Psychotherapy of Childhood Anxiety Disorders: A Meta-Analysis

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Key Words
Child psychotherapy evaluation · Childhood anxiety disorders · Treatment evaluation

Abstract
Background: The present study compared the efficacy of psychotherapy for childhood anxiety disorders (excluding trials solely treating post-traumatic stress disorder or obsessive-compulsive disorder). Methods: The meta-analysis included studies that met the basic CONSORT (consolidated standards of reporting trials) criteria. Several outcome variables (e.g. effect sizes, percentage of recovery) were analyzed using completer and intent-to-treat analyses during post-treatment and follow-up assessment. Twenty-four studies published by March 2005 were included in this meta-analysis. Results: In all the included studies, the active treatment condition was cognitive-behavioral. The overall mean effect of treatment was 0.86. No differences in outcome were found between individual and group treatments or child- and family-focused treatments. Follow-up data demonstrated that treatment gains were maintained up to several years after treatment. Conclusions: These findings provide evidence that anxiety disorders in children can be treated efficaciously. The gathered data support the clinical utility of cognitive-behavioral therapy in this regard. Randomized controlled trial studies investigating treatments other than cognitive-behavioral therapy are missing.

Introduction
Anxiety disorders are the most prevalent mental disorders not only of adulthood but also of childhood and adolescence [1, 2]. Trait anxiety seems to have increased greatly [3] and new research shows that childhood anxiety disorders are important risk factors for the development of mental disorders in adulthood, including anxiety disorders, depression and substance abuse [4–6]. The development and dissemination of efficacious treatments are, therefore, essential.

Clinical research on psychotherapy for anxiety disorders in children has advanced considerably in recent years. Different research groups have conducted several randomized controlled trials (RCTs) and substantial progress has been made in treating anxiety disorders in children and adolescents. However, these studies need to be summarized and discussed.

Meta-analyses represent an empirical approach to evaluating psychotherapy research. For a review of strengths and limitations of meta-analyses, see Sensky [7]. First meta-analyses have been published examining the efficacy of child and adolescent psychotherapies in general; yet further specific analyses of anxiety disorders were not the aim of these studies. Casey and Berman [8] reported an overall outcome effect size of 0.74 for child psychotherapy (n = 75; studies published from 1952 to 1983) and an average treatment versus control effect size of 1.16 for phobias (n = 9 studies). Weisz et al. [9] aggre-
gated the effects of 39 treatment-control comparisons and reported a mean overall effect size of 0.79 for child psychotherapy (n = 163; studies published from 1958 to 1984) and a mean effect size of 0.74 for phobias and anxiety.

A meta-analysis by Weisz et al. [10] produced an effect size of 0.60 (n = 16 studies) for phobias and anxiety and an overall effect size of 0.54 (n = 150; studies published from 1967 to 1993). An examination of the different types of therapies used in these studies showed that most were behavioral therapies. In addition, effects were more robust for behavioral than nonbehavioral treatments.

A major limitation of previously published meta-analyses is the inclusion of studies which did not require the establishment of diagnoses on the basis of DSM or ICD criteria. Furthermore, these meta-analyses did not include studies published after 1993. However, in the last 10 years, substantial progress has been made regarding psychotherapy research in the area of childhood anxiety disorders.

In order to improve the quality of reports using RCTs and to comprehend the results of RCTs, the consolidated standards of reporting trials (CONSORT) were developed. The CONSORT criteria checklist contains recommendations for four stages of a trial (enrolment, intervention allocation, follow-up, and analysis). According to current guidelines [11, 12] (http://www.consort-statement.org) and the fact that in recent years, several methodologically rigorous child and adolescent anxiety treatment outcome studies have been completed, the purpose of this paper was to provide a comprehensive meta-analysis of child and adolescent psychotherapy outcomes for anxiety disorders, including only studies that meet basic methodological criteria. Just recently, an extensive review by Compton et al. [13] summarized RCTs on anxiety and depression and concluded that cognitive-behavioral therapy (CBT) is currently the treatment of choice for anxiety and depression in children and adolescents [see also 14]. However, this review did not include any statistical analyses summarizing the results of these studies and did not analyze different treatment settings, which could be relevant regarding treatment outcome. Therefore, the present study means to close this gap and summarize the overall effect sizes of recently published RCTs, including analyses of different treatment settings in regard to treatment outcome. The data are aggregated from published studies in the field and multiple outcome variables are reported including effect sizes of several treatment settings (e.g., individual vs. group treatment or child- vs. family-focused treatment), recovery rates (both completer and intent-to-treat), sustained recovery rates and the results of follow-up assessments.

**Method**

**Literature Search**

In this meta-analysis, we included only published, peer-reviewed psychotherapy outcome studies in English and German. The literature search was conducted in PsycINFO and Medline (= PubMed) using the following key words: ‘childhood’, ‘children’, ‘anxiety disorders’, ‘specific phobia’, ‘social phobia’, ‘panic disorder’, ‘separation anxiety disorder’, ‘generalized anxiety disorder’, ‘overanxious disorder’, ‘avoidant disorder’, ‘therapy’, ‘outcome’, and ‘treatment’. We also included psychotherapy studies that were listed in the reference sections of the papers we collected after the initial computer search and conducted an Internet search. This literature search produced a total of 36 treatment studies. Obsessive-compulsive disorder (OCD) and post-traumatic stress disorder (PTSD) will not be addressed in this meta-analysis because both basic research and psychotherapy research of these two disorders differ substantially from childhood anxiety disorders mentioned above. In this regard, there is an ongoing discussion whether OCD should even be assigned to anxiety disorders (see ICD-10). For separate reviews of psychotherapy research in OCD and PTSD, see Cartwright-Hatton et al. [14], Cohen et al. [15], Franklin et al. [16] and Simons et al. [70].

**Inclusion Criteria**

All studies included in this meta-analysis were required to have investigated the efficacy of a specific treatment for anxiety disorders in children against a control condition or alternative credible psychotherapeutic treatment. Participants (i.e., the children who were treated) had to have met DSM or ICD diagnostic criteria for a principal anxiety disorder, and participants had to have been randomly assigned to either treatment or control conditions (i.e., an RCT). In accordance with the CONSORT guidelines, the authors of the studies were required to have written a standard treatment protocol. In order to facilitate computation of effect sizes, studies were included only if they reported means and standard deviations of the outcome measures as well as sample sizes at each assessment time point. We excluded studies in which treatment groups had fewer than 10 patients (lack of power), single case studies, subclinical cases and pharmacological studies. Studies had to have been published by March 2005. All decisions were made a priori, before examining any individual studies. These several steps produced a pool of 24 studies for further analyses. An overview of all the studies is in Appendix 1. The treatment orientation of the excluded studies was primarily CBT, except the chart review of Target and Fonagy [18], which involved a psychodynamic approach.

**Effect Size**

The effect size (Cohen’s d) is an index of the size and direction of treatment effects. Cohen [19] suggests that an effect size of 0.20 may be considered a small effect, 0.50 a medium effect, and 0.80 a large effect.
We computed different effect sizes for both treatment and control conditions including a global effect size and separate effect sizes for the three most commonly used outcome measures (see below). Effect sizes were calculated for pre-/post-treatment and for pre-follow-up assessments. Within each study, effect size values were averaged across all outcome variables to an overall effect size (one effect size per study). In a second step, these effect sizes were averaged across all studies (global effect size). Specific effect sizes were calculated separately for the most commonly used measurements, i.e. Revised Children’s Manifest Anxiety Scale (RCMAS), Fear Survey Schedule for Children-Revised (FSSC-R), and Children’s Depression Inventory (CDI) within each study and then averaged across all studies. In addition, 95% confidence intervals were computed for the effect sizes of each category. When the associated confidence intervals do not include 0, this indicates that the average effect size is significantly greater than zero at p < 0.05. Q statistic was calculated to determine whether the effect size data from the studies are homogeneous. If the studies share a common effect size, the value of Q will be nonsignificant. Fail-safe number statistics were calculated to check the robustness of the findings and whether significant mean effect sizes might have been inflated by a publication bias [20]. The fail-safe number informs about the number of file drawer studies required to bring the mean effect size down to a defined level. Orwin [21] adopted Rosenthal’s formula for probabilities to effect sizes d: 

\[ n_{fs} = \frac{[1/2]^2}{d^2} \]

In additional analyses, we distinguished between child- and family-focused treatments. Child-focused treatment was defined as the treatment which focuses on the children and in which the parents are either not included at all or minimally involved (i.e., in three sessions or fewer). Family-focused treatment was defined as the treatment in which parents are actively involved in the treatment process for more than three sessions, the nature and extent of parental involvement being clearly described in the treatment procedure section of the study.

Recovery
Percent recovered is an essential index of a clinically significant change. However, most of the studies reported recovery rates based on the percentage recovered compared to those who completed treatment rather than those who entered treatment (i.e., intent-to-treat). Intent-to-treat analyses produce a more conservative estimate of recovery by answering the question: ‘Of those patients who entered in and began treatment, what percentage is likely to recover?’ We will report recovery rates for both completer and intent-to-treat analyses.

Follow-Up Data
Follow-up data provide an accurate picture of the efficacy of treatment and its lasting effects.

Measures
In order to establish diagnoses before and after treatment and at follow-up and to determine diagnostic recovery, most studies used the Anxiety Disorders Interview Schedule for Children (ADIS) [22]. The ADIS is a structured diagnostic interview consistent with the DSM-IV criteria. It assesses child functioning (anxiety, mood, and externalizing disorders) in separate interviews: one with the child and one with the parents. The ADIS contains a Clinician Severity Rating, a scale ranging from 0 (absent) to 8 (very severe) that represents the degree of symptom severity and interference of diagnosis assigned by the interviewer based on clinical information provided by the interview. The interview has good interrater reliability for the child and the parent interview, adequate test-retest reliability [23], and is sensitive to treatment effects in studies with youths with anxiety disorders [24].

Child Measures
The RCMAS [25] measures chronic anxiety and is one of the most frequently used anxiety self-report measures for children. The RCMAS consists of 37 items, each one rated as true or false. There has been extensive work supporting the RCMAS’s validity and reliability [25]. Significant correlations have been found between RCMAS scores and other self-report measures of anxiety and related constructs such as depression [26].

The FSSC-R consists of 80 items and assesses specific fears in children [27]. The child rates his or her level of fear on a three-point Likert scale. Ollendick [27] reported solid internal consistency and adequate test-retest reliability.

The CDI consists of 27 items and assesses cognitive, affective, and behavioral symptoms of depression. For each item, children are given three choices from which they select the one which best describes themselves over the past 2 weeks. The CDI has good internal consistency, moderate test-retest reliability [28] and correlates in the expected direction with measures of related constructs such as self-esteem, negative cognitive attributes, and hopelessness [29].

Procedure
We assessed the following major variables from each study: sample size, means and standard deviations of outcome measures, percentage of children who completed treatment, pre-/post-treatment and treatment-control effect sizes, and percentage of children who recovered after treatment, both for those who entered (intent-to-treat) and for those who completed treatment. For follow-up, we assessed the same variables provided that such data were available.

Effect Size Calculation
The pre-/post-treatment and pre-follow-up effect sizes were calculated as follows:

\[ d_{es} = \frac{M_1 - M_2}{\sqrt{\frac{(N_{t1} - 1)SD_{t1}^2 + (N_{t2} - 1)SD_{t2}^2}{N_{t1} + N_{t2} - 2}}} \]

where \( \bar{M} \) is the mean of the measurement, \( t1 \) represents the pretreatment assessment, \( t2 \) represents the post-treatment or follow-up assessment, \( N \) is the sample size, and \( SD \) the standard deviation [30].

To investigate the treatment-control effect size we used the following formula:

\[ d_{es} = \frac{\left( M_{TG} - M_{CG} \right)}{SD_{pooled pre}} \]

where \( M \) is the mean of the measure, \( TG \) represents the treatment group, \( CG \) represents the control group, and \( SD_{pooled pre} \) is the pooled within-group standard deviation.
Results

Data from 24 clinical trials were included in this meta-analysis. In all 24 studies, the active treatment condition was cognitive-behavioral. Twelve studies included individual treatment and fifteen included group therapy. Seventeen studies included child-focused treatment and 14 included family-focused treatment. Sixteen studies had a waiting list control condition; the remaining included an alternative therapy as a control condition. The length of treatment ranged from 3 to 18 sessions, with an average length of 12.3 sessions. The total number of patients across all studies was 1,275, and the mean age of participants was 10.9 years (range: 6–18 years). The vast majority of studies evaluated the efficacy of psychotherapy in children with anxiety disorders. Only the study of Ginsburg and Drake [31] examined the feasibility and effectiveness of a school-based group CBT for anxiety disorders with low-income African-American adolescents. The study conducted by Baer and Garland [32] evaluated a CBT program in a community psychiatric setting.

Interventions Used by the Studies

Most programs were designed to target the child’s anxiety using a mixture of exposure techniques (90.5%), cognitive restructuring strategies (66.7%), relaxation techniques (52.4%), and positive self-talk (38.1%). In regard to parents, the majority of family-focused treatment studies used the following interventions: teaching parents to deal and cope with child anxiety and its related behaviors, teaching communication and problem-solving skills, and managing the parents’ own anxiety. Interventions used in the attention placebo control condition were designed to provide the child and the parents with a broad range of information about anxiety (education support), but no encouragement or instructions were offered to confront feared situations [33–36].

Exclusion Criteria Used by the Studies

Many studies excluded children with psychotic symptoms (72.2%) and mental retardation (50%). Most studies (77.8%) excluded children who were currently involved in psychosocial or psychopharmacological treatment. However, more recent studies [32, 37–39] did include children who were on psychopharmacological medica-

1 The exclusion criteria for each study included in this meta-analysis can be requested from the authors.

Completion Rates

Most children who entered treatment (n = 1,275) completed it (n = 1,080; 84.7%). Completion rates were 85.8% for group treatment, 84.9% for individual therapy, 82.9% for family-focused treatment, and 84.9% for child-focused treatment. Thus, very similar completion rates were found across all modes of treatment.

Effect Sizes

Active Treatment vs. Waiting List (Global and Specific Effect Sizes)

The mean overall pre-/post-treatment effect size across all treatments was d = 0.86 (95% CI = 0.69–1.03; n = 24), whereas the effect size for the waiting list control group was d = 0.13 (95% CI = 0.03–0.24; n = 16). The specific effect sizes for the three most common treatment outcome measures (RCMAS, FSSC-R, and CDI) were as follows: pre-/post-treatment effect size in the treatment condition was 0.83 (95% CI = 0.64–1.02) for trait anxiety (RCMAS) and 0.85 (95% CI = 0.45–1.24) for phobic anxiety (FSSC-R). In the control condition, effect sizes were 0.22 (95% CI = 0.08–0.35; RCMAS) and 0.32 (95% CI = 0.11–0.54; FSSC-R). Furthermore, symptoms of depression (CDI) improved significantly (d = 0.70; 95% CI = 0.54–0.86) in the treatment condition compared to the waitlist control condition (d = 0.20; 95% CI = 0.07–0.33). The mean overall treatment versus control effect size across all active treatments averaged 0.66 (95% CI = 0.36–0.96) at the post-treatment assessment. The t test with independent samples revealed a significant difference between treatment versus control effect size [t(37) = 6.92, p = 0.00].

Homogeneity of effect sizes for post-test data was tested. The resulting χ² value was nonsignificant [χ²(24) = 25.08, p = 0.20], indicating that the assumption of homogeneity for the post-test effect sizes cannot be rejected.

The fail-safe number of this study suggests that 79 file drawer studies with effect sizes of zero are necessary to reduce the effect size of 0.86 to a mean effect size of 0.20. To reduce the effect size of 0.86 to a mean effect size of 0.50, 17 studies of zero effect would be necessary.

Follow-Up

Of the 24 outcome studies, 16 reported short-term follow-up assessment data. On average, short-term follow-up assessments occurred 10.4 months after the end of treatment. The effect size for pre-follow-up was 1.36 (95%
There are 3 long-term follow-up studies: the 6-year follow-up study by Barrett et al. [40], the 3.4-year study by Kendall and Southam-Gerow [41] and the 7.4-year study by Kendall et al. [42]. The long-term effect size for the study by Kendall and Southam-Gerow [41] was 0.61. The effect size for the 6-year follow-up study of Barrett et al. [40] was 0.82. Finally, the effect size for the 7.4-year follow-up study by Kendall et al. [42] was 1.13 for child report and 1.54 for parent report.

Treatment Setting (Global Effect Sizes)

Table 1 displays the mean overall pre-/post-treatment and pre-follow-up effect sizes for the outcome studies by treatment type. Group and individual treatments were equally effective in reducing children’s symptoms. The mean overall treatment versus control pre-/post-treatment effect sizes were 0.52 (95% CI = 0.04–0.99) for individual therapy and 0.61 (95% CI = 0.44–0.79) for group therapy. The t test with independent samples revealed a nonsignificant difference [t(11) = 0.33, p = 0.75].

As shown in table 1, the pre-/post-treatment effect size for child-focused treatments was similar to the effect size of treatments involving parents. The mean overall treatment versus control pre-/post-treatment effect sizes were 0.53 (95% CI = 0.30–0.77) for child-focused therapy and 0.63 (95% CI = 1.3–0.58) for family-focused therapy. This difference was not significant [t(21) = 0.26, p = 0.79].

Placebo Condition vs. Waiting List (Global Effect Sizes)

We compared waiting list control conditions with active, nonspecific control conditions, and the pre-/post-treatment effect sizes were 0.13 (95% CI = 0.03–0.24; n = 16) for waiting list and 0.58 (95% CI = -1.6 to 1.3; n = 4) for attention placebo control.

Percent Recovered

Table 2 displays the percentage of patients who recovered from those who completed and those who entered treatment. Across all active treatments, 68.9% of children who completed therapy no longer met the diagnostic criteria for their principal pretreatment anxiety disorder compared to only 12.9% of children who were assigned to a waiting list. The t test with independent samples re-
revealed a significant difference between treatment versus control group after treatment with respect to children without anxiety diagnosis \( [t(34) = 15.06, p < 0.00] \). At the follow-up assessment, the recovery rate of children who completed treatment increased slightly to 72%.

Individual and group therapies showed comparable recovery rates of 72.1% and 66%, respectively. For the intent-to-treat sample, the overall recovery rate after treatment was 55.4%. Intent-to-treat recovery rates were similar for individual (57.3%) and group (53.2%) therapy. Child- and family-focused treatment showed recovery rates of 64.1% and 76.9%, respectively. For the intent-to-treat sample, the recovery rate after treatment for child-focused treatment was 54.4% and 65.2% for family-focused treatment. Because of a lack of the majority of treatment studies to report the number of dropouts in the waiting list conditions, the intent-to-treat recovery rate could not be investigated.

**Discussion**

In the present study, we examined 24 RCTs out of a pool of 36 treatment outcome studies on anxiety disorders in children and adolescents. The results of the present study indicate that treatment produces acute effects, which reflect substantial symptom improvement. The current findings are consistent with previous meta-analyses and provide convergent evidence that CBT is effective for children with anxiety disorders.

The robust effects of therapy are reflected in the pre-/post-treatment effect sizes of 0.86 in the treatment and 0.13 in the waiting list control conditions. Of the children included in our study who completed treatment, 68.9% recovered to the extent that they no longer met criteria for their principal pretreatment anxiety diagnosis. In comparison, only 12.9% of waiting list participants recovered. Furthermore, the fail-safe number statistic shows that 79 file drawer studies with effect sizes of zero are needed to reduce this effect size to a mean effect size of 0.20.

Symptom improvements following treatment occurred not only for anxiety but also for depression. The treatment versus control effect size was 0.66. Previous meta-analytic estimates reported similar \([10]\) or higher pre-/post-treatment effect sizes \([8, 9]\) in general child and adolescent psychotherapy.

Interestingly, the attention placebo control condition reached a pre/post-treatment effect size of 0.58 \((n = 4)\). Compared to the overall pre-/post-treatment effect size of the active treatment condition \((0.86)\), the effect size for the active control condition is quite high. Four studies included in our meta-analysis compared their active interventions with an attention placebo control condition \([33–36]\). However, only the study of Beidel et al. \([33]\) and Muris et al. \([36]\) used a strict placebo condition consisting of study skills, test-taking strategies and emotional disclosure, which led to a significant treatment vs. placebo effect. The two other studies included psychoeducation on anxiety comparable to information given in CBT programs, which were equally effective as the active treatments. Thus, these results could imply that improvement in childhood anxiety disorder can be achieved purely with psychoeducation on anxiety, without explicit instructions for exposures. This interesting issue requires further investigation.

Treatment outcome studies comparing the efficacy of child-alone therapy to interventions involving both children and parents have found conflicting results. As in the meta-analysis of Casey and Berman \([8]\), we did not find a difference between the two types of treatment. The mean treatment versus control effect size was 0.53 for child-focused therapy and 0.63 for family-focused treatment. To date, there is no clear empirical evidence indicating who might benefit more from one type of therapy over the other. Some data suggest that younger children and children with parents who have an anxiety disorder themselves may benefit more from a combined child and parent treatment than from a child-alone therapy \([43, 44]\). However, in the study by Barrett et al. \([43]\), the relative superiority of CBT plus family anxiety management condition compared to CBT alone at the 12-month post-treatment assessment disappeared at the 6-year follow-up assessment \([40]\). Likewise, the results of other studies \([45–47]\) suggest that for some children, a child-alone approach may be sufficient for anxiety symptom reduction. Further research must examine the role of parental involvement in treatment for children with different anxiety disorders and identify child and family characteristics that will enable clinicians to assign anxious children to either a child-alone intervention or a combined parent-child intervention.

While earlier studies established the efficacy of individual therapy in treating child anxiety disorders, the results of a number of recent trials suggest that group treatment may be equally efficacious \([38, 48]\). In the present study, the mean effect sizes for group and individual therapy were comparable. These results should nevertheless be interpreted with caution. A number of group treatment studies in our meta-analysis included family-focused in-
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interventions (n = 7), making it difficult to disentangle the effects of group therapy and family involvement. Though Muris et al. [38] have recently demonstrated the similarity in outcomes of group and individual treatment without parental involvement in either condition, future research should continue to investigate this question while controlling the overlap of family-focused and group treatment settings.

Weisz et al. [9, 10] and Kazdin et al. [49] reported in their meta-analyses that only around one third of all published treatment outcome studies reported data from follow-up assessments. Recently published studies seem to have improved in this regard, as 16 (66.7%) of the 24 outcome studies included in our meta-analysis reported follow-up data. In our meta-analysis, the mean pre-follow-up effect size was 1.36, suggesting that treatment gains were well maintained. Furthermore, 72% of the children did not meet the criteria for their principal pretreatment diagnosis at follow-up. These results are consistent with three long-term follow-up studies [40–42], which have demonstrated that the effects of child anxiety treatments are relatively stable and long-lasting (up to 7.4 years).

With the exception of five studies that investigated the efficacy of psychotherapy for social phobia [32, 33, 46, 50, 51], none of the other treatment studies differentiated between the different types of anxiety disorders. In fact, these studies tested general treatments for anxiety disorders in groups of children with different anxiety disorders (separation anxiety disorder, generalized anxiety disorder, specific phobia, social phobia). However, RCT studies with a focus on specific anxiety disorders, in particular separation anxiety disorder, generalized anxiety disorder and specific phobia, are needed to complement the previous global approaches.

All studies included in the meta-analysis investigated the efficacy of CBT. Thus, non-CBT outcome studies that fulfill the methodological criteria required by CONSORT are still missing. In the literature search for the present study, we found a chart review of child psychoanalysis and psychotherapy of children with emotional disorders [18], which reported an improvement rate of 47.2% in a large child sample (n = 352). However, this study did not meet the inclusion criteria for our meta-analysis (see Appendix 2).

It has to be considered that the studies included in this meta-analysis demonstrate efficacy of psychotherapy for anxious children; however, effectiveness has to be proven. Therefore, now that CBT trials are found to be successful, it is important to extend studies to the clinical settings. Baer and Garland [32] investigated a pilot study of community-based cognitive-behavioral group therapy for adolescents with social phobia. Although the sample size (n = 12) was small, the study provides support for the use of simplified cognitive-behavioral interventions in community psychiatric settings. Similar conclusions can be drawn from the study of Ginsburg and Drake [31] that evaluated the feasibility and effectiveness of a school-based group treatment for anxiety disorders with African-American adolescents. For the evaluation of the presented results, general limitations regarding meta-analyses have to be considered. The reported effect sizes only provide information about how much the patients improved after therapy compared to their values before therapy. Thus, the effect sizes do not provide information regarding symptom severity and functional level after therapy compared to healthy control participants.

Reported effect sizes are based entirely on self-report measures, due to the fact that parent and teacher reports were not regularly assessed. However, the gold standard in assessing efficacy of psychotherapy in children and adolescents should be the inclusion of self-reports as well as parent and teacher reports [71]. Further psychotherapy research should take this into consideration in order to get a sound and accurate description of therapy outcome.

Knowing how to treat childhood anxiety disorders is important. Such knowledge both helps ease the suffering of children and contributes to the reduction and prevention of future suffering as adults. As we work toward improving psychotherapy for childhood anxiety disorders, we must especially focus on helping those children for whom existing therapies continue to produce less than optimal outcomes.

Acknowledgements

This research was supported by a Swiss National Science Foundation scholarship to Tina In-Albon (PBBSI-102350) and by a grant of the Swiss National Science Foundation to Silvia Schneider (SNF PP001-68701). The authors thank David A. Moscovitch, Michael K. Suvak, Department of Psychology, Boston University, and Andrea H. Meyer, University of Basel.
## Appendix 1

### Studies Included in the Meta-Analysis

<table>
<thead>
<tr>
<th>Study</th>
<th>n</th>
<th>Age</th>
<th>Mode</th>
<th>Diagnosis</th>
<th>Weeks</th>
<th>Com-</th>
<th>FU</th>
<th>Pre-/post-treatment effect size (d) per treatment condition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Barrett et al. [43]</td>
<td>79</td>
<td>7–14</td>
<td>Ind, 1, 2</td>
<td>SAD, OAD, SoP</td>
<td>12</td>
<td>94</td>
<td>6, 12 months; Barrett et al. [40]: 6 years</td>
<td>CBT: 0.59, CBT+Fam: 0.89</td>
</tr>
<tr>
<td>Barrett [48]</td>
<td>60</td>
<td>7–14</td>
<td>Gr, 1, 2</td>
<td>SAD, OAD, SoP</td>
<td>12</td>
<td>83</td>
<td>12 months</td>
<td>FSSC-R Ch: 1.54, FSSC-R Fam: 2.37</td>
</tr>
<tr>
<td>Beidel et al. [33]</td>
<td>67</td>
<td>8–12</td>
<td>Gr, 1, 3</td>
<td>SoP</td>
<td>12</td>
<td>75</td>
<td>6 months</td>
<td>SPAI: 1.21; SPAI Tbl: 0.18</td>
</tr>
<tr>
<td>Cobham et al. [44]</td>
<td>67</td>
<td>7–14</td>
<td>Ind, act tx1, 2</td>
<td>SAD, OAD, GAD, SPP, SoP</td>
<td>10</td>
<td>91</td>
<td>6, 12 months</td>
<td>CBT Ch anx: 0.77, CBT+PAM Ch anx: 0.49, CBT+PAM Ch+Pa anx: 0.40, CBT Ch+Pa anx: 0.66</td>
</tr>
<tr>
<td>Flannery-Schröder et al. [52]</td>
<td>37</td>
<td>8–14</td>
<td>Ind, Gr1</td>
<td>GAD, SAD, SoP</td>
<td>18</td>
<td>76</td>
<td>3 months</td>
<td>Ind: 1.26, Gr: 0.73</td>
</tr>
<tr>
<td>Gallagher et al. [51]</td>
<td>23</td>
<td>8–11</td>
<td>Gr</td>
<td>SoP</td>
<td>3</td>
<td>50</td>
<td>3 weeks</td>
<td>0.36</td>
</tr>
<tr>
<td>Ginsburg et al. [31]</td>
<td>12</td>
<td>14–17</td>
<td>Gr</td>
<td>GAD, SPP, AG, SoP</td>
<td>10</td>
<td>75</td>
<td>–</td>
<td>SCARED: 0.27</td>
</tr>
<tr>
<td>Hayward et al. [50]</td>
<td>35</td>
<td>14–18</td>
<td>Gr</td>
<td>SoP</td>
<td>16</td>
<td>81</td>
<td>12 months</td>
<td>SPAI: 1.10</td>
</tr>
<tr>
<td>Heyne et al. [45]</td>
<td>61</td>
<td>7–14</td>
<td>Ind, act tx1, 2</td>
<td>SAD, SoP, GAD, PD, OCD, CD</td>
<td>8</td>
<td>–</td>
<td>4.5 months</td>
<td>Ch: 0.51, Pa+Te: 1.58, Ch+Pa+Te: 0.78</td>
</tr>
<tr>
<td>Kendall et al. [53]</td>
<td>47</td>
<td>9–13</td>
<td>Ind</td>
<td>SAD, OAD, AD</td>
<td>16</td>
<td>78</td>
<td>12 months; Kendall and Southam-Gerow [41]: 3.4 years</td>
<td>1.32</td>
</tr>
<tr>
<td>Kendall et al. [54]</td>
<td>94</td>
<td>9–13</td>
<td>Ind</td>
<td>SAD, OAD, AD</td>
<td>16</td>
<td>80</td>
<td>12 months; Kendall et al. [42]: 7 years</td>
<td>1.08</td>
</tr>
<tr>
<td>King et al. [55]</td>
<td>34</td>
<td>5–15</td>
<td>Ind</td>
<td>SAD, OAD, SoP, SPP, CD</td>
<td>4</td>
<td>100</td>
<td>3 months</td>
<td>1.37</td>
</tr>
<tr>
<td>Last et al. [34]</td>
<td>56</td>
<td>6–17</td>
<td>Ind, 1, 3</td>
<td>Schoolpho</td>
<td>12</td>
<td>73</td>
<td>1 month</td>
<td>STAIC-T CBT: 0.57, STAIC-T ES: 1.33</td>
</tr>
<tr>
<td>Manassis et al. [37]</td>
<td>78</td>
<td>8–12</td>
<td>Ind, Gr2</td>
<td>GAD, SAD, SPP, SoP, PD</td>
<td>12</td>
<td>–</td>
<td>–</td>
<td>SASC Gr: 0.21, SASC Ind: 0.43</td>
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<tr>
<td>Mendlewitz et al. [39]</td>
<td>62</td>
<td>7–12</td>
<td>Gr, act tx1, 2</td>
<td>–</td>
<td>12</td>
<td>91</td>
<td>–</td>
<td>Ch: 0.26, Pa: 0, Ch+Pa: 0.33</td>
</tr>
<tr>
<td>Muris et al. [38]</td>
<td>36</td>
<td>8–13</td>
<td>Gr+Ind</td>
<td>GAD, SAD, SoP</td>
<td>12</td>
<td>100</td>
<td>–</td>
<td>STAIC Ind: 0.82, STAIC Gr: 1.08</td>
</tr>
<tr>
<td>Muris et al. [36]</td>
<td>30</td>
<td>9–12</td>
<td>Gr, 1, 3</td>
<td>GAD, SAD, SoP</td>
<td>12</td>
<td>80</td>
<td>–</td>
<td>STAIC: 1.38</td>
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<tr>
<td>Nauta et al. [47]</td>
<td>79</td>
<td>7–18</td>
<td>Ind, 1, 2</td>
<td>SAD, SoP, GAD, PD</td>
<td>12</td>
<td>96</td>
<td>3 months</td>
<td>SCAS Ch: 1.13, SCAS Pa: 0.65</td>
</tr>
<tr>
<td>Rapee [56]</td>
<td>95</td>
<td>7–16</td>
<td>Gr</td>
<td>SAD, GAD, SoP</td>
<td>12</td>
<td>–</td>
<td>12 months</td>
<td>0.81</td>
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<tr>
<td>Silverman et al. [69]</td>
<td>56</td>
<td>6–16</td>
<td>Gr</td>
<td>OAD, SoP, GAD</td>
<td>–</td>
<td>73</td>
<td>3, 6, 12 months</td>
<td>0.64</td>
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<tr>
<td>Silverman et al. [35]</td>
<td>81</td>
<td>6–16</td>
<td>Ind, 2, 3</td>
<td>SPP, SoP, AG</td>
<td>10</td>
<td>78</td>
<td>3, 6, 12 months</td>
<td>SC: 0.91, CM: 0.57, ES: 0.27</td>
</tr>
<tr>
<td>Spence et al. [46]</td>
<td>50</td>
<td>7–14</td>
<td>Gr, act tx1, 2</td>
<td>SoP</td>
<td>–</td>
<td>–</td>
<td>6, 12 months</td>
<td>Ch: 0.97, Ch+Pa: 0.75</td>
</tr>
<tr>
<td>Toren et al. [57]</td>
<td>24</td>
<td>6–13</td>
<td>Gr</td>
<td>SAD, OAD</td>
<td>10</td>
<td>–</td>
<td>12, 36 months</td>
<td>0.58</td>
</tr>
</tbody>
</table>

SAD = Separation anxiety disorder; GAD = generalized anxiety disorder; OAD = overanxiety disorder; SoP = social phobia; SPP = specific phobia; PD = panic disorder; AG = agoraphobia; CD = conduct disorder; Schoolpho = school refusal; Gr = group treatment; Ind = individual treatment; act tx = active placebo treatment; SPAI = Social Phobia and Anxiety Inventory; SCARED = Screen for Childhood Anxiety Related Emotional Disorders; SCAS = Spence Child Anxiety Scale; STAIC-T = State-Trait Anxiety Inventory for Children; SASC = Social Anxiety Scale for Children; ES = education support; SC = exposure-based cognitive self-control; CM = exposure-based contingency management; PAM = parental anxiety management; Tb = testbuster.

1 Family-focused treatment.
2 Child-focused treatment.
3 Attention-placebo control group.
4 Effect sizes are indicated for outcome measure RCMAS, when not otherwise specified.
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