JAMA | US Preventive Services Task Force | EVIDENCE REPORT

Primary Care Interventions to Prevent Child Maltreatment Updated Evidence Report and Systematic Review for the US Preventive Services Task Force

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IMPORTANCE Child maltreatment, also referred to as child abuse and neglect, can result in lifelong negative consequences.

OBJECTIVE To update the evidence on interventions provided in or referable from primary care to prevent child maltreatment for the US Preventive Services Task Force.

DATA SOURCES PubMed, Cochrane Library, EMBASE, and trial registries through December 18, 2017; references; experts; literature surveillance through July 17, 2018.

STUDY SELECTION English-language fair- and good-quality randomized clinical trials that (1) included children with no known exposure to maltreatment and no signs or symptoms of current or past maltreatment, (2) evaluated interventions feasible in a primary care setting or that could result from a referral from primary care, and (3) reported abuse or neglect outcomes or proxies for abuse or neglect (eg, injury with a specificity for abuse, visits to the emergency department, hospitalization).

DATA EXTRACTION AND SYNTHESIS Two reviewers independently assessed titles/abstracts, full-text articles, and study quality; a third resolved conflicts when needed. When at least 3 similar trials were available, random-effects meta-analyses were conducted.

MAIN OUTCOMES AND MEASURES Direct measures (including reports to child protective services and removal of the child from the home) or proxy measures of abuse or neglect; behavioral, emotional, mental, or physical well-being; and harms.

RESULTS Twenty-two trials (33 publications) were included (N = 11132). No significant association was found between interventions and reports to child protective services within 1 year of intervention completion (10.6% vs 11.9%; pooled odds ratio [OR], 0.94 [95% CI, 0.72-1.23]; 10 trials [n = 2444]) or removal of the child from the home within 1 to 3 years of follow-up (3.5% vs 3.7%; pooled OR, 1.09 [95% CI, 0.16-7.28]; 4 trials [n = 609]). No statistically significant associations were observed between interventions and outcomes for emergency department visits in the short term (<2 years), hospitalizations, child development, school performance, and prevention of death. Nonsignificant results from single trials led to a conclusion of insufficient evidence for injuries, failure to thrive, failure to immunize, school attendance, and other measures of abuse or neglect. Inconsistent results led to a conclusion of insufficient evidence for long-term (≥2 years) outcomes for reports to child protective services (ORs range from 0.48 to 1.13; 3 trials [n = 1690]), emergency department visits (1 of 2 trials reported significant differences) and internalizing and externalizing behavior symptoms (3 of 6 trials reported reductions in behavior difficulties). No eligible trials on harms of interventions were identified.

CONCLUSIONS AND RELEVANCE Interventions provided in or referable from primary care did not consistently prevent child maltreatment. No evidence on harms is available.

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Corresponding Author: Meera Viswanathan, PhD, RTI International, 3040 E Cornwallis Rd, Research Triangle Park, NC 27709 (viswanathan@rti.org). hild abuse and neglect, also referred to as child maltreatment—words or actions that cause or fail to protect children from harm, potential harm, or the threat of harm¹—is associated with negative physical and emotional health outcomes that persist and can lead to serious disorders throughout the life course. ²⁻⁶ Injuries may include brain injuries, blindness, and fractures⁷ and can lead to disability or death. ⁸ The goal of interventions to prevent child maltreatment is to reduce or eliminate exposure to abuse and neglect, improve well-being, and reduce mortality.

This review was conducted to inform the US Preventive Services Task Force (USPSTF) in its update of the 2013 recommendation on interventions relevant to US primary care to prevent child maltreatment. In 2013, the USPSTF concluded that the evidence was insufficient (I statement) to assess the balance of benefits and harms.

Methods

Scope of the Review

The analytic framework and key questions (KQs) that guided the review are shown in **Figure 1**. Detailed methods, evidence tables, sensitivity analyses, and contextual information are available in the full evidence report at http://www.uspreventiveservicestaskforce.org/Page/Document/UpdateSummaryFinal/child-maltreatment-primary-care-interventions1.

Data Sources and Searches

PubMed, the Cochrane Library, and EMBASE were searched for English-language articles published from November 1, 2011, through December 18, 2017. ClinicalTrials.gov, Cochrane Clinical Trials Registry, and the World Health Organization International Clinical Trials Registry Platform were also searched. To supplement systematic electronic searches (eMethods 1 in the Supplement), reference lists of pertinent articles and studies suggested by reviewers were reviewed. Since December 18, 2017, ongoing

surveillance was conducted through article alerts and targeted searches of journals to identify major studies published in the interim that may affect the conclusions or understanding of the evidence and the related USPSTF recommendation. The last surveillance was conducted on July 17, 2018, and identified no eligible studies.

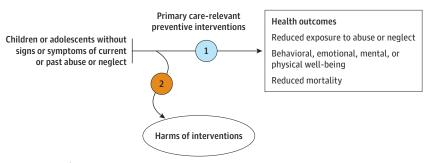
Study Selection

Two investigators independently reviewed titles, abstracts, and full-text articles using prespecified inclusion criteria for each KQ (eTable 1 in the Supplement); disagreements about inclusion were resolved by discussion.

To be eligible, study samples had to have a majority of children (>50%) without known exposure to maltreatment and no signs or symptoms of current or past maltreatment. Studies of symptomatic children undergoing diagnostic evaluation for conditions related to abuse or neglect or asymptomatic children with known exposure to child maltreatment were excluded. All interventions that were feasible in a primary care setting or for which a primary care clinician could give a referral were eligible, with the exception of community-wide programs. Eligible comparators included usual care, delayed interventions, and active interventions that allow for assessment of the independent contribution of the primary care–feasible or primary care–referable preventive intervention.

Studies needed to report at least 1 direct or proxy measure of abuse or neglect to be eligible. Direct measures included those reflecting physical, sexual, or emotional abuse perpetrated by a parent or caregiver; physical (eg, failure to thrive), emotional, dental or medical (eg, lack of immunizations or well-child visits), or educational neglect; reports to Child Protective Services (CPS); and removal of the child from the home. Proxy measures included injuries (eg, fractures, bruises, burns, nonaccidental injuries [injuries with a high specificity for abuse, 12 regardless of whether the injury was intentional]), visits to the emergency department, and hospitalizations. Parent-reported measures of abuse or neglect were summarized in response to

Figure 1. Analytic Framework: Primary Care Interventions to Prevent Child Maltreatment

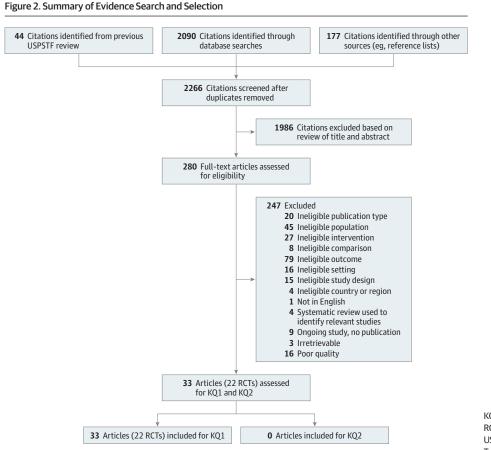


Key questions

Do primary care-feasible or -referable interventions to prevent child maltreatment reduce exposure to abuse or neglect; improve behavioral, emotional, physical, or mental well-being; or reduce mortality among children and adolescents without obvious signs or symptoms of abuse or neglect?

What are the harms of primary care-feasible or -referable interventions to prevent child maltreatment?

Evidence reviews for the US Preventive Services Task Force (USPSTF) use an analytic framework to visually display the key questions that the review will address to allow the USPSTF to evaluate the effectiveness and safety of a preventive service. The questions are depicted by linkages that relate interventions and outcomes. Refer to the USPSTF Procedure Manual for further details. ¹¹ KQ indicates key question.



KQ indicates key question; RCT, randomized clinical trial; USPSTF, US Preventive Services Task Force.

a contextual question (eResults 2 in the Supplement). For studies that reported direct or proxy measures of abuse or neglect (other than self-report), behavioral, emotional, mental, or physical wellbeing outcomes were also assessed.

Randomized clinical trials (RCTs) and systematic reviews were eligible for all KQs; observational study designs were also eligible for harms of screening (KQ2). Only studies that had been published in English and conducted in the 49 countries categorized as very highly developed by the 2015 Human Development Index were included.¹³

Data Extraction and Quality Assessment

For each included study, 1 investigator extracted information about design, population, intervention, and outcomes, and a second investigator reviewed the information for completeness and accuracy. Two independent investigators assessed the quality of each study as good, fair, or poor, using predefined criteria developed by the USPSTF (eMethods 2 in the Supplement)¹⁴ and others¹⁵ for assessing the risk of bias of trials. Individual study quality ratings are reported in eTables 2 through 6 in the Supplement.

Data Synthesis and Analysis

Findings were qualitatively synthesized for each KQ in tabular and narrative formats. Studies were included in the main analysis if they met all study selection criteria and were fair or good quality; this

included studies from the prior review that informed the USPSTF 2013 recommendation that continued to meet the study selection criteria for this update.¹⁶ Qualitative and quantitative sensitivity analyses were conducted by adding poor-quality studies. When at least 3 independent and similar RCTs were available, randomeffects models using the inverse-variance-weighted method of DerSimonian and Laird was used to estimate pooled effects for pooled odds ratios (ORs). Statistical heterogeneity was assessed using the *I*² statistic.¹⁵ All quantitative analyses were conducted using Comprehensive Meta Analysis (Version 3.3) software.¹⁷

All testing was 2-sided and assumed statistical significance if the 95% confidence intervals of pooled estimates did not cross the null value. The strength of evidence for each outcome was assessed as high, moderate, low, or insufficient using methods developed for the USPSTF (and the Evidence-based Practice Center program), based on the quality of studies, consistency of results between studies, precision of findings, and risk of reporting bias for each intervention comparison and major outcome of interest. ^{14,18}

Results

Twenty-two RCTs (N = 11132) of good or fair quality, from 33 publications, were included (Figure 2). Table 1 summarizes study characteristics, and eTables 7 through 11 in the Supplement provide

Table 1. Characteristics of Studies of Interventions to Prevent Child Maltreatment

Characteristics and Subcharacteristics	No. of Studies (%) (N = 22)
Study Quality	
Good	4 (18)
Fair	18 (82)
Population Characteristics	
Timing of enrollment	
Enrolled in prenatal period or immediately after birth	13 (59)
Mixed enrollment	1 (5)
Enrolled after the perinatal period	8 (36)
Maltreatment reported at baseline	
Yes	6 (27)
No	16 (73)
Risk status	
Parent identified to be at risk	12 (55)
Child identified to be at risk because of birth status (premature or low birth weight)	2 (9)
Participants not specifically identified to be at risk	7 (32)
Age of mothers	
Most or all younger than 20 y	7 (32)
≥20 y on average	15 (68)
Intervention Characteristics	
Home visit component	
Yes	21 (95)
No	1 (5)
Clinical personnel involved in care	
Yes	17 (77)
No	5 (23)
Usual-Care Comparator	
Yes	19 (86)
No	3 (14)
Geographic Setting	
United States of America	16 (73)
United Kingdom	3 (14)
Canada	1 (5)
Australia	1 (5)
New Zealand	1 (5)

details. The evidence base spans more than 3 decades; the earliest included study recruited participants in 1976¹⁹ and the most recent through 2010.²⁰ Twenty-one of 22 trials (95%) had a home visiting component; some trials present results from attempting a similar model of home visiting interventions in different settings (eg, the Healthy Families intervention in Alaska^{21,22} and New York^{23,24}; replications^{20,25} of the Nurse Family Partnership²⁶⁻³²). Fifteen trials (68%) recruited mothers younger than 20 years, on average. Nineteen trials (86%) included a usual-care comparator. Sixteen trials (73%) were conducted in the United States. In other respects, however, the evidence base is heterogenous in study populations (time of enrollment, age of infant, experience of maltreatment) and interventions (number, intensity, and duration of components; use of clinical personnel).

Regarding enrolled populations, 13 trials enrolled mothers before or immediately after birth; the mean age of children in the 9 other trials ranged from younger than 6 months to 8 years. One study enrolled fathers. Six trials (27%) reported on whether participants had experienced maltreatment at baseline; other trials either did not specify or enrolled women during pregnancy.

Twelve trials (55%) identified participants or children to be at risk. Risk factors varied across studies and included infant health status or child emotional or behavioral concerns, maternal depression or other mental health problems, presence or history of intimate partner violence, parental substance abuse history, and demographic and socioeconomic characteristics of the mother or family (eg, adolescent parent, poverty, marital status).

Regarding interventions, only 1 trial³³ did not feature home visits. It focused on behavioral therapy for male patients entering outpatient alcohol treatment who had legal guardianship of at least 1 child aged 8 to 12 years. For the trials with home visit components, the content, use of other components, personnel, intensity, and duration varied. Although the specific intervention goals of the home visiting program varied by trial, trials described the following activities: assessing family needs; developing a relationship between the home visitor and the client; providing information, referral, and parenting education; promoting child health, safety, and development; providing clinical care; enhancing family functioning and supporting positive child-parent interactions; building supportive social networks; and creating family plans to support parental life course development and self-sufficiency. Five trials reported home visit services without a clinical component or personnel. 21-24,34-37 Sixteen trials provided home visits in the context of clinical support, which included nurses or mental health professionals supervising home visitors, serving as home visitors, or providing comprehensive pediatric service as an intervention component. 19,20,25-28,38-50

Of the 21 home visit trials, 8 had nurses as home visitors, ^{20,25-28,39,41,42,45,46,49} 1 had mental health clinicians as home visitors, ⁴⁷ 6 had paraprofessional home visitors, ^{19,21-24,36,38,40} 3 included home visitors with early childhood expertise, ^{43,45-47} 3 included social workers, ^{25,41,42,45,46} and 1 had peer home visitors. ⁴⁸ The remaining trials did not specify the training of the home visitors. ^{34,35,37,44,50,51}

The duration of the intervention ranged from 3 months^{19,33} to 3 years.^{41,42,45,46} The planned number of home visits, when reported, ranged from 5 sessions⁴⁹ to 41 sessions.^{34,35}

Benefits of Interventions

Key Question 1. Do primary care–feasible or primary care–referable interventions to prevent child maltreatment reduce exposure to abuse or neglect; improve behavioral, emotional, physical, or mental well-being; or reduce mortality among children and adolescents without obvious signs or symptoms of abuse or neglect?

Direct or Proxy Measures of Abuse or Neglect

Reports to Child Protective Services | Thirteen trials (14 publications)^{19,21,23,25,28,33,34,36,37,39,41,42,44,47} reported on reports to CPS, and 1 trial²⁰ reported on safeguarding actions (eTable 12 and eTable 13 in the Supplement). All eligible trials reported their first results during the intervention (1 year from baseline), at the end

Figure 3. Pooled Results: Reports to Child Protective Services and Removal of the Child From the Home

A Reports to Child Protective Services

		Follow-up	No. of Reports/ Total No. of Children		Odds Ratio	Favors		Favors	
Source	Comparison		Intervention	Control	(95% CI)	Intervention	Control		
Brooten et al, ³⁹ 1986	Home visits	18 mo	2/39	4/40	0.49 (0.08-2.82)				
Duggan et al, ²¹ 2007	Home visits	1 y	30/151	25/158	1.31 (0.73-2.35)		-	_	
DuMont et al, ²³ 2008	Home visits	1 y	41/524	32/536	1.35 (0.84-2.17)		-	-	
Fergusson et al, ⁴¹ 2005	Home visits	3 y	36/184	44/207	0.90 (0.55-1.47)		-	}	
Finello et al, ⁴⁴ 1998	Home health	6 mo	1/20	0/20	3.15 (0.12-82.16))		-	
Lam et al, ³³ 2009	Combined	12 mo	3/20	3/10	0.39 (0.04-3.77)				
Lowell et al, ⁴⁷ 2011	Home visits	2 y	NA	NA	0.59 (0.25-1.39)		_	_	
Olds et al, ²⁸ 1986	Combined	2 y	12/181	16/161	0.61 (0.22-1.65)			_	
Sadler et al, ²⁵ 2013	Home visits	2 y	0/44	2/34	0.15 (0.01-3.15)		-		
Silovsky et al, ³⁷ 2011	Home visits	2 y	10/48	18/57	0.57 (0.23-1.39)		-	_	
Pooled estimate			135/1211	144/1223	0.94 (0.72-1.23)		<	>	
Random-effects meta-analysis	s: $I^2 = 6.3\%$; $P = .64$					0.04		11111111	100
						0.01	0.1 Odds Ratio	l 10 o (95% CI)	100

B Removal of child from home

		No. of Children Removed/ Total No. of Children		Odds Ratio	Favors	Favors	
Source	Time Point, mo	Intervention	Control	(95% CI)	Intervention	Control	
Brayden et al, ³⁸ 1993	36	5/141	1/122	4.45 (0.51-38.61)	_		
Brooten et al, ³⁹ 1986	12	0/39	2/40	0.19 (0.01-4.19)	←	 	
McIntosh et al, ³⁵ 2009	12	4/68	0/63	8.86 (0.47-167.97) —		→
Quinlivan et al, ⁴⁹ 2003	12	2/65	8/71	0.25 (0.05-1.22)		÷	
Pooled estimate Random-effects meta-analysis: I ² = 61.8%; P	=.93	11/313	11/296	1.09 (0.16-7.28)			
,				1	0.01 0.1	1 10 io (95% CI)	100

The size of the boxes indicates the relative contribution of each study to the pooled estimate; error bars, 95% CIs for each study's estimate of effect; centers of diamonds, pooled estimates; width of diamonds, 95% CIs for pooled estimates. A, For studies with multiple groups, the pooled estimates average the treatment effect from active groups and are reported as "combined."

Finello et al reported results from a combination of home health and home visits, home health alone, and home visits alone. Because that trial reported cases only for the home health group, the pooled analysis includes the home health group only. Lowell et al did not report number of events per group; the total N analyzed for both groups is 117. NA indicates not available.

of the intervention, or within 1 year of completion of the intervention. Four trials reported outcomes at 1 or more time points after the first analysis of results. The timing of these reports varied: within 6 months of the initial results, 44 1 year 23,47 to 2 years 47 after the initial results, or over the longer term (6 years after the initial results, 23 when the child was 7 years of age 24 ; 13 years after the initial results, when the child was 15 years of age $^{30-32}$). The pooled OR from 10 trials, all having reported results within 1 year of completion, suggested association between the intervention and reports to CPS (OR, 0.94 [95% CI, 0.72-1.23]; l^2 = 6.3%) (Figure 3). Trials reporting additional results within 6 months 44 or 1 year 23,47 of the original results also reported no statistically significant differences between the groups.

Trials measuring outcomes for later time points provided mixed results. ^{24,30-32,47} No patterns emerged regarding features of the intervention or characteristics of the study sample across the trials showing benefit.

Removal of Child from Home | Five trials^{34,35,38,39,48,49} reported on outcomes related to removal of the child from the home. Four of 5 trials contributed to a pooled analysis of removal of the child

from the home across time points ranging from 12 months to 3 years after baseline (eTable 14 in the Supplement). 34,35,38,39,49 The results show no statistically significant associations between study groups and the outcome (3.5% [11/313] vs 3.7% [11/296]; pooled OR, 1.09 [95% CI, 0.16-7.28]; $I^2 = 61.8\%$; 4 trials [n = 609]) (Figure 3).

Other Measures of Abuse or Neglect | Two RCTs 38,40 reported on study-specific measures of abuse (eTable 15 and eTable 16 in the Supplement). These measures included physical abuse, 38 neglect, 38 and results from the Framingham Safety Survey about household hazards. 40 One trial reported no statistically significant differences, finding 13 of 141 cases (9.2%) of physical abuse in the intervention group vs 8 of 122 (6.6%) in the comparator group (risk ratio [RR], 1.4 [95% CI, 0.58-3.62]). 38 The same study 38 reported 15 of 141 cases (10.6%) of neglect in the intervention group vs 5 of 122 (4.1%) in the comparator group (RR, 2.79 [95% CI, 0.98-7.91]). 38 The other trial reported a statistically significant difference (P = .03), but the clinical importance of differences in the Framingham Safety Survey score is unclear. 40 That trial reported mean values on the Framingham Safety Survey

score of 1.72 (intervention) vs 1.68 (comparator); higher scores represent greater safety.

Injuries With a High Specificity for Abuse or Neglect | One trial found no statistically significant differences in the rates of nonaccidental injuries (O/64 for the intervention vs 1/71 for the comparator; calculated RR, 0.37 [95% CI, 0.02-8.91]) (eTable 17 in the Supplement).

Emergency Department Visits | Eleven trials reported on emergency department visits (eTable 18 and eTable 19 in the Supplement). 19-22,26-32,34,35,41-46,50,51 The timing of outcome measurement varied substantially across trials, ranging from 6 months to more than 4 years. In addition, outcome measures varied and included (1) mean number of all-cause emergency department visits; (2) mean number of emergency department visits for accidents, injuries, and ingestions; (3) number of children seen in the emergency department; (4) number of children seen for accidents or injuries; (5) number of children seen for injuries or ingestions, and (6) total emergency department visits. These variations precluded pooling. Several trials presented outcomes at multiple periods. The results were generally inconsistent in direction of effect.

Hospitalization: Findings | Twelve trials reported on hospitalization outcomes (eTable 20 and eTable 21 in the Supplement). 19-22,26,29,34,39,41,42,44,46,49,51 Because of substantial heterogeneity in outcome definitions and periods of interest, results could not be pooled. Outcomes varied in degree of specificity to child abuse and neglect and included (1) number of children with hospital admission as a result of an injury that was "referred for independent investigation by the Family and Children's Services staff and concluded to have arisen as a result of a nonaccidental injury to the neonate"49; (2) number of children hospitalized because of child abuse and neglect⁴¹; (3) proportion of children hospitalized because of injury or ingestion²⁰; (4) number of children hospitalized for ambulatory care-sensitive conditions; (5) number of children rehospitalized (at 14 days and 18 months, the original cause was not specified); (6) number of children with all-cause hospitalization; (7) mean number of all-cause hospitalizations; (8) total counts of hospital visits; (9) mean number of hospital days; and (10) types of injuries reported among those hospitalized. Results were not statistically significant for 7 outcomes. One of 5 trials that reported on the number of children with all-cause hospitalization demonstrated a significant reduction, but only for 1 of 4 outcome measures.⁴⁴ One of 2 trials found a lower mean number of hospital days, ²⁹ and 1 trial found lower overall rates of hospital admission for unintentional injury at a 9-year follow-up. 42 These results suggest an overall lack of benefit for the active intervention group(s).

Failure to Thrive | One trial reported on failure to thrive. It found no statistically significant differences between study groups for this outcome (0% [0/39] vs 2.5% [1/40]; RR, 0.34 [95% CI, 0.01-8.14]) (eTable 22 in the Supplement).³⁹

Failure to Immunize | One trial reported on failure to immunize. It found no statistically significant differences between study groups in the rate of no vaccinations at 6 months (calculated RR, 0.49 [95% CI, 0.16-1.52]) (eTable 23 in the Supplement).⁴⁹

Behavioral, Emotional, Mental, or Physical Well-being

Internalizing and Externalizing Behavior | Six trials reported on internalizing (depression, anxiety) and externalizing (disruptive, aggressive, or delinquent) behavioral outcomes in children (eTable 24 and eTable 25 in the Supplement). 21-24,26,27,41,42,45-47 The evidence included substantial heterogeneity in the timing and type of outcome measurement, and findings were inconsistent. Three of 6 trials found a reduction in behavioral difficulties in children in primary care interventions to prevent child maltreatment. 21,22,41,42,47

Social, Emotional, and Other Developmental Outcomes Not Otherwise Categorized | Five trials evaluated discrete social, emotional, or other developmental outcomes separately from overall measures of externalizing or internalizing problems (eTable 26 and eTable 27 in the Supplement). ^{24,27,34,35,45-47} The heterogeneity of outcomes precluded meta-analysis, but all trials reported results that were not statistically significant.

Child Development as Measured by the Bayley Scales of Child

Development | Four trials^{21,22,26-32,34} reported on child development as measured by the Bayley Scales of Child Development (eTable 28 and eTable 29 in the Supplement). The results generally indicated no statistically significant differences between intervention and control groups, with the exception of 1 trial. This trial found a statistically significant difference in the Bayley Mental Development Index among children in the experimental group (mean score, 88 vs 84.8; difference, 3.2 [95% CI, 1.2-5.2]; <85 is the threshold for mild delay). The mean difference between the 2 groups was not statistically significant for the psychomotor index.

Other Developmental Outcomes | Three trials reported no consistent differences in other developmental outcome measures (infant development tests, parent concerns about infant development) (eTable 30 and eTable 31 in the Supplement). 20,28,46

School Performance and Attendance | One trial found no statistically significant differences in school performance at 9 years of age. One trial reported on school attendance; children at age 7 years in the intervention group were less likely to report skipping school than children in the usual-care group (2.35% [9/388] vs 6.47% [26/405]; RR, 0.36 [95% CI, 0.17-0.76]) (eTable 32 and eTable 33 in the Supplement). The same study reported no statistically significant differences using maternal reports of skipping school.

Death | Four trials of fair quality reported on the outcome of child death (eTable 34 in the Supplement); outcome specification and follow-up times varied and precluded quantitative synthesis. 26.27.34.39.49 No study reported statistically significant differences in the rates of child death between intervention and usual-care groups. One trial reported mortality at the 6-month follow-up, 49 1 at 12 months, 34 and 1 at 18 months. 39 Another trial reported child deaths at the 9-year follow-up. 26.27 One trial included deaths attributed to sudden infant death syndrome. 39 One trial included only those deaths for which a child protection concern was known and an open

Table 2. Summary of Evidence of Interventions to Prevent Child Maltreatment^a

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Outcome	Population, Intervention	No. of Studies (No. of Observations)	Summary of Findings by Outcome	Consistency and Precision	Body of Evidence Limitations (Including Reporting Bias)	Strength of Evidence	Applicability
KQ1: Benefits of Primary	y Care-Feasible or Pri	mary Care-Refera	ble Interventions				
Reports to CPS	Caregivers of children at risk of maltreatment	14 (4958)	Reports at or within 1 y of trial completion: 11.1% vs 11.8% ; OR, 0.94 (95% CI, $0.72-1.23$); I^2 =6.3%; 10.2434 participants Mixed results for long-term follow-up ^b	Consistent, imprecise (short-term outcomes) Inconsistent, imprecise (long-term outcomes)	Heterogeneity across studies in type of intervention; no evidence of reporting bias	Low for no benefit for short-term outcomes Insufficient for long-term outcomes	Unclear whether findings apply to subgroups defined by parent risk factors
Removal of the child from home	Infants and toddlers <3 y	5 (885)	At 0-3 y: 11/313 (3.51%) vs 11/296 (3.71%); OR, 1.09 (95% CI, 0.16-7.28); I^2 = 61.8%; 4609 participants At birth (for intervention started in pregnancy) in 1 study: calculated OR, 1.55 (95% CI, 0.61-3.94)°; 225 participants	Inconsistent, imprecise	Heterogeneity across studies in timing of outcome; no evidence of reporting bias	Low for no benefit	Unclear whether findings apply to subgroups defined by parent risk factors
Other measures of abuse or neglect	Caregivers (mothers or families)	2 (461)	Abuse ^d : 13/141 (9.2%) vs 8/122 (6.6%); RR, 1.4 (95% CI, 0.58-3.62); 1263 participants Neglect: e 15/141 (10.6%) vs 5/122 (4.1%); RR, 2.79 (95% CI, 0.98-7.91); 1 trial, 263 participants Significantly higher safety scores in intervention group; 1 trial, 147 participants	Inconsistent, imprecise	Heterogeneity across studies in outcome measures; no evidence of reporting bias	Insufficient	Unclear whether findings apply to subgroups defined by parent risk factors
Injuries with a high specificity for abuse	Adolescent mothers	1 (136)	Nonaccidental injuries: 0/64 (0%) vs 1/71 (1.4%); calculated RR, 0.37 (95% CI, 0.015-8.91)	Consistency unknown (single trial), imprecise	Single small trial; no evidence of reporting bias	Insufficient	Unclear whether findings apply to subgroups defined by parent risk factors
Visits to the ED	Children	11 (5732)	2 of 7 studies reported a statistically significant difference in mean number of all-cause ED visits the first 2 y of follow-up; all other studies reported results that are not statistically significant? 1 of 2 studies reported statistically significant results at 2- to 4-y follow-up for each of the following: mean number of all-cause ED vists; mean number of ED visits for accidents, injuries, and ingestions; and number of children seen for accidents or injuries 2 studies found no statistically significant differences for number of children seen in the ED; 1 study found no statistically significant difference in the proportion of children seen for injuries and ingestions 1 of 2 studies reported statistically significant differences at long-term follow-up	Inconsistent, imprecise	Heterogeneity across studies in outcome measures; no evidence of reporting bias	Low for no benefit for short-term outcomes, insufficient for long-term outcomes	Unclear whether findings apply to subgroups defined by parent risk factors

(continued)

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Clinical Review & Education US Preventive Services Task Force

Table 2. Summary of Evidence of Interventions to Prevent Child Maltreatment^a (continued)

Outcome	Population, Intervention	No. of Studies (No. of Observations)	Summary of Findings by Outcome	Consistency and Precision	Body of Evidence Limitations (Including Reporting Bias)	Strength of Evidence	Applicability
Hospitalization	Infants	12 (5491)	1 of 5 studies showed a reduction in number of children with all-cause hospitalization, but only for 1 of 4 outcome measures 1 of 2 studies found a lower mean number of hospital days 1 trial found lower overall rates of hospital admission for unintentional injury at 9-y follow-up All other outcomes not statistically significantly different9	Consistent, imprecise for results less than 3 y; inconsistent, imprecise for long-term follow-up	Heterogeneity outcome measures; each outcome/timing only presented in a single study; no evidence of reporting bias	Low strength of evidence of no benefit	Unclear whether findings apply to subgroups defined by parent risk factors
Failure to thrive	Infants	1 (79)	0/39 (0%) vs 1/40 (2.5%); RR, 0.34 (95% CI, 0.01-8.14)	Consistency unknown (single trial), imprecise	Single small trial; no evidence of reporting bias	Insufficient	Unclear whether findings apply to subgroups defined by parent risk factors
Failure to immunize	Adolescent mothers	1 (136)	No vaccinations at 6 mo: 4/71 (5.6%) vs 9/65 (13.8%); calculated RR, 0.49 (95% CI, 0.16-1.52)	Consistency unknown (single trial), imprecise	Single small trial; no evidence of reporting bias	Insufficient	Unclear whether findings apply to subgroups defined by parent risk factors
Internalizing and externalizing behavior symptoms	Caregivers of children at risk of maltreatment	6 (5529)	3 of 6 trials reported reductions in behavior difficulties ^h Other outcomes not statistically significantly different ⁱ	Inconsistent, imprecise	Small number of trials; heterogeneity of outcome measures; no evidence of reporting bias	Insufficient	Home-based intervention targeting high-risk families may be effective in decreasing behavior problems
Other social, emotional, and developmental outcomes	Infants and toddlers <3 y	4 (3965) children	O of 5 studies reported statistically significant differences on a variety of social, emotional, and developmental measures ^j	Consistent, imprecise	Heterogeneity outcome measures; each outcome and timing only presented in a single study; no evidence of reporting bias	Low strength of evidence of no benefit for children <3 y	Unclear whether findings apply to subgroups defined by parent risk factors; one intervention may not be readily generalizable to other (pediatric practice) settings
Bayley Scales of Development	Caregivers and families	4 (1638) caregivers and families	1/4 trials reported higher scores in the intervention group (mean difference between groups, 3.2 [95% CI, 1.2-5.2])	Consistent, imprecise	Outcomes measured at different ages; no evidence of reporting bias	Low for no benefit	All studies focused on at-risk caregivers and families
Other measures of development	Pregnant mothers	3 (3204)	1 of 3 trials reported statistically significant differences on other development outcomes, but for subset of reported outcome measures and timing	Consistent, imprecise	Heterogeneity in outcome measures; no evidence of reporting bias	Low for no benefit	Unclear whether findings apply to subgroups defined by parent risk factors
School performance	School-aged children	1 (1139)	1 study found no statistically significant difference on various school performance measures	Consistency unknown (single trial), imprecise	Single trial; no evidence of reporting bias	Low for no benefit	Single study, applicability to other settings and ages unclear
School attendance	School-aged children and their families	1 (1184)	Self-reported school attendance at age 7 y: 9/388 (2.35%) vs 26/405 (6.47%); RR, 0.36 (95% CI, 0.17-0.76) No statistically significant difference in maternal reports of skipping school	Consistency unknown (single trial), imprecise	Single trial; inconsistent self-report and maternal report; no evidence of reporting bias	Insufficient	Single study, applicability to other settings and ages unclear

(continued)

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Table 2. Summary of Evidence of Interventions to Prevent Child Maltreatment^a (continued)

Outcome	Population, Intervention	No. of Studies (No. of Observations)	Summary of Findings by Outcome	Consistency and Precision	Body of Evidence Limitations (Including Reporting Bias)	Strength of Evidence	Applicability		
Death	Pregnant or postpartum women; 3 studies included only women at risk for maltreatment; all studies included home visiting	4 (1065)	0/4 trials reported statistically significant differences in death	Consistent, imprecise	Heterogeneity in included studies; no evidence of reporting bias	Low for lack of effect on outcome of death	Unclear whether findings apply to subgroups defined by parent risk factors		
Composite maltreatment outcome ^k	Mothers of newborns	1 (136)	2/65 (3.1%) vs 9/71 (12.7%); RR, 0.24 (95% CI, 0.05-1.08) Adjusted RR, 0.22 (95% CI, 0.02 to 0.98); P = .04	Consistency unknown (single trial), imprecise	Single small trial; no evidence of reporting bias	Insufficient	Unclear whether findings apply to subgroups other than teenaged first-time mothers		
KQ2: Harms of Primary C	KQ2: Harms of Primary Care-Feasible or Primary Care-Referable Interventions								
Harms	NA	0 (0)	No eligible studies	NA	NA	Insufficient	NA		

Abbreviations: CPS, child protective services; ED, emergency department; KQ, key question; NA, not applicable; OR, odds ratio; RR, relative risk.

(1 study), proportion of children with ED visits for injuries and ingestions (1 study), number of children using the ED (2 studies), and total ED visits (1 study).

- ^h One study reported statistically significant differences on each of the following: mean and proportion of children with higher externalizing behaviors at 12 months; internalizing behaviors at 2 years and 3 years; behavior problems at 5, 6, and 9 years; and more maternal concerns on the child behavior checklist.
- i Outcomes with no statistically significant results include internalizing behaviors at 6 and 12 months (1 study); child behavior at 2 years (1 study), 30 to 33 months and 5.5 years (1 study), and 7 years (1 study); and internalizing and externalizing behaviors at 9 years (1 study).
- ^j Outcomes included dysregulation, sleep problems, social skills, attention and social problems, school-related conduct outcomes, and infant social and emotional adjustment.

^a All bodies of evidence were rated as fair quality.

^b Long-term CPS reports: adjusted OR, 0.48 (95% CI, 0.23-1.00) in one study (3-year follow-up, 157 participants); adjusted OR, 1.13, P > .10 in second study (5-year follow-up, 1173 participants); P = .04 in another study (13-year follow-up, 216 participants, no effect size provided).

^c Calculations based on number randomized.

^d Defined as hitting with the hand or objects, biting, burning with objects or by immersion, twisting, shaking, throwing or pushing so as to cause a fall, or hair pulling; identified from review of public agency documents from the Tennessee Department of Human Services.

e Defined as abandonment, leaving a child with an inappropriate caretaker, gross failure to seek medical care, failure to provide shelter or nutrition, or gross failure to provide for normal intellectual development; identified from review of public agency documents from the Tennessee Department of Human Services.

f Outcomes with no statistically significant results include mean number of ED visits for accidents and injuries

^g Outcomes with no statistically significant results include number of hospitalizations attributable to nonaccidental injury to the neonate (1 study), number of children hospitalized because of child abuse and neglect (1 study), proportion of children hospitalized for injuries and ingestions (1 study), number of children hospitalized for ambulatory-care sensitive conditions (1 study), number of children rehospitalized (1 study), mean number of all-cause hospitalizations (3 studies), and total count of hospital stays (1 study).

^k Defined as infant death, severe nonaccidental injury, and involuntary foster care placement.

verdict was reached.³⁴ The other 2 trials included all deaths in the period specified for follow-up. Overall rates of death were low (0% to 3.0%), even though eligible studies included at-risk children.

Composite Outcome | One trial reported on a composite outcome comprising infant death, severe nonaccidental injury, and involuntary foster care placement (eTable 35 in the Supplement). ⁴⁹ The investigators did not find a statistically significant risk before adjusting for covariates (3% [2/65] vs 12.7% [9/71]; RR, O.24 [95% CI, O.05-1.08]). When adjusted for baseline covariates, the RR was 0.22 (95% CI, 0.02-0.98). ⁴⁹

Harms of Interventions

Key Question 2. What are the harms of primary care–feasible or primary care–referable interventions to prevent child maltreatment? No studies met the inclusion criteria.

Discussion

The findings for this evidence review are summarized in Table 2. The evidence for the effect on short-term outcomes of primary carefeasible and primary care-referable interventions to prevent child maltreatment, reports to CPS, emergency department visits, and hospitalizations showed no statistically significant benefit in terms of child maltreatment outcomes or proxy measures. Few studies reported long-term outcomes; results from these studies are not consistent. Sensitivity analyses that included poor-quality studies did not change the results (eResults 1, eFigure 1 and eFigure 2, eTables 37-55 in the Supplement).

Furthermore, interpretation of some outcomes can be challenging. Lower rates of all-cause emergency department visits or hospitalization may represent changes in patterns of health care utilization as a result of the intervention rather than lower rates of abuse or neglect.

The evidence was also inconclusive for other outcomes, based primarily on the limited number of trials reporting on each outcome and lack of statistically significant results. These include injuries, failure to thrive, failure to immunize, internalizing and externalizing behavior symptoms, school attendance, and other measures of abuse or neglect. The evidence also suggests no statistically significant benefit for removal of the child from the home, child development, school performance, and prevention of death. No eligible studies on harms of interventions were identified. Results for parent-reported measures of exposure to abuse or neglect (eResults 2, eTables 56-71 in the Supplement) also did not suggest consistent benefit.

The evidence consisted entirely of RCTs and almost entirely of interventions that included home visits. Trials generally focused on young mothers and drew from vulnerable populations. Some interventions, such as the Nurse Family Partnership (Memphis, Tennessee; Elmira, New York) and Healthy Families, were tested in multiple settings. Nevertheless, the 21 included home-visit trials differed substantially in other respects, such as the populations of interest, baseline risk of maltreatment, intervention intensity and duration, and outcomes measured. These underlying characteristics may explain variations in the effectiveness of the intervention, but the evidence base for each outcome

was not extensive enough to identify any patterns. Additionally, the unaccounted effects of enhanced surveillance in preventive interventions, particularly in the context of home visiting programs, can result in increased identification of maltreatment. This is an important source of bias inherent in child maltreatment prevention research.

Limitations

This study had several limitations. First, inclusion in the review was restricted to studies focused on preventive interventions for children who had not yet experienced maltreatment and to individual rather than community-wide interventions. Therefore, the conclusions of this systematic review do not apply to the effectiveness of interventions to prevent the recurrence of child maltreatment for children who have experienced maltreatment or to community-wide interventions.

Second, in general, information about the type and severity of parent mental health problems was underreported. The resulting gap in the evidence precludes a fuller understanding of the salient factors affecting the effectiveness of the interventions.

Third, the focus of this review was on direct or proxy measures of abuse or neglect and their sequelae. It does not address the effectiveness of home visiting and other interventions for the other outcomes also measured in many of the included trials (eg, maternal outcomes, family functioning).

Fourth, there are limitations in the evidence that relate primarily to the considerable diversity of the interventions and the uncertainties stemming from such heterogeneity. Although all but 1 of the included trials had a home visiting component, several aspects of this particular activity differed. These aspects included the nature and theoretical basis of the interventions delivered during the home visits, credentials of the home visiting staff, and intensity and duration of the intervention.

Fifth, all trials involved implementing multiple components. Complex, multicomponent interventions need to report the theoretical foundation for the intervention to help interpret the results and reproduce successful interventions. Study authors generally did not provide a theory of change or logic model that identified components essential to the success of the intervention. Without theoretical or contextual information on critical intervention components, the review cannot comment on how successful interventions are different from unsuccessful interventions. ⁵²

Sixth, the question of applicability of the findings to other pediatric or caregiver populations (eg, with lower or higher risk profiles) and other settings (eg, with fewer resources) remains uncertain.

Seventh, although some child maltreatment prevention studies reported positive outcomes, the results do not consistently indicate benefit when sorted by outcome. Chance positive findings cannot be ruled out. Some studies reported on outcomes that may be important but are distal or unrelated to child maltreatment outcomes (such as interpregnancy interval).

Conclusions

Interventions provided in or referable from primary care did not consistently prevent child maltreatment. No evidence on harms is available.

ARTICLE INFORMATION

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Author Contributions: Dr Viswanathan had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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Statistical analysis: Viswanathan, Pan, Hart. Obtained funding: Viswanathan, Jonas. Administrative, technical, or material support: Viswanathan, Fraser, Pan, McKeeman, Forman-Hoffman, Hart, Zolotor, Patel. Supervision: Viswanathan.

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