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SpringerPlus

DOI: 10.1186/s40064-016-2826-9

Published: 20/07/2016

Peer reviewed version

Cyswllt i'r cyhoeddiad / Link to publication

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Health Related Quality of Life for Young People receiving Dialectical Behaviour Therapy (DBT): A routine outcome-monitoring pilot

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Running Head: Health Related Quality of Life in Dialectical Behaviour Therapy
Abstract

Purpose: Adults presenting with Borderline Personality Disorder (BPD) score poorly on measures of Health Related Quality of Life (HRQoL). Little is known about HRQoL in adolescents with BPD type presentations and how treatment impacts quality of life. Our primary aim was to use routinely collected quality-of-life outcome measures pre and post-treatment in Dialectical Behaviour Therapy (DBT) for adolescents to address this gap. Secondary aims were to benchmark these data against EuroQol 5 dimensions (EQ-5D™) outcomes for clients treated in clinical trials and to assess the potential of the EQ-5D™ as a benchmarking tool.

Method: Four adolescent DBT teams, routinely collecting outcome data using a pseudonymised secure web-based system, supplied data from consecutive discharges.

Results: Young people in the DBT programmes (n=43) had severely impaired HRQoL scores that were lower at programme admission than those reported in published studies using the EQ-5D™ in adults with a BPD diagnosis and in one study of adolescents treated for depression. 40% of adolescents treated achieved Reliable Clinical Change. HRQoL improved between admission and discharge with a large effect size. These results were not statistically significant when clustering in programme outcomes were accounted for.

Conclusion: Young people treated in NHS DBT programmes for BPD type presentations had poorer HRQoL than adults with a BPD diagnosis and adolescents with depression treated in published clinical trials. The EQ-5D™ detected reliable change in this group of adolescents. Programme outcome clustering suggests that both the measure and the web-based monitoring system provide a mechanism for benchmarking clinical programmes.

Key Words: Adolescents; borderline personality disorder; health-related quality-of-life; routine outcome monitoring
### Background

Whilst borderline personality disorder (BPD) is most commonly diagnosed in adults, more recently clinicians and researchers have begun to consider the assessment and identification of personality disorders in adolescents [1, 2]. Studies investigating BPD traits in young people report that these presentations predict the presence of personality disorders in adulthood and are also linked to other psychiatric disorders, impaired long-term functioning and to increased mortality [3-5]. Currently, there are no established effective treatments for young people with BPD-type presentations. More typically, adolescent research has focussed on interventions for repeated self-harm, one of the diagnostic criteria for BPD. A meta-analysis of psychological and social interventions for suicide attempts and self-harm [6] reviewing 19 trials comprising 2176 young people concluded that the selected interventions appeared to be effective overall for self-harm. Dialectical behaviour therapy (DBT) [7,8], cognitive-behavioural therapy (CBT) [9] and mentalization-based therapy (MBT) [10] demonstrated the largest effect-sizes; however, these results require independent replication.

DBT is an effective treatment for BPD in adults with robust evidence from randomised controlled trials demonstrating significant impacts on a number of important outcomes, including reduced suicidal and self-harming behaviours and service utilisation [11-18]. The recent Cochrane Review concluded that DBT was the only psychological treatment for BPD with sufficient data to pool into a meta-analysis [19]. Results demonstrated moderate to large statistically significant effects for DBT over treatment as usual in reductions in suicidal and self-harm behaviours (SMD -0.54, 95% CI -0.92 to -0.16); improvements in mental health (SMD 0.65, 95% CI 0.07 to 1.24) and decreases in anger (SMD -0.83, 95% CI -1.43 to -0.22). The National Institute for Health and Care Excellence (NICE) Guidelines for the treatment and management of BPD recommend DBT particularly where reduction in self-harm is a
clinical priority. An earlier review reported that DBT had the potential for cost-effectiveness [20].

As a direct consequence of its success in treating suicidal and self-harm behaviours with adults diagnosed with BPD, DBT was adapted for the treatment of adolescents (DBT-A) presenting with suicidal and self-harm behaviours who were also demonstrating features of a developing BPD [21]. A recent RCT of the adapted form of DBT conducted in Norway replicated the findings of earlier studies in adult populations. Mehlum et al. [8] randomly allocated 77 adolescents presenting with suicidal and self-harming behaviour with at least two other BPD characteristics to either DBT-A or Enhanced Treatment as Usual (ETAU). After 16 weeks of treatment DBT-A was significantly superior to ETAU in terms of decreases in self-harm behaviour and depression. Several non-randomised studies conducted in the UK and elsewhere also indicate that DBT maybe a promising intervention with adolescents presenting with self-harm behaviour in the context of BPD [22-25].

Adults diagnosed with BPD have severe impairments in Health Related Quality of Life (HRQoL) [26-28]. In addition to the significant personal burden of a diagnosis of BPD [29], clients with BPD are typically “treatment seeking” with associated high utilisation of health services [30-33], leading researchers and policy makers alike to highlight the importance of investigating and implementing clinically and cost-effective treatments for this population [27,9]. Establishing cost-effectiveness (specifically cost-utility analysis) requires generic preference based measures that can be used to calculate the cost per additional Quality-Adjusted Life Year (QALY). QALYs are a generic measure of disease burden that includes both the quality and length of life and allow for comparison across conditions to help inform decision makers where best to invest scarce health care resources. NICE [34] recommend the EuroQol 5 Dimensions (EQ-5D™) as the most appropriate measure for health economic
evaluations of new technologies. This standardised and validated self-report measure describes an individual’s current health status and can be used to identify changes occurring over time. The construct validity and test-retest reliability of the EQ-5D™ have also been supported [35].

As the measure is generic rather than condition-specific, the EQ-5D™ provides a common denominator for different evaluations allowing comparison of new technologies with each other. Such generic measures assess broad levels of functioning in contrast to symptomatic measures that may address a single clinical outcome e.g. self-harm or depression. Whilst individual symptom measures are an important measure of clinical outcome, clinical guidance recommends considering broader measures of functioning and quality of life rather than simply symptomatic improvement [34]. Use of generic measures maybe of particular importance in the case of BPD where clients’ problems impinge on a wide range of health domains.

We therefore sought to examine the routine effectiveness of DBT as delivered in the NHS, with a focus solely on the EQ-5D™ as the Health Related Quality of Life (HRQoL) outcome. Using the EQ-5D has an additional advantage above addressing functional outcomes more broadly; measures that enable the calculation of QALYs across a number of services will provide data about cost-effectiveness of DBT as delivered in routine clinical practice, and, subject to sufficient variation across programmes, allow for the development of a national benchmarking system. Transferring evidence-based treatments established as efficacious in randomised-controlled clinical trials to routine clinical practice is fraught with difficulty [36]. Only by measuring outcomes routinely in clinical settings can the effectiveness of such treatments be established. Once programmes that do not deliver good clinical outcomes can
be identified, organisational interventions to address poor outcomes can be developed and implemented.

The primary aim of the pilot study was to evaluate the HRQoL outcomes of adolescents receiving DBT in routine clinical practice. Our secondary aim, given the absence of studies reporting on HRQoL in adolescents, was to compare the findings with published RCT data on adults with BPD and adolescents with other mental health conditions. Finally, we wished to assess the potential of the EQ-5D™ as a benchmarking measure.

Methods

Procedure.

All DBT programmes (N=9) with a subscription to the DBT pseudonymised outcome benchmarking (DBT-POB) website at www.dbt.uk.net were invited to participate in the study. The pseudonymised outcome website was developed to assist teams to collect routine outcome data based on the potential value of such data for implementation of DBT programmes [37]. To reduce the administrative burden on participating programmes and hopefully maximise the success of data collection, the amount of data required was kept to an absolute minimum using a single outcome measure, the EQ-5D™. Consistent with keeping the demands on busy programmes low, only pre- and post-treatment data were required for entry on the website. Programmes were asked to assess clients on the EQ-5D™ at admission to the programme and on discharge, regardless of whether discharge was planned or unplanned. Assessing all entrants to the programme, regardless of whether they complete treatment or not, provides a more conservative test of the effectiveness of a treatment programme. Only including data from treatment completers may overestimate the
effectiveness of treatment in routine practice. Pseudonyms were selected according to the gender of the client so number of male and female clients in the sample as a whole is known; otherwise no demographic data is available at the individual client level.

Seven of the subscribed nine teams were working with adolescents. A census of DBT clients in treatment on a specified date established that only four of these programmes were using the website to routinely collect data with sufficient accuracy for benchmarking purposes, namely that the website was reporting the correct number of clients in treatment that day. These four programmes consented for their data to be downloaded from the system on the predefined census date and for their programme data to be included in the multi-site data analysis. All participating programmes had broadly similar admission criteria (ages 14-18; five or more BPD criteria of which one must be the recent occurrence of self-harm behaviour).

All programmes had permission from their information governance officers to upload pseudonymised routine outcomes to the website. Ethical Approval was sought and received from the Ethics Committee of the School of Psychology, Bangor University. The Local NHS Research Ethics Committee was asked for an opinion on whether the study required NHS approval and considered that the study qualified as service evaluation. No patient identified information was submitted to or held by the outcome benchmarking website.

**Measures**

**EQ-5D™**

The EQ-5D™ is a generic outcome measure that asks participants to rate their current general health status on 5 dimensions: mobility, self-care, usual activities, pain or discomfort and anxiety & depression. The participant rates each of these dimensions at one of three different levels (no problems, some problems or extreme problems). The scoring system then classifies
the individual into one of 243 possible health states. Each health state provides a summary of
the participants rating of their health status on each dimension. For example, the health state
‘11111’ represents perfect health on all 5 dimensions – a state that 56% of the UK population
and 70% of the Spanish population will report on any one day [38,39]. In contrast, a health
status of ‘33333’ indicates the worst possible health status with extreme problems on all 5
dimensions. These scores can be transformed into a utility score that ranges between -0.59
and 1 (with death anchored at 0 and 1 representing perfect health) by applying societal
weights to each level that are based on the values of these health states in adult general
population samples derived from a choice based method such as Time Trade-Off (TTO) [40].
Currently there are no weights derived from valuations of health states by children and
adolescents. Negative scores indicating health states judged as ‘worse than death’ are
possible. The utility scores can be used to calculate QALYs. The sole adolescent study
utilising the EQ-5D™ provided tentative support for its use in this client group [41].

Data collection and entry

Each DBT team had its own protocol for administering and entering the data onto the website.
In most cases young people scored the questionnaires themselves. On some occasions
clinicians completed the questionnaires on their behalf. Proxy completion of the EQ-5D™ is
a recognised method of data collection. In a recent study of the EQ-5D™ with children and
adolescents from a community study, where parents acted as proxies, high levels of
agreement were found between the self-report and proxy versions of the EQ-5D-Y [42]. Each
DBT team uploaded EQ-5D scores for all patients treated in their DBT programme at
admission and discharge. An audit on a random selection of inputs from each team was
conducted to ensure that the data had been accurately entered into the website. To conduct
the audit one of the research team (LB) telephoned the DBT team and asked them to provide
the EQ-5D™ scores stored on paper in the clinical notes for each client for a set of randomly chosen patients and time points. These scores were checked for accuracy with the data already entered into the website. No inconsistencies between the data entered onto the system and the original paper copies of the data were detected during this audit.

**Data analysis strategy**

Utility scores at or below zero might be anticipated in clinical samples for which achieving a ‘life worth living’ [43] remains a daily struggle. In a sample of adolescents in treatment for suicidal and self-harm behaviours a measure that captures health states considered ‘worse than death’ displays considerable face and criterion validity; these scores were retained. Difference scores were calculated by subtracting EQ-5D™ utility scores on discharge from those at admission.

In an endeavour to establish whether the change in EQ-5D™ scores represented a robust change at the individual level, the Reliable Change Index for the EQ-5D was calculated using the Jacobson method [44]. This formula takes into account the reliability of the measure and variance in measurement to generate a change score in excess of which we can essentially be 95% certain that the change in score is a real (hence, reliable) change over time. Ideally, in order to exclude sources of systematic error in this calculation, an estimate of test-retest reliability derived from a clinical sample whose clinical characteristics have not changed over a period of time would be preferred. Since clinical samples are, by definition, in treatment and therefore on a change trajectory and the EQ-5D™ is not yet used widely in mental health settings, such estimates are hard to obtain. Hurst et al. [45 Table IX], using an intraclass correlation coefficient (ICC) rather than Pearson correlation because of concerns about over-estimating reliability, reported a reliability coefficient for the EQ-5D™ of 0.78 (0.6-0.96)
over 2 weeks in a clinical sample of 31 rheumatoid arthritis sufferers where there was no change in rheumatoid arthritis. More recently Sonntag and colleagues have tabulated ICCs for n=106 social phobics at an interval of 6 months and n=60 at 12 months [46 Table 4] anchored by no change on the Liebowitz Social Anxiety Scale which are all also 0.78. In view of the replication of this estimate and albeit limited clinical similarities in the study populations, this value was used in the calculation of Reliable Change Index in the DBT programmes in this study.

**Results**

**Description of the sample**

On the census date, 76 sets of client data were downloaded from four programmes of which 66 had female pseudonyms and 10 male. Of these 76 clients, 43 (38 female and 5 male) had been discharged from their respective programme and 33 were still in treatment. Admission EQ-5D™ utility scores for the whole sample (n=76) were 0.236 (SD 0.32, range -0.594 - 0.848). The admission EQ-5D scores of the 43 adolescents who had completed treatment prior to the census date were not significantly different from the 33 young people who were still in treatment. [Mean admission utility score still in treatment (n=33) = 0.244, mean admission utility score discharged (n=43) = 0.230, t(75)=0.19, p=0.85]. Average length of stay of adolescents consecutively discharged from programmes (n=43) was 177 days (SD 116, range 23-462).

**Comparison of Admission and Discharge HRQoL scores**

Admission and discharge scores for the 43 consecutive discharges in the dataset were compared. Mean admission utility scores were 0.230 (SD 0.345, range -0.590 to 0.883) and
mean discharge utility scores were 0.554 (SD 0.376, range -0.008 to 1.000). Fourteen clients reported health states as worse than death at admission and nine at discharge (Table 1). These data were not normally distributed and were tested for significance using the Wilcoxon signed-rank test. Utility scores between admission and discharge were significantly different (z=-4.26, p<.001).

Variability between DBT programmes in baseline health status on admission (p<0.001) and their ability to generate changes in EQ-5D scores between admission and discharge (p<0.0001) was apparent in the dataset. This was indicative of clustering in the data that would exaggerate the significance levels in analyses if not accounted for. An intra-programme correlation coefficient was calculated from the difference scores (treating DBT programme as a random effect) and used to inflate the width of the estimated confidence interval for mean differences to account for between-programme variability in outcomes. The high-level of clustering, indicated by an ICC of 0.71, increased the variance of the average difference score by a factor of approximately 12 according to the exact method of calculation for unequal cluster sizes given by Donner, Birkett and Buck [47]. The average difference in EQ-5D™ utility scores of 0.32 between admission and discharge failed to attain statistical significance once this adjustment was made (t=1.56, p=0.13).

The Reliable Change Index calculation indicated that EQ-5D™ difference scores of at least 0.45 could be considered reliable in the present study. Seventeen clients in the sample of 43 (40%) experienced reliable change between admission and discharge. The ability of clients to achieve change on the EQ-5D™ was constrained by an obvious ceiling effect in that over 25% in the DBT sample had a utility score >0.55 on admission and so could not achieve a change of >0.45.
Comparison of HRQoL scores with other published data sets

Ishak et al. [48] identified only 10 studies of psychotherapeutic interventions for BPD incorporating the use of HRQoL measures, of which only three report the EQ-5D™ as an outcome measure (McMain et al. [17] analyse the Euroqol VAS thermometer but do not report the EQ-5D utility scores) and, of these remaining three, only two represent unique datasets (van Asselt et al. [26] re-analyse the Giesen-Bloo 2006 dataset [49]). Both are studies in adult BPD. There is currently limited, but promising evidence to support the use of the EQ-5D instrument in CAMHS settings as a HRQoL measure [41].

Baseline and discharge EQ-5D™ scores for this clinical sample of admissions are presented in Table 1, alongside pre-and post-intervention EQ-5D™ scores from the few available RCTs in mental health for comparison purposes. As the present study is a small-uncontrolled study and primarily of clinical interest in relation to benchmarking outcomes across DBT programmes, effect sizes using Cohen’s d are also reported. The EQ-5D™ scores at treatment commencement are much lower in this study than in the studies of BPD in adult populations (Nadort et al t (103) = 2.15, p<.05, [50]; van Asselt et al t (89) = 2.70, p<0.005 [51]), both of which were of Schema-Focussed therapy. The adolescents in this pilot study end treatment at an EQ-5D™ score average commensurate with the starting EQ-5D™ scores of adolescents in the RCT for depression [41] (t (241) = 0.88, p>.02, ns). The effect size for this pilot study is higher than the two adult studies [50,51] but a similar size to the adolescent depression study [41].

INSERT Table 1 ABOUT HERE
Discussion

This study is the first to report on HRQoL of adolescents with BPD-type presentations in routine clinical practice. Participants in the DBT programmes in this study had significantly impaired health related quality of life (HRQoL) on admission, scoring significantly lower than data available from a published RCT using the EQ-5D with BPD participants [26] and from a study reporting on the treatment of adolescents with depression [41]. These data therefore support the often-expressed view of clinicians that clients in clinical services are more severe than those participating in clinical trials. Indeed, this view has been highlighted as a barrier to the implementation of evidence-based practice [52].

Alternatively the experience of BPD in adolescence compared to adulthood may have a particularly significant effect on HRQoL. Typically adolescents presenting with developing BPD in adolescence have experienced high levels of adversity in a context of genetic vulnerability over many years [2]. In addition to significant mental health difficulties with high rates of comorbidity [53, 54], they encounter major problems at school and with familial relationships and friends [55-57] that persist often into adulthood even when some of the more impulsive behaviours may have subsided. In clinical practice the broad ranging impact on almost all aspects of development is striking. Poor HRQoL in this context is perhaps unsurprising.

Despite the severity of impairment demonstrated in the patients treated in the DBT programmes, their HRQoL improved between admission and discharge with 40% of the sample achieving Reliable Clinical Change. This finding is particularly interesting when considering the fact that patients were discharged for multiple reasons – including both
planned (e.g. they completed the DBT programme and no longer needed DBT) and unplanned
discharges (e.g. the patient chose to stop attending treatment). This study reported on the
changes between admission and discharge for all patients who had received DBT regardless
of whether they completed the treatment programme or not. This finding also runs counter to
the views clinicians express [52] that increased clinical severity might prevent significant and
reliable clinical change.

Despite the change in HRQoL scores for patients in the DBT programmes, discharge scores
remained considerably lower than the general population norm and comparable to the scores
of adolescents commencing treatment in an RCT for depression [41]. These findings
underscore the view, reported in the literature, that individuals with BPD have significant
impairment of HRQoL [26-28] even following what is an effective treatment. The results
suggest that further research should be directed towards further enhancing clinical outcomes.

Typically, adaptations of DBT for adolescents have shortened the programme duration from 1
year (typical in adult services) to 16 weeks [8, 20]. The final HRQoL outcomes in this study
would argue against this given the low level of functioning of adolescents at treatment end
and that adolescents in this study had a longer average treatment length than that described in
the literature [8, 21]. In the shorter forms of DBT-A adolescents entering the programme
typically have fewer BPD symptoms (typically 2-3) whereas in the clinical programmes
studied here inclusion criteria to the programme required 5 or more BPD criteria which may
account for the longer treatment duration. These results suggest that for adolescents at this
level of severity longer treatment durations may be necessary.

Not all of the programmes were equally successful in producing good HRQoL outcomes.

When this variability was accounted for outcomes were not significantly different at treatment
end. The level of clustering in the dataset (ICC of 0.71) is unusually high, beyond the range
commonly encountered in naturally occurring biological and disease-related phenomena. Such variability is perhaps not uncommon in pilot data, especially with a team-based treatment delivered within highly variable organisational contexts. Whether the high levels of clustering were driven by differences in programme fidelity or therapist competence or are simply the artefact of small numbers of participants with a condition with highly variable outcomes is unknown. Future studies with more programmes with larger numbers of teams and treated clients will be necessary to tease out this finding. Paradoxically, the differential success of the DBT programmes in producing HRQoL outcomes, with some programmes performing better than others, augurs well for the intended use of the website for future benchmarking between programmes.

Whilst these findings in this pilot study are of interest, aspects of the data collection indicate that they should be interpreted with a significant degree of caution. Firstly, only four out of seven CAMHS teams with a subscription to the website were successfully using the system to collect routine data. The teams that were collecting data may have been especially motivated and thus potentially have been more likely to produce improved outcomes. Resolving problems in routinely collecting outcomes using the system would be essential for future meaningful use of the system to collect national outcome data or to benchmark programmes systematically. Secondly, these data were collected under routine clinical practice conditions and so the research team did not control data collection and entry. Each individual DBT programme operated their own data administration, collection and entry procedures for their own clinical purposes and this may have led to different protocols for data collection. In some teams the proportion of EQ-5Ds completed by clinicians as a proxy, may have been higher and in some cases clinicians recorded the measure retrospectively, particularly in circumstances where patients left the programme prematurely. Both of these practices may have resulted in biases in the data. However, as all teams were collecting data in real time
(i.e. they were administering scores as and when patients were admitted and discharged) before consenting to participate in the research study, opportunities for systematic bias were reduced. Secondly, the absence of any additional data on either the participants or other outcomes limits generalizability. Thirdly, the absence of a control group means that change in the sample cannot be attributed to the treatment that they received. The absence of a control group and the high-level of clustering means that these data cannot make any definitive statement about whether DBT is effective in adolescents with BPD treated in routine clinical practice.

In conclusion, although further research is necessary to unpick the findings of this pilot study, the DBT outcome monitoring website has demonstrated its potential to collect data in routine practice and in real-time and thus may be a promising tool to benchmark what is gained and lost following the implementation of DBT in clinical practice settings.
Acknowledgements

We would like to extend out thanks to the DBT teams and their patients who participated in this study. Thanks to Dr. Barbara Baragwanath for tracking down references and proofreading the manuscript.

Compliance with Ethical Standards

**Funding:** The study was funded by a grant to Bangor University from the Knowledge Economy Skills Scholarships (KESS) programme (Bangor University KESS Mini 016), and by Integral Business Support Ltd.

**Conflicts of Interest:** MS is the Director of the British Isles DBT Training Team that trains practitioners in DBT with a licensed training programme and receives fees for training in DBT. She also receives royalties from books and training products in DBT. MS is married to the managing director and principal shareholder of Integral Business Support Ltd that delivers licensed training in DBT. RABH is the Managing Director and principal shareholder of Integral Business Support Ltd that delivers licensed training in DBT. RPH’s spouse is a member of the British Isles DBT Training Team and receives income from DBT training.

**Ethical approval:** All procedures were in accordance with the ethical standards of the institutional and national research committees and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

**Informed consent:** The Ethical Committees to whom the study was submitted did not require informed consent for individuals as the data was pseudonymised and was collected as part of routine clinical data collection.
Authors’ Contributions

MS, RABH, & RPH designed the study; LB recruited participants and supervised data collection; MS & RPH supervised the study on a day-to-day basis; RABH & LB analyzed the data; MS & LB wrote a first draft of the paper; all authors contributed to the final draft of the manuscript.
References


Table 1: Comparison of EQ-5D scores across studies.

<table>
<thead>
<tr>
<th>Sample</th>
<th>EQ-5D at t1</th>
<th>EQ-5D at t2</th>
<th>t2-t1 (mean length of treatment)</th>
<th>Average Difference</th>
<th>effect size d</th>
</tr>
</thead>
<tbody>
<tr>
<td>Present study: n=43 routine admissions to DBT programmes</td>
<td>0.23 (0.12 – 0.34)</td>
<td>0.55 (0.44 – 0.67)</td>
<td>10 months</td>
<td>0.32 (0.20 – 0.44)</td>
<td>1.00</td>
</tr>
<tr>
<td>RCTs in Adults</td>
<td></td>
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</tr>
<tr>
<td>Nadort et al. [50] n=62 SFT v modified SFT for adult BPD</td>
<td>0.44 (0.31 – 0.56)</td>
<td>0.56 (0.47 – 0.65)</td>
<td>18 months</td>
<td>0.12 (0.02 – 0.22)</td>
<td>.39</td>
</tr>
<tr>
<td>van Asselt et al. [51] n=48 transference v SFT for adult BPD</td>
<td>0.50 (0.41 – 0.59)</td>
<td>0.69 (0.61 – 0.77)</td>
<td>3 years</td>
<td>0.20 (0.09 – 0.29)</td>
<td>.64</td>
</tr>
<tr>
<td>RCTs in CAMHS</td>
<td></td>
<td></td>
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<td></td>
</tr>
<tr>
<td>Byford et al. [41] n=199 CBT/SSRI v SSRI for adolescent depression</td>
<td>0.50 (0.46 – 0.54)</td>
<td>0.76 (0.72 – 0.80)</td>
<td>28 weeks</td>
<td>0.26 (not reported)</td>
<td>.90</td>
</tr>
</tbody>
</table>

1 Includes 9 clients (21%) admitted in state 11233 whose utility remained unchanged at -0.008
2 Relative to baseline variance which is the effect size commonly used in power calculations, Nadort et al. [50] calculate effect sizes relative to pooled variance (pre/post) and report a lower estimate of 0.35