.66          | 31.7 (2)                      | 1741 (460)        | PA, USA    | Lab            | Mechanical ventilation > 7 d; BPD; CNS malformations; IVH > 2; progressive hydrocephalus; meningitis; cyanotic cardiac malformations; multiple congenital abnormalities; ROP; strabismus; nystagmus; optic atrophy. |
Table 1. Characteristics of the studies included in the meta-analyses

| Study | Included attention measure/s | Age in months | GA<sub>PT</sub> (weeks) | BW<sub>PT</sub> (g) | Cohort | Setting | Exclusion criteria |
|-------|-------------------------------|---------------|--------------------------|-------------------|--------|---------|------------------|
| Ortiz-Mantilla et al. 2008 | Novelty preference | 8.15<sup>T1</sup> (T1: 6.65; T2: 9.65) | 26.9 (2.2; 23–32) | 976 (245) | New Jersey, USA 1996–1999 | Lab | Primarily classified as low-risk infants. |
| Palmer et al. 1982 | Visual following | Neonates | 31.5 (2.3; 27–35) | 765–2490 | London, UK 1979 | Hospital | |
| Paludetto et al. 1982 | Visual following | Neonates | 33 (27–36) | 1723 | Naples, Italy | Hospital | GA at birth > 37 weeks; abnormal BW for GA; Apgar 5' < 8; hyperbilirubinemia; abnormal glycaemia; neurological damage; serious attacks of apnea; sepsis; serious respiratory distress. |
| Pel et al. 2016 | Latency to fixate | 12.1 | 27.2 (1.1; < 29) | Rotterdam, Netherlands 2010–2013 | Lab | Congenital malformations; white matter or gray matter damage (detected by MRI); visual acuity < 0.15; ROP > 3. |
| Petkovic et al. 2016 | Visual following | 5 | 35.0 (1.3; 33–36) | 2261 (382) | Zagreb, Croatia | Lab | IVH > 1; visual or auditory impairment; major neurological complications. |
| Petrie Thomas et al. 2012 | Focused attention | 8.06 | 29.2 (2.7; 23–32) | 1322 (461) | Vancouver, Canada 2001–2004 | Lab | Major congenital anomalies; IVH > 2; PVL; maternal antenatal drug use; abnormal BW. |
| Pridham et al. 2000 | Focused attention | 8 | 29.2 (2.7; < 33) | 1220 (92) | WI, USA 1995–1998 | Home | RDS group; IVH > 2; BPD |
| Provasi et al. 2017 | Habituation | 4.27 | 34.4 (0.9; 31–36) | 1974 (372) | Paris, France | Lab | History of medical complications; prolonged ventilation; hemorrhagic and ischemic brain lesions; serious infections; visual impairments. |
eTable 1. Characteristics of the studies included in the meta-analyses

| Study | Included attention measure/s | Age in months | GA\textsubscript{PT} (weeks) M (SD; Range) | BW\textsubscript{PT} (g) M (SD) | Cohort | Setting | Exclusion criteria |
|-------|-------------------------------|---------------|------------------------------------------|-------------------------------|--------|---------|-------------------|
| ball on a computer screen; computerized eye tracking. **Visual following** Total looking time on ball course according to areas of interest; only synchronized trials considered. | | | | | | | |
| Reuner et al\textsuperscript{37} 2015 | **Focused attention** Time of intense looking and/or manipulating the toy, with high interest (based on Ruff et al\textsuperscript{15} 1990) during one minute of free play with three toys (20 sec each). | 7.90 | 31.9 (3.1; 23–36) | 1860 (698) | Heidelberg, Germany 2008–2009 | Lab | Congenital anomalies; major sensory impairment; PVL; IVH > 2; other neurological complications; maternal antenatal drug use. |
| Ricci et al\textsuperscript{38} 2008 | **Visual following** Quality of eye tracking of an object; items 3–7 (i.e., horizontal, vertical, arc and tracking a colored stimulus); FT group assessed at 48 hours after birth; PT group assessed at GA of 35 and 40 weeks (only second assessment was considered). | Neonates | 28.2 (1.5; 25–31) | 1155 (262) | Rome and Milan, Italy 2004–2007 | Hospital | GA at birth other than 25–31; abnormal cranial ultrasound; unstable clinical condition at GA of 35; oxygen-dependence at term; major congenital malformations; genetic chromosomal abnormality; metabolic disorders; congenital infection; signs of encephalopathy or seizures during neonatal course; jaundice requiring phototherapy; ROP > 2. |
| Riese\textsuperscript{39} 1987 | **Focused attention** "Attentiveness": indicating more focused or sustained attention (infant was alert to and maintained attention on the toy; 9-point scale); offline coding of a videotaped 2 min episode of free play with a mechanical toy. | 24 | 34.0 (29–37) | | KY, USA | Lab | Twins study |
| Rose et al\textsuperscript{40} 1978 | **Novelty preference** Ratio of looking at the novel stimulus to the looking time at the familiar and novel stimuli (only intramodal/visual tasks considered); stimuli: objects; simultaneous presentation of familiar and novel stimuli (3 trials; 20 sec each) after a 30 sec familiarization trial (without habituation criterion). | 12.4 | 32.6 (2.5) | 1651 (303) | New York, USA | Lab | Visual or neurological abnormalities. |
| Rose et al\textsuperscript{41} 1979 | **Novelty preference** Ratio of looking at the novel stimulus to the looking time at the familiar and novel stimuli (only visual task considered); stimuli: objects in different shapes; simultaneous presentation of familiar and novel stimuli for 20 sec. **Habituation** Time to reach criterion (fixed fixation time); criterion: accumulated 20 sec of looking at the stimulus. | 6 m: 6.72 12 m: 12.4 | 6 m: 33.4 (1.6) 12 m: 33.2 (2.4) | 6 m: 1610 (271) 12 m: 1738 (250) | New York, USA | Lab | Visual or neurological abnormalities. |
| Rose\textsuperscript{42} 1980 | **Novelty preference** Difference in time looking at the novel vs. the familiar stimulus; stimuli: abstract patterns, multidimensional variations of black lines on a white background and naturalistic faces (3 trials; 10 sec each); simultaneous presentation of familiar and novel stimuli. **Habituation** Time to reach criterion (fixed fixation time); criterion: accumulated 5, 10 or 20 sec of looking at the stimulus. | 6.33 | 33.0 (2.4) | 1633 (347) | New York, USA 1974–1976 | Lab | Only non-intervention group: visual or neurological abnormalities. |
| Rose\textsuperscript{43} 1983 | **Novelty preference** | 6 m: | 6 m: 34.5 (1.9) | 6 m: | New York, Lab | | Visual or neurological abnormalities. |

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| Study | Included attention measure/s | Age in months | GA <sub>RT</sub> (weeks) | BW <sub>RT</sub> (g) | Cohort | Setting | Exclusion criteria |
|-------|-----------------------------|---------------|--------------------------|------------------|--------|---------|-------------------|
| Rose et al, 1988 | Novelty preference | 6.49 | 12 m: 12.9 | 1802 (450) | USA | New York, USA 1979–1981 | Congenital, neurological or physical abnormalities; BW > 1500 g. |
| Rose et al, 2001 | Novelty preference | 7 | 31.4 | 1184 (211) | New York, USA | Lab | Congenital, neurological or physical abnormalities; BW > 1500 g. |
| Rose et al, 2002 | Latency to fixate | 8.49 | (T1: 5.24; T2: 7.80; T3: 12.9) | 1108 (283) | New York, USA 1994–1997 | Lab | Congenital, neurological or physical abnormalities; BW > 1750 g. |
| Ross et al, 1992 | Novelty preference | 10 | 30.35 | 1463 (242) | New York, USA | Lab | Congenital malformations; moderate to severe neurosensory deficits; maternal antenatal drug use; abnormal weight for GA. |
| Ross-Sheehy et al, 2017 | Latency to fixate | 7.47 | (range: 2–14) | 1963 | Iowa, USA | Lab | |
| Ruff et al, 1982 | Visual following | 1.49 | (T1: 1.02; T2: 1.96) | <1500 | New York, USA | Lab | Both preterm groups considered; BW > 1500 g. |
| Ruff et al, 1984 | Focused attention | 8.96 | 31.7 (26–34) | 1217 | New York, USA | Lab | Only low-risk group; BW > 1500 g. |
| Study | Included attention measure/s | Age in months | GA (weeks) M (SD; Range) | BW (g) M (SD) | Cohort | Setting | Exclusion criteria |
|-------|------------------------------|---------------|--------------------------|--------------|--------|---------|--------------------|
| Ruff,50 1986 | **Focused attention** Frequency of "examining" (i.e., looking accompanied by fingerling or turning the object with an intent facial expression; 6 trials; each with a different toy; 1 min each). | 7.22 | 31.6 | 1198 | USA | | |
| Soares et al,51 2012 | **Focused attention** Med.1 Frequencies of exploratory actions (based on Ruff et al,9 1984; fingerling, transferring, rotating and alternating were considered); two trials (120 sec each) of free play with three soft rubber toys. | 5.56 (T1: 4.62; T2: 5.57; T3: 6.48) | 35.6 (0.5; 32–36) | 2960 (320) | São Paulo, Brazil | Lab | Prenatal complications; risk of cerebral palsy (i.e., PVL, IVH, alterations in cerebral ultrasound); cardiopulmonary problems; hyperbilirubinemia; ROP; genetic syndromes; BW < 1750 g. |
| Spungen et al,52 1985 | **Novelty preference** The difference in the amount of time spent looking at the familiar (target 1, last trial) vs. the novel stimulus (target 2, first trial); stimuli: abstract patterns; FT group assessed between the first and the third PND; PT group assessed at a mean GA of 41.3 weeks. | Neonates | 32.0 (3.1; 28–40) | 1219 (354) | New York, USA | Hospital | Neurological abnormalities and sickness at the time of testing. |
| Stjernqvist & Svenningsen,53 1990 | **Visual following** BNBAS: orienting to an inanimate visual stimulus (9-point scale); FT group assessed at a mean GA of 40.6 weeks; PT group assessed at a mean GA of 41.3 weeks. | Neonates | 25.4 (24–31) | 755 (109) | Lund, Sweden 1984–1986 | Hospital | Oxygen or medications needed during the time of testing; BW > 900 g. |
| Strand-Brodd et al,54 2011 | **Visual following** PD Gain of smooth pursuit (Gain), indicating the proportion of the object's movement that is followed by smooth pursuit; eye movement measured with EOG; stimuli: small happy face sketch; four test trials (35 sec each) indicating 4 conditions of object's movement (sinusoidal/triangular movement modulation x small/large amplitude) | 3 (T1: 2; T2: 4) | 28.5 (22–32) | 1198 | Uppsala, Sweden 2004–2007 | Lab | Down syndrome; Bartter's syndrome; GA at birth > 32 weeks. |
| Sun & Buys,55 2012 | **Focused attention** "Sustained attention": persistence of effort, infant leaned in towards and fixated on an object while manipulating it (based on Ruff et al,56 1990; 5 trials, each with a different toy; 1 min each). | 8 | 28.0 (1.9; < 32) | 1008 (246) | Brisbane, Australia 1998–1999 | Hospital | Health problems. |
| Wilcox et al,56 1996 | **Habituation** Time to reach criterion/trial length (fixed fixation time); criterion; accumulated 30 sec of looking at the stimulus or looking away from the display three times (2 sec each) after accumulated 10 sec of looking (odd-numbered trials). | 4.56 (T1: 2.54; T2: 4.79; T3: 6.56) | 34.1 (0.8; 32–35) | 2150 (240) | Texas, USA | Lab | Twins; abnormal BW for GA; known neurological insult; genetic and chromosomal abnormalities, infection, or disease. |
| Zuccarini et al,57 2017 | **Focused attention** MOE: exploration of the object using the hands or mouth while having continuous contact with it (based on Ruff et al,8 1984); 5 min session of free play with age-appropriate toys, with the mother's presence. | 7.6 (T1: 6.1; T2: 9.1) | 25.9 (1.5; 23–28) | 803 (191) | Bologna, Italy | Hospital /Lab | GA at birth > 28; major cerebral damages detected by ultrasound and MRI (PVL; IVH > 2; hydrocephalus); congenital malformations; visual impairment (ROP > 2; blindness); hearing impairment. |

**Note.** The measures obtained from each study are described. The age measure comprises both the full-term and the preterm groups. GA, BW and exclusion criteria refer to the preterm group, as the
Table 1. Characteristics of the studies included in the meta-analyses

| Study       | Included attention measure/s | Age in months | GA<sub>PT</sub> (weeks) | BW<sub>PT</sub> (g) | Cohort | Setting | Exclusion criteria |
|-------------|-------------------------------|---------------|-------------------------|---------------------|--------|---------|-------------------|
|             |                               |               | M (SD; Range)           | M (SD)              |        |         |                   |
| full-term groups were sampled from typically developing and healthy infants. BPD = bronchopulmonary dysplasia; BW = birthweight; GA = gestational age; IVH = intraventricular hemorrhage; PND = postnatal day; PT = preterm; PVL = periventricular leukomalacia; RDS = respiratory distress syndrome; ROP = retinopathy of prematurity; RT = response time; T = Time. Longitudinal study: measurements from all time points were pooled together to maintain an accurate sample size for each study. Cor Manuscript's authors have supplied additional data via personal correspondence. Med.1 Median and interquartile range were reported and were transformed into means and standard deviations according to equations 14 and 15 described by Wan et al. Med.2 Median, minimum and maximum were reported and were transformed into mean and standard deviation according to equations 3 and 7 described by Wan et al. PD Plot digitizer<sup>61</sup> was employed to extract data from the manuscript. # Both studies include the same sample, but report on different outcome measures.

Table 2. Preterm Birth and the Development of Visual Attention during the First Two Years of Life: A Systematic Review and Meta-analysis (because of the size of the table we are hosting it here: https://data.mendeley.com/datasets/y6v85b5pny/2)
eAppendix 1. Modified Newcastle-Ottawa Scale for assessment of studies quality and risk of bias

The following coding guidelines were followed:

Selection
1) Representativeness of the preterm cohort
+ Truly or somewhat representative of the average characteristics of preterm-born infants in the community.
- Selected group of preterm-born infants (e.g., specific hospital/clinic); no description of the derivation of the cohort.
2) Selection of the full-term cohort
+ Drawn from the same community as the preterm cohort.
- Drawn from a different source; no description of the derivation of the full-term cohort.
3) Ascertainment of prematurity
+ Secure record (e.g., hospital record).
- Self report; no description.

Comparability
1) Comparable age
+ Study controls for corrected age.
- No description of group ages / a statistic test for comparing group ages / age correction for prematurity.
2) Other comparable factors
+ Study controls for gender, socioeconomic status, ethnicity or parental education.
- Significant differences between the cohorts in these factors.

Outcome
1) Independent assessment
+ Independent coders with an excellent inter-rater reliability statistic; computerized assessment.
- No description; non-excellent inter-rater reliability.
2) Blind assessment
+ Experimenters/coders were blind to the infant’s group (preterm/full-term) or to the test condition.
- No description; experimenters/coders aware of the infant’s group.
3) Data loss
+ Complete data from all subjects accounted for; subjects’ data loss unlikely to introduce a bias – less than 20% of the data, or if an adequate reason for data loss is provided.
- More than 20% of the data lost and no description of those lost; no statement.
| Study               | Selection | Control sample | Ascription of prematurity | Comparability | Outcome | Overall |
|--------------------|-----------|----------------|---------------------------|---------------|---------|---------|
| Bonin et al,1 1998 | -         | +             | +                         | -             | +       | +       | 5      |
| Butcher et al,2 2002 | -         | +             | -                         | -             | +       | -       | 3      |
| Cherkes-Julkowski,3 1998 | -         | -             | -                         | +             | +       | +       | 2      |
| de Jong et al,5 2015 | -         | +             | +                         | +             | +       | +       | 6      |
| De Schuymer et al,6 2012 | -         | +             | +                         | -             | +       | +       | 5      |
| DiPietro et al,7 1992 | -         | +             | -                         | -             | +       | +       | 5      |
| Ferrari et al,9 1983 | -         | +             | -                         | -             | +       | +       | 4      |
| Foreman et al,10 1991 | -         | -             | -                         | -             | -       | -       | 0      |
| Forslund & Bjerre,11 1983 | +         | -             | +                         | +             | -       | +       | 5      |
| Friedman et al,12 1981 | -         | +             | -                         | +             | +       | +       | 6      |
| Harel et al,13 2011 | -         | -             | -                         | +             | +       | +       | 4      |
| Hodel et al,14 2017 | -         | +             | +                         | +             | +       | +       | 7      |
| Hunnius et al,16 2008 | -         | +             | -                         | +             | +       | +       | 5      |
| Kooiker et al,17 2019 | -         | -             | +                         | +             | -       | +       | 6      |
| Landry & Chapieski,18 1988 | -         | +             | -                         | -             | +       | +       | 3      |
| Landry et al,19 1985 | -         | +             | -                         | -             | -       | +       | 4      |
| Landry et al,20 1996 | -         | +             | -                         | -             | -       | +       | 3      |
| Lawson et al,21 1984 | -         | +             | +                         | -             | +       | +       | 4      |
| Leijon,22 1982 | -         | +             | +                         | -             | +       | +       | 4      |
| Lejeune et al,23 2019 | -         | +             | +                         | +             | -       | +       | 6      |
| Study                        | Selection |          |          | Comparability |          |          | Outcome | Overall |
|-----------------------------|-----------|----------|----------|---------------|----------|----------|---------|---------|
| Lobo et al,24 2015          | -         | -        | +        | +             | -        | +        | +       | 5       |
| Loi et al,25 2017           | -         | -        | +        | +             | +        | +        | +       | 6       |
| Mash et al,26 1998          | -         | -        | -        | -             | -        | +        | +       | 3       |
| Ortiz-Mantilla et al,27 2008| -         | -        | +        | +             | +        | +        | +       | 6       |
| Palmer et al,28 1982        | -         | +        | +        | +             | -        | -        | +       | 4       |
| Paludetto et al,29 1982     | -         | +        | +        | -             | +        | -        | +       | 5       |
| Pel et al,30 2016           | -         | -        | +        | +             | -        | +        | +       | 5       |
| Petkovic et al,31 2016      | -         | -        | +        |               | -        | -        | +       | 2       |
| Petrie Thomas et al,32 2012 | -         | +        | +        | +             | -        | +        | -       | 5       |
| Pridham et al,35 2000       | -         | -        | +        |               | +        | +        | +       | 5       |
| Provasi et al,36 2017       | -         | -        | -        |               | +        | +        | -       | 2       |
| Reuner et al,37 2015        | -         | -        | +        |               | +        | +        | -       | 5       |
| Ricci et al,38 2008         | +         | +        | +        |               | +        | -        | +       | 6       |
| Riese,39 1987               | -         | -        | +        |               | +        | -        | +       | 4       |
| Rose et al,40 1978          | -         | +        | +        | +             | -        | +        | +       | 6       |
| Rose et al,41 1979          | -         | -        | -        |               | +        | +        | +       | 5       |
| Rose,42 1980                | -         | -        | -        |               | +        | -        | +       | 4       |
| Rose,43 1983                | -         | -        | -        |               | +        | +        | +       | 5       |
| Rose et al,44 1988          | -         | +        | +        | -             | -        | +        | +       | 5       |
| Rose et al,45 2001          | -         | +        | +        | +             | -        | -        | +       | 6       |
| Study                          | Selection | Comparability | Outcome |
|-------------------------------|-----------|---------------|---------|
|                              | Representative | Corrected age controlled | Blind assessment | Data loss | Overall |
| Rose et al,46 2002           | -         | +            | +       | +         | +         | 6       |
| Ross et al,47 1992           | -         | +            | -       | +         | +         | 5       |
| Ross-Sheehey et al,48 2017   | -         | -            | +       | -         | +         | 4       |
| Ruff et al,49 1982           | -         | +            | -       | +         | +         | 5       |
| Ruff et al, 8 1984           | -         | +            | +       | -         | +         | 5       |
| Ruff,50 1986                 | -         | -            | +       | -         | -         | 2       |
| Soares et al,51 2012         | -         | +            | +       | -         | +         | 5       |
| Spungenen et al,52 1985      | -         | +            | +       | +         | +         | 6       |
| Stjernqvist & Svenningsen,53 1990 | +          | -            | -       | -         | +         | 5       |
| Strand-Brodd et al,54 2011   | +         | -            | +       | +         | -         | 5       |
| Sun & Buys,55 2012           | -         | +            | +       | +         | -         | 5       |
| Wilcox et al,56 1996         | -         | +            | +       | +         | +         | 6       |
| Zuccarini et al,57 2017      | -         | +            | +       | +         | +         | 6       |
eAppendix 2. Description of the moderation analyses

Here we present a description of all of the factors that were considered in the moderation analyses. All of the significant moderators were conveyed in the main text.

Factors weighed in all analyses:
Mean age at test time (months); % of female infants; mean GA at birth of the preterm group; GA age group classification of the preterm group\(^6\) (i.e., extreme, very or moderate/late preterm birth); birthweight of the preterm group; publication year; location (USA vs. other); setting (i.e., lab, hospital or home); cohort's birth era (before the antenatal corticosteroids and surfactant era [i.e., year < 1990; Era I], after the introduction of antenatal corticosteroids and surfactants [i.e., 1990<>2000; Era II], or after the introduction of further advancement in neonatal thermal, emotional and physical care strategies [i.e., year > 2000; Era III]); NOS score (either as a continuous variable or a categorical one, with scores larger than 5 classified as high quality studies\(^6\)); % of intraventricular hemorrhage in the preterm group.

Additional factors weighed in the visual following meta-analysis:
Age group (either neonates \([k = 7]\) or infants \([k = 4]\)); gaze tracking technology (either manual/videotaped examination by an experimenter or computerized eye tracking system \([k = 2]\)); method of assessment in neonates (either the Brazelton Neonatal Behavioral Assessment Scale [BNBAS; \(k = 5\)] or other standardized tests \([k = 2]\).

Additional factors weighed in the latency to fixate meta-analysis:
Gaze tracking technology (either manual/videotaped \([k = 7]\) or computerized eye tracking system \([k = 3]\)); type of measure (either response time \([k = 5]\) or response time + % of correct fixations \([k = 5]\)).

Additional factors weighed in the habituation meta-analysis:
The type of outcome measure (either time \([k = 10]\) or trials \([k = 3]\) to reach the habituation criterion); criterion type (either accumulating a predetermined fixation time \([k = 7]\) or personally-customized, as a given frequency of decrement in looking time from baseline \([k = 6]\)).

Additional factors weighed in the novelty preference meta-analysis:
The type of outcome measure (either the ratio \([k = 11]\) of looking time at the novel stimulus to the total looking time \([i.e., \text{novel + familiar}]\) or the raw difference in looking time between the two stimuli \([k = 6]\)); the method of stimuli presentation (either simultaneous presentation of the novel and familiar stimuli \([k = 14]\) or presentation of the novel stimulus in a subsequent trial \([k = 3]\)).
eAppendix 3. Moderation analysis in the latency to fixate meta-analysis

A moderation analysis was conducted to assess whether differences in latency to fixate between the preterm and term groups were associated with the cohort’s birth era.

The analysis indicated that the cohort’s birth era classification was associated with the differences in latency to fixate between the preterm-term groups (Q_M = 6.38, df = 2, p = 0.041). This suggests that when compared to studies with infants born prior to the corticosteroids and surfactant era (β = -0.442, z = -2.34, p = 0.019), in studies conducted with cohorts that were born between 1990 and 2000 better outcomes were more likely to be observed (β = 0.598, z = 2.41, p = 0.016). Regarding cohorts from Era III, neither increased risk nor advantage were observed when compared to cohorts born at Era I (β = 0.231, z = 1.10, p = 0.27); however, increased risk for deficits was found compared to cohorts from Era II (β = -0.367, z = -1.98, p = 0.048). A possible explanation for this finding is the inclusion of two studies\textsuperscript{17,30} that involved infants born extremely preterm in Era III, while no such preterm GA group was examined in studies from previous eras. Indeed, upon removal of these studies from the analysis no increased risk for deficits was found in preterm infants born in Era III, when compared to Era II (β = -0.282, z = -1.42, p = 0.15).

A further inspection of studies involving cohorts from Era III, suggests that the differences between the preterm-term groups are still statistically significant in this era (d = -0.211; 95% CI, -0.39 to -0.03, z = -2.28, p = 0.023; k = 6), reflecting increased risk for deficits in latency to fixate ability following preterm birth. However, there are two factors that might explain this finding. First, as articulated above, Era III was the only Era including cohorts of infants born extremely preterm-- a descriptive examination of the studies in Era III suggests the possibility of larger effect sizes in studies involving such populations (i.e., d = -0.53, 30, -0.32\textsuperscript{17}) when compared to studies involving very-to-late preterm born populations (i.e., d = 0.12, 16, -0.01, 14, -0.18, 48, -0.42\textsuperscript{6}). Upon removal of these two studies from the analysis no significant differences were observed between the preterm-term groups (d = -0.126; 95% CI, -0.35 to 0.10, z = -1.11, p = 0.27; k = 4). A second factor that could explain the findings is advancements in gaze assessment technologies, involving the implementation of computerized gaze tracking systems that were not utilized in previous eras (half of the studies in Era III utilized such equipment). A descriptive examination of the effect sizes in studies from Era III suggests the possibility of larger effect sizes in studies utilizing computerized gaze tracking systems (i.e., d = -0.42, 6, -0.53, 30 and -0.32\textsuperscript{17}) rather than manual/videotaped assessments (i.e., d = 0.12, 16, -0.01, 14 and -0.18\textsuperscript{48}). Taken together, it could be conjectured that the differences between the preterm-term groups in Era III are at least partially related to the inclusion of extremely preterm cohorts and the implementation of more sensitive methods for gaze assessment. The possibility of assessing the interaction between these two factors and birth era classification was not viable, as no studies from Eras I and II included them.

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eAppendix 4. Moderation analysis in the habituation meta-analysis

A moderation analysis was conducted to assess whether differences in habituation between the preterm and term groups were associated with the cohort’s birth era.

The analysis indicated that when considering all three birth era groups, only a trend towards significance was found for the differences in effect sizes ($Q_M = 4.98, df = 2, p = 0.083$). However, when considering the difference between studies conducted before 2000 to all other studies a significant moderation effect was found ($Q_M = 4.77, df = 1, p = 0.029; k = 13$; see eFigure 4), suggesting increased risk for deficits in habituation in preterm infants born before 2000 ($\beta = -0.186, z = -2.3, p = 0.016; d = -0.186, 95\% CI, -0.34 to -0.03, p = 0.016; k = 10$), and lesser likelihood for such deficits in infants born in Era III ($\beta = 0.318, z = 2.18, p = 0.029; d = 0.132, 95\% CI, -0.11 to 0.37, p = 0.29; k = 3$). Further, this association remained significant even when the only study involving a group of infants born extremely preterm was removed from the analysis ($Q_M = 3.91, df = 1, p = 0.048; k = 12$).

eFigure 4. Graphical depiction of the moderation of the differences between the preterm and full-term groups in habituation by birth era

**eFigure 4.** Each circle represents an effect size of a distinct study, conducted either in cohorts born before ($k = 10$; depicted in blue) or after ($k = 3$; depicted in green) the year 2000. Diamonds represent the standardized total mean differences between the term-preterm groups at each birth era. The asterisk represents significant difference between the groups in studies involving cohorts born before the year 2000; no significant difference was observed in studies from the current era. Circle sizes are proportional to studies’ weight. * $p < 0.05$
eAppendix 5. Additional analyses in the focused attention meta-analysis

Screening for outliers in the focused attention meta-analysis
A strong evidence of heterogeneity in effect sizes between studies was found in the focused attention meta-analysis ($Q_{1} = 36.46, p = 0.004; I^2 = 53.4\%$). Additional screening has pinpointed two potential outliers with studentized residuals of $-2.53$ and $-3.45$. A possible explanation for the deviant effect size in the study conducted by Holly Ruff^50 (a pioneer in the research of focused attention and exploratory behavior in infants) could be related to a partial report of the outcome measures. In the study by Sun & Buys^55 a possible explanation for the deviant effect size is that the study included a cohort of infants born extremely preterm (see the subsequent moderation analysis for a further evaluation). Removal of the two potential outliers, slightly attenuated the pooled effect size, but did not annulled it ($d = -0.17; 95\% CI, -0.28$ to $-0.05; z = -2.88, p = 0.004$). However, these studies were responsible for the heterogeneity, as no heterogeneity was evident following their removal ($Q_{15} = 10.4, p = 0.79; I^2 = 0.0\%$).

Moderation analysis assessing whether birth era and GA group are associated with focused attention outcomes
In the focused attention meta-analysis, sufficient data were available to assess the interaction between cohort’s birth era classification (i.e., birth before or after 2000; as no studies with cohorts born before 1990 included infants born extremely preterm, only a comparison between these two eras was applicable) and GA group^62 (i.e., extreme preterm [GA < 28 weeks], very preterm [GA of 28 to 32 weeks] or moderate/late preterm [GA of 32 to 37 weeks]). Classifications of the GA groups were based on the reported means and ranges for GA at birth.

The analysis indicated a significant interaction between cohort’s birth era and GA group ($Q_M = 11.6, df = 5, p = 0.04$). To test whether birth after the year 2000 was associated with better focused attention performance in infants born extremely preterm, a post hoc analysis using a Dunnett’s test (using the Bonferroni correction) with the before 2000-extreme preterm as the control group was conducted. The analysis indicated that infants born extremely preterm after 2000, had a lesser risk to show deficits in focused attention when compared to infants born extremely preterm before 2000 ($\beta = 1.73, z = 2.87, p = 0.021$). Further, in cohorts born before 2000, both the very- ($\beta = 1.99, z = 3.52, p = 0.002$) and moderate-late-preterm ($\beta = 1.82, z = 3.11, p = 0.009$) cohorts also had a lesser risk for deficits when compared to the extreme-preterm cohort that was born before 2000 (see eFigure 5 for a graphical depiction of the interaction).

Taken together the moderation analysis suggests that in cohorts born in Era III following extreme preterm birth, the risk for deficits in focused attention has attenuated– plausibly due to further advancements in neonatal care.

eFigure 5. Graphical depiction of the moderation of the differences between the preterm and full-term groups in focused attention by birth era and GA group

eFigure 5. Asterisks represent significant differences in effect sizes. ** $p < 0.01$ * $p < 0.05$
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