

Implementation of routine physical and psychological symptom distress screening in cancer care

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Prevalence of psychological distress in patients with cancer

- Active phase of illness trajectory
 - Major depression (16%)
 - Anxiety (10%)
- Survivorship (at least 2 years post-diagnosis)
 - Major depression (11.6%)
 - Healthy controls (10.2%)
 - Anxiety (17.9%)
 - Healthy controls (13.9%)
- Palliative settings
 - Major depression (14.3%)
 - Anxiety (9.6%)



Mitchell, Chan, Bhatti, et al., 2011

Mitchell, Ferguson, Gill, et al., 2013

Mitchell, Chan, Bhatti, et al., 2011



Psychological and Physical Adjustment to Breast Cancer Over 4 Years: Identifying Distinct Trajectories of Change

> Vicki S. Helgeson, Pamela Snyder, and Howard Seltman Carnegie Mellon University

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IDENTIFICATION OF DISTINCT DEPRESSIVE SYMPTOM TRAJECTORIES IN WOMEN FOLLOWING SURGERY FOR BREAST CANCER

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Trajectories of psychological distress among Chinese

women diagnosed with breast cancer Wendy W. T. Lam¹*, George A. Bonanno², Anthony D. Mancini², Samuel Ho^{1,3}, Miranda Chan⁴, Wai Ka Hung⁴, Amy Or⁴ and Richard Fielding¹ ¹Centre for Psycho-Oncology Research & Training, Department of Community Medicine & Unit for Behavioural Sciences, The University of Hong Kong, Hong Kong

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The evolution of psychological distress trajectories in women diagnosed with advanced breast cancer: a longitudinal study

W. W. T. Lam¹*, I. Soong², T. K. Yau², K. Y. Wong³, J. Tsang⁴, W. Yeo⁵, J. Suen⁵, W. M. Ho⁵, W. K. Sze⁶, A. W. Y. Ng⁶, ^{© 2010} A. Kwong⁷, D. Suen⁷ and R. Fielding¹ ^(278-6133/0612.00) (10.1017)⁽³⁷⁾

Health Psychology 2010, Vol. 29, No. 2, 160–168

> Identification and Prediction of Distress Trajectories in the First Year After a Breast Cancer Diagnosis

Inge Henselmans University Medical Center Groningen Vicki S. Helgeson Carnegie Mellon University

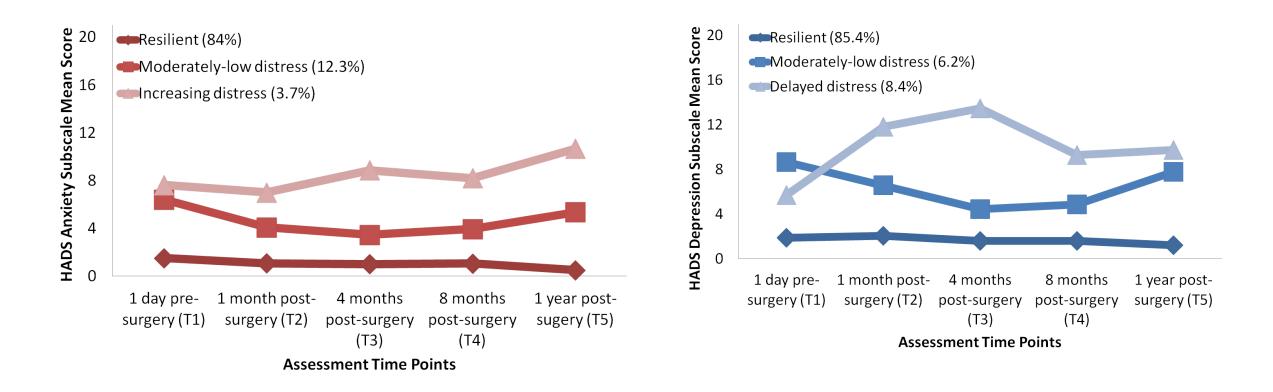
Howard Seltman Carnegie Mellon University Jakob de Vries University Medical Center Groningen

Robbert Sanderman and Adelita V. Ranchor University Medical Center Groningen





Trajectories of psychological distress among Chinese patients with colorectal cancer (n=232)



Trajectory patterns of distress from pre-surgery to 1 year post-surgery



Due distance for American	Moderately-low distress group			Persistent distress group				
Predictors for Anxiety	Resilient group as a reference group							
	OR	95% CI	SE	P-value	OR	95% CI	SE	P-value
Pre-surgical negative cancer- related intrusive thoughts	1.231	1.093; 1.385	0.060	0.001	1.303	1.021; 1.663	0.125	0.034
Pre-surgical physical symptom intrusiveness	1.095	0.996; 1.204	0.048	NS	1.216	0.976; 1.514	0.112	NS
Dispositional Optimism	0.974	0.925; 1.034	0.114	NS	0.531	0.292; 0.965	0.305	0.038
	Moderately-low distress group			Delayed distress group				
Predictors for Depression	Resilient group as a reference group							
	OR	95% CI	SE	P-value	OR	95% CI	SE	P-value
Pre-surgical negative cancer- related intrusive thoughts	1.158	1.006; 1.332	0.072	0.041	1.040	0.873; 1.239	0.089	NS
Pre-surgical physical symptom intrusiveness	1.241	1.112; 1.385	0.056	<0.001	1.118	1.013; 1.394	0.081	0.034
Dispositional Optimism	1.065	0.822; 1.380	0.132	NS	0.625	0.419; 0.931	0.204	0.021
Stage of disease								
Stage 0-II	1.214	0.352; 4.189	0.632	NS	0.126	0.021; 0.740	0.904	0.022
Stage III-IV	Referent			Referent				

⁺Insignificant predictors were excluded from the table, and these models were adjusted for covariates (including age, gender, marital status, educational level and occupation).



Psychological distress in cancer survivors

- Most patients were psychologically resilience in response to cancer diagnosis
- Chronic distress
 - Breast cancer 10% to 15%
 - Colorectal cancer 4% to 20%
 - Predictors
 - Poor social support
 - Poor personal resources (e.g. pessimism, low self-esteem, negative intrusive thoughts)
 - Unmanaged physical symptom distress
 - Poor satisfaction with treatment decision making



Impacts of chronic distress on long-term survivorship

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Distress trajectories at the first year diagnosis of breast cancer in relation to 6 years survivorship

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Multiple regression modeling of predictors of 6-year psychosocial outcome measures

ß values	HADS-A	HADS-D	CIES-R Intrusive	CIES-R Avoidance	CIES-R Hyper- arousal	ChSAS Family	ChSAS Self- image	ChSAS Appearance & Sexuality
Distress Trajectory Resilient [†]	Referent	Referent	Referent	Referent	Referent	Referent	Referent	Referent
$Recovered^{\dagger}$	0.02	-0.04	0.14*	0.10	0.08	0.03	-0.17*	-0.15*
Delay Recovery [†]	0.17**	0.05	0.15*	0.06	0.13	0.01	-0.05	-0.22**
Chronic Distress [†]	0.31**	0.22**	0.40***	0.25***	0.32***	-0.17*	-0.36***	-0.31***
C-LOT-R	-0.29**	-0.33***	-0.19**	-0.17*	-0.12	0.21**	0.20**	0.23**
GSES	-0.17**	-0.09	-0.07	0.05	-0.14*	0.09	-0.01	-0.01
EORTC Arm Breast	0.06 0.22**	0.03 0.19**	0.06 0.19**	- 0.20**	- 0.21**	-	- -0.04	- -0.14*
Age Marital Status Married	-0.22	20**	-0.03	-	-	-	0.07	- 0.17**
Occupation Housewife Employed Unemployed Retired	-0.22** Referent -0.04 -0.05	-0.22** Referent 0.01 -0.09	-0.35*** Referent Referent -0.20**	-0.19** Referent Referent 0.01	-0.19** Referent Referent -0.12	0.06 0.05 Referent -0.04	0.14* -0.06 Referent 0.11	0.10 Referent -0.12 0.01
Breast Cancer recurrence	-	-	-	-	-	-	-0.15*	-0.14*

Distress trajectory: 1-8 months post-surgery. All variable measured concurrently at 6 years, except † based on 1-8 months distress trajectory. Significance: *<0.05, **<0.01, ***<0.001

Implications



- Cancer patients who experienced chronic distress during the acute phase reported the worst longer-term outcomes
- Interventions should be targeted to differentiate those who are at risk of chronic distress during the acute phase of illness trajectory
 - Ensuring optimal communications and decision-making support are essential
 - Assessing symptom distress and optimizing symptom management should be implemented at early post-operative phase
 - Screening patients with poor social and personal resources





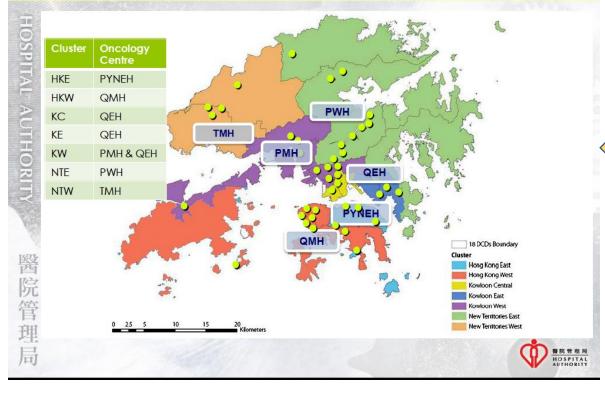
IPOS International Standard of Quality Cancer Care July 2010 (Revised October 2014)

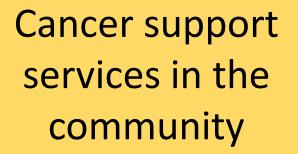
- 1. Psychosocial cancer care should be recognised as an universal human right
- 2. Quality cancer care must integrate the psychosocial domain into routine care



Psychosocial cancer care in Hong Kong

Distribution of Public Hospitals and 6 Clinical Oncology Centres in HA





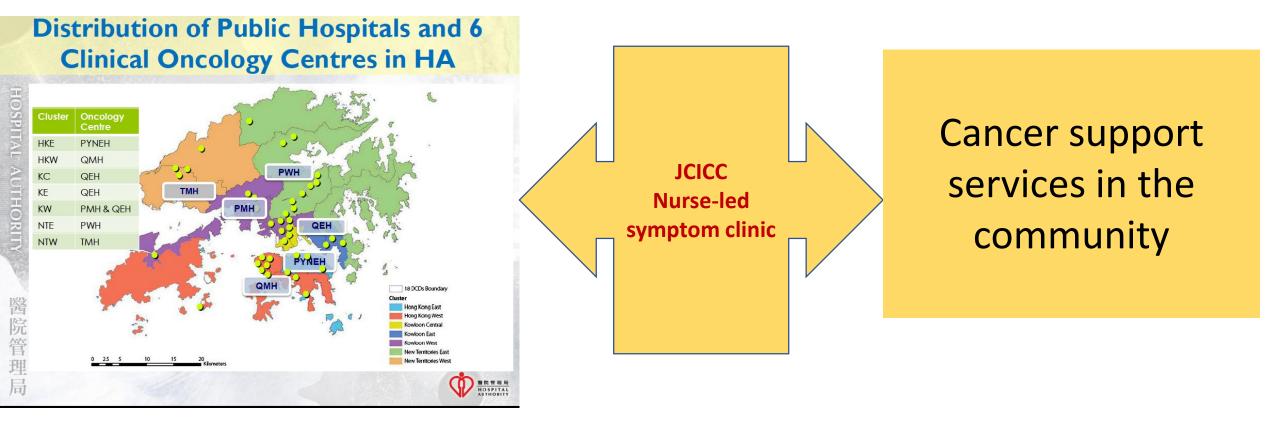


HKU Jockey Club Institute of Cancer Care (ICC)

- A critical platform to provide supportive care in collaboration with clinical oncology units and non-government organizations.
- Screening for psychosocial health need and symptom management services
 - Interventions addressing specific symptom distress
 - Cancer survivorship programme aiming to regain normalcy

New Initiatives



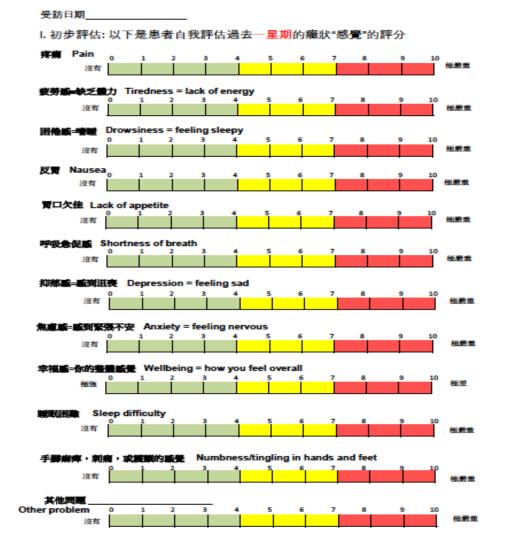


Step-Up Personalized Supportive Care (SUPer Care)



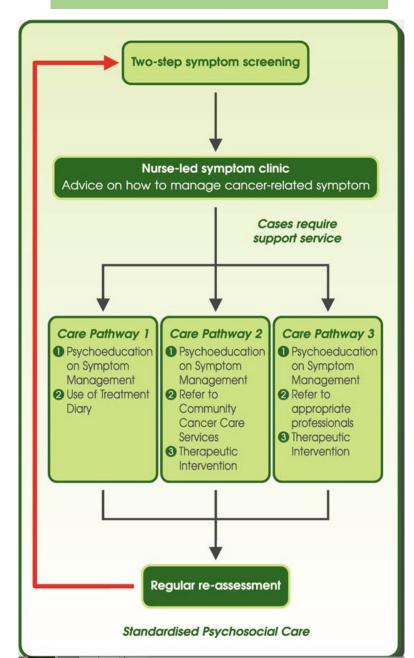
HKU Jockey Club Institute of Cancer Care (JCICC) 香港大學賽馬會癌症綜合關鍵中心

Edmonton Symptom Assessment System (ESAS)



SUPer Care Model









August 2018 – Nov 2020

Assessment	Number of patients to be screened	Number of patients who met the cut-off scores
ESAS assessment	2234	629 Non-case (Score <4) (28%) 779 Sub-clinical case (scores 4 -6) (35%) 826 cases (scores ≥7) (37%) - 590 (71.4%) consent for service
PSQI (Pittsburg sleep Quality Inventory)	339	$318 (94\%) (scores \ge 5 greater)$
BFI (Brief fatigue inventory)	209	$37 (17.7\%) (\text{score} \ge 7)$
ID Pain – neuropathy	220	$71 (32.3\%) (\text{score} \ge 3)$
Pain (Brief pain inventory)	120	112 (93.3%) (current pain severity score \geq 5)
HADS (Hospital anxiety and depression scale)	403	A: 94 (score 8-10– borderline) (23.3%) A: 94 (score >11 – clinical case) (23.3%) D: 80 (score 8-10 –borderline) (20%) D: 98 (score >11 – clinical case) (24.3%)
FCR (Fear of cancer recurrence)	137	$88 \text{ (score } \geq 13) (63.8\%)$
Appetite (Council of Nutrition Appetite Questionnaire)	65	$31 (\text{score} \le 28) (47.7\%)$

Patient Characteristics



	Non case/refuse for service [n=1644]	Consent to service (Case) [n=590]
Age (mean)	60.42±17.97	59.1±11.57
Gender		
Male	414 (25.2%)	156 (26.4%)
Female	730 (44.4%)	433 (73.4%)
Missing	500 (30.4%)	1 (0.2%)
Marital Status		
Single	149 (9.1%)	63 (10.7%)
Married/Cohabited	826 (50.2%)	299 (50.7%)
Divorced/Separated	88 (5.4%)	48 (8.1%)
Widowed	77 (4.7%)	25 (4.2%)
Missing	504 (30.7%)	155 (26%)
Educational level		
No formal education or primary education	460 (28%)	137 (23.2%)
Secondary education	731 (44.5%)	270 (45.8%)
Tertiary education	428 (26%)	170 (28.8%)
Missing	25 (1.5%)	13 (2.2%)



Patient Characteristics

	Non case or refuse for service [n=1644]	Consent to service (Case) [n=590]
Cancer type		
Breast	448 (27.3%)	226 (38.3%)
CRC	598 (36.3%)	157 (26.6%)
Lung	215 (13.2%)	91 (15.4%)
Gynaecology	85 (5.2%)	58 (9.5%)
Head and neck	124 (7%)	27 (4.7%)
Gastrointestinal / Hepatobiliary	51 (3.2%)	10 (1.7%)
Sarcoma	20 (1.3%)	7 (1.2%)
Neurological	15 (1.1%)	4 (0.7%)
Others	73 (4.4%)	10 (1.8%)
Missing	15 (1%)	0 (0%)

Patients who refused for service



Focus assessment	Number of patients who met screening (ESAS ≥ 7) cut off	No. of patients who met the cut-off scores
PSQI (Pittsburg sleep Quality Inventory)	114	74 (score ≥ 5) (65%)
Fatigue – BFI	67	0 (score ≥ 7)
ID Pain - neuropathy	63	3 (score ≥ 3) (5%)
Pain – BPI	24	10 (#3 score ≥ 5) (42%)
Anxiety and Depression – HADS	129	Anxiety: 4 (score 8-10 – borderline) (3%) 6 (score≥ 11 – clinical case) (5%) Depression: 5 (score 8-10 – borderline) (4%) 5 (score ≥ 11 – clinical case) (4%)
Fear of cancer recurrence - FCR	64	2 (score ≥ 13) (3%)
Appetite – Council of Nutrition Appetite Questionnaire	18	2 (score ≤28) (11%)



ESAS score distribution by symptoms for cases (n=590)

Symptoms/ESAS score	0-3	4-6	≥7	Total
Pain	336 (56.9%)	134 (22.7%)	120 (20.3%)	590
Tiredness	148 (25.1%)	233 (39.5%)	<mark>209 (35.4%)</mark>	590
Drowsiness .	199 (33.8%)	202 (34.3%)	<mark>188 (31.9%)</mark>	589
Nausea	492 (83.5%)	72 (12.2%)	25 (4.2%)	589
Lack of appetite	401 (68.1%)	136 (23.4%)	52 (8.8%)	589
Shortness of breath	388 (66%)	150 (25.5%)	50 (8.5%)	588
Depression	272 (46.1%)	185 (31.4%)	<mark>133 (22.5%)</mark>	590
Anxiety	268 (45.4%)	175 (29.7%)	<mark>147 (24.9%)</mark>	590
Wellbeing	263 (44.8%)	216 (36.8%)	108 (18.4%)	587
Sleep difficulty	205 (34.7%)	150 (25.4%)	<mark>235 (39.8%)</mark>	590
Numbness	228 (38.8%)	140 (23.8%)	<mark>220 (37.4%)</mark>	588



JCICC Triage Pathway (n=590)

Pathway 1 Provide psychoeducation for self-management	337	57.1%
Pathway 2 Group activity (outside or in-house)	41	7%
Pathway 3 In depth one to one specialty consultation	212	35.9%

Summary



- 1 in 3 cancer survivors screened were identified as cases for symptom distress.
- The most common symptoms included sleep disturbance, numbness, fatigue and anxiety.
- Most cases consented for receiving interventions and follow-up in managing the unmet needs.
- 1 in 3 who are identified as distressed but decline professional help.
- Implementation research is needed to assess the impact of screening for symptom distress on the well-being of patients.



