- An umbrella review of the benefits and risks associated with youths' interactions with 1 electronic screens
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Abstract

The influence of electronic screens on children and adolescents' health and education is not 34 well understood. In this prospectively registered umbrella review (PROSPERO; 35 CRD42017076051), we harmonised effects from 102 meta-analyses (2,451 primary studies; 36 1,937,501 participants) on screen time and outcomes. 43 effects from 32 meta-analyses met 37 our criteria for statistical certainty. Meta-analyses of associations between screen use and 38 outcomes showed small-to-moderate effects (range: r = -0.14 - 0.33). In education, results 39 were mixed; for example, screen use was negatively associated with literacy (r = -0.14, 95%40 confidence interval [CI] -0.20 to -0.09,  $p = \langle 0.001, k = 38, N = 18,318 \rangle$ , but this effect was 41 positive when parents watched with their children (r = 0.15, 95% CI 0.02 to 0.28, p = 0.028, 42 k=12, N=6,083). In health, we found evidence for several small negative associations; for example, social media was associated with depression (r = 0.12, 95% CI 0.05 to 0.19, p =<0.001, k = 12, N = 93,740). Limitations include a limited number of studies for each outcome, medium-to-high risk of bias in 95/102 included meta-analyses and high heterogeneity (17/22 in education and 20/21 in health with  $I^2 > 50\%$ ) We recommend that

caregivers carefully weigh the potential harms and benefits of specific types of screen use.

49 Keywords:

Word count: 5300

An umbrella review of the benefits and risks associated with youths' interactions with electronic screens

## Introduction

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In the 16th century, hysteria reigned around a new technology that threatened to be

"confusing and harmful" to the mind. The cause of such concern? The widespread

availability of books brought about by the invention of the printing press. In the early 19th

century, concerns about schooling "exhausting the children's brains" followed, with the

medical community accepting that excessive study could be a cause of madness. By the

20th century, the invention of the radio was accompanied by assertions that it would distract

children from their reading (which by this point was no longer considered confusing and

harmful) leading to impaired learning.

Today, the same arguments that were once levelled against reading, schooling, and radio are being made about screen use (e.g., television, mobile phones, and computers).<sup>4</sup>
Excessive screen use is the number one concern parents in Western countries have about their children's health and behaviour, ahead of nutrition, bullying, and physical inactivity.<sup>5</sup>
Yet, the evidence to support parents' concerns is inadequate. A Lancet editorial<sup>6</sup> suggested that, "Our understanding of the benefits, harms, and risks of our rapidly changing digital landscape is sorely lacking."

While some forms of screen use (e.g., television viewing) may be detrimental to health and wellbeing, <sup>7,8</sup> evidence for other forms of screen exposure (e.g., video games or online communication, such as Zoom<sup>TM</sup>) remains less certain and, in some cases, may even be beneficial. <sup>9,10</sup> Thus, according to a Nature Human Behaviour editorial, research to determine the effect of screen exposure on youth is "a defining question of our age". <sup>11</sup> With concerns over the impact of screen use including education, health, social development, and psychological well-being, an overview that identifies potential benefits and risks is needed.

Citing the negative effects of screens on health (e.g., increased risk of obesity) and 76 health-related behaviours (e.g., sleep), guidelines from the World Health Organisation<sup>12</sup> and 77 numerous government agencies<sup>13,14</sup> and statements by expert groups<sup>15</sup> have recommended that young people's time spent using electronic media devices for entertainment purposes should be limited. For example, the Australian Government guidelines regarding sedentary behaviour recommend that young children (under the age of two) should not spend any time 81 watching screens. They also recommend that children aged 2-5 years should spend no more than one hour engaged in recreational sedentary screen use per day, while children aged 5-12 and adolescents should spend no more than two hours. However, recent evidence suggests that longer exposures may not have adverse effects on children's behaviour or mental health—and might, in fact, benefit their well-being—as long as exposure does not reach extreme levels (e.g., 7 hours per day)<sup>16</sup>. Some research also indicates that content (e.g., video games vs television programs) plays an important role in determining the potential benefit or harm of youths' exposure to screen-based media. <sup>17</sup> Indeed, educational screen use is positively related to educational outcomes. 18 This evidence has led some researchers to argue that a more nuanced approach to screen use guidelines is required. 19

In 2016, the American Academy of Pediatrics used a narrative review to examine the 92 benefits and risks of children and adolescents' electronic media<sup>20</sup> as a basis for updating their 93 guidelines about screen use. 15 Since then, a large number of systematic reviews and meta-analyses have provided evidence about the potential benefits and risks of screen use. While there have been other overviews of reviews on screen use, these have tended to focus on a single domain (e.g., health<sup>21</sup>), focus on a particular exposure (e.g., social media<sup>22,23</sup>) or provide only a narrative summary of the literature.<sup>24</sup> Focusing on a single domain or exposure makes it difficult to understand what trade-offs are involved in any guidelines around screen use. For example, prohibiting screen use might reduce exposure to advertising 100 but may also thwart learning opportunities from interactive educational tools. Reviews on 101 either of these exposures or outcomes would likely miss being able to quantify these 102

trade-offs. Overviews are one method of evidence synthesis that helps address these
trade-offs, by providing 'user-friendly' summaries of a field of research.<sup>25</sup> These overviews
provide a reference point for the field and allow for easier comparison of risks and benefits
for the same behaviour. By analogy, reading is a sedentary behaviour, and only by
comparing the health risks against the educational benefits can researchers and policymakers
make clear recommendations about what young people should do.

In order to synthesise the evidence and support further evidence-based guideline 109 development and refinement, we reviewed published meta-analyses examining the effects of 110 screen use on children and youth. This review synthesises evidence on any outcome of 111 electronic media exposure. We deliberately did not pre-specify outcomes, in order to get a 112 list of areas where there is meta-analytical evidence. Adopting this broad approach allowed 113 us to provide a holistic perspective on the influence of screens on children's lives. By 114 synthesising across life domains (e.g., school and home), this review provides evidence to 115 inform guidelines and advice for parents, teachers, pediatricians and other professionals in 116 order to maximise human functioning. 117

118 Results

The searches yielded 50,649 results, of which 28,675 were duplicates. After screening titles and abstracts, we assessed 2,557 full-texts for inclusion. Of those, 217 met the inclusion criteria and we extracted the data from all of these meta-analyses. Figure 1 presents the full results of the selection process.

The most frequently reported exposures were physically active video games (n = 31), 123 general screen use (n = 27), general TV programs and movies (n = 20), and screen-based 124 interventions to promote health (n = 14). Supplementary File 1 provides a list of all 125 exposures identified. The most frequently reported outcomes were body composition (n =126 30), general learning (n=24), depression (n=13), and general literacy (n=12). Of the 273 127 unique exposure/outcome combinations, 241 occurred in only one review, with 23 appearing 128 twice, and 9 appearing three or more times. Full characteristics of the included studies are 129 provided in Supplementary File 2. After removing reviews with duplicate exposure/outcome 130 combinations, our process yielded 252 unique effect/outcome combinations (retaining 131 multiple effects for different age groups or study designs) contributed from 102 reviews. 132 These effects represent the findings of 2,451 primary studies, involving 1,937,501 participants. 133 The characteristics of the included effects are available in Supplementary File 3. 134

## TABLE 1

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The quality of the included meta-analyses was mixed (see Table 1). Most assessed heterogeneity (n low risk = 93/102, 91% of meta-analyses), reported the characteristics of the included studies (n low risk = 86/102, 84%), and used a comprehensive and systematic search strategy (n low risk = 71/102, 70%). Most reviews did not clearly report if their eligibility criteria were predefined (n unclear = 71/102, 70%). Many papers also did not complete dual independent screening of abstracts and full text (n high risk = 20/102, 20%) or did not clearly report the method of screening (n unclear = 37/102, 36%). A similar trend

was observed for dual independent quality assessment (n high risk = 52/102, 51%; n high risk = 19/102, 19%). Overall, only 7 meta-analyses were graded as low risk of bias on all criteria.

#### 145 Education Outcomes

There were 88 unique effects associated with education outcomes, including general 146 learning outcomes, literacy, numeracy, and science. We removed 28 effects that did not 147 provide individual study-level data, 19 effects with samples < 1,000, and 19 effects with a 148 significant Egger's test or insufficient studies to conduct the test. Effects not meeting one or 149 more of these standards are presented in Supplementary File 4. The remaining 22 effects met 150 our criteria for statistical credibility and are described in Figure 2. These 22 effects came 151 from 17 meta-analytic reviews analysing data from 337 empirical studies with 262,497 152 individual participants. 153

Among the statistically credible effects, general screen use (r = -0.11, 95%) confidence 154 interval [CI] -0.24 to 0.01, p = 0.071, k = 18, N = 13,100), television viewing (r = -0.10, 1.00)155 95% CI -0.15 to -0.04, p = <0.001, k = 18, N = 62,135), and video games (r = -0.08, 95%156 CI -0.12 to -0.04,  $p = \langle 0.001, k = 10, N = 4,276 \rangle$  were all negatively associated with 157 learning. E-books that included narration (r = 0.11, 95% CI 0.05 to 0.17, p = <0.001, k =158 50, N=2,288), as well as touch screen education interventions (r=0.21,95% CI 0.15 to 159 0.28, p = < 0.001, k = 79, N = 5.810, and augmented reality education interventions (r =160 0.33, 95% CI 0.25 to 0.42, p = <0.001, k = 15, N = 1,474) were positively associated with 161 learning. General screen use was negatively associated with literacy outcomes (r = -0.14, 162 95% CI -0.20 to -0.09, p = < 0.001, k = 38, N = 18,318). However, if the screen use involved co-viewing (e.g., watching with a parent; r = 0.15, 95% CI 0.02 to 0.28, p = 0.028, k = 12, N=6,083), or the content of television programs was educational (r=0.13, 95% CI 0.03 to 165 0.23, p = 0.012, k = 13, N = 1,955), the association with literacy was positive and 166 significant at the 95% confidence level (weak evidence). Numeracy outcomes were positively 167 associated with screen-based mathematics interventions (r = 0.27, 95% CI 0.21 to 0.33, p =168

<0.001, k = 85, N = 36,793) and video games that contained numeracy content (r = 0.32, 95% CI 0.21 to 0.43, p = <0.001, k = 25, N = 2,008).

As shown in Figure 2, most of the credible results (13 of 22 effects) showed statistically 171 significant associations, with 99.9% confidence intervals not encompassing zero (strong 172 evidence). The remaining six associations were significant at the 95% confidence level (weak 173 evidence). All credible effects related to education outcomes were small-to-moderate. 174 Screen-based interventions designed to influence an outcome (e.g., a computer based 175 program designed to enhance learning; r = 0.21, 95% CI 0.15 to 0.28, p = <0.001, k = 79176 N=5,810) tended to have larger effect sizes than exposures that were not specifically 177 intended to influence any of the measured outcomes (e.g., the association between television 178 viewing and learning; r = -0.10, 95% CI -0.15 to -0.04, p = <0.001, k = 18, N = 62,135). 170 The largest effect size observed was for augmented reality-based education interventions on 180 general learning (r = 0.33, 95% CI 0.25 to 0.42, p = <0.001, k = 15, N = 1,474). Most 181 effects showed high levels of heterogeneity (17 of 22 with  $I^2 > 50\%$ ). 182

#### 183 Health-related Outcomes

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We identified 163 unique outcome-exposure combinations associated with health or 184 health-related behaviour outcomes. We removed 39 effects that did not provide individual 185 study-level data, 50 effects with samples < 1,000, and 53 effects with a significant Egger's 186 test or insufficient studies to conduct the test. No remaining studies had statistically 187 significant tests for excess significance. Effects not meeting one or more of these standards 188 are presented in Supplementary File 5. The remaining 21 meta-analytic associations met our 189 criteria for credible evidence and are described below (see also Figure 3). These 21 effects 190 came from 15 meta-analytic reviews analysing data from 344 empirical studies with 859,562 191 individual participants. 192

Digital advertising of unhealthy foods—both traditional advertising (r = 0.23, 95% CI

0.10 to 0.37,  $p = \langle 0.001, k = 13, N = 1,756 \rangle$  and video games developed by a brand for 194 promotion (r = 0.18, 95% CI 0.10 to 0.25, p = <0.001, k = 15, N = 3,842)—were associated 195 with higher unhealthy food intake. Social media use and sexual content were positively 196 associated with risky behaviors (e.g., social media and risky sexual behaviour; r = 0.21, 95%197 CI 0.14 to 0.28, p = <0.001, k = 14, N = 23,096). Television viewing was negatively 198 correlated with sleep duration, but with stronger evidence only observed for adolescents (r =199 -0.06, 95% CI -0.10 to -0.01, p = 0.018, k = 10, N = 9,798). Both television and video games 200 were associated with body composition (e.g., television r = 0.06, 95% CI 0.03 to 0.10, p =201  $<0.001, k = 12, N = 3{,}196$ ). Screen-based interventions which target health behaviours 202 appeared mostly effective. 203

Across the health outcomes, most (14 of 21) effects were statistically significant at the 99.9% confidence interval level, with the remaining four significant at 95% confidence. However, most of the credible effects exhibited high levels of heterogeneity, with all but two having  $I^2 > 75\%$ . Additionally, most effects were small, with the association between internet use and depression the largest at r = 0.25 (95% CI 0.22 to 0.27, p = <0.001, k = 118, N = 527,696). Most of the effect sizes (17/21) had an absolute value of r < 0.2.

210 Discussion

The primary goal of this review was to provide a holistic perspective on the influence of screens on children's lives across a broad range of outcomes. We found that when meta-analyses examined general screen use, and did not specify the content, context or device, there was strong evidence showing potentially harmful associations with general learning, literacy, body composition, and depression. However, when meta-analyses included a more nuanced examination of exposures, a more complex picture appeared.

As an example, consider children watching television programs—an often cited form of screen use harm. We found evidence for a small association with poorer academic

performance and literacy skills for general television watching<sup>27</sup>. However, we also found 219 evidence that if the content of the program was educational, or the child was watching the 220 program with a parent (i.e., co-viewing), this exposure was instead associated with better 221 literacy. Thus, parents may play an important role in selecting content that is likely to 222 benefit their children or, perhaps, interact with their children in ways that may foster 223 literacy (e.g., asking their children questions about the program). Similar nuanced findings 224 were observed for video games. The credible evidence we identified showed that video game 225 playing was associated with poorer body composition and learning. 27,29 However, when the 226 video game were designed specifically to teach numeracy, playing these games showed 227 learning benefits.<sup>30</sup> One might expect that video games designed to be physically active 228 could confer health benefits, but none of the meta-analyses examining this hypothesis met 229 our thresholds for statistical credibility (see Supplementary Files 4 & 5) therefore this hypothesis could not be addressed. 231

Social media was one type of exposure that showed consistent associations with poor health, with no indication of potential benefit. Social media showed strong evidence of harmful associations with risk taking in general, as well as unsafe sex and substance abuse. These results align with meta-analytic evidence from adults indicating that social media use is also associated with increased risk of depression. Recent evidence from social media companies themselves suggest there may also be negative effects of social media on the mental health of young people, especially teenage girls. 4

One category of exposure appeared to be consistently associated with benefits:

screen-based interventions designed to promote learning or health behaviours. This finding

indicates that interventions can be effectively delivered using electronic media platforms, but

does not necessarily indicate that screens are more effective than other methods (e.g.,

face-to-face, printed material). Rather, it reinforces that the content of the screen use may

be the most important aspect. The way that a young person interacts with digital screens

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may also be important. We found evidence that touch screens had strong evidence for benefits on learning, <sup>26</sup> as did augmented reality. <sup>35</sup> 246

Largely owing to a small number of studies or missing individual study data, there 247 were few age-based conclusions that could be drawn from reviews which met our criteria for 248 statistical certainty. Given the differences in development across childhood and adolescence 249 and the different ways children of various ages use screens, further examination of age-based 250 differences is needed. However, in the absence of this work, our study has shown how 251 children are affected by screens in general. 252

Among studies that met our criteria for statistical certainty heterogeneity was high, 253 with almost all effects having  $I^2 > 50\%$ . Much of this heterogeneity is likely explained by differences in measures across pooled studies, or in some cases, the generic nature of some of the exposures. For example, "TV programs and movies" covers a substantial range of content, which may explain the heterogeneous association with education outcomes.

Our results have several implications for policy and practice. Broadly, our findings 258 align with the recommendations of others who suggest that current guidelines may be too 259 simplistic, mischaracterise the strength of the evidence, or do not acknowledge the important 260 nuances of the issue. 36-38 Our findings suggest that screen use is a complex issue, with 261 associations based not just on duration and device type, but also on the content and the environment in which the exposure occurs. Many current guidelines simplify this complex 263 relationship as something that should be minimised. 12,13 We suggest that future guidelines need to embrace the complexity of the issue, to give parents and clinicians specific information to weigh the pros and cons of interactions with screens.

Given our results, we support the continuing trend of guidelines moving away from 267 recommendations to reduce 'screen use', and instead focusing on the type of screen use. For 268 example, we suggest that guidelines should discourage high levels of social media and 269

internet use. Guidelines may also consider adapting recommendations that promote the use
of educational apps and video games, although these recommendations need to be balanced
against the (very small) risks to adiposity.<sup>39</sup>

Our results also have implications for future research. Screen use research is extensive, 273 varied, and rapidly growing. Reviews tended to be general (e.g., all screen use) and even 274 when more targeted (e.g., social media) nuances related to specific content (e.g., Instagram 275 vs Facebook) have not been meta-analysed or have not produced credible evidence. Fewer 276 than 20% of the effects identified met our criteria for statistical credibility. Most studies 277 which did not meet our criteria failed to provide study-level data (or did not provide 278 sufficient data, such as including effect estimates but not sample sizes). Newer reviews were 270 more likely to provide this information than older reviews, but it highlights the importance 280 of data and code sharing as recommended in the PRISMA guidelines. 40 When study level 281 data was available, many effects were removed because the pooled sample size was small, or 282 because there were fewer than ten studies on which to perform an Egger's test. It seems that 283 much of the current screen use research is small in scale, and there is a need for larger, high-quality studies.

Our results highlight the need for the field to more carefully consider if the term 'screen use' remains appropriate for providing advice to parents. Instead, our results suggest that more nuanced and detailed descriptions of the behaviours to be modified may be required.

Rather than suggesting parents limit 'screen use', for example, it may be better to suggest that parents promote interactive educational experiences but limit exposure to advertising.

Screen use research has a well-established measurement problem, which impacts the individual studies of this umbrella review. The vast majority of screen use research relies on self-reported data, which not only lacks the nuance required for understanding the effects of screen use, but may also be inaccurate. In one systematic review on screen use and sleep, 66 of the 67 included studies used self-reported data for *both* the exposure and outcome variable.

It has been established that self-reported screen use data has questionable validity. In a meta-analysis of 47 studies comparing self-reported media use with logged measures, Parry 297 et al<sup>41</sup> found that the measures were only moderately correlated (r = 0.38), with 298 self-reported problematic usage fairing worse (r = 0.25). Indeed, of 622 studies which 299 measured the screen use of 0—6 year-olds, only 69 provided any sort of psychometric 300 properties for their measure, with only 19 studies reporting validity. 42 While some 301 researchers have started using newer methods of capturing screen behaviours—such as 302 wearable cameras<sup>43</sup> or device-based loggers<sup>44</sup>—these are still not widely adopted. It may be 303 that the field of screen use research cannot be sufficiently advanced until accurate, validated, 304 and nuanced measures are more widely available and adopted. 305

There were a number of strengths and limitations to our work. Our primary goal for
this umbrella review was to provide a high-level synthesis of screen use research, by
examining a range of exposures and the associations with a broad scope of outcomes. Our
results represent the findings from 2,451 primary studies comprised of 1,937,501 participants.
To ensure findings could be compared on a common metric, we extracted and reanalysed
individual study data where possible.

Our high-level approach limits the feasibility of examining fine-grained details of the 312 individual studies. For example, we did not examine moderators beyond age, nor did we rate 313 the risk of bias for the individual studies. Thus, our assessment of evidence quality was 314 restricted to statistical credibility, rather than a more complete assessment of quality (e.g., 315 GRADE<sup>45</sup>). As such, we made decisions regarding the credibility of evidence, where others may have used different thresholds or metrics. In addition, when faced with duplicate 317 outcome/exposure combinations we chose to keep the one with the largest pooled sample 318 size, assuming that this would capture the most comprehensive and most recent review. 319 Inspection of the excluded effect sizes suggests that this decision was not that impactful: our 320 results would have been almost exactly the same has we used the number of included studies 321

(k) or the most recent review by publication year. However, we provide the complete results in Supplementary Files 4 & 5, along with the dataset (Supplementary File 6) for others to consider alternative criteria.

Our high-level approach also means that we could not engage with the specific 325 mechanisms behind each association, and as such, we cannot make claims on the directions 326 of causality. These likely depend on the specific exposure and outcome. It is tempting to 327 draw inferences that the associations are due to screen use causing these outcomes, but we 328 cannot rule out reverse causality, a third variable, or some combination of influences. Many 329 of the individual reviews go into more detail about the strength of the evidence for causal 330 associations, but those judgements were difficult to synthesise across more than 200 reviews. 331 Readers who wish to more deeply understand one specific relationship are directed to the 332 cited review for that effect, where the authors could engage more deeply with the 333 mechanisms.

We converted all effect sizes to a common metric (Pearson's r) to allow for comparisons 335 of magnitude, but acknowledge that this assumes a linear relationship between the variables. 336 Some previous research suggests that associations are typically linear. <sup>18</sup> However, others 337 have identified instances where non-linear relationships exist, especially for very high levels 338 of screen use. 17,46,47 Additionally, our conversion may not always adequately account for differences in study design or measures of exposures and outcomes. Care is needed, therefore, when interpreting the effect sizes. In addition, reviews provide only historical evidence which may not keep up with the changing ways children can engage with screens. While our synthesis of the existing evidence provides information about how screens might have 343 influenced children in the past, it is difficult to know if these findings will translate to new 344 forms of technology in the future. 345

Screen use is a topic of significant interest, as shown by the wide variety of academic domains involved, parents' concerns, and the growing pervasiveness into society. Our

findings showed that screen use is associated with both positive (e.g., educational video games were associated with improved literacy) and negative (e.g., general screen use was 340 associated with poorer body composition) outcomes. Based on our findings, we recommend 350 that parents, teachers, and other caregivers need to carefully weigh the pros and cons of each 351 specific activity for potential harms and benefits. However, our findings also lead us to 352 suggest that in order to aid caregivers to make this judgement, researchers need to conduct 353 more careful and nuanced measurement and analysis of screen use, with less emphasis on 354 measures that aggregate screen use and instead focus on the content, context, and 355 environment in which the exposure occurs. 356

Methods 357

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We prospectively registered our methods on the International Prospective Register of 358 Systematic Reviews (PROSPERO; CRD42017076051) in October 2017. We followed the 359 Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) 360 guidelines.<sup>40</sup> 361

Population: To be eligible for inclusion, meta-analyses needed Eligibility criteria. to include meta-analytic effect sizes for children or adolescents (age 0-18 years). We included 363 meta-analyses containing studies that combined data from adults and youth if meta-analytic effect size estimates specific to participants aged 18 years or less could be extracted (i.e., the highest mean age for any individual study included in the meta-analysis was < 18 years). A meta-analysis was still included if the age range exceed 18 years, provided that the mean age was less than 18. We excluded meta-analyses that only contained evidence gathered from adults (age >18 years).

Exposure: We included meta-analyses examining all types of electronic screens 370 including (but not necessarily limited to) television, gaming consoles, computers, tablets, 371 and mobile phones. We also included analyses of all types of content on these devices, 372 including (but not necessarily limited to) recreational content (e.g., television programs, 373

movies, games), homework, and communication (e.g., video chat). In this review we focused 374 on electronic media exposure that would be considered typical for children and youth. That 375 is, exposure that may occur in the home setting, or during schooling. Consistent with this 376 approach, we excluded technology-based treatments for clinical conditions. However, we 377 included studies examining the effect of screen exposure on non-clinical outcomes (e.g., 378 learning) for children and youth with a clinical condition. For example, a meta-analysis of 379 the effect of television watching on learning among adolescents diagnosed with depression 380 would be included. However, a meta-analysis of interventions designed to treat clinical depression delivered by a mobile phone app would be excluded.

Outcomes: We included all reported outcomes on benefits and risks.

Publications: We included meta-analyses (or meta-regressions) of quantitative evidence. 384 To be included, meta-analyses needed to analyse data from studies identified in a systematic 385 review. For our purposes, a systematic review was one in which the authors attempted to 386 acquire all the research evidence that pertained to their research question(s). We excluded 387 meta-analyses that did not attempt to summarise all the available evidence (e.g., a 388 meta-analysis of all studies from one laboratory). We included meta-analyses regardless of 380 the study designs included in the review (e.g., laboratory-based experimental studies, 390 randomised controlled trials, non-randomised controlled trials, longitudinal, cross-sectional, 391 case studies), as long as the studies in the review collected quantitative evidence. We 392 excluded systematic reviews of qualitative evidence. We did not formulate 393 inclusion/exclusion criteria related to the risk of bias of the review. We did, however, employ 394 a risk of bias tool to help interpret the results. We included full-text, peer-reviewed meta-analyses published or 'in-press' in English. We excluded conference abstracts and meta-analyses that were unpublished. 397

Information sources. We searched records contained in the following databases:
Pubmed, MEDLINE, CINAHL, PsycINFO, SPORTDiscus, Education Source, Embase,

Cochrane Library, Scopus, Web of Science, ProQuest Social Science Premium Collection, and ERIC. We conducted an initial search on August 17, 2018 and refreshed the search on September 27, 2022. We searched reference lists of included papers in order to identify additional eligible meta-analyses. We also searched PROSPERO to identify relevant protocols and contacted authors to determine if these reviews have been completed and published.

Search strategy. The search strategy associated with each of the 12 databases can
be found in Supplementary File 7. We hand searched reference lists from any relevant
umbrella reviews to identify systematic meta-analyses that our search may have missed.

Selection process. Using Covidence software (Veritas Health Innovation,
Melbourne, Australia), two researchers independently screened all titles and abstracts. Two
researchers then independently reviewed full-text articles. We resolved disagreements at each
stage of the process by consensus, with a third researcher employed, when needed.

Data items. From each included meta-analysis, two researchers independently
extracted data into a custom-designed database. We extracted the following items: First
author, year of publication, study design restrictions (e.g., cross-sectional, observational,
experimental), region restrictions (e.g., specific countries), earliest and latest study
publication dates, sample age (mean), lowest and highest mean age reported, outcomes
reported, and exposures reported.

Study risk of bias assessment. For each meta-analysis, two researchers independently completed the National Health, Lung and Blood Institute's Quality
Assessment of Systematic Reviews and Meta-Analyses tool<sup>48</sup> (see Table 1). We resolved disagreements by consensus, with a third researcher employed when needed. We did not assess risk of bias in the individual studies that were included in each meta-analysis.

Effect measures. Two researchers independently extracted all quantitative meta-analytic effect sizes, including moderation results. We excluded effect sizes which were reported as relative risk ratios or odds ratios, as meta-analyses did not contain sufficient information to meaningfully convert to a correlation. We also excluded effect size estimates
when the authors did not provide a sample size. Where possible, we also extracted effect
sizes from the primary studies included in each meta-analysis.

To facilitate comparisons, we converted effect sizes to Pearson's r using established formulae. Fifect sizes on the original metric are provided in Supplementary File 6. Throughout the results section we interpret the size of the effects using Funder and Ozer's guidelines: very small (0.05 < r <= 0.1), small (0.1 < r <= 0.2), medium (0.2 < r <= 0.2), large (0.3 < r <= 0.4), and very large (r >= 0.4). These are similar to other interpretations based on empirical data.

Synthesis methods. After extracting data, we examined the combinations of
exposure and outcomes and removed any effects that appeared multiple times (i.e., in
multiple meta-analyses, or with multiple sub-groups in the same meta-analysis), keeping the
effect with the largest total sample size. In instances where effect sizes from the same
combination of exposure and outcome were drawn from different age-groups (e.g., children vs
adolescents), or were drawn using different study designs (e.g., cross-sectional vs
longitudinal) we retained both estimates in our dataset.

We descriptively present the remaining meta-analytic effect sizes. To remove the 443 differences in approach to meta-analyses across the reviews, we reran the effect size estimate 444 using a random effects meta-analysis via the metafor package<sup>53</sup> in R<sup>54</sup> (version 4.3.0) when 445 the meta-analysis's authors provided primary study data associated with these effects. When 446 required, we imputed missing sample sizes using mean imputation from the other studies within that review. From our reanalysis we also extracted  $I^2$  values. To test for publication bias, we conducted Egger's test<sup>55</sup> when the number of studies within the review was ten or more, <sup>56</sup> and conducted a test of excess significance. <sup>57</sup> We contacted authors who did not 450 provide primary study data in their published article. Where authors did not provide data in 451 a format that could be re-analysed, we used the published results of their original 452

453 meta-analysis.

Evidence assessment criteria. Statistical Credibility: We employed a statistical
classification approach to grade the credibility of the effect sizes in the literature. To be
considered 'credible' an effect needed to be derived from a combined sample of >1,000
participants<sup>58</sup> and have non-significant tests of publication bias (i.e., Egger's test and excess
significance test). We performed these analyses, and therefore the review needed to provide
usable study-level data in order to be included.

Consistency of Effect within the Population: We also examined the consistency of the effect size using the  $I^2$  measure. We considered  $I^2 < 50\%$  to indicate effects that were relatively consistent across the population of interest.  $I^2$  values of > 50% were taken to indicate an effect was potentially heterogeneous within the population.

Direction of Effect: Finally, we examined the extent to which significance testing suggested screen exposure was associated with benefit, harm, or no effect on outcomes. We used thresholds of P < .05 for weak evidence (i.e., 95% confidence intervals did not cross zero) and  $P < 10^{-3}$  (i.e., 99.9% confidence intervals did not cross zero) for strong evidence. An effect with statistical credibility but with P > .05 (i.e., 95% confidence intervals included zero) was taken to indicate no association of interest.

Deviations from protocol. As described above, we have summarised the
meta-analytic findings from all included systematic reviews. In our protocol, we originally
planned to also conduct a narrative synthesis of all systematic reviews, even those without
meta-analyses. However, we determined that combining results from the meta-analyses alone
allow readers to compare relative strength of associations more easily. Readers interested in
the relevant systematic reviews (i.e., without meta-analysis) can consult the list of references
in Supplementary File 8.

We altered our evidence assessment plan when we identified that, as written, it could not classify precise evidence of null effects (i.e., from large reviews with low heterogeneity

and low risk of publication bias) as 'credible' because a highly-significant *P*-value was a

criteria. This would have significantly harmed knowledge gained from our review as it would

have restricted our ability to show where the empirical evidence strongly indicated that there

was no association between screen use and a given outcome.

## Data availability statement

All data for this review are available from the authors' GitHub repository

(https://github.com/motivation-and-Behaviour/screen\_umbrella) or from the Open Science

Foundation (https://osf.io/3ubqp/).

# <sup>487</sup> Code availability statement

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All code used in these analyses are available on the authors' GitHub repository

(https://github.com/motivation-and-Behaviour/screen\_umbrella).

# Acknowledgements

The authors received no specific funding for this work.

## Author contributions

TS, MN, PP, and CL conceptualised the review and drafted the manuscript. TS, MN, and PP conducted the analyses. All authors contributed to data extraction, interpretation, and editing of the manuscript.

#### Competing interests

The authors declare no conflicts of interest.

## 498 Tables

Table 1: Review characteristics and quality assessment for meta-analyses providing unique effects

# Figure legends

Figure 1: PRISMA flow diagram.

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- Figure 2: Education outcomes. Forest plot for 22 unique effect sizes related to
  educational outcomes which met the criteria for statistical certainty. Findings are presented
  as correlations (two-sided) with both 95% and 99.9% confidence intervals.
- Figure 3: Health and health-related behaviour outcomes. Forest plot for 21 unique effect sizes related to health and health-related behaviour outcomes which met the criteria for statistical certainty. Findings are presented as correlations (two-sided) with both 95% and 99.9% confidence intervals.

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 $\label{thm:continuous} \begin{tabular}{ll} Table 1 \\ Quality \ assessment \ for \ studies \ providing \ unique \ effects \end{tabular}$ 

		Quality Assessment							
First Author	Year	Elig. Crit. <sup>1</sup>	Lit. Search <sup>2</sup>	Dual Screen <sup>3</sup>	Dual Qual.4	Studies Listed $^5$	Pub. Bias <sup>6</sup>	Hetero. <sup>7</sup>	
Abrami	2020	U	U	L	Н	L	L	L	
Adelantado-Renau	2019	L	${ m L}$	L	L	L	L	L	
Andrade	2019	U	${ m L}$	L	U	L	Н	L	
Arztmann	2022	U	Н	Н	Н	Н	L	L	
Aspiranti	2020	U	L	L	Н	L	Н	L	
Bartel	2015	L	L	U	U	L	U	U	
Beck Silva	2022	L	${ m L}$	L	L	L	Н	L	
Benavides-Varela	2020	U	Н	L	Н	L	${ m L}$	L	
Blok	2002	U	${ m L}$	Н	Н	L	Н	L	
Bossen	2020	U	${ m L}$	L	L	L	Н	L	
Boyland	2016	Н	${ m L}$	L	U	L	L	L	
Byun	2018	U	U	U	Н	Н	Н	Н	
Cao	2020	U	Н	U	Н	L	L	L	
Champion	2019	L	L	L	L	L	L	L	
Chan	2014	U	Н	Н	Н	L	${f L}$	L	
Chauhan	2017	U	${ m L}$	U	Н	Н	${ m L}$	L	
Chen	2020	U	Н	U	Н	Н	Н	L	
Cheung	2012	U	${ m L}$	L	Н	Н	L	L	
Cheung	2013	L	Н	Н	U	L	${ m L}$	L	
Cho	2018	U	Н	U	Н	L	${f L}$	L	

Table 1

Quality assessment for studies providing unique effects (continued)

First Author	Year	Elig.	Lit.	Dual	Dual	Studies	Pub.	Hetero. <sup>7</sup>
		Crit. <sup>1</sup>	Search <sup>2</sup>	Screen <sup>3</sup>	Qual. <sup>4</sup>	Listed <sup>5</sup>	Bias <sup>6</sup>	
Claussen	2022	U	L	U	Н	L	Н	L
Clinton	2019	U	Н	U	U	L	L	L
Comeras-Chueca	2021	L	U	L	U	L	Н	L
Comeras-Chueca	2021	L	L	L	U	L	Η	${ m L}$
Coyne	2018	L	L	L	Н	L	L	L
Cunningham	2021	U	L	L	Н	L	L	L
Cushing	2010	U	L	Н	Н	L	L	L
Darling	2017	U	L	U	U	L	Н	Н
Eirich	2022	U	L	L	L	L	L	L
Feng	2021	L	L	L	L	L	Н	L
Ferguson	2017	U	L	L	Н	L	L	L
Ferguson	2020	L	U	L	L	L	L	L
Folkvord	2018	U	L	L	U	L	Н	L
Furenes	2021	Н	Н	L	U	L	L	L
Gardella	2017	U	L	L	U	L	L	L
Garzón	2019	U	Н	U	Н	Н	L	L
Graham	2015	U	L	Н	Н	L	L	L
Hammersley	2016	L	L	Н	L	L	Н	L
Нао	2021	U	L	L	L	L	Н	L
Hassan-Saleh	2019	U	L	U	U	Н	Н	L
Не	2021	L	L	L	L	L	L	L

Table 1

Quality assessment for studies providing unique effects (continued)

First Author	Year	Elig.	Lit.	Dual	Dual	Studies	Pub.	Hetero. <sup>7</sup>
		Crit. <sup>1</sup>	Search <sup>2</sup>	Screen <sup>3</sup>	Qual. <sup>4</sup>	Listed <sup>5</sup>	Bias <sup>6</sup>	
Hernandez-Jimenez	2019	U	L	Н	L	L	L	${f L}$
Hurwitz	2018	L	L	Н	Н	L	L	${f L}$
Ivie	2020	U	L	L	L	L	L	${ m L}$
Janssen	2020	U	${f L}$	L	L	L	U	L
Kates	2018	U	Н	L	Н	Н	L	L
Kim	2021	U	L	U	L	L	L	${f L}$
Kroesbergen	2003	U	L	U	Н	L	Η	${ m L}$
Kucukalkan	2019	U	L	U	U	Н	L	${ m L}$
Li	2010	U	L	L	U	L	Н	L
Li	2022	L	Н	L	L	L	Н	L
Li	2022	U	Н	L	Н	L	L	${f L}$
Liao	2008	L	Н	Н	L	Н	Н	Н
Liao	2014	U	L	Н	L	L	L	${ m L}$
Liu	2019	U	L	U	Н	L	L	L
Liu	2022	U	Н	U	Н	Н	L	L
Lu	2021	U	L	U	L	L	L	${f L}$
Madigan	2020	U	L	L	U	L	L	${f L}$
Major	2021	U	L	L	Н	L	L	${f L}$
Mallawaarachchi	2022	L	L	L	L	L	L	L
Mares	2005	U	L	Н	Н	L	Н	Н
Mares	2013	U	Н	Н	Н	L	Н	L

Table 1

Quality assessment for studies providing unique effects (continued)

First Author	Year	Elig.	Lit.	Dual	Dual	Studies	Pub.	Hetero. <sup>7</sup>
		Crit. <sup>1</sup>	Search <sup>2</sup>	Screen <sup>3</sup>	Qual. <sup>4</sup>	$\mathrm{Listed}^5$	$\mathrm{Bias}^6$	
Marker	2022	U	${ m L}$	Н	L	L	L	L
Marshall	2004	U	L	Н	Н	Н	Н	L
Martins	2019	U	${ m L}$	U	Н	L	L	L
Martins	2022	L	L	L	L	L	Н	L
Mazeas	2022	L	L	L	L	L	L	L
McArthur	2012	L	L	L	L	L	L	L
McArthur	2018	L	L	L	L	L	L	L
Mei	2018	U	Н	U	L	L	Н	L
Merchant	2014	U	L	Н	Н	Н	Н	L
Neitzel	2022	U	L	Н	Н	L	Н	Н
Oldrati	2020	U	L	U	Н	L	L	L
Paik	1994	U	Н	U	Н	Н	L	Н
Pearce	2016	U	L	Н	Н	Н	L	L
Peng	2011	U	L	U	U	L	Н	L
Powers	2013	U	L	U	Н	L	L	L
Prescott	2018	U	L	U	Н	L	L	L
Reynard	2022	Н	L	L	L	L	L	L
Rodriguez-Rocha	2019	U	L	L	L	L	L	L
Sadeghirad	2016	Н	L	L	L	L	L	L
Scherer	2020	U	Н	U	Н	L	L	L
Schroeder	2013	L	L	U	Н	L	L	L

Table 1

Quality assessment for studies providing unique effects (continued)

First Author	Year	Elig.	Lit.	Dual	Dual	Studies	Pub.	Hetero. <sup>7</sup>
		Crit. <sup>1</sup>	Search <sup>2</sup>	$Screen^3$	Qual. <sup>4</sup>	$\mathrm{Listed}^5$	$\mathrm{Bias}^6$	
Scionti	2019	L	${f L}$	L	Н	${ m L}$	L	L
Shin	2019	U	L	L	L	L	Н	L
Shin	2022	L	Н	L	L	${ m L}$	L	${ m L}$
Slavin	2014	U	Н	Н	Н	L	Н	Н
Strouse	2021	U	L	U	Н	Н	L	L
Takacs	2014	Н	L	U	Н	L	L	L
Takacs	2019	L	L	U	Н	L	L	${f L}$
Tekedere	2016	U	Н	U	U	L	L	${ m L}$
Tokac	2019	U	Н	L	Н	L	L	L
Vahedi	2018	L	L	U	U	L	L	L
van Ekris	2016	U	L	L	L	L	Н	L
Vannucci	2020	U	L	U	Н	L	L	L
Williams	1982	U	U	Н	U	${ m L}$	Н	Н
Wouters	2013	U	Н	U	Н	L	L	L
Xie	2018	U	L	L	Н	L	L	L
Yin	2019	U	Н	U	Н	L	L	L
Zhou	2020	U	L	U	Н	L	L	L

Zucker 2009 L L U H L H L

Note: Items are from the National Health, Lung and Blood Institute's Quality Assessment of Systematic Reviews and Meta-Analyses tool. Note that we excluded the first item of the tool. U = Unclear; L = Low; H = High <sup>1</sup> Eligibility criteria predefined and specified <sup>2</sup> Literature search strategy comprehensive and systematic <sup>3</sup> Dual independent screening and review <sup>4</sup> Dual independent quality assessment <sup>5</sup> Included studies listed with important characteristics and results of each <sup>6</sup> Publication bias assessed <sup>7</sup> Heterogeneity assessed