

# CORE-10 USER MANUAL

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#### **PREAMBLE**

### **Acknowledgements & enquires**

The development of the CORE-10 and manual was funded by the Artemis Trust and Priorities & Needs R&D funding via Leeds Mental Health Teaching NHS Trust. Enquires regarding CORE-10 itself should be sent to CORE-IMS at <a href="mailto:admin@coreims.co.uk">admin@coreims.co.uk</a>

#### Citing the manual

This manual should be referenced as follows:

Connell, J. & Barkham, M. (2007). *CORE-10 User Manual, Version 1.1.* CORE System Trust & CORE Information Management Systems Ltd.

### Copyright

Paper versions of all CORE measures are copyleft: that is, all measures can be freely photocopied but cannot be changed in any way or used for commercial gain. Services who wish to incorporate CORE-questionnaires into locally developed/commissioned software should seek permission from the CORE-System Trustees at <a href="mailto:cst-iapt@psyctc.org">cst-iapt@psyctc.org</a>.

### IT support for IAPT pathfinder sites

The CORE System Trust (CST) and CORE User Network have asked for the following information to be available to potential Improving Access to Psychological Therapy pathfinder sites.

#### **Reproduction of CORE measures in Trust software**

CST recognises that pathfinder sites may prefer to develop/commission local software to capture, store and report on data for all the measures in the Minimum Data Set (MDS) or to incorporate such functionality into an existing software application. A service wishing to incorporate the CORE questionnaires into locally developed/commissioned software (or to incorporate them into an existing software application) should seek prior permission from the Trustees of CST because this would otherwise constitute a breach of copyright. The CST Trustees can be contacted at <a href="mailto:cst-iapt@psyctc.org">cst-iapt@psyctc.org</a>. CORE questionnaires may be reproduced freely on paper.

#### **CORE-Net software**

CST and the CORE User Network welcome the Improving Access to Psychological Therapies Pathfinder programme and are supportive of the central role given to the measurement and evaluation. The pathfinder programme sets a demanding schedule which involves capture and reporting of data from a wide range of measures, including CORE, starting in September 2007 for a period of 6-7 months. At the request of services who are already using CORE and who are considering bidding as a pathfinder site, CORE Information Management Systems Ltd (CORE IMS) have agreed to develop the existing CORE-Net software to allow data capture for the key measures included in the Minimum Data Set. For further details of available support, see Section V.

#### SECTION I

#### Introduction

The CORE-10 is a brief outcome measure comprising 10 items drawn from the CORE-OM which is a 34-item assessment and outcome measure. The CORE-OM has been widely adopted in the evaluation of counselling and the psychological therapies in the UK.

Where resources and time allow, the CORE-OM remains the preferred version for the assessment and measurement of outcome. This is because it offers a wider range of items for clients to identify to practitioners at assessment and provides more precise estimates of outcomes and change following an intervention. In addition, there is to date a far larger amount of accumulated data on the CORE-OM from which to derive benchmarks and comparisons.

However, there is a need for a shorter measure in such settings as busy practices where information is needed quickly and which places minimum demands on clients and practitioners. In addition, further measures tapping other outcomes may be needed from services such that some trade-offs have to be made. For the pathfinder sites taking part in the Improving Access to Psychological Therapies initiative, the CORE-10 has been selected as part of a minimum data set that will enable services to meet the requirements of the Balanced Score Card with the least burden to clients and services.

#### The CORE family of measures

The CORE-10 is one instrument within the CORE family of measures which, together with the CORE System, are briefly summarised in Section V. The parent measure is the CORE-OM, the properties of which have been reported widely in the literature. [1-10] In addition it has been used widely in benchmarking applications within NHS settings. [11-14] The CORE-OM is part of a broader CORE System which has also been reported in the literature and provides a context within which to evaluation outcomes. [15,16]

Acknowledging that one outcome measure is not fit for all purposes, shorter versions derived from the CORE-OM have also been developed. These include CORE Short Forms A&B for session-by-session use in research settings<sup>[3, 7]</sup> and the CORE-5 for session-by-session use in practice settings.<sup>[17]</sup> Population-specific versions of CORE also exist for the general population (GP-CORE)<sup>[18]</sup> and learning disabilities (LD-CORE).<sup>[19]</sup> A range of translations are also available (see Section V). Training and IT support for CORE is available and has been documented together with a resulting national research database.<sup>[20, 21]</sup>

#### Overview

The CORE-10 has 10 items on one side of A4, making it quick and easy to administer. It can easily be scored by hand. The CORE-10 has been designed to tap into a pan-theoretical 'core' of users' distress, including commonly experienced symptoms of anxiety and depression and associated aspects of life and social functioning. In addition, there is a key item on risk to self.

The CORE-10 taps global distress and is, therefore, suitable for use as an initial quick screening tool and also as an outcome measure. Like most self report measures, it cannot be used to gain a diagnosis of a specific disorder. A *clinical score* can be derived directly by summing the items and used as a global index of distress. The risk item should be regarded as a clinical flag and some services may wish to use it to trigger more discussion of risk at assessment. A non-response to the risk item should also be explored further.

It is important to remember that no service can be fully evaluated through the use of one outcome measure alone. Hence, the CORE-10 should be viewed as one component of an evaluation system to be used either with the CORE System tools (see Section V) or with whatever local or theory-specific methods and clinical supervision/audit practices are in place.

#### **Development**

The CORE-10 was derived from the 34 item CORE-OM using a number of specific requirements to determine item selection. First, we sought to devise a tool that met the following characteristics: item coverage, format, range of intensity, and content. Secondly, we sought to find a balance between the role of statistical procedures, face and clinical validity.

<u>Item coverage</u>: We aimed to ensure that the measure provided a balance in terms of being short and easily scored but still giving sufficient scope to tap as far as possible the 10 item clusters in the CORE-OM: (1) subjective well-being, (2) anxiety, (3) depression, (4) physical, (5) trauma, (6) general functioning, (7) close relationships (functioning); (8) social relationships (functioning), (9), risk to self, and (10) risk to others.

<u>Format</u>: Although our aim of restricting the item number to 10 was driven by brevity, the specificity of 10 items yielded additional advantages in terms of simplicity of scoring. Simple summing of the individual item totals would directly yield a 'clinical' score for the CORE measure as has been used in recent reports and would therefore not require practitioners to make any further calculations.<sup>[9]</sup>

<u>Range of intensity</u>: We sought to draw on both high and low intensity items. Excluding the 6 risk items, the CORE-OM is balanced between high and low intensity items (i.e., 14 each). Hence, we sought a structure that reflected 6 high and 4 low intensity items.

Content: Although our aim of 10 items tapping 10 item clusters would appear to fit a one-item per cluster assignment, we were cognisant of the need to sample both high and low items in relation to *depression* and *anxiety* due to their high prevalence in the population. Hence, this combination alone would require a total of 4 items. In terms of functioning, we elected to select one item from each of the three clusters – *general, social, and close* relationships yielding a total of 3 items. In addition, we selected single items from *physical* and from *trauma* – a total of 2 items. To keep to 10 items, we selected a single *risk* item as a key component in screening with 'risk to self' taking priority over 'risk to others' which has a relatively low incidence rate in people presenting in primary care. To accommodate this structure, we elected to omit items from the Subjective Well-being domain due to their having a high correlation with the problem domain, higher than the correlation between any other two domains<sup>[3]</sup>. In addition, we aimed to exclude symptom and functioning items which had been found to be most frequently omitted by clients in previous reports on the CORE-OM.<sup>[3,5]</sup>

In order to achieve the above requirements, we employed a method comprising four steps:

<u>Step 1: Omission of items</u> First, we omitted 'Subjective Well Being' items [4, 14, 17, 31] and 'Risk to Others' items [6 and 22]. Second we omitted frequently missed items from the symptom and functioning dimensions [19, 20, 21, 30, and 32]. [3, 5]

<u>Step 2: Item coverage and mapping</u> The omission of items in step 1 resulted in some items being 'forced choices' in order to retain items from specific clusters and the balance of 6/4 high and low intensity items. This resulted in the selection of items **2** (*low anxiety*), **7** (*high general functioning*), **27** (*low depression*), and **3** (low *close relationship*) with the subsequent omission of items 12, 1, 26, 25, 29, 9, 24. There remained 6 paired items in which one item from each pair needed to be selected.

<u>Step 3: Regression analyses</u> We then employed regression analyses to identify items which best predicted item cluster scores on the CORE-OM. This resulted in the selection of items **23** (*high depression*), **10** (*high social functioning*), **18** (*low physical*), and **16** (*high risk to self*).

<u>Step 4: Clinical judgement</u> Where decisions on the basis of regression analyses were marginal, we invoked clinical judgement to inform item selection. For *trauma* item **28** was more focused specifically on trauma than item 13. For *high anxiety* there was a clear preference for item **15** rather than item 11.

The data used for the development of the CORE-10 measure consisted of 6610 clients drawn from 33 primary care services. Using SPSS 12.01 the data was split into 3 random groups with each data set drawing from the 33 services. One data set was used for item selection, as detailed above. The second dataset was used as a means of replicating results obtained from the first sample. The third sample, together with other datasets, was used as an independent sample to determine a range of psychometric properties of the extracted items.

Table 1: CORE-10 development - Selection of items

Item	Domain: Cluster	Item	Inten- sity	Step Omitted/ Included
4	Subj Wellbeing	I have felt O.K. about myself	Lo	1 (omit)
14	Subj Wellbeing	I have felt like crying	Hi	1 (omit)
17	Subj Wellbeing	I have felt overwhelmed by my problems	Hi	1 (omit)
31	Subj Wellbeing	I have felt optimistic about my future	Lo	1 (omit)
2	P: Anxiety	I have felt tense, anxious or nervous	Lo	2 (incl)
20	P: Anxiety	My problems have been impossible to put to one side	Lo	1 (omit)
11	P: Anxiety	Tension and anxiety have prevented me doing important things	Hi	3 (omit)
15	P: Anxiety	I have felt panic or terror	Hi	3 (incl)
5	P: Depression	I have felt totally lacking in energy and enthusiasm	Hi	3 (omit)
23	P: Depression	I have felt despairing or hopeless	Hi	3 (incl)
27	P: Depression	I have felt unhappy	Lo	2 (incl)
30	P: Depression	I have thought I am to blame for my problems and difficulties	Lo	1 (omit)
8	P: Physical	I have been troubled by aches, pains or other physical problems	Lo	3 (omit)
18	P: Physical	I have difficulty getting to sleep or staying asleep	Lo	3 (incl)
13	P: Trauma	I have been disturbed by unwanted thoughts and feelings	Hi	3 (omit)
28	P: Trauma	Unwanted images or memories have been distressing me	Hi	3 (incl)
1	F: Close Rel	I have felt terribly alone and isolated	Hi	2 (omit)
3	F: Close Rel	${\bf I}$ have felt ${\bf I}$ have someone to turn to for support when needed	Lo	2 (incl)
19	F: Close Rel	I have felt warmth and affection for someone	Lo	1 (omit)
26	F: Close Rel	I have thought I have no friends	Hi	2 (omit)
7	F: General F	I have felt able to cope when things go wrong	Hi	2 (incl)
12	F: General F	I have been happy with the things I have done	Lo	2 (omit)
21	F: General F	I have been able to do most things I needed to	Lo	1 (omit)
32	F: General F	I have achieved the things I wanted to	Hi	1 (omit)
33	F: Social Rel	I have felt humiliated or shamed by other people	Hi	3 (omit)
10	F: Social Rel	Talking to people has felt too much for me	Hi	3 (incl)
25	F: Social Rel	I have felt criticised by other people	Lo	2 (omit)
29	F: Social Rel	I have been irritable when with other people	Lo	2 (omit)
22	R: Harm to O	I have threatened or intimidated another person	Hi	1 (omit)
6	R: Harm to O	I have been physically violent to others	Hi	1 (omit)
9	R: Harm to S	I have thought of hurting myself	Lo	2 (omit)
16	R: Harm to S	I made plans to end my life	Hi	3 (incl)
24	R: Harm to S	I have thought it would be better if I were dead	Lo	2 (omit)
34	R: Harm to S	I have hurt myself physically or taken dangerous risks with my Health	Hi	3 (omit)

**Domains:** Subj Wellbeing = Subjective Wellbeing; P = Problems; F = Functioning; R = Risk

**Clusters:** Close Rel = Close Relationships; General F = General Functioning; Social Rel = Social Relationships Harm to O = Harm to Others; Harms to S = Harm to Self

#### Structure

Item selection for the CORE-10 was determined by a combination of clinical utility, coverage of item clusters, and statistical procedures (i.e., regression analyses). The 10 items comprise the following: Depression (2 items), Anxiety (2 items), Functioning (3 items, 1 each for 'general' 'social', and 'close'); Trauma (1 item); Physical (1 item); and Risk (1 item). The measure includes high and low intensity items to increase sensitivity and is loaded slightly in favour of higher intensity items (6 items) in order to reduce any ceiling effect. Two of the items are 'positively' framed. The items are presented in Table 2. They are presented here grouped according to problem areas (i.e., item clusters).

Table 2: CORE-10 items

Item	Problem Area	Item	Positive/ Negative	Intensity
1	Anxiety	I have felt tense, anxious or nervous	-	Low
5	Anxiety	I have felt panic or terror	-	High
9	Depression	I have felt unhappy	-	Low
8	Depression	I have felt despairing or hopeless	-	High
7	Physical	I have had difficulty getting to sleep or staying asleep	-	Low
10	Trauma	Unwanted images or memories have been distressing me	-	High
2	Close relationships	I have felt I have someone to turn to for support when needed	+	Low
4	Social relationships	Talking to people has felt too much for me	-	High
3	General Functioning	I have felt able to cope when things go wrong	+	High
6	Risk	I made plans to end my life	-	High

#### **Format**

The actual sequence of items in the CORE-10 follows the order they appear in the CORE-OM and is shown in the item number in Table 2. The final version of the CORE-10 is presented in Appendix I.

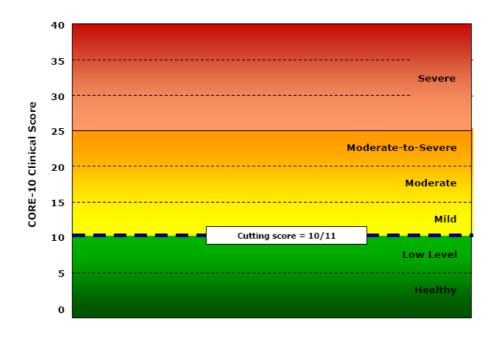
#### Scoring procedures and meaning

Key points in the scoring of the CORE-10 are as follows:

- 1. Each item within the CORE-10 is scored on a 5-point scale ranging from 0 ('not at all') to 4 ('most or all the time').
- 2. The clinical score is calculated by adding the response values of all 10 items.
- 3. Where there are missing data the clinical score is derived by calculating the total mean score (dividing the total score by the number of completed items) and multiplying by 10.
- 4. We do not recommend re-scaling the clinical score if more than one item is missing.
- 5. The minimum score that can be achieved is 0 and the maximum is 40.
- 6. The measure is problem scored, that is, the higher the score the more problems the individual is reporting and/or the more distressed they are.

A score of 10 or below denotes a score within the non-clinical range and of 11 or above within the clinical range. Within the non-clinical range we have identified two bands called 'healthy' and 'low' level distress. People may score on a number of items at any particular time but still remain 'healthy'. Similarly, people may score in the 'low' range which might be a result of raised pressures or particular circumstances but which is still within a non-clinical range. Within the clinical range we have identified the score of 11 as the lower boundary of the 'mild' level, 15 for the 'moderate' level, and 20 for the 'moderate-to-severe' level. A score of 25 or over marks the 'severe' level.





#### Normative data

Normative data is available for three samples, two clinical and one non-clinical population (Table 3). For samples 1 and 3 the normative data is for the CORE-10 items as embedded in the CORE-OM (i.e., the CORE-OM has been administered to the sample and the scores for the items making up the CORE-10 subsequently extracted).

### Non-clinical sample: General population (sample 1)

The sample comprised 553 adults who participated in the follow-up to the psychiatric morbidity survey and returned valid CORE-OM forms was used. [22] This data was weighted to be representative of a general population taking into account design factors, non-response and sampling procedures from the original psychiatric morbidity sample to the subsequent follow-up sample. [23] This resulted in an effective general population sample of 535 cases consisting of 268 males (50.2%) and 267 females (49.8%) with a mean age of 43.4 years (SD = 15.3).

### **Clinical sample: Primary care routine practice (sample 2)**

The sample comprised one third, randomly selected, of a dataset of 5831 people presenting in 33 primary care services who had completed a CORE-OM. The other two thirds of the dataset were used in the development and psychometric analysis of the measure. This clinical sample consisted of 1835 clients comprising 1319 females (72%) and 516 males (28%). The average age was 38 (SD = 12.6) ranging from 16 to 84 years. Ethnicity data was available for 1606 (87.5%) people. Of these, 92% were White/European and the remainder were Asian (4%), African/Caribbean (2%), and Mixed/Other (2.3%). The majority of clients presented with anxiety (71%) and/or depression (65%). Half of the sample reported experiencing interpersonal problems (50.9%).

#### Clinical sample: GP practice (sample 3)

This data was drawn from a naturally occurring sample from a GP practice in the north of England who completed the CORE-10 consistent with the Quality Outcomes Framework agenda. A total of 323 measures were completed. In the majority of cases (n=271;84%) the client completed the measure directly onto computer. The ten items comprising the CORE-10 were presented individually to the client on the computer monitor. A response was required before the next item was offered. The remaining clients (n=52;16%) completed a pen and paper version of the CORE-10. Data was collected between April and October 2006. A reliable measure (no more than 1 missing item) was completed by 321 patients; 209 (65%) females and 112 (35%) males. Age data was unavailable.

Table 3: Means, standard deviations, and confidence intervals for clinical and non-clinical sample populations

	Sample 1 General Population (n=535)			ral Population Primary Care		Sample 3 GP Practice (n=321)			
	Mean	SD	CI	Mean	SD	CI	Mean	SD	CI
Males	4.8	4.6	4.3-5.4	18.5	8.1	17.8-19.2	19.8	7.6	18.3-21.2
Females	4.7	4.9	4.1-5.2	20.1	7.5	19.7-20.5	20.4	8.0	19.3-21.5
Total	4.7	4.8	4.3-5.2	19.7	7.7	19.3-20.0	20.2	7.9	19.3-21.0

#### **Gender differences**

There was no significant difference between males and females in the General Practitioner (stand alone/non-embedded, clinical) sample (19.8 vs. 20.4; t=.68, p=.50) or the General Population (non-clinical) sample (4.8 vs. 4.7; t=.46, p=.64). There was, however, a statistically significant difference between males and females for the Primary Care Counselling (embedded, clinical) sample (18.5 vs. 20.1; t=-4.1, p<.001). More data is required to establish whether gender should be taken into account when relating individual scores to clinical distribution data.

### **Section II**

### **Psychometric qualities: Is the measure valid and reliable?**

A measure should be reliable, valid, and responsive to the clinical change that occurs over time.

- Validity is the extent to which the measure measures what it intends to measure;
   that is, is it asking the right questions?
- Reliability is how uniformly the test can be repeated when administered on more than one occasion or by more than one rater.
- Responsiveness or sensitivity is the ability of the measure to detect true change in clients' status over time; that is, is it sensitive to the subtle changes patients make?

#### Discrimination between clinical and non-clinical populations

One aspect of validity is the requirement of the measure to show a statistically and clinically significant difference between the clinical population for which it has been designed and a non-clinical sample. A comparison of the Primary Care Counselling (clinical) and the General Population (non-clinical) samples revealed a large and clinically significant difference between the two populations (19.7 vs. 4.7; t = 37.3; p < .001). This is illustrated in the box plot of the data (Figure 2). The boxes cover the middle 50% of the scores in each group. The (red) dot in the box indicates the median and the dark blue 'notched' area around the line is the confidence interval (CI) around the median. Neither the boxes nor the confidence intervals overlap, indicating the statistically significant difference between the two populations. It also shows that there are very few cases in the clinical sample scoring zero and very few outliers in the non-clinical sample scoring very high, a further indication of the difference in scores between the two populations.

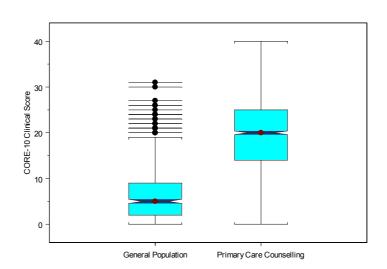


Figure 2: Box-plot showing difference between clinical and non-clinical populations

#### Discrimination between clients with and without a diagnosis of depression

Whether a measure can discriminate between those diagnosed and not diagnosed with a problem refers to the sensitivity and specificity of the measure. Sensitivity and specificity relate to the proportion of people correctly 'diagnosed' by a measure when compared with a well established 'gold standard' diagnostic tool. The sensitivity of the measure is the proportion of people with a positive diagnosis on both tests and specificity the proportion of people with a negative diagnosis on both tests. A perfect test is one that achieves 100% sensitivity and 100% specificity. However this state is virtually unobtainable and there is always some trade off between the rates. The lower the trade off between sensitivity and specificity the better the measure performs as a diagnostic test.

Table 4 shows a range of cut off scores on the CORE-10 against a diagnosis of DSM-IV depression using the SCID. [24] A cut-off score for depression of 13 on the CORE-10 yields sensitivity and specificity values of .92 (CI = .83-1.0) and 0.72 (CI = .60-.83) respectively.

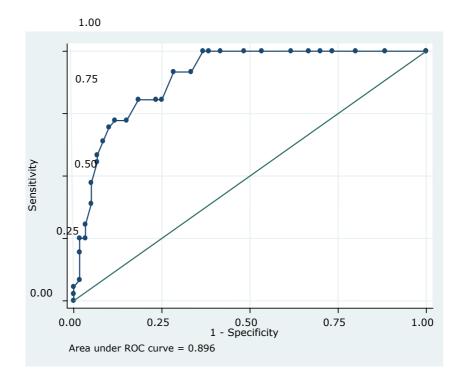
Table 4: Sensitivity and specificity of the CORE-10 against a SCID diagnosis of depression

Cut	Sensitivity	Specificity
point	(true positives)	(true negatives)
≥ 13	0.92	0.72
≥ 14	0.81	0.75
≥ 15	0.81	0.77
≥ 16	0.81	0.82
≥ 17	0.72	0.85
≥ 18	0.72	0.88
≥ 19	0.69	0.90
≥ 20	0.64	0.92
≥ 21	0.58	0.93

As outlined above a good diagnostic test is one that has small false positive (high sensitivity) and small false negative (high specificity) rates across a reasonable range of cut off values. This can be tested using a ROC (Receiver Operating Characteristic) curve which is a graphical representation of the trade off between the false negative and false positive rates for every possible cut off (see Figure 3).

A rapid climb towards the upper left hand corner of the graph signifies that the false negative rate is high and the false positive rate is low. A diagonal path from the lower left hand corner to the upper right hand corner means that every improvement in false positive rate is matched by a corresponding decline in the false negative rate. To quantify how quickly the graph climbs towards the left hand corner the area under the curve is measured. The closer the area is to 1.0 the better the test. The area under the curve for the cut off values of the CORE-10 compared with a DSM-IV SCID diagnosis was 0.896 (SE = 0.031, 95%CI = 0.835 to 0.957).

Figure 3: ROC curve of the CORE-10 against the SCID diagnosis of moderate depression



#### **Correlation with other measures**

It is expected that a measure of psychological distress would correlate with other well-established mental health measures examining similar constructs. We identified reports in which the CORE-OM had been correlated with other outcome measures to which we had access to the original data. [3, 7, 23, 25] We recalculated the correlations using the 10 CORE items drawn from the CORE-OM. The correlation between the CORE-10 items and other measures is shown in Table 5. These results indicate a good correlation between the CORE-10 and other measures of anxiety, depression and overall mental health.

Table 5: Correlation of the CORE-10 with other measures

Measure	CORE-10	Reference for dataset used
Symptom CheckList-90-R	.81	[3]
Brief Symptom Inventory	.75	[3]
Beck Depression Inventory	.77	[3]
Beck Depression Inventory-II	.75	[7]
Beck Depression Inventory-II	.76	[3]
Beck Anxiety Inventory	.65	[3]
Patient Health Questionnaire-9	.56	[25]
Clinical Interview Schedule-R	.74	[23]

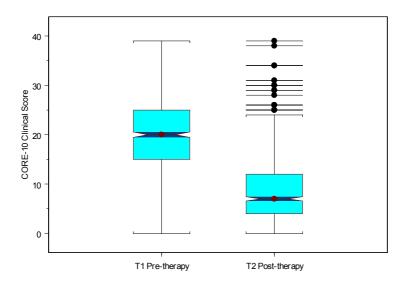
### Internal reliability: Do all the items measure the same trait?

One minimal requirement of any measure is that it shows high internal reliability. Internal reliability tells you whether the items are all measuring the same trait by examining the consistency of responses. Thus a series of random unrelated questions would have a low internal consistency and conversely a series of related items (e.g., known symptoms of depression) should have a high internal consistency. Internal consistency is measured on a scale of 0-1. A very high score (e.g., .9) may indicate redundant items, a low score (e.g., .2) indicates inconsistent or random responses. A measure does not need to be unidimensional to achieve a high internal consistency as multiple dimensions can be interrelated. Internal reliability is measured using Cronbach's coefficient alpha which is the percentage of the variance in a score that is covariance between items. An alpha of 0.82 (CI .79 to .85) provides evidence of very good internal consistency. For women the alpha was .82 (CI .78 to .86) and for men .81 (CI .76 to .86).

### Sensitivity to change

The most important requisite of an outcome measure is that it is responsive to change over time. To test this we used the pre- and post-therapy data within the Primary Care Counselling (clinical) sample. The sample consisted of 780 clients of whom 27% were male and 73% female with an average age of 40. The clients had an average of 6 therapy sessions. The pre-therapy score for those clients with both a pre- and post-therapy measure was 19.5 (SD = 7.6) compared with a post-therapy score of 8.6 (SD = 6.7) yielding a change score of 10.9 (SD = 8.0). The difference was statistically significant (t = 38.1, p<.001). The difference between the pre- and post-therapy scores is illustrated in Figure 4.

Figure 4: Difference between CORE-10 pre- and post-therapy scores



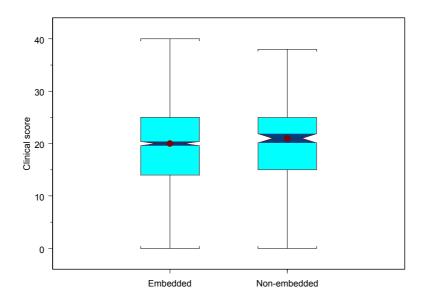
### **Acceptability**

One aspect of acceptability can be judged by the rates of completion and non-completion of items. One of the criteria for inclusion of items in CORE-10 was a low rate of missing data for each of the chosen 10 items from the original CORE-OM. From the Primary Care Counselling (clinical) sample there was a mean of .08 (SD = .35) missing items. All ten items were completed by 94.4% of the sample. This analysis was repeated on the General Population (non-clinical) sample. When clients who had obviously not turned over the page were omitted (n = 4), for the CORE-10 items there was a mean of .06 (SD = .46) missing items with 520 (96.7%) completing all 10 items. The analysis could not be repeated on the General Practitioner sample as this was administered using computer software where it was mandatory to complete one item before the next item became available.

#### Comparison of embedded vs. stand alone version

The majority of the psychometric testing of the CORE-10 has been performed on the CORE-10 items as embedded within the CORE-OM. This is a logical step in measure development. Although presenting items outside the context of the other items may yield slight differences in how clients respond, it would be expected that these differences will be relatively small. Indeed, a comparison of the Primary Care (embedded) and General Practitioner (non-embedded) samples found the CORE-10 measure yielded a significantly higher score than from the embedded items (19.7 vs. 20.2 t=41.3, p<.001). However the difference was small and clinically insignificant as illustrated in Figure 5.

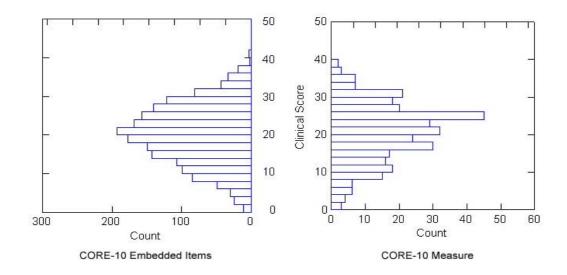
Figure 5: Notched box plot of embedded and non-embedded CORE-10 items



The boxes cover the middle 50% of the scores in each group. The (red) dot in the box indicates the median and the dark blue 'notched' area around the line is the confidence interval (CI) of the median. The similarity between the two samples is illustrated by the comparable range of scores for the two samples. The narrow confidence interval for the Primary Care Counselling sample is an indication of the precision of the median score as a result of the large number of clients in this dataset. This is the reason for the non-overlap of the confidence intervals and hence the statistically significant difference between the two samples.

We also plotted the distribution of scores for the CORE-10 measure against that of the CORE-10 embedded items which are shown in Figure 6. The shapes of the plots are similar. The clinical scores ranged from 0 to 38. Of the total sample of 321 people, 85.7% scored 11 or above (i.e., scored above the cut-off as indicating psychological distress) while the percentage of people presenting with depression was 81.7% as determined by use of a cutting score of 13. But this value would be lower if a higher cutting score was adopted.

Figure 6: Back-to-back histograms of embedded and non-embedded CORE-10 items



#### **Section III**

#### Reliable and clinically significant change

Researchers traditionally compare group means for treatments using statistical methods to suggest whether change is a sampling artefact or chance finding. Practitioners are more concerned with changes in particular individuals and often dichotomise outcome as 'successful' or 'not successful'. Jacobson and colleagues proposed methods bridging these languages by presenting changes in the group under investigation at the level of the individual. <sup>[26]</sup> The methods rest on two questions being asked of each client's data:

- Has the client's score changed sufficiently to be confident that the change is not attributable to measurement error? This question relates to 'reliable change'.
- How does the end state score of the client compare with the scores observed in socially
  and clinically meaningful comparison groups? And more precisely, is the end-state score
  better represented by a non-clinical or general population rather than a clinical
  population? This question relates to 'clinically significant change'.

To achieve reliable and clinically significant change a client's score must fulfil both these criteria.

#### Reliable change

Reliable change refers to the extent to which change falls beyond that likely based on the measurement variability of the measure. This is calculated using the following formulae:

$$(SE_{diff} = SD_1\sqrt{2}\sqrt{1-r}) \times 1.28$$

where  $SD_1$  is the standard deviation of the baseline (pre-therapy) observations and, r is the reliability (coefficient alpha) of the measure. Multiplying by 1.28 yields a value which is unlikely to occur more than 10% of the time due to the unreliability of the measure alone. The value of 1.28 is recommended for use with short outcome measures as opposed to the traditional value of 1.96. $^{[27,28]}$ 

The resulting values for the CORE-10 ( $SD_1$ = 7.9 r=.82) gives us a reliability figure of 5.9 which we would round up to 6.0 for ease of measurement. The client must therefore improve by 6.0 or more from pre- to post-therapy to be able to say that they have made reliable improvement.

# Cutting score between clinical and non-clinical populations

The cutting score to discriminate between a clinical and non-clinical population was calculated using the Primary Care Counselling (clinical) and General Population (non-clinical) samples and the formulae proposed by Jacobson and Truax:<sup>[26]</sup>

$$\frac{mean_{clin}sd_{norm} + mean_{norm}sd_{clin}}{sd_{norm} + sd_{clin}}$$

Replacing this with the values for the CORE-10 (see Table 3) yielded a cutting score for the sample as a whole of 11.0. The difference between men (10.6) and women (11.1) was less than 1 full point.

It is therefore recommended that a cut off of 10/11 is used where 10 is in the non-clinical range and 11 in the clinical range.

In summary, to achieve reliable improvement, a client must improve by 6.0 or more points from pre- to post-therapy. To achieve clinically significant change they must change from a pre-therapy score of 11 or above to a post-therapy score of 10 or below. To achieve reliable and clinically significant change both criteria must be fulfilled. Note that those clients whose pre-therapy score is below the cutting score (average 13% of total clients), while it is possible to achieve reliable improvement it is not possible to achieve clinical improvement because their score is already below the cutting score. For this reason it is often necessary to calculate reliable and clinical improvement rates for (a) only for those clients above the cut off value and (b) for all clients, particularly if a service has a high rate of clients with lower client scores at intake. Table 6 shows comparative improvement rates for the Primary Care Counselling sample using four mutually exclusive groups: (1) reliable and clinically significant improvement, (2) reliable improvement only, (3) no reliable change, and (4) reliable deterioration.

Table 6: Rates of reliable and clinically significant change for Primary Care Counselling

	All cli (n=3		Clients above cut off only (n=263)	
	N	%	N	%
Reliable and clinically significant improvement	439	54.8	439	63.4
Reliable improvement only	170	21.2	146	21.1
No reliable change	173	21.6	95	13.7
Reliable deterioration	19	2.4	12	1.7

#### **Section IV**

# Comparison of CORE-10 with CORE-OM

#### Correlation with the CORE-OM

CORE-OM and CORE-10 clinical scores were calculated for the clinical (Primary Care Counselling) and the non-clinical (General Population) samples. For the clinical sample the two scores correlated at r=0.94 (CI 0.93 to 0.95). For the non-clinical sample the two scores correlated at r=0.92 (CI 0.91 to 0.93). This high correlation is an indication of the close relationship between the clinical scores of the CORE-10 and the CORE-OM.

#### **Correlation with other measures**

The differences between correlations of the CORE-10 and CORE-OM with other common mental health measures are shown in Table 7. Despite the considerable reduction in items from the CORE-OM to the CORE-10, the correlations with mental health measures measuring similar constructs remains high.

Table 7: Comparison of the CORE-10 and CORE-OM correlation with other measures

Measure	CORE-10	CORE-OM	Difference
Symptom CheckList-90-R	.81	.88	.07
Brief Symptom Inventory	.75	.81	.06
Beck Depression Inventory	.77	.85	.08
Beck Depression Inventory-II	.75	.75	.00
Beck Depression Inventory-II	.76	.81	.05
Beck Anxiety Inventory	.65	.65	.00
Patient Health Questionnaire-9	.56	.63	.07
Clinical Interview Schedule-R	.74	.77	.03

#### **Internal reliability**

Despite the considerable reduction in the number of items, which will almost invariably result in a lowering of the coefficient alpha, the internal consistency for the CORE-10 is good at 0.82 (CI .79-.85). By comparison, the coefficient alpha for the CORE-OM is .94 (CI .93-.95) for both clinical and non-clinical populations.<sup>[3]</sup> Such a high value might suggest redundancy of items.

### Reliable and clinical change

Using the Primary Care Counselling sample a comparison of the rates of reliable and clinical change for the CORE OM and CORE-10 was carried out (Table 8). For the CORE-OM a cut off value of 10 and an RCI of 5 were used while the values for the CORE-10 a cutting score of 11 and an RCI of 6 were used. [22] The difference between the two measures was small with a slightly higher proportion of clients achieving reliable and clinical change on the CORE-OM who did not on the CORE-10 and a corresponding higher rate of clients achieving reliable change, but not clinical change, on the CORE-10.

Table 8: Comparison of rates of reliable and clinical change for CORE-10 and CORE-OM

	COR	E-10	CORE-OM		
	N	%	N	%	
Reliable and clinically significant improvement	439	54.8	452	56.1	
Reliable improvement only	170	21.2	154	19.1	
No reliable change	173	21.6	186	23.1	
Reliable deterioration	19	2.4	14	1.7	

# **Section V**

### The CORE family of outcome measures

The CORE-10 is one of a family of CORE outcome measures which have been developed to be used under different circumstances and for different populations. These measures are outlined below.

CORE Measures	Thumbnail summary
CORE-OM: 'Parent' outcome measure	The CORE-OM is a 34-item generic measure of psychological distress which is pan-theoretical (i.e., not associated with a school of therapy), pan-diagnosis (i.e., not focused on a single presenting problem), and draws upon the views of what practitioners considered to be the most important aspects of mental health to measure. The CORE-OM comprises 4 domains Well-being (4 items); Symptoms (12 items - depression x 4, anxiety x 4, trauma x 2 physical x 2); Functioning (12 items general x 4, social x 4, & close x 4); and Risk (6 items - to self x 4 or to others x 2). It takes between 5-10 minutes to complete.
Short Forms A&B:  Session-by-session repeated administration (research)	Two parallel 18 item psychometrically balanced measures for use at alternate therapy sessions and which together make up the CORE-OM. The use of two short forms at alternate sessions rather than the CORE-OM measure at every session reduces memory effects. Due to administrative complexities, repeated administration of the two short forms is usually used only in research studies.
CORE-10:  Review or quick initial assessment	A short 10 item version of the CORE-OM to be used as a screening tool and outcome measure when the CORE-OM is considered too long for routine use. Items cover anxiety (2 items), depression (2 items), trauma (1 item), physical problems (1 item) functioning (3 items – day to day, close relationships, social relationships) and risk to self (1 item). The measure has 6 high intensity/severity (e.g., and 4 low intensity/severity items.
CORE-5: Session-by-session monitoring	The CORE-5 comprises 5 items drawn from the CORE-OM and was designed to provide a brief tool for practitioners to monitor ongoing progress session by session. Items cover anxiety, depression, and functioning.

# The CORE family of population-specific outcome measures

CORE Measures	Thumbnail summary
GP-CORE:	A short 14 item measure derived from the CORE-OM suitable
General Population	for use with general populations, including students. In contrast to the CORE-OM, the GP-CORE does not comprise
For use with general or student populations	items denoting high-intensity presenting problems or risk and over half the items are positively keyed. These aspects increase its acceptability in a non-clinical population.
YP-CORE:	A 10-item measure derived from the CORE-OM and designed
Young Person's	for use in the 11-16 age range. Structure is similar to that of the CORE-OM but with items rephrased to be more easily understood by the target age group.
LD-CORE:	LD-CORE is being developed by therapists and adults with LD
Learning Disability	in Scotland and England. The measure includes simplified items from the CORE-OM selected by therapists and adults with LD, and also includes new items designed to cover the major issues they face that are not in the CORE-OM.
CORE Translations	Approved translations now exist for Gujarati, Norwegian, Italian, Slovak, Swedish, Icelandic, Albanian and Greek. Kannada, Tamil and Welsh versions are nearing completion. Referential and psychometric data for the Slovak, Italian, Norwegian and Greek translations should be available in 2008. People seeking other translations or wishing to help produce them should contact core-trans@psyctc.org.

### **The CORE System**

The CORE System was developed by a multidisciplinary group of practitioners and researchers and the content of the system was informed by extensive collaboration with practitioners, managers, and service commissioners.<sup>[15]</sup> The system comprises three tools, sharing the onus of evaluation data provision equally between clients completing a CORE outcome measure pre- and post-therapy and practitioners completing the CORE Assessment Form at pre-therapy and End of Therapy Form at post-therapy.

CORE System	Thumbnail summary
A CORE measure	See section above.
Therapy Assessment Form	The CORE Therapy Assessment Form captures a 'core' set of contextual information that aids the quality of both client assessment and overall service development. To enhance client assessment, the form collects important contextual information including client support, previous/concurrent attendance for psychological therapy, medication, as well as a categorisation system to record presenting difficulties, their impact on day-to-day functioning, and any associated risk. To aid the development of service quality, the form collects data on critical assessment audit items that profile the accessibility and appropriateness of service provision. These include client demographics, waiting times, and the suitability of referral.
End of Therapy Form	The CORE End of Therapy Form complements the other components and captures a 'core' set of treatment descriptors that aid the interpretation of CORE-OM scores to help contextualize therapy outcomes and inform service development. The form collects profile information that includes therapy length, type of intervention, modality, and frequency. To enhance the development of service quality, the form collects data on critical discharge audit items that profile the effectiveness and efficiency of service provision. These include problem and risk review, therapy benefits, session attendance rates, and therapy ending (i.e., planned or unplanned).

#### **Currently available software**

The CORE-Net software currently available from CORE IMS:

- Allows data capture, storage, analysis and reporting facilities of data for CORE measures
- Allows data capture of most of the routine data specified in the MDS (nature of intervention/treatment offered; number of sessions; type of therapist; therapist identifier; gender; age; ethnicity)
- Is being used by a rapidly growing number of staff (currently estimated at about 100) spread across a number of services, including services operating in the stepped care environment. About 600 new patients are currently being added per month in CORE-Net
- Allows data to be entered either by therapists or administrative staff spread across multiple locations (an essential requirement if data entry is carried out in surgeries)
- Allows any PC with internet access to be used without the installation of any local software since CORE-Net requires only a standard internet browser
- CORE-Net server software can either be mounted on a local server or, to minimise
  potential delays arising from local IT installation and support tasks can be used on a
  currently available secure hosted server
- Provides a high level of security in order to protect data confidentiality

#### **Development**

In order to provide a more complete data capture and reporting system for IAPT pathfinder sites in CORE-Net:

- The data capture of sexuality and disability is under development
- Data capture for the PHQ-9 and IAPT Inclusion & Employment Questionnaire is in development
- Development of data capture for the other primary measures included in the IAPT
   Outcome Framework and Data Collection Minimum Data Set (GAD-7; PEQ1; PEQ2;
   Work & Social Adjustment Scale) is subject to agreement with pathfinder sites,
   agreement of copyright holders of the measures and availability of development
   funding
- Development of data capture for disorder specific measures (e.g., SPIN; OCI; IES;
   HAI; Mobility Inventory; HADS; BDI; Fear Questionnaire; Penn State Worry) and other measures (e.g., CSQ8; SF6-D; EQ5-D) is subject to specific requests from

pathfinder sites, agreement of copyright holders of the measures and availability of funding for software development.

- Reporting the aggregate data at monthly intervals
- CORE-Net provides a facility to download all data to an Excel spreadsheet for analysis and reporting purposes. This facility will be extended to allow downloading of data captured from the additional measures (PHQ-9, etc) to assist pathfinder sites in meeting the aggregate reporting requirements specified in Appendix 1, IAPT Reporting Template 1, Service Activity Data and Appendix 3, IAPT Reporting Template 3, Data Collection Framework & Key Findings.

Availability of the software supporting additional measures is in all cases subject to the agreement of the copyright owner for the specific measure. CORE-Net can provide interoperability with a local EPR system but it is **not** intended to provide EPR functionality within CORE-Net. There is a license fee payable for use of the CORE-Net software (but there is no charge for use of CORE measures). CORE-IMS can be contacted for more information at <a href="mailto:admin@coreims.co.uk">admin@coreims.co.uk</a>

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# Appendix I: CORE-10

CLINICAL OUTCOMES in ROUTINE EVALUATION CORE-10 v.1	Site ID  Client ID  letters only numbers only  Sub codes  Therapist ID numbers only (1)  Date form given D D M M Y Y	Stage Completed S Screeening R Referral A Assessment F First Therapy Session P Pre-therapy (unspecified) D During Therapy (review) L Last therapy session X Follow up 1 Y Follow up 2  Episode Stage  numbers only (2)  Gender Y Male Age Female	
	ANT - PLEASE READ T		
	ents about how you have beer ement and think how often yo		
Thei	n tick the box which is closest	to this.	
Please use a dair	k pen (not pencil) and tick clea	•	
Over the last week		World of the control	
1 I have felt tense, anxious or ne	rvous	0 1 2 3 4	
2 I have felt I have someone to to	ırn to for support when neede	d	
3 I have felt able to cope when th	nings go wrong	4 3 2 1 0	
4 Talking to people has felt too m	nuch for me	0 1 2 3 4	
5 I have felt panic or terror		0 1 2 3 4	
6 I made plans to end my life		0 1 2 3 4	
7 I have had difficulty getting to s	leep or staying asleep	0 1 2 3 4	
8 I have felt despairing or hopele	SS	0 1 2 3 4	
9 I have felt unhappy		0 1 2 3 4	
10 Unwanted images or memorie	es have been distressing me	0 1 2 3 4	
Total (Clinical Score*)			
then multiply by 10 to get the Clinical	Score.	uestions completed to get the mean score, the item scores to get the Clinical Score.	
Thank you for yo	ur time in completing	g this questionnaire	
, ,	Conveight CODE System Trust		



# Appendix II: CORE-OM

CLINICAL OUTCOMES ROUTINE EVALUATION OUTCOM MEASU	Ietters Client ME Therap JRE Sub c	only numbers only  t ID  ist ID numbers only (1) number	Agors only (2)	Stage C S Scree R Refer A Asset F First P Pre-ti D Durin	rral ssment Therapy (u herapy (u ig Therapy therapy s w up 1	Session nspecified) y	Stage  Episode
		IMPORTANT - PLEASE RE	AD THIS FI	RST			
	Please read e	statements about how you ha each statement and think how on Then tick the box which is see a dark pen (not pencil) and	often you fel closest to th	t that w nis.	ay last v	veek.	
	ne last week		Not at all	Occasions		Most or	
1 I have fe	It terribly alone an	d isolated	<b>U</b> °	<b>1</b>	<u> </u>   2	3 4	. ∐ F
2 I have fe	lt tense, anxious o	or nervous	<b>□</b> ∘	<u> </u>	2	3 4	. <u> </u>
3 I have fe	It I have someone	to turn to for support when needed	j 4	_ з	_ 2 _	1 0	F
4 I have fe	lt O.K. about myse	elf	<u> </u>	В	2	1 🗆 0	w
5 I have fe	lt totally lacking in	energy and enthusiasm	<u> </u>	□ 1	_ 2 _	3 🔲 4	. P
6 I have b	een physically viol	ent to others	□ ∘	<u> </u>	_ 2 [	3 🔲 4	- R
7 I have fe	It able to cope wh	en things go wrong	<b>4</b>	_ 3	_ 2 _	1 🔲 0	F
8 I have b	een troubled by ac	thes, pains or other physical proble	ms 🔲 o	<u> </u>	2	3 🔲 4	Р
9 I have th	ought of hurting n	nyself	<u> </u>	<b>1</b>	_ 2 _	3 4	R
10 Talking t	o people has felt t	oo much for me	□ ∘	<u></u> 1	2	3 4	- F
11 Tension and anxiety have prevented me doing important things		ıs 🗌 o	<u> </u>	_ 2 _	3 4	. P	
12 I have b	een happy with the	e things I have done	<u> </u>	Вз	2 [	1 🔲 0	F
13 I have b	een disturbed by u	inwanted thoughts and feelings	<u> </u>	□ 1	_ 2 _	3 🔲 4	. 🗌 Р
14 I have fe	lt like crying		□ ∘	□ 1	2	3 4	. W
		Please turn ov	/er				

# CORE-OM (cont'd)

	Over the last week	Vot at all   Occasionally   Sometimes   Offen   All the time   OFFICE USE   ONLY   SE	
15	I have felt panic or terror	0 1 2 3 4 P	
16	I made plans to end my life	0 1 2 3 4 R	
17	I have felt overwhelmed by my problems	0 1 2 3 4 W	
18	I have had difficulty getting to sleep or staying asleep	0 1 2 3 4 P	
19	I have felt warmth or affection for someone	43210F	
20	My problems have been impossible to put to one side	0 1 2 3 4 P	
21	I have been able to do most things I needed to	43210F	
22	I have threatened or intimidated another person	0 1 2 3 4 R	
23	I have felt despairing or hopeless	0 1 2 3 4 P	
24	I have thought it would be better if I were dead	0 1 2 3 4 R	
25	I have felt criticised by other people	0 1 2 3 4 F	
26	I have thought I have no friends	0 1 2 3 4 F	
27	I have felt unhappy 0 1 2 3 4 P		
28	Unwanted images or memories have been distressing me 0 1 2 3 4 P		
29	I have been irritable when with other people 0 1 2 3 4 F		
30	I have thought I am to blame for my problems and difficulties 0 1 1 2 3 4 P		
31	I have felt optimistic about my future	43210W	
32	I have achieved the things I wanted to	43210F	
33	I have felt humiliated or shamed by other people	0 1 2 3 4 F	
34			
	my health		
	THANK YOU FOR YOUR TIME IN COMPL	ETING THE QUESTIONNAIDS	
	THANK YOU FOR YOUR TIME IN COMPL	ETING THIS QUESTIONNAIRE	
Tot	al Scores		
Ме	an Scores		
dim	(Total score for each dimension divided by number of (W) (P) (F) (R) All items All minus R items completed in that dimension)		

# **Appendix III: CORE Practitioner-completed measures**

# **Therapy Assessment Form**

CLINICAL OUTCOMES IN ROUTINE EVALUATION THERAPY ASSESSMEN FORM v.2	Site ID
Referral date First assessment date attended Last assessment date	DD MM YYYY Previously seen for therapy Yes in this service? No DD MM YYYY Months since last episode
Living alon Living with Caring for Caring for Living wiht	e (not including dependents)
Please tick as a Primary GP Secondary In a In a In a Hos Specialist Psy Atta	bus use of services for psychological problems?  many boxes as appropriate or other member of primary care team (eg practice nurse, counsellor)

# Therapy Assessment Form (cont'd)

Is the client currently prescribed medication to help with their psychological problem(s)? Yes No
If yes, please indicate type of medication:  Anti-psychotics
Brief description of reason for referral
Depression
Risk
What has the client done to cope with/aviod their problems? Please tick, and then specify actions Positive actions  Negative actions
Assessment outcome (tick one box only)  Assessment/one session only  Accepted for therapy  Accepted for trial period of therapy

# **End of Therapy Form**

CLINICAL				
OUTCOMES in	Site ID			Number of
ROUTINE	OI: . ID	letters numbers		sessions planned
EVALUATION	Client ID			
END OF	Sub Codes	Therapist ID SC4 nu	mbers SC5 numbers	Number of
THERAPY	Sub Codes			sessions attended
FORM v.2	Date therapy commenced	D D M M Y Y	/ Y Y 	Number of
	Date therapy	DD MM Y	/ Y Y	Number of sessions
	completed			unattended
What type of thera	py was undertak	en with the client? F	Please tick as manv bo	oxes as appropriate
Psychodynar			Person-centred	,, -,
Psychoanaly		H	Integrative	H I
Cognitive		Ħ	Systemic	H H
Behavioural		Ħ	Supportive	H H
Cognitive/Behavioural Art			T I	
Structured/Brief Other (specify below)			ow) 🗌 📗	
				l
What madality of th	arany was unds	rtakan with the aliant?	Places tiek as many	havaa aa annranriata
Individual	ierapy was unde	rtaken with the client?		boxes as appropriate
111111111111			Family Marital/Couple	H
Group			wantai/Couple	
What was the frequ	ency of therapy	with the client?		_
What was the frequ		with the client?	Less than once we	· =
		with the client?	Less than once we	· =
More than or		with the client?		· =
More than or Weekly	nce weekly	with the client?	Not at a fixed frequ	· =
More than or Weekly	nce weekly	es the ending of thera	Not at a fixed frequ	· =
More than or Weekly  Which of the follow	nce weekly	es the ending of thera	Not at a fixed frequency?	· =
More than or Weekly  Which of the follow Unplanned	ing best describ	es the ending of therap	Not at a fixed frequency?	· =
Which of the follow Unplanned Due to crisis Due to loss of	ing best describ	es the ending of therapy PI	Not at a fixed frequency?  anned anned from outset	lency
Which of the follow Unplanned Due to crisis Due to loss of	ing best describ	es the ending of theral PI PI Age Age	Not at a fixed frequency?  anned anned from outset greed during therapy	ency
More than or Weekly  Which of the follow Unplanned Due to crisis Due to loss of Client did not	ing best describ	es the ending of theral PI PI Age Age	Not at a fixed frequency?  anned anned from outset greed during therapy greed at end of therap	ency

# **End of Therapy Form (cont'd)**

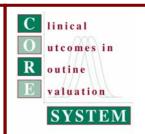
Review of Identified Problems/Concerns
Depression   Trauma/Abuse   Anxiety/Stress   Bereavement/Loss   Self esteem   Interpersonal/relationship   Living/Welfare   Physical Problems   Work/Academic   Other (specify below)   Addictions
Contextual Factors   Contextual Factors   Suicide   Contextual Factors   Contextual Factor   Contex
Improved Yes No Yes N
Has contact with this service resulted in a change of medication? Yes No Not applicable If yes, is this change likely to be of benefit to the client? Yes No Details of change: Started Discontinued Increased Decreased Modified Has the client been given a follow-up appointment? Number of months until appointment

#### **Appendix IV: CORE Publications**

# The CORE Measures & CORE System

#### **Publications**

Up to June 2007



This is a chronological listing of publications which focuses on the following:

- A: CORE System and its constituent measures (i.e., CORE-OM, CORE-SF, GP-CORE)
- B: Use of the CORE System or its components in trials and routine settings
- C: Outcome measurement, implementation, & infrastructure (e.g., other outcome measures, core battery, practice research networks, outcomes policy)
- D: Broader issues of practice-based evidence and evidence-based practice
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