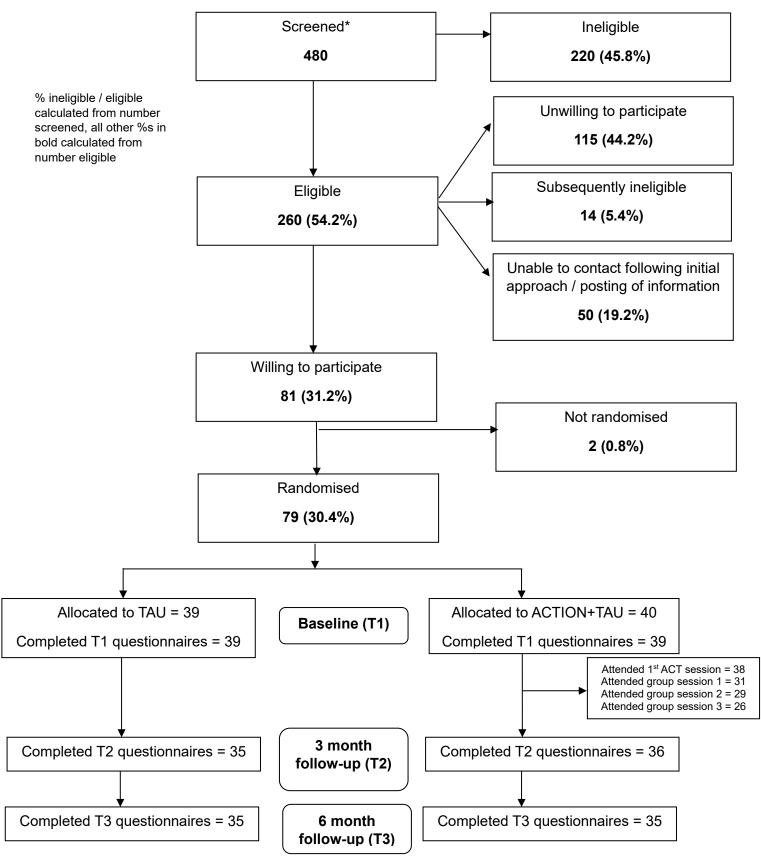
## **Participant flow**



\* Sites conducted screening of hospital records to assess for patient eligibility. Three sites provided data on the number of records screened but one did not. The total screened is therefore the number screened for the three sites who returned this data plus the number of patients eligible for approach from the fourth site.

## **Baseline characteristics**

	TAU	ACTION + TAU (n=40)		
	(n=39)			
Age				
Mean (SD)	60.2 (10.17)	58.5 (10.62)		
Median (Range)	62.0 (43.0, 84.0)	59.0 (39.0, 84.0)		
Missing	0	0		
Which best describes your ethnic background?				
White British	38 (97.4%)	38 (95.0%)		
Any other White background	1 (2.6%)	1 (2.5%)		
Vixed White and Black Caribbean	0 (0.0%)	1 (2.5%)		
Any other mixed background	0 (0.0%)	0 (0.0%)		
Asian/Asian British Indian	0 (0.0%)	0 (0.0%)		
Black/Black British African	0 (0.0%)	0 (0.0%)		
Any other Black background	0 (0.0%)	0 (0.0%)		
Any other ethnic group	0 (0.0%)	0 (0.0%)		
Not stated	0 (0.0%)	0 (0.0%)		
What is your marital status?				
Married	24 (61.5%)	27 (67.5%)		
Single	3 (7.7%)	1 (2.5%)		
Living with partner	5 (12.8%)	1 (2.5%)		
Divorced or separated	3 (7.7%)	6 (15.0%)		
Widowed	3 (7.7%)	3 (7.5%)		
Missing	1 (2.6%)	2 (5.0%)		
What is your employment status?				
Full time	11 (28.2%)	10 (25.0%)		
Part time	8 (20.5%)	9 (22.5%)		
Not currently working	5 (12.8%)	4 (10.0%)		
Retired	14 (35.9%)	11 (27.5%)		
Other	1 (2.6%)	4 (10.0%)		
Missing	0 (0.0%)	2 (5.0%)		

	TAU			
	(n=39)	ACTION + TAU (n=40)		
What is the highest level of education or professional qualification you have obtained?				
Postgraduate qualification	6 (15.4%)	3 (7.5%)		
Degree level education	12 (30.8%)	9 (22.5%)		
Higher educational qualifications (below degree level)	5 (12.8%)	9 (22.5%)		
Vocational Qualifications (NVQ1+2)	5 (12.8%)	4 (10.0%)		
A-Level or equivalent	3 (7.7%)	5 (12.5%)		
GCSE/O-Level/CSE	7 (17.9%)	7 (17.5%)		
No formal qualifications	1 (2.6%)	0 (0.0%)		
Missing	0 (0.0%)	3 (7.5%)		
Do you have children?				
Yes	30 (76.9%)	37 (92.5%)		
No	9 (23.1%)	2 (5.0%)		
Missing	0 (0.0%)	1 (2.5%)		
How many children?				
1	9 (30.0%)	7 (18.9%)		
2	15 (50.0%)	18 (48.6%)		
3	4 (13.3%)	11 (29.7%)		
4	2 (6.7%)	0 (0.0%)		
6	0 (0.0%)	1 (2.7%)		
How many children?				
Mean (SD)	2.0 (0.85)	2.2 (0.95)		
Median (Range)	2.0 (1.0, 4.0)	2.0 (1.0, 6.0)		
Missing	0	0		
Recruitment Route				
Recently completed treatment	17 (43.6%)	15 (37.5%)		
Experiencing medication problems	1 (2.6%)	0 (0.0%)		
Retrospective screening of hospital records	21 (53.8%)	25 (62.5%)		

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	TAU	ACTION + TAU (n=40)		
	(n=39)			
Year of cancer diagnosis				
2020	21 (53.8%)	19 (47.5%)		
2021	18 (46.2%)	21 (52.5%)		
Cancer Incidence at randomisation				
First primary	37 (94.9%)	37 (92.5%)		
Second primary	1 (2.6%)	1 (2.5%)		
Recurrence of previous primary	1 (2.6%)	0 (0.0%)		
Missing	0 (0.0%)	2 (5.0%)		
Current stage of cancer				
Stage IA	16 (41.0%)	16 (40.0%)		
Stage IB	2 (5.1%)	2 (5.0%)		
Stage IIA	15 (38.5%)	16 (40.0%)		
Stage IIB	6 (15.4%)	4 (10.0%)		
Stage IIIA	0 (0.0%)	2 (5.0%)		
Missing	0 (0.0%)	0 (0.0%)		
Breast cancer treatment received*				
Surgery: Lumpectomy	30 (76.9%)	25 (62.5%)		
Surgery: Unilateral mastectomy	10 (25.6%)	14 (35.0%)		
Surgery: Double Mastectomy	0 (0.0%)	0 (0.0%)		
Radiotherapy	19 (48.7%)	22 (55.0%)		
Chemotherapy	6 (15.4%)	9 (22.5%)		
Other Treatment	0 (0.0%)	3 (7.5%)		
Hormone therapy regimen*				
Tamoxifen	15 (38.5%)	12 (30.0%)		
Anastrozole	22 (56.4%)	27 (67.5%)		
Raloxifene	0 (0.0%) 0 (0.0%			
Exemestane	2 (5.1%)	2 (5.0%)		
Letrozole	0 (0.0%)	1 (2.5%)		

	TAU			
	(n=39)	ACTION + TAU (n=40)		
Supportive therapies currently receiving or received within the last month*				
Occupational therapy	0 (0.0%)	0 (0.0%)		
Physiotherapy	3 (7.7%)	3 (7.5%)		
Other Supportive therapy	8 (20.5%)	7 (17.5%)		
Other therapies received / undertaken following HT prescription*				
Mindfulness	3 (7.7%)	0 (0.0%)		
Other therapy	2 (5.1%)	2 (5.0%)		
Menopausal status				
Pre-menopausal (no change in menstrual frequency in the previous 12 months)	5 (12.8%)	6 (15.0%)		
Peri-menopausal (changes in menstrual frequency in the previous 12 months or 3-11 months of amenorrhoea)	5 (12.8%)	4 (10.0%)		
Post-menopausal (no menstrual cycles for 12 months or more)	27 (69.2%)	26 (65.0%)		
Unsure	2 (5.1%)	4 (10.0%)		
Charlson comorbidities (Present)*				
Myocardial infarction	1 (2.6%)	0 (0.0%)		
Congestive heart failure	0 (0.0%)	0 (0.0%)		
Peripheral vascular disease	0 (0.0%)	0 (0.0%)		
Cerebrovascular disease	1 (2.6%)	0 (0.0%)		
Dementia	0 (0.0%)	0 (0.0%)		
Chronic pulmonary disease	0 (0.0%)	0 (0.0%)		
Connective tissue disease	4 (10.3%)	3 (7.5%)		
Peptic ulcer disease	1 (2.6%)	0 (0.0%)		
Liver disease				
Mild	0 (0.0%)	0 (0.0%)		
Moderate to severe	0 (0.0%)	0 (0.0%)		
None	39 (100.0%)	40 (100.0%)		

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	TAU			
	(n=39)	ACTION + TAU (n=40		
Diabetes Mellitus				
End-organ damage	0 (0.0%)	0 (0.0%)		
No/diet controlled	37 (94.9%)	37 (92.5%)		
Uncomplicated	2 (5.1%)	3 (7.5%)		
Hemiplegia	0 (0.0%)	0 (0.0%)		
Moderate to severe chronic kidney disease	1 (2.6%)	0 (0.0%)		
Solid tumor				
Localised	0 (0.0%)	0 (0.0%)		
Metastatic	0 (0.0%)	0 (0.0%)		
None	39 (100.0%)	40 (100.0%)		
Leukaemia	0 (0.0%)	0 (0.0%)		
Lymphoma	0 (0.0%)	0 (0.0%)		
AIDS	1 (2.6%)	0 (0.0%)		
Charlson Comorbidity Index				
0	30 (76.9%)	34 (85.0%)		
1	6 (15.4%)	6 (15.0%)		
2	1 (2.6%)	0 (0.0%)		
3	1 (2.6%)	0 (0.0%)		
6	1 (2.6%)	0 (0.0%)		
Charlson Comorbidity Index				
Mean (SD)	0.4 (1.12)	0.2 (0.36)		
Median (Range)	0.0 (0.0, 6.0)	0.0 (0.0, 1.0)		
Missing	0	0		

\* More than one response can be provided

## Outcome measures

	Baseline			3 months			6 months		
	TAU N (%)	ACTION + TAU N (%)	Total N (%)	TAU N (%)	ACTION + TAU N (%)	Total N (%)	TAU N (%)	ACTION + TAU N (%)	Total N (%)
Randomised	39	40	79	39	40	79	39	40	79
Any questionnaire data received	39 (100%)	39 (100%)	78 (98.7%)	35 (89.7%)	36 (90.0%)	71 (89.9%)	35 (89.7%)	35 (87.5%)	70 (88.6%)
Participant withdrew from questionnaires	0 (0.0%)	0 (0.0%)	0 (0.0%)	0 (0.0%)	2 (5.0%)	2 (2.5%)	0 (0.0%)	2 (5.0%)	2 (2.5%)
No questionnaire data received	0 (0.0%)	1 (2.5%)	1 (1.3%)	4 (10.3%)	2 (5.0%)	6 (7.6%)	4 (10.3%)	3 (7.5%)	7 (8.9%)

## Adverse events

Only related and unexpected serious adverse events (RUSAEs) were reportable in this study. No RUSAEs were reported.