

Institute of Psychiatry



at The Maudsley

# Cognitive behavioural interventions to enhance SDM

Rona Moss-Morris Professor of Psychology as Applied to Medicine

# Outline

#### What is CBT

#### **MS** Fatigue

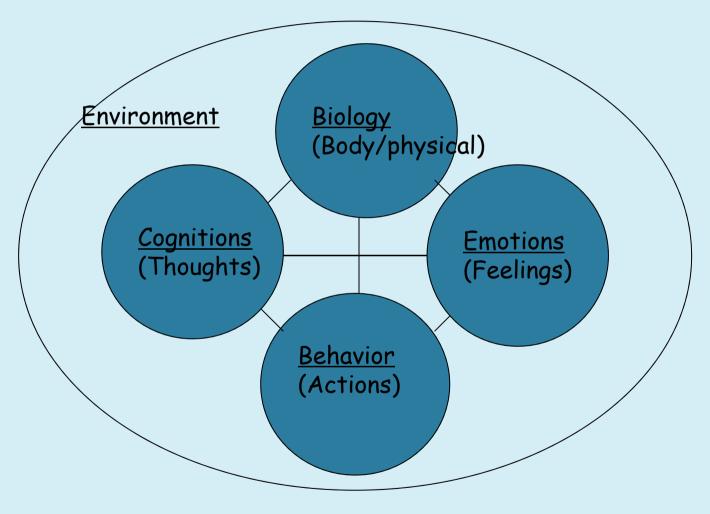
- CB Model of MS fatigue
- Therapist driven CBT for MS fatigue
- CCBT for MS fatigue

#### **Adjustment to MS**

- CB model of adjustment to MS
- Nurse-led CBT for MS

2

### **The five part CBT Model**

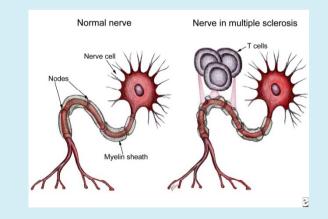


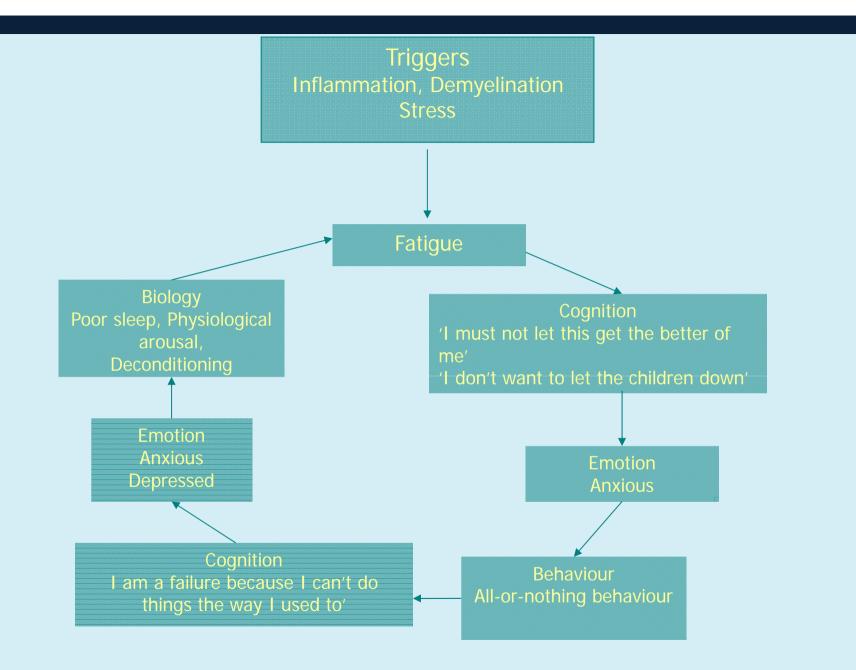
# **Core Therapeutic processes**

- Formulation
- Collaborative approach
- Guided discovery
- Behavioural change e.g. reduce
  avoidance
- Identifying unhelpful thoughts
- Alternate thoughts

# **Fatigue in MS**

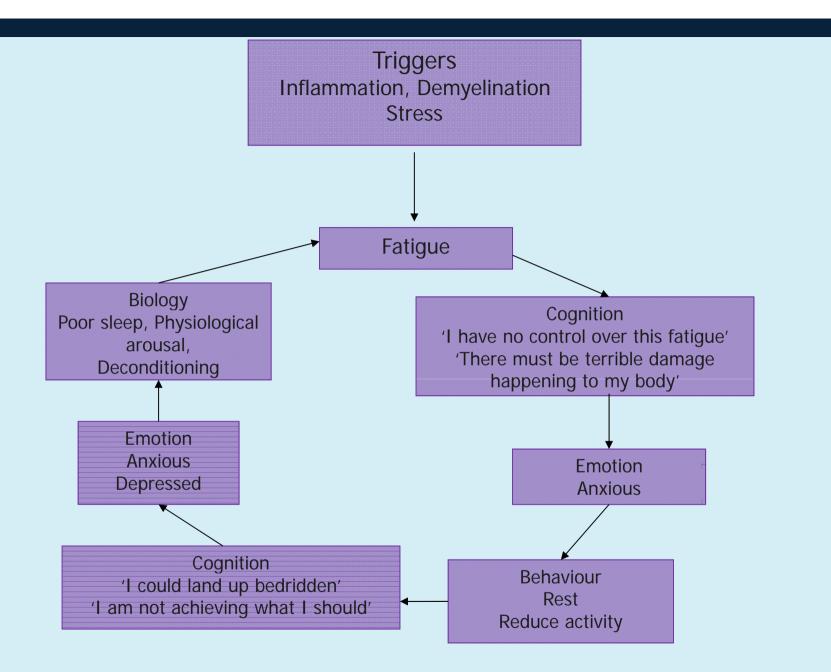
- Fatigue is among the most common, and least understood, symptoms of MS.
- 76% to 97% of MS patients report fatigue.
- Fatigue has a substantial impact on daily activities.





Van Kessel, & Moss-Morris. (2006) Understanding multiple sclerosis fatigue: A synthesis of biological and psychological factors, JPR

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Van Kessel, & Moss-Morris. (2006) Understanding multiple sclerosis fatigue: A synthesis of biological and psychological factors, JPR

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# **Can CBT reduce fatigue in MS?**

# Cognitive behaviour therapy compared to relaxation training for multiple sclerosis fatigue: A randomized controlled trial.



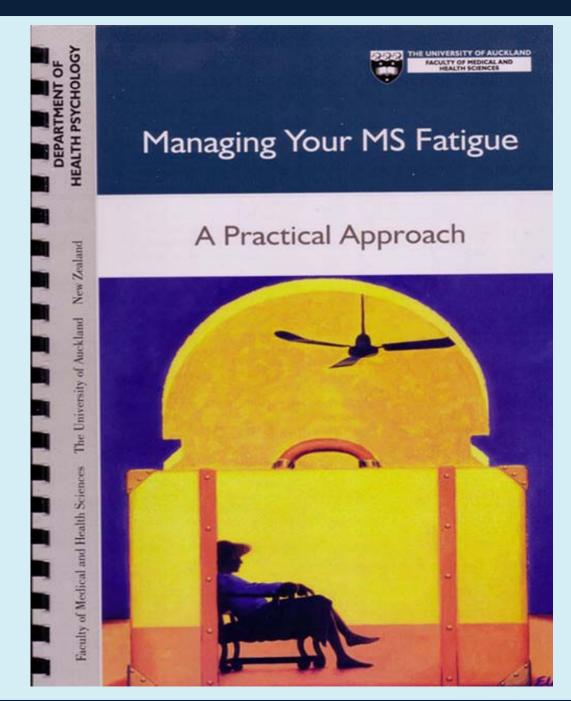
van Kessel, K., Moss-Morris, R, Willoughby, E, Chalder, T. Johnson, M.H., Robinson, E. (2008). *Psychosomatic Medicine*, 70, 205-213.



8 weekly manualised sessions

Combination of face-to-face and telephone sessions.

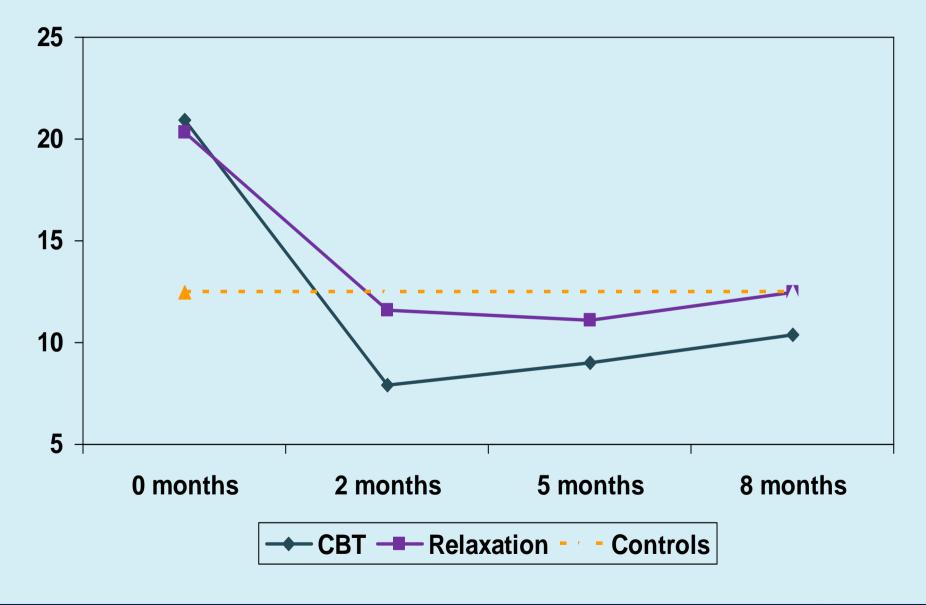
n=72



# The programme

- Develop personal model of fatigue
- Changing activity patterns
- Challenging unhelpful thoughts
- **Stress and sleep management**
- Managing difficult emotions

#### Fatigue across groups



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Psychological Medicine (2012), 42, 205–213. © Cambridge University Press 2011 doi:10.1017/S0033291711000924

# Which cognitions and behaviours mediate the positive effect of cognitive behavioural therapy on fatigue in patients with multiple sclerosis?

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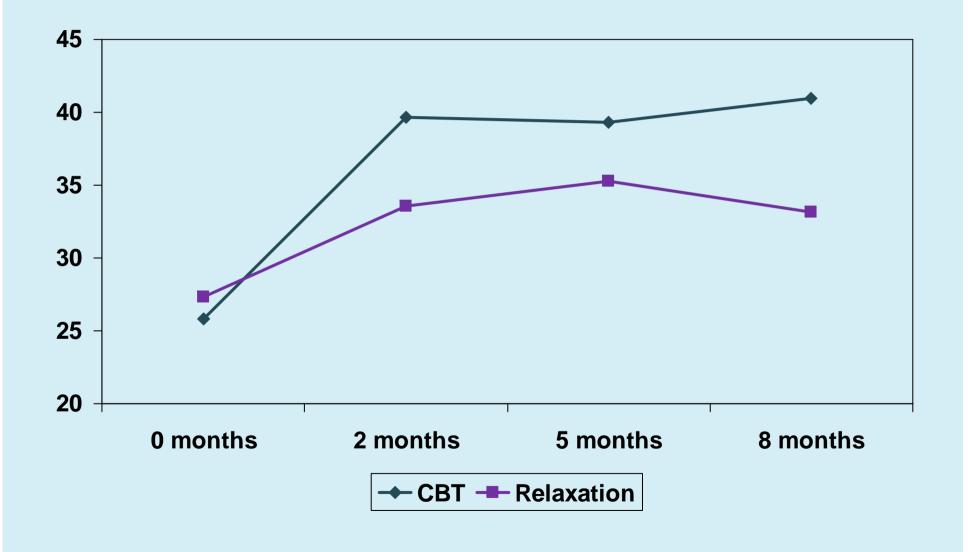
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<sup>8</sup> School of Psychology, University of Southampton, Highfield, Southampton, UK

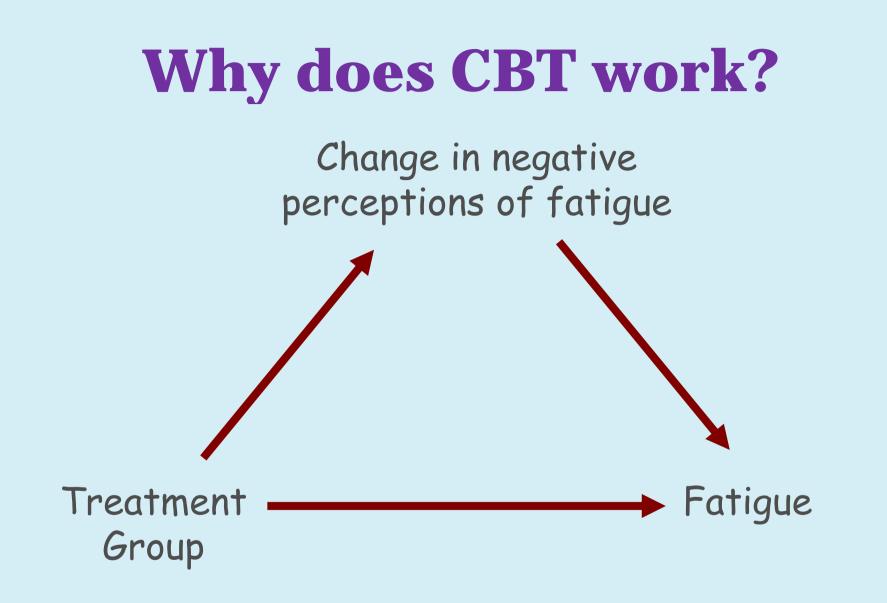
**Background.** Chronic fatigue is a common symptom of multiple sclerosis (MS). A randomized controlled trial (RCT) showed that cognitive behavioural therapy (CBT) was more effective in reducing MS fatigue than relaxation training (RT). The aim of the current study was to analyse additional data from this trial to determine whether (1) CBT compared to RT leads to significantly greater changes in cognitions and behaviours hypothesized to perpetuate MS fatigue; (2) changes in these variables mediate the effect of CBT on MS fatigue; and (3) these mediation effects are independent of changes in mood.

Method. Seventy patients (CBT, n=35; RT, n=35) completed the Cognitive and Behavioural Responses to Symptoms Questionnaire (CBSQ), the Brief Illness Perception Questionnaire (B-IPQ) modified to measure negative representations of fatigue, the Hospital Anxiety and Depression Scale (HADS), and the Chalder Fatigue Questionnaire (CFQ), pre- and post-therapy. Multiple mediation analysis was used to determine which variables mediated the change in fatigue.

#### **Total Positive Perceptions of fatigue**



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Shorter communication

#### A pilot randomised controlled trial of an Internet-based cognitive behavioural therapy self-management programme (MS Invigor8) for multiple sclerosis fatigue

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#### ABSTRACT

The majority of people affected by Multiple Sclerosis (paMS) experience severe and disabling fatigue. A recent randomised controlled trial (RCT) showed that cognitive behaviour therapy with a clinical psychologist was an effective treatment for MS fatigue. An Internet-based version of this intervention, MS Invigor8, was developed for the current study using agile design and input from paMS. MS Invigor8 includes eight tailored, interactive sessions. The aim was to test the feasibility and potential efficacy and cost-effectiveness of the programme in a pilot RCT. 40 patients were randomised to MS Invigor8 (n = 23) or standard care (n = 17). The MS Invigor8 group accessed sessions over 8–10 weeks and received up to three 30–60 min telephone support sessions. Participants completed online standardised questionnaires assessing fatigue, mood, quality of life and service use at baseline and 10 weeks follow-up. Large between group treatment effects were found for the primary outcomes of fatigue severity (d = 1.19) and impact (d = 1.02). The MS Invigor8 group also reported significantly greater improvements in anxiety, depression and quality-adjusted life years. These data suggest that Internet-based CBT may be a clinically and cost-effective treatment for MS fatigue. A larger RCT with longer term follow-up is warranted.

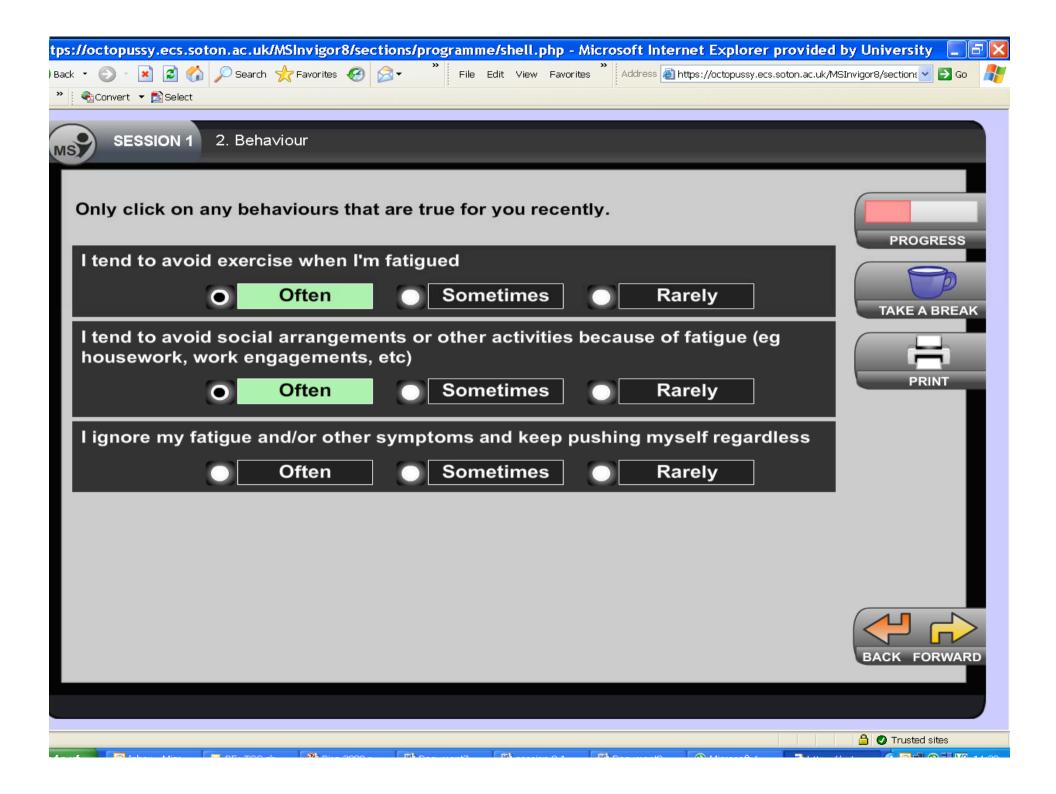
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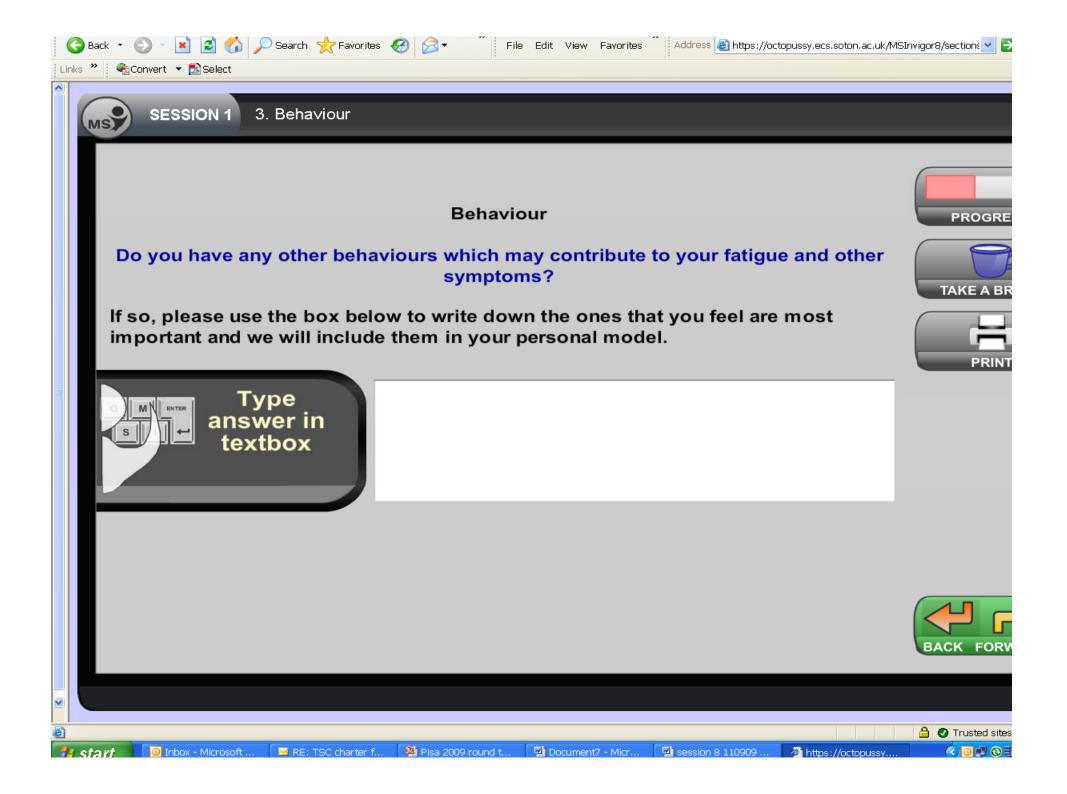
## MSInvigor8 Design team

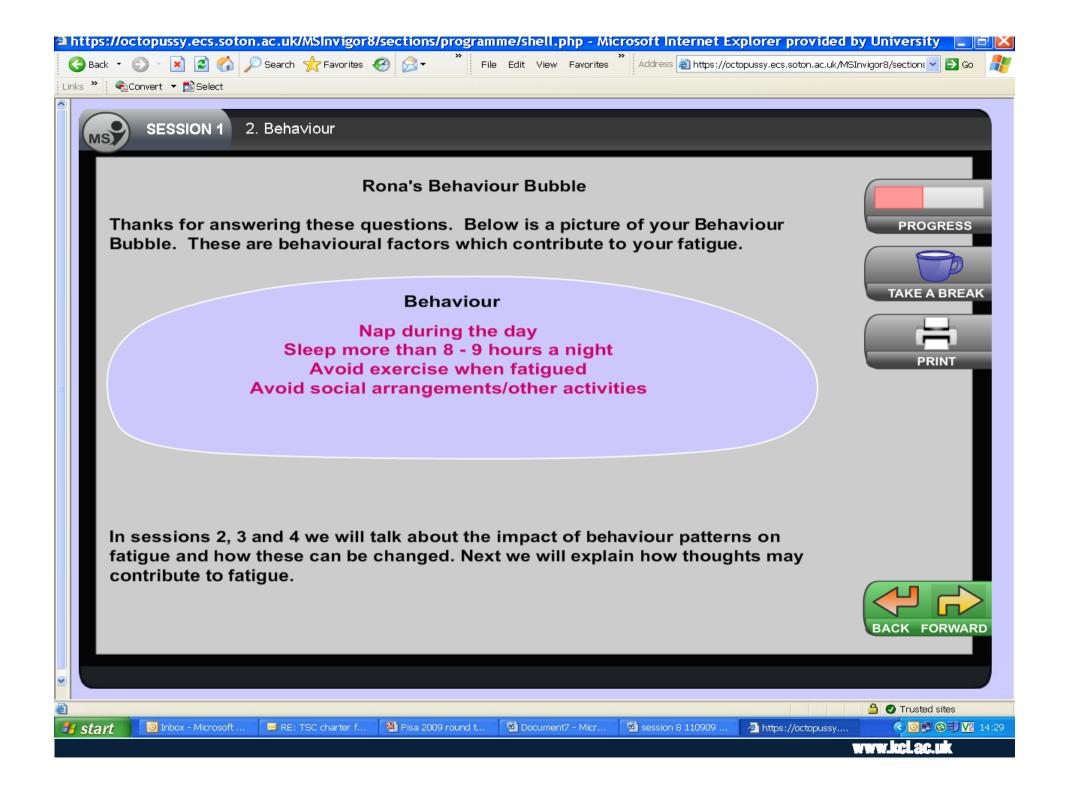


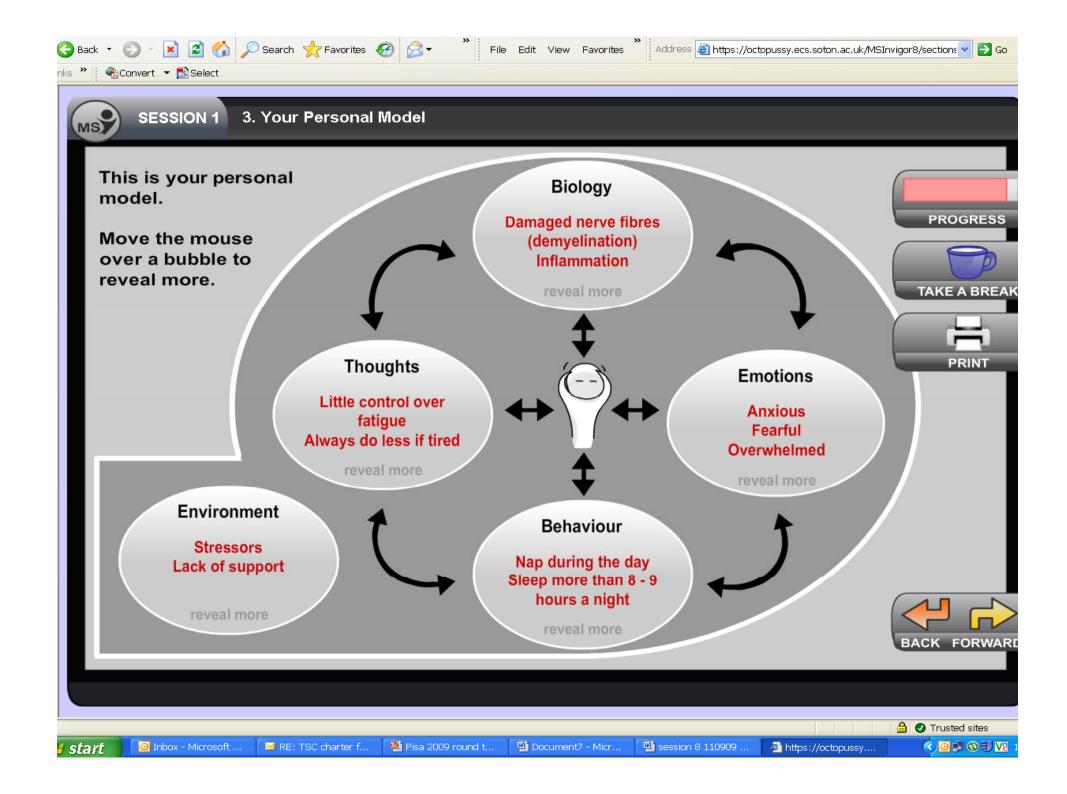


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	SESSION 1 2. Behaviour									
		Only click on any b	pehaviours that	t are true fo	r you recer	ntly.				
									PROGRESS	
		I nap during the d	ay to manage	my fatigue						
			Often	Som	etimes	R	arely			
			Onteri		letimes		arery		TAKE A BREAK	
		I tend to overdo t	hings when I h	ave energy						
		$\Box$	Often	Som	etimes	R	arely			
								_	PRINT	
≡		I regularly sleep n	nore than 8 - 9	hours a nigl	nt					
		•	Often	Som	etimes	R	arely			
								_		
		I find myself rushi	ng to get thing	gs done befo	re I crash					
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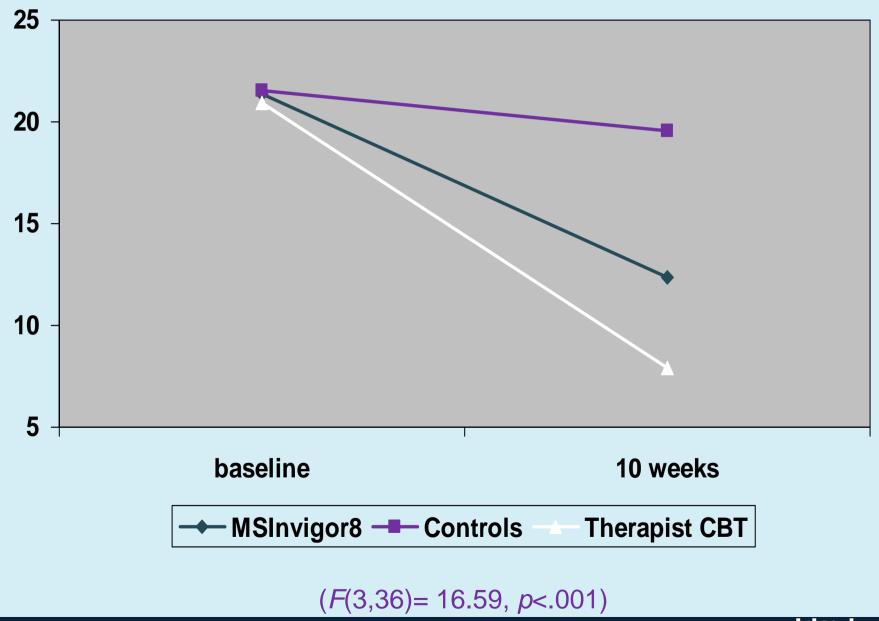




MSI Pilot trial				
	MSInvigor8 (n=23)	Control group (n=17)		
Age: M (SD)	40 (17.8)	42 (11.43)		
Gender: % female	73%	94%		
Time since diagnosis	21 (9.05)	16 (7.9)		

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#### Fatigue Severity Across Groups



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### Within group Effect Sizes for CBT

- MSInvigor8 study = 1.49 and 1.40
- Therapist driven CBT = 3.03



# Summary of economic data

- MSInvigor8
  - produced more QALYs
  - did not have an impact on non-intervention service costs.
- To achieve NICE recommended cost per QALY of £20,000
  - intervention costs need to be no more than £300 per person
- If 300 people use the intervention then this would cover a £90,000 development cost which is above the actual costs that were accrued.



Supportive adjustment for Multiple Sclerosis

#### **ACADEMIC TEAM**

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University Hospitals NHS Trust



Institute of Psychiatry at the Maudsley

#### **CLINICAL TEAM**

**Sarah Morton Sally Baines Suzanne Roche** 



iniversity of London



# **Modelling phase**

Systematic review of the empirical literature on adjustment to MS (n=72)

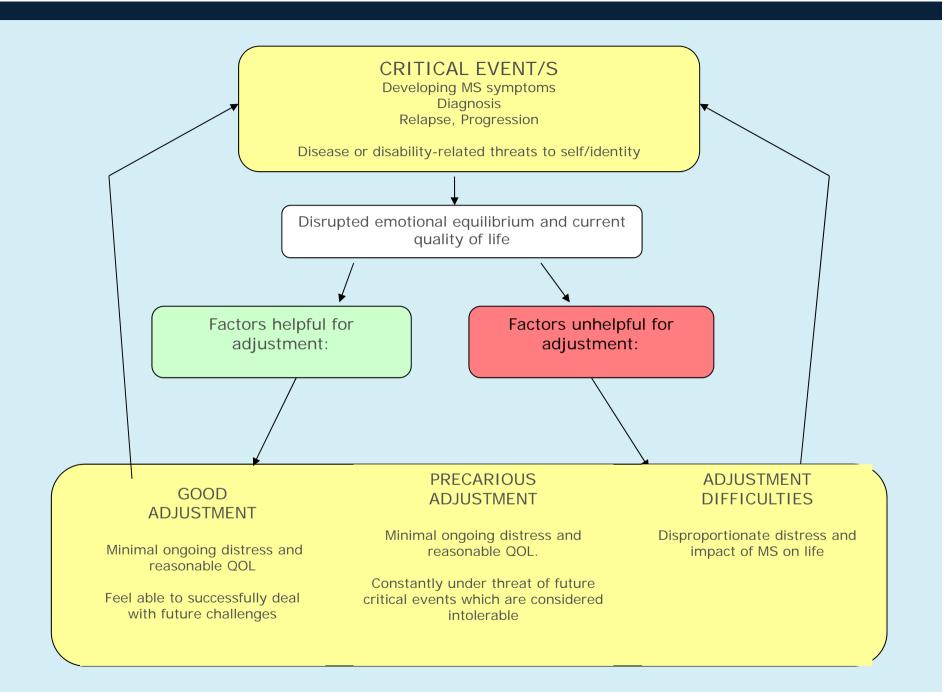
Dennison et al., (2009). Clinical Psychology Review. 29, (2), 141-153.



In-depth interviews with pwMS (n=30) and partners of pwMS (n=15)

Bogosian et al., (2009). Multiple Sclerosis. 15 (7), 876-884.

Dennison et al. (2010) Journal of Health Psychology, 16(3) 478–488.



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An eight-week CBT programme manual

Supportive Adjustment for Multiple Sclerosis (saMS)

> Written by Professor Rona Moss-Morris, Laura Dennison and Professor Trudie Chalder

This information is the result of MS Society-funded research at

H KING'S LONDON

Southampton and



Supportive Adjustment for Multiple Sclerosis (saMS)

page

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First Edition December 2010 Institute of Psychiatry



at The Maudsley

#### Coping when somebody close to you has MS

A booklet for partners, relatives or friends

Written by Professor Rona Moss-Morris, Laura Dennison and Professor Trudie Chalder This information is the result of MS Society-funded research at:



and Southampton



### Dealing with feeling helpless

A diagnosis of MS, and the realisation of what this means, often generates feelings of helplessness in partners and those close to people with MS. Jenny provides a good example of these feelings:

> "I feel a bit lost at times, because you can't help someone suffering with this disease, it's impossible, it's out of your hands and you feel sort of... like... I don't know, helpless, frustrated, and very angry."

Men in particular seemed to struggle with feeling out of control, as Gary describes:

"As the actual man, I am supposed to protect, to lay down my life, and do whatever it takes to protect my partner, my write. But in this situation there is nothing you can do. It's very difficult to describe, you feel like you're just powerless to actually do anything. It is like the control has been taken from you. So it's emasculating really, it's difficult to describe more than that..."

It's not surprising that MS provokes such strong feelings of helplessness – it is unpredictable and ourrently incurable and it is distressing to see somebody you care about having to deal with this. There is no magio wand available for you to put things back to how they were before. But there are ways to manage symptoms, limitations and emotional reaponses related to MS. Your partner or friend will look at these in his/ her sessions with the nurse. Also, from what the people with MS we interviewed told us, there is a whole range of important ways in which support people help, including:

- just being there!
- listening and talking
- finding out information
- taking an interest and taking part in managing MS (for example, going to doctors' appointments, MS society meetings)
- making adaptations to lifestyle, activities, roles, expectations
- providing different types of support sometimes emotional rather than practical

### How to best support somebody with MS

Most people don't want to see their partner or friend upset and struggling and want to help and support them in some way. However, it is difficult to know what to do in a situation where you can't get rid of the main problem. We found that often there is a mismatch between the amount and type of support that is given and what the person with MS wants and needs. Mike and Sandra are a typical example:

Mike (husband) reported how, when Sandra was diagnosed with MS, he didn't know what to do to help the situation. He was shocked and upset and also wanted to ease her distress. He thought that maybe in the future her mobility would get worse and she might need to use a wheelchair. So he focussed all his energy into making changes to the house and garden and building wheelchair ramps.

Sandra (wife with MS) talked about how, when she was first diagnosed, what she really wanted was for Mike to give her a hug! At that time she wasn't ready to make plans for how she would cope with potential disability in the future. She just wanted emotional support and someone to talk to.

# **CBT versus Supportive Listening**

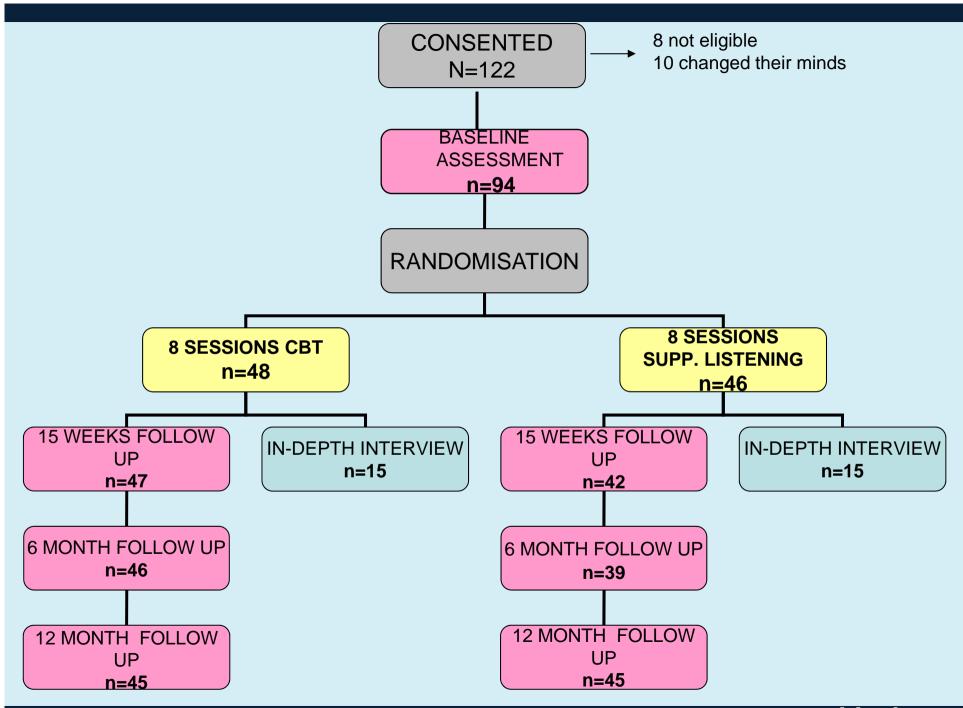
- Moss-Morris et al. (2009). Protocol for the saMS trial. BMC Neurology, 2009, 9:45
- Delivered by 2 general nurses
- Trained specifically for this project

-MS

-CBT

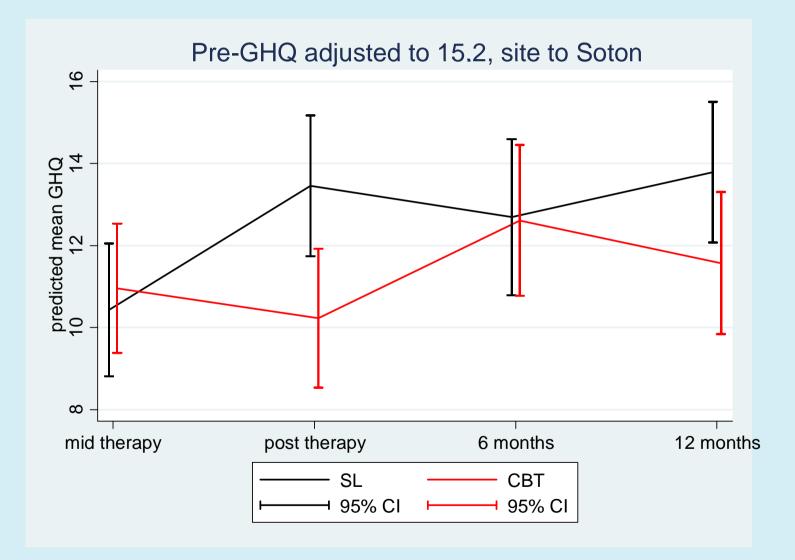
-Supportive Listening





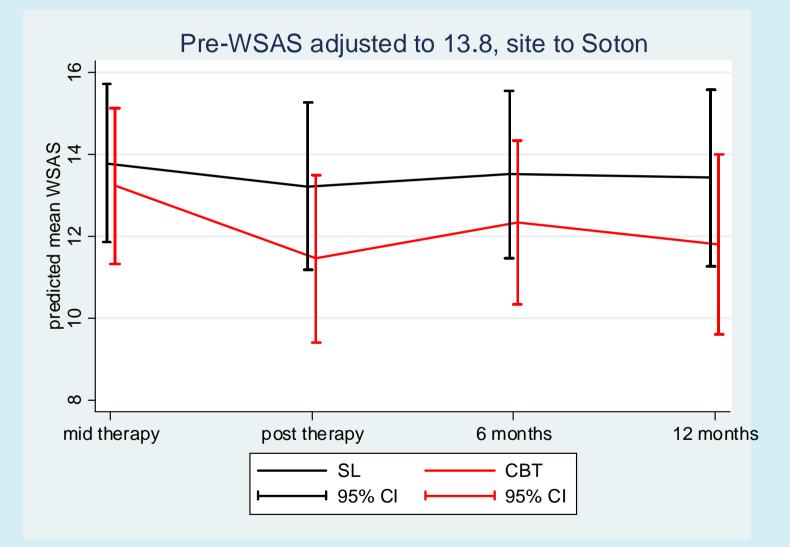
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## **Distress (GHQ)**



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# Functional Impairment (WSAS)



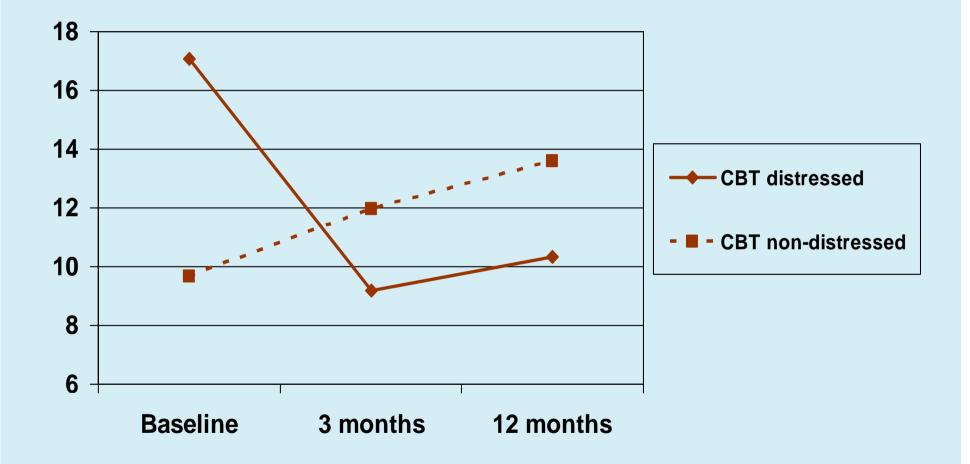
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Global improvement					
	CBT	SL			
Improved	67.4%	31%			
Not improved or worse	32.6%	69%			
Therapy Satisfaction					
Satisfied	84.8%	57.1%			
Neutral or dissatisfied	15.2%	42.9%			

## **Moderation analyses**

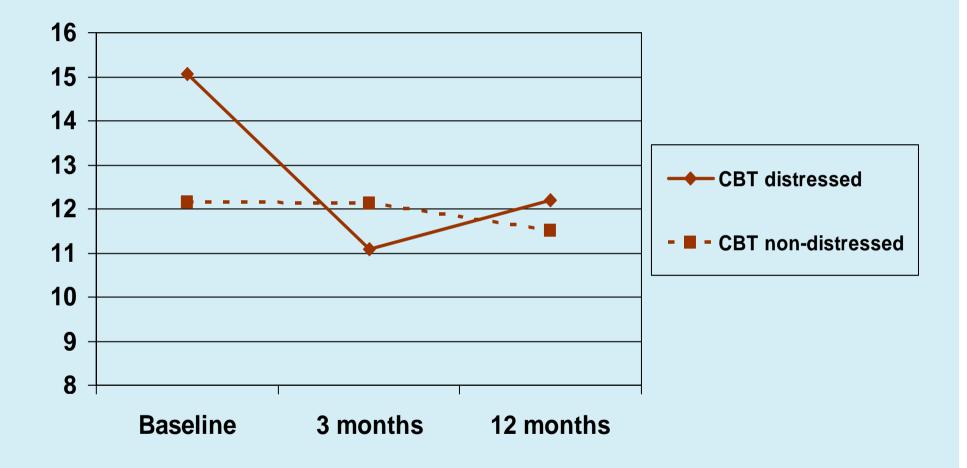
- GHQ caseness (score of 3 or greater)
  - 60.6% of sample clinically distressed
- Social Support Satisfaction Significant Others Scale Power & Champion (1988)
  - 72% dissatisfied

#### Change in GHQ moderated by Clinical Distress



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#### **Change in WSAS moderated by Clinical Distress**



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# **Summary of saMS**

- CBT for adjustment to MS is more effective at reducing distress than SL up to one year post initiation of treatment.
- There were no significant differences between SL and CBT on WSAS.
- Patients who are 'clinically' distressed and/ or dissatisfied with their social support benefit more from CBT both in terms of reduced distress and reduced impact of MS on life roles.

# **CBT for MS**

- Appears to be an effective therapy for reducing MS related distress and managing symptoms like fatigue
- Challenge for services
  - How do provide these treatments (cost issues)
  - Training staff in these approaches ongoing supervision
  - CCBT