

Agreement for depression diagnosis between DSM-IV-TR criteria, three validated scales, oncologist assessment, and psychiatric clinical interview in elderly patients with advanced ovarian cancer

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Background: Depression, a major outcome in cancer patients, is often evaluated by physicians relying on their clinical impressions rather than patient self-report. Our aim was to assess agreement between patient self-reported depression, oncologist assessment (OA), and psychiatric clinical interview (PCI) in elderly patients with advanced ovarian cancer (AOC).

Methods: This analysis was a secondary endpoint of the Elderly Women AOC Trial 3 (EWOT3), designed to assess the impact of geriatric covariates, notably depression, on survival in patients older than 70 years of age. Depression was assessed using the Geriatric Depression Scale-30 (GDS), the Hospital Anxiety Depression Scale, the distress thermometer, the mood thermometer, and OA. The interview guide for PCI was constructed from three validated scales: the GDS, the Hamilton Depression Rating Scale, and the Montgomery Asberg Depression Rating Scale (MADRS). *The Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised* (DSM) criteria for depression were used as a gold standard.

Results: Out of 109 patients enrolled at 21 centers, 99 (91%) completed all the assessments. Patient characteristics were: mean age 78, performance status ≥ 2 : 47 (47%). Thirty six patients (36%) were identified as depressed by the PCI versus 15 (15%) identified by DSM. We found moderate agreement for depression identification between DSM and GDS ($\kappa=0.508$) and PCI ($\kappa=0.431$) and high agreement with MADRS ($\kappa=0.663$). We found low or no agreement between DSM with the other assessment strategies, including OA ($\kappa=-0.043$). Identification according to OA (yes/no) resulted in a false-negative rate of 87%. As a screening tool, GDS had the best sensitivity and specificity (94% and 80%, respectively).

Conclusion: The use of validated tools, such as GDS, and collaboration between psychologists and oncologists are warranted to better identify emotional disorders in elderly women with AOC.

Keywords: depression, elderly, cancer, screening, geriatric assessment

Introduction

Depression is a major outcome in cancer patients, with an estimated frequency of 16%,¹ and this frequency is reported to be higher for elderly patients (45%).² Depression is the most common psychiatric illness in patients with cancer and it is known to reduce patients' quality of life and to decrease their adherence to medical treatments.³⁻⁵ Depressive symptoms (asthenia, sleep disturbances, anorexia, etc) have been reported to significantly affect patients and their families.^{3,6} Moreover, recent studies have

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shown that depression is an independent predictive factor for cancer-related mortality,^{7–9} even though depression usually responds well to treatment (nonpharmacologic and pharmacologic).^{10–12}

The evaluation of depression has been a challenge because symptoms cover a broad spectrum, from sadness to major depressive disorder, and because mood change is often difficult to assess when patients are confronted by repeated threats to life or are experiencing pain.⁶

For a depression diagnosis, clinicians typically rely on their clinical impression rather than on patient self-report or structured assessment.^{13,14} A systematic routine screening for depression has been implemented in some cancer care settings. Nevertheless, a substantial amount of data shows that depression remains underdiagnosed and undertreated in older patients, especially in the context of cancer,^{15–19} with low agreement between physicians and patients about the prevalence of mild and moderate/severe depression (33% and 13%, respectively).^{20–22} Similar results have been reported for nurses.^{14,23} Several reasons have been highlighted to explain those barriers to diagnosis and thus to adequate treatment.

First, many cancer patients are not treated for depression due to assessment difficulties, especially with older patients.^{6,19} This population often develops symptoms that differ from those experienced by younger patients, with more somatic complaints and fewer emotional symptoms (sadness, guilt, anhedonia).^{18,19} Patients in later adulthood are more reluctant to report depressive symptoms such as sadness or suicidal ideations.¹² Furthermore, they might consider these symptoms as belonging to aging processes.¹²

Those difficulties may be due to nondisclosure by the patients themselves, who may feel they are wasting the doctor's time or that they are in some way to blame for their distress and therefore choose to hide their feelings.²⁴ There is also the wrong belief among some professionals that terminal illness invariably causes depression,²⁵ and there are disagreements about the meaning of the term “depression” within psychiatric classification.²⁶ We also know that some symptoms of depression (eg, fatigue, loss of appetite) might be expected consequences of physical illness.^{27,28} It is possible that some of the difficulties faced when attempting to identify depression might be decreased by the use of appropriate screening scales (adapted to the population and to the clinical setting).¹²

This study aimed to assess agreement between patient self-reported depression using different scales,¹⁸ oncologist assessment (OA) based on the oncologist's clinical impressions, and a psychiatric clinical interview (PCI) conducted by

a psychologist using *the Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised* (DSM)²⁹ among elderly patients with advanced ovarian cancer (AOC).

Methods

This analysis was a secondary endpoint of the Groupe d'Investigateurs Nationaux pour l'Etude des Cancers Ovariens (GINECO) Elderly Women AOC Trial 3 (EWOT3) Phase II multicenter trial, designed to assess the impact of geriatric covariates, notably depression, on survival in patients older than 70 years of age receiving six courses of carboplatin as first-line chemotherapy for stage 3–4 ovarian cancer. The local ethics committee and the Institutional Review Board at the Hospices Civils de Lyon approved this study, and all patients gave written informed consent.

Participants

To be included, patients had to have advanced (International Federation of Gynecology and Obstetrics stage 3 or 4) ovarian cancer and be older than 70 years of age. They could not have received chemotherapy previously and should have had a life expectancy of at least 3 months. Inclusion and exclusion criteria and patient characteristics have been previously reported.³⁰

Assessments

Emotional disorders – and particularly depression – were assessed using patient self-report, OA, and PCI. Chronologically, oncologists were asked to provide their clinical impression before the self-reports. The PCI was performed within 10 days of enrollment according to the trial design, and the interviewer was blinded to the OA and to the self-report results.

Self-reports

All self-reports used were validated in their French form and included the Geriatric Depression Scale-30 (GDS),³¹ the Hospital Anxiety Depression Scale (HADS),^{32,33} the distress thermometer (DT), and the mood thermometer (MT).³⁴ The GDS is a 30-item scale. This scale has been developed and validated especially for depression screening among elderly patients^{35,36} and has also been used in cancer patients.³⁷ Scores of 0–9 are considered normal, 10–19 indicate mild depression, and 20–30 indicate severe depression.³⁵ The HADS is a 14-item scale with separate subscales for anxiety (HADS-A) and depression (HADS-D). It has been validated in cancer patients for depression and anxiety diagnosis³⁸ and can be administered or used as a self-assessment tool. The DT is a

one-item self-report screening tool for assessing psychological distress in cancer patients.³⁹ The DT asks patients to rate the intensity of their psychological distress using an eleven-point visual analog scale from 0 (no distress) to 10 (extreme distress),⁴⁰ with a score of 4 used to indicate a significant level of distress.⁴¹ The MT is a one-item self-report screening tool for measuring mood in cancer patients.³⁴ Patients are asked to answer the question “How depressed have you been today and over the last week?” by rating their answer on a 0–10 scale (0 = normal mood; 10 = extreme depression). A score of 4 is used to indicate a significant level of depression.³⁴ All of these self-report assessment tools were transformed into binary variables with the following cutoff scores: $\geq 10/30$ for GDS-30, $> 14/42$ for HADS, $> 4/10$ for MT, and $> 4/10$ for DT.

Oncologist assessment

Oncologists were invited to give their clinical impression regarding depression (yes/no). This item was previously shown to be correlated to survival in two previous GINECO studies of elderly patients with AOC.⁴²

Psychiatric clinical interview

A PCI was conducted by psychologists within 10 days of enrollment. The interview guide for the PCI was constructed and adapted from the DSM criteria and three validated scales: GDS, Hamilton Depression Rating Scale (HDRS), and Montgomery Asberg Depression Rating Scale (MADRS), according to a structured interview guide for evaluating depression in elderly patients developed by Tison.⁴³ Psychologists were to assess depression using DSM criteria (DSM major depressive disorder, yes/no) and to give their clinical impression (PCI, yes/no). The DSM was considered to be the gold standard.¹⁸

Procedure and outcomes

Demographic information (ie, age) and clinical data (ie, performance status, albumin level, weight, cancer diagnosis, and metastatic status) were collected.

Patient functioning was assessed using two validated tools: basal Activities of Daily Living (ADL) and Instrumental Activities of Daily Living (IADL).^{44–47}

On the day of enrollment, patient assessments included a clinical evaluation, laboratory results, and psychological self-reports (HADS, DT, MT, and GDS-30). In the 10 days after enrollment, a PCI based on MADRS and HDRS questionnaires was conducted.

Survival data were collected from the medical chart, starting from the time of enrollment to the patient’s death. Patients

alive at the time of data collection were included in the analysis as censored cases using the date of last follow-up.

Statistical considerations

The number of patients to include was defined in the first aim of the GINECO EWOT3 study, a secondary endpoint of which is examined in the current study, using previous findings from our group.⁴² A single analysis requires a sample of 105 patients, assuming that depression triples the risk of death (two-tailed test), with a significance level of $\alpha=5\%$ and a power of 90% ($\beta=10\%$).³⁰ This calculation was based on the assumption, on the basis of results from the EWOT1 and EWOT2 studies, that a death rate of 85% is expected at 2 years (86.4%, according to EWOT1 and EWOT2).

Categorical variables were presented by their frequencies and percentages and continuous variables by their means and standard deviations if they were normally distributed. When not normally distributed, data were reported by their median and interquartile range (Q1–Q3) and analyzed using nonparametric methods. Mann–Whitney *U*-tests and chi-square tests were performed to determine factors associated with depression according to the DSM. Agreements between depression diagnosis and DSM, PCI, and self-reported assessments were evaluated using Cohen’s kappa coefficient (κ).

P-values < 0.05 were considered statistically significant. All statistical analyses were performed using SPSS v17 (SPSS, Inc., Chicago, IL, USA).

Results

Patient characteristics

One hundred and nine patients had a psychological assessment; of these, 99 had a complete set of psychological data and were included in the analysis. Patients’ characteristics are presented in Table 1.

Depression assessments

Out of the 99 patients included, 15 (15%) were depressed according to DSM and 39 (39%) were depressed according to PCI. We found different levels of agreement between DSM – considered as the gold standard – and other assessments (Table 2):

- no agreement (kappa coefficient between 0.0 and 0.2): with the DT ($\kappa=-0.060$), MT ($\kappa=0.058$), HADS ($\kappa=0.127$), HDRS ($\kappa=0.151$), and OA ($\kappa=-0.043$),
- moderate agreement (kappa coefficient between 0.4 and 0.6): with GDS ($\kappa=0.508$) and with PCI ($\kappa=0.431$),
- high agreement (kappa coefficient between 0.6 and 0.8): with MADRS ($\kappa=0.663$).

Table 1 Patient characteristics, N=99

| Variables | N (%) |
|---|------------|
| Age in years, mean (SD) | 78 (5) |
| ≥ 80 | 38 (38) |
| ≥ 85 | 11 (11) |
| PS, N (%) | |
| 0–1 | 52 (53) |
| 2–3 | 47 (47) |
| ADL, median (Q1–Q3) | 5.5 (5–6) |
| Number of patients with an ADL ≤ 4 (moderate impairment), N (%) | 10 (10) |
| IADL, median (Q1–Q3) | 22 (17–26) |
| Number of patients with an IADL ≤ 24 , N (%) | 68 (69) |
| Depression according to DSM criteria, N (%) | 15 (15) |
| Depression according to PCI, N (%) | 39 (39) |
| Overall survival in months, mean (SD) | 15.7 (9.2) |

Abbreviations: ADL, Activities of Daily Living scale; DSM, *Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised*; IADL, Instrumental Activities of Daily Living scale; N, number; PCI, psychiatric clinical interview; PS, performance status; Q1–Q3, first through third quartiles; SD, standard deviation.

Considering the agreement between CPI and other assessments, results are the following:

- no agreement (kappa coefficient between 0.0 and 0.2): with the DT ($\kappa=-0.060$), MT ($\kappa=0.193$), HADS ($\kappa=-0.051$), and OA ($\kappa=-0.061$),
- low agreement (kappa coefficient between 0.2 and 0.4): with HDRS ($\kappa=0.389$),
- moderate agreement (kappa coefficient between 0.4 and 0.6): with DSM ($\kappa=0.431$), with GDS ($\kappa=0.517$), and with MADRS ($\kappa=0.494$).

The tool giving the best combination of sensitivity/specificity for depression screening among elderly patients was the GDS (cutoff score ≥ 10), with a sensitivity of 0.94 and a specificity of 0.80 as compared to DSM (Table 3). The DT, MT, and HADS had very poor sensitivity/specificity for depression identification (0.50/0.40, 0.50/0.57, and

Table 3 Sensitivity and specificity of the different assessments to DSM

| Variables | Sensitivity | Specificity | PPV | NPV |
|-----------------|-------------|-------------|------|------|
| DT ≥ 4 | 0.50 | 0.40 | 0.14 | 0.81 |
| MT > 4 | 0.50 | 0.57 | 0.18 | 0.86 |
| GDS ≥ 10 | 0.94 | 0.80 | 0.47 | 0.99 |
| HADS ≥ 16 | 0.50 | 0.67 | 0.22 | 0.88 |
| HDRS ≥ 7 | 0.94 | 0.42 | 0.23 | 0.97 |
| MADRS ≥ 16 | 0.88 | 0.91 | 0.63 | 0.97 |
| PCI | 1.00 | 0.71 | 0.38 | 1.00 |
| OA | 0.13 | 0.82 | 0.12 | 0.84 |

Abbreviations: DSM, *Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised*; DT, Distress Thermometer; GDS, Geriatric Depression Scale; HADS, Hospital Anxiety Depression Scale; HDRS, Hamilton Depression Rating Scale; MADRS, Montgomery Asberg Depression Rating Scale; MT, Mood Thermometer; NPV, Negative Predictive Value; OA, oncologist assessment; PCI, psychiatric clinical interview; PPV, Positive Predictive Value.

0.50/0.67, respectively). Among clinical assessments, a good sensitivity/specificity (1.00/0.71) was obtained with the PCI, whereas OA resulted in a very low sensitivity (0.13).

Depression and overall survival

Mean survival was 15.7 months (standard deviation = 9.2 months). We did not find any significant difference regarding patients' characteristics between depressed and nondepressed patients according to DSM (Table 4). Patients with major depressive disorder according to the DSM tended to have a shorter overall survival than did nondepressed patients, at 14.9 months (3.0–18.3 months) versus 15.2 months (9.4–22.8 months); however, this difference was not statistically significant ($P=0.281$).

Discussion

The main objective of this study was to assess the level of agreement between patient self-reported depression rated on different scales, oncologist clinical impression, and

Table 2 Comparison of depression diagnosis using different strategies between depressed and nondepressed patients according to DSM diagnosis, N=99

| Variables | Depressed patients | Nondepressed patients | κ coefficient | P-value* |
|-----------------|--------------------|-----------------------|------------------------|----------|
| DT ≥ 4 | 50 (60) | 8 (53) | −0.060 (−0.189; 0.069) | <0.001 |
| MT > 4 | 36 (43) | 8 (53) | 0.058 (0.096; 0.212) | <0.001 |
| GDS ≥ 10 | 17 (20) | 15 (100) | 0.508 (0.326; 0.690) | <0.001 |
| HADS > 14 | 28 (33) | 8 (53) | 0.127 (−0.051; 0.305) | 0.01 |
| HDRS ≥ 7 | 49 (58) | 15 (100) | 0.151 (0.051; 0.251) | <0.001 |
| MADRS ≥ 16 | 8 (10) | 14 (94) | 0.663 (0.473; 0.853) | 0.109 |
| PCI | 24 (29) | 15 (100) | 0.431 (0.268; 0.594) | <0.001 |
| OA | 15 (18) | 2 (13) | −0.043 (−0.228; 0.223) | 0.851 |

Note: *McNemar's test.

Abbreviations: κ , Cohen's kappa coefficient; DSM, *Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised*; DT, Distress Thermometer; GDS, Geriatric Depression Scale; HADS, Hospital Anxiety Depression Scale; HDRS, Hamilton Depression Rating Scale; MADRS, Montgomery Asberg Depression Rating Scale; MT, Mood Thermometer; N, number; OA, oncologist assessment; PCI, psychiatric clinical interview.

Table 4 Associations between depression diagnosis according to DSM and patient characteristics, N=99

| Variables | No depression (DSM) N=84 | Depression (DSM) N=15 | P-value |
|------------------------------------|-----------------------------|--------------------------|---------|
| Age (years), median (Q1–Q3) | 78 (73–82) | 78 (76–82) | 0.261* |
| PS 2–3, N (%) | 37 (44) | 10 (67) | 0.106‡ |
| ADL, median (Q1–Q3) | 6 (5–6) | 6 (5–6) | 0.737* |
| IADL, median (Q1–Q3) | 22 (16–26) | 22 (17–26) | 0.357* |
| Survival in months, median (Q1–Q3) | 15.2 (9.4–22.8) | 14.9 (3.0–18.3) | 0.281* |

Notes: *Mann–Whitney U-test; ‡chi-square test.

Abbreviations: ADL, Activities of Daily Living scale; DSM, *Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised*; IADL, Instrumental Activities of Daily Living scale; N, number; PS, performance status; Q1–Q3, first through third quartiles.

a PCI conducted by a psychologist using the DSM criteria among elderly patients with AOC. We found moderate agreement for depression identification between DSM and GDS ($\kappa=0.508$) and PCI ($\kappa=0.431$) and high agreement with MADRS ($\kappa=0.663$). We found low or no agreement between DSM and the other assessments strategies, including OA ($\kappa=-0.043$). One explanation may be the difficulty, according to the DSM classification, to connect somatic symptoms with depression, considering that ovarian cancer may lead to appetite change, fatigue, and libido or sleep disruption.

This result confirms the importance of the National Comprehensive Cancer Network guidelines and the work of Jacobsen regarding the implementation of systematic strategies for distress screening among cancer patients.^{48–50} Indeed, OA results in a false-negative rate of more than 80%, leading to the recommendation that health care professionals without specific training for depression diagnosis should be encouraged to use validated tools in their daily clinical practice.

The screening tool with the best combination of sensitivity/specificity for depression among elderly patients with AOC was the GDS, with a sensitivity of 0.94 and a specificity of 0.80. These results suggest that the GDS is a good tool, as part of a screening “routine” among elderly patients with AOC, and should be preferred to more classic and unspecific tools such as the DT or the HADS. Our results might be partially explained by the fact that most of the scales for depression screening/diagnosis were validated among middle-aged patients.⁵¹

The systematic use of this type of tool (eg, GDS) has been widely advocated to improve the identification of depression in this frail population.^{2,20,52} However, the vast majority of cancer centers do not have a strategy for the identification of depression in patients with late-stage cancer. Our results suggest that some existing tools are consistent with the DSM diagnosis of depression that is still considered as the gold

standard. Nonetheless, our findings suggest that we should distinguish first-line tools (screening) from second-line tools (diagnostic).^{11,26,50,53} Indeed, even if these tools allow easy identification of depression, in case of treatment failure and/or severe risk factors, these patients should be referred to a mental health specialist to confirm the diagnosis.⁵⁴

Our group has recently published a report proposing a geriatric vulnerability score in elderly patients with AOC.³⁰ In this previous work, we used the HADS for the development of the vulnerability score. Our findings suggest that this score is less accurate than the GDS for the identification of depression. More research is necessary on the best way to screen for depression among this specific population, so as to allow simple screening in daily practice. It is important to recognize that in this population of patients with severe physical and/or psychosocial distress, only very brief assessments are likely to be adopted in clinical practice.

We did not find significant difference in overall survival between depressed patients and nondepressed patients (Table 4, Figure S1). These results are in contradiction with our previous results, which reported a significant impact of OA and HADS on survival.^{42,55} These results were recently confirmed in a pooled analysis of EWOT1, EWOT2, and EWOT3 (submitted) data, leading to the idea that emotional disorders in general, including cancer distress, might have a more significant impact on survival than authentic depression as defined by DSM. Such results are in line with the current debate on HADS performance, considered as poor in depression identification^{56,57} but powerful in predicting various medical outcomes.⁵⁸ More research is necessary to better characterize the association between both depression and survival among elderly cancer patients.

The main limitation of this study was that this analysis was a secondary endpoint of the GINECO EWOT3 Phase II multicenter trial, designed to assess the impact of geriatric covariates, notably depression, on survival in elderly patients with AOC. As a result, the sample size was not calculated

regarding sensitivity and specificity of the different tools as compared to DSM, limiting generalization of our results.

Another limitation is that all the threshold scores used for this study come from existing literature and it is known that sensitivity and specificity of the instrument depends on cutoff points, which might need to be changed in different patient populations (eg, inpatients, outpatients, and elderly patients, or by stage and type of disease).⁵⁹ This is probably true for all these tools, and future research should try to explore this issues using the same tool among different populations. Finally, our results suggest that depression systematic screening using specific tool (eg, GDS) is feasible and efficient, and that these tools should be selected according to the population and the clinical setting. More research is also necessary to better understand what should be done in case of positive screening for depression. Research that includes prospective longitudinal studies will help to characterize depression and develop optimal treatment strategies.

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Disclosure

The authors report no conflicts of interest in this work.

References

1. Mitchell AJ, Chan M, Bhatti H, et al. Prevalence of depression, anxiety, and adjustment disorder in oncological, haematological, and palliative-care settings: a meta-analysis of 94 interview-based studies. *Lancet Oncol*. 2011;12(2):160–174.
2. Bellera CA, Rainfray M, Mathoulin-Pélissier S, et al. Screening older cancer patients: first evaluation of the G-8 geriatric screening tool. *Ann Oncol*. 2012;23(8):2166–2172.
3. DiMatteo MR, Lepper HS, Croghan TW. Depression is a risk factor for noncompliance with medical treatment: meta-analysis of the effects of anxiety and depression on patient adherence. *Arch Intern Med*. 2000;160(14):2101–2107.
4. Block S. Assessing and managing depression in the terminally ill patient. ACP-ASIM End-of-Life Care Consensus Panel. American College of Physicians – American Society of Internal Medicine. *Ann Intern Med*. 2000;132(3):209–218.
5. Goodwin JS, Zhang DD, Ostir GV. Effect of depression on diagnosis, treatment, and survival of older women with breast cancer. *J Am Geriatr Soc*. 2004;52(1):106–111.
6. Weinberger MI, Bruce ML, Roth AJ, Breitbart W, Nelson CJ. Depression and barriers to mental health care in older cancer patients. *Int J Geriatr Psychiatry*. 2011;26(1):21–26.
7. Lloyd-Williams M, Shiels C, Taylor F, Dennis M. Depression – an independent predictor of early death in patients with advanced cancer. *J Affect Disord*. 2009;113(1–2):127–132.
8. Satin JR, Linden W, Phillips MJ. Depression as a predictor of disease progression and mortality in cancer patients: a meta-analysis. *Cancer*. 2009;115(22):5349–5361.
9. Pirl WF, Traeger L, Greer JA, et al. Depression, survival, and epidermal growth factor receptor genotypes in patients with metastatic non-small cell lung cancer. *Palliat Support Care*. 2013;11(3):223–229.
10. Rayner L, Price A, Evans A, Valsraj K, Hotopf M, Higginson IJ. Antidepressants for the treatment of depression in palliative care: systematic review and meta-analysis. *Palliat Med*. 2011;25(1):36–51.
11. Rayner L, Price A, Hotopf M, Higginson IJ. The development of evidence-based European guidelines on the management of depression in palliative cancer care. *Eur J Cancer*. 2011;47(5):702–712.
12. Evans M, Mottram P. Diagnosis of depression in elderly patients. *Adv Psychiatr Treat*. 2000;6:49–56.
13. Söllner W, DeVries A, Steixner E, et al. How successful are oncologists in identifying patient distress, perceived social support, and need for psychosocial counselling? *Br J Cancer*. 2001;84(2):179–185.
14. Little L, Dionne B, Eaton J. Nursing assessment of depression among palliative care cancer patients. *J Hosp Palliat Nurs*. 2005;7(2):98–106.
15. Klap R, Unroe KT, Unützer J. Caring for mental illness in the United States: a focus on older adults. *Am J Geriatr Psychiatry*. 2003;11(5):517–524.

16. Morley JE. The top 10 hot topics in aging. *J Gerontol A Biol Sci Med Sci*. 2004;59(1):24–33.
17. Flaherty JH, McBride M, Marzouk S, et al. Decreasing hospitalization rates for older home care patients with symptoms of depression. *J Am Geriatr Soc*. 1998;46(1):31–38.
18. Nelson CJ, Cho C, Berk AR, Holland J, Roth AJ. Are gold standard depression measures appropriate for use in geriatric cancer patients? A systematic evaluation of self-report depression instruments used with geriatric, cancer, and geriatric cancer samples. *J Clin Oncol*. 2010;28(2):348–356.
19. Weinberger MI, Roth AJ, Nelson CJ. Untangling the complexities of depression diagnosis in older cancer patients. *Oncologist*. 2009;14(1):60–66.
20. Passik SD, Dugan W, McDonald MV, Rosenfeld B, Theobald DE, Edgerton S. Oncologists' recognition of depression in their patients with cancer. *J Clin Oncol*. 1998;16(4):1594–1600.
21. Mitchell AJ, Vaze A, Rao S. Clinical diagnosis of depression in primary care: a meta-analysis. *Lancet*. 2009;374(9690):609–619.
22. Mitchell AJ, Rao S, Vaze A. International comparison of clinicians' ability to identify depression in primary care: meta-analysis and meta-regression of predictors. *Br J Gen Pract*. 61(583):e72–e80.
23. Rhondali W, Hui D, Kim SH, et al. Association between patient-reported symptoms and nurses' clinical impressions in cancer patients admitted to an acute palliative care unit. *J Palliat Med*. 2012;15(3):301–307.
24. Maguire P. Improving the detection of psychiatric problems in cancer patients. *Soc Sci Med*. 1985;20(8):819–823.
25. Lloyd-Williams M, Friedman T, Rudd N. A survey of antidepressant prescribing in the terminally ill. *Palliat Med*. 1999;13(3):243–248.
26. Hotopf M, Chidgey J, Addington-Hall J, Ly KL. Depression in advanced disease: a systematic review Part 1. Prevalence and case finding. *Palliat Med*. 2002;16(2):81–97.
27. Yennurajalingam S, Kwon JH, Urbauer DL, Hui D, Reyes-Gibby CC, Bruera E. Consistency of symptom clusters among advanced cancer patients seen at an outpatient supportive care clinic in a tertiary cancer center. *Palliat Support Care*. 2013;11(6):473–480.
28. Yennurajalingam S, Urbauer DL, Casper KL, et al. Impact of a palliative care consultation team on cancer-related symptoms in advanced cancer patients referred to an outpatient supportive care clinic. *J Pain Symptom Manage*. 2011;41(1):49–56.
29. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition, Text Revision (DSM-IV-TR)*. 4th ed. Washington, DC: American Psychiatric Association; 2000.
30. Falandry C, Weber B, Savoye AM, et al. Development of a geriatric vulnerability score in elderly patients with advanced ovarian cancer treated with first-line carboplatin: a GINECO prospective trial. *Ann Oncol*. 2013;24(11):2808–2813.
31. Clément JP, Nassif RF, Léger JM, Marchan F. [Development and contribution to the validation of a brief French version of the Yesavage Geriatric Depression Scale]. *Encephale*. 1997;23(2):91–99. French.
32. Lépine JP, Godchau M, Brun P, Lempérière T. [Evaluation of anxiety and depression among patients hospitalized on an internal medicine service]. *Ann Med Psychol (Paris)*. 1985;143(2):175–189. French.
33. Lépine JP, Godchau M, Brun P. Anxiety and depression in inpatients. *Lancet*. 1985;2(8469–8470):1425–1426.
34. Gil F, Grassi L, Travado L, Tomamichel M, Gonzalez JR; Southern European Psycho-Oncology Study Group. Use of distress and depression thermometers to measure psychosocial morbidity among southern European cancer patients. *Support Care Cancer*. 2005;13(8):600–606.
35. Yesavage JA, Brink TL, Rose TL, et al. Development and validation of a geriatric depression screening scale: a preliminary report. *J Psychiatr Res*. 1982–1983;17(1):37–49.
36. Burke WJ, Nitcher RL, Roccaforte WH, Wengel SP. A prospective evaluation of the Geriatric Depression Scale in an outpatient geriatric assessment center. *J Am Geriatr Soc*. 1992;40(12):1227–1230.
37. Crawford GB, Robinson JA. The geriatric depression scale in palliative care. *Palliat Support Care*. 2008;6(3):213–223.
38. Lloyd-Williams M, Spiller J, Ward J. Which depression screening tools should be used in palliative care? *Palliat Med*. 2003;17(1):40–43.
39. Hegel MT, Collins ED, Kearing S, Gillock KL, Moore CP, Ahles TA. Sensitivity and specificity of the Distress Thermometer for depression in newly diagnosed breast cancer patients. *Psychooncology*. 2008;17(6):556–560.
40. Keir ST, Calhoun-Eagan RD, Swartz JJ, Saleh OA, Friedman HS. Screening for distress in patients with brain cancer using the NCCN's rapid screening measure. *Psychooncology*. 2008;17(6):621–625.
41. Jacobsen PB, Donovan KA, Trask PC, et al. Screening for psychologic distress in ambulatory cancer patients. *Cancer*. 2005;103(7):1494–1502.
42. Trédan O, Geay JF, Touzet S, et al; Groupe d'Investigateurs Nationaux pour l'Etude des Cancers Ovariens. Carboplatin/cyclophosphamide or carboplatin/paclitaxel in elderly patients with advanced ovarian cancer? Analysis of two consecutive trials from the Groupe d'Investigateurs Nationaux pour l'Etude des Cancers Ovariens. *Ann Oncol*. 2007;18(2):256–262.
43. Tison P. [Structured interview guide for evaluating depression in elderly patients, adapted from DSM IV and the GDS, HDRS and MADRS scales]. *Encephale*. 2000;26(3):33–43. French.
44. Katz S, Akpom CA. A measure of primary sociobiological functions. *Int J Health Serv*. 1976;6(3):493–508.
45. Lawton MP, Brody EM. Assessment of older people: self-maintaining and instrumental activities of daily living. *Gerontologist*. 1969;9(3):179–186.
46. Katz S, Ford AB, Moskowitz RW, Jackson BA, Jaffe MW. Studies of illness in the aged. The Index of ADL: a standardized measure of biological and psychosocial function. *JAMA*. 1963;185:914–919.
47. Cohen ME, Marino RJ. The tools of disability outcomes research functional status measures. *Arch Phys Med Rehabil*. 2000;81(12 Suppl 2):S21–S29.
48. Donovan KA, Jacobsen PB. Progress in the implementation of NCCN guidelines for distress management by member institutions. *J Natl Compr Canc Netw*. 2013;11(2):223–226.
49. Holland JC, Bultz BD; National Comprehensive Cancer Network (NCCN). The NCCN guideline for distress management: a case for making distress the sixth vital sign. *J Natl Compr Canc Netw*. 2007;5(1):3–7.
50. Jacobsen PB, Ransom S. Implementation of NCCN distress management guidelines by member institutions. *J Natl Compr Canc Netw*. 2007;5(1):99–103.
51. Stukenberg KW, Dura JR, Kiecolt-Glaser JK. Depression screening scale validation in an elderly, community-dwelling population. *Psychol Assess*. 1990;2(2):134–138.
52. Jadoon NA, Munir W, Shahzad MA, Choudhry ZS. Assessment of depression and anxiety in adult cancer outpatients: a cross-sectional study. *BMC Cancer*. 2010;10:594.
53. Mackenzie LJ, Carey ML, Sanson-Fisher RW, D'Este CA, Paul CL, Yoong SL. Agreement between HADS classifications and single-item screening questions for anxiety and depression: a cross-sectional survey of cancer patients. *Ann Oncol*. 2014;25(4):889–895.
54. Colasanti V, Marianetti M, Micacchi F, Amabile GA, Mina C. Tests for the evaluation of depression in the elderly: a systematic review. *Arch Gerontol Geriatr*. 2010;50(2):227–230.
55. Freyer G, Geay JF, Touzet S, et al. Comprehensive geriatric assessment predicts tolerance to chemotherapy and survival in elderly patients with advanced ovarian carcinoma: a GINECO study. *Ann Oncol*. 2005;16(11):1795–1800.
56. Cosco TD, Doyle F, Ward M, McGee H. Latent structure of the Hospital Anxiety And Depression Scale: a 10-year systematic review. *J Psychosom Res*. 2012;72(3):180–184.
57. Coyne JC, van Sonderen E. No further research needed: abandoning the Hospital and Anxiety Depression Scale (HADS). *J Psychosom Res*. 2012;72(3):173–174.
58. Doyle F, Cosco T, Conroy R. Why the HADS is still important: reply to Coyne and van Sonderen. *J Psychosom Res*. 2012;73(1):77–78.
59. Ibbotson T, Maguire P, Selby P, Priestman T, Wallace L. Screening for anxiety and depression in cancer patients: the effects of disease and treatment. *Eur J Cancer*. 1994;30A(1):37–40.

Supplementary material

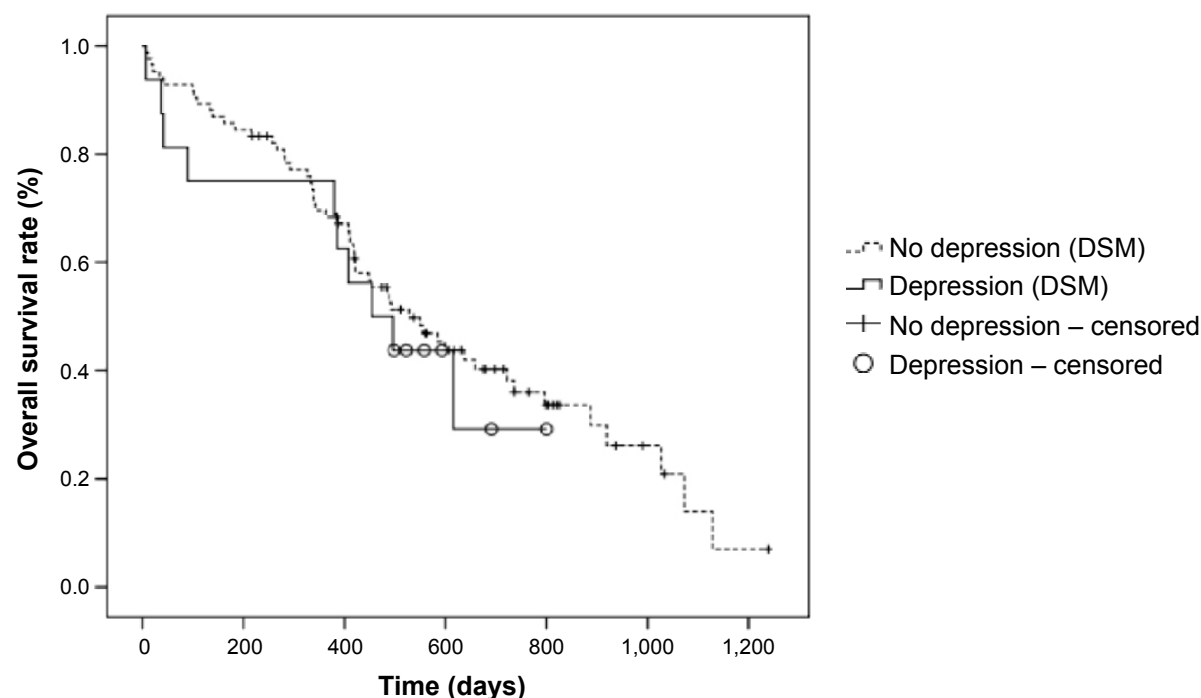


Figure S1 Kaplan–Meier estimates of overall survival according to the presence or the absence of depression using DSM criteria.

Abbreviation: DSM, *Diagnostic and Statistical Manual of Mental Disorders, fourth edition, revised*.

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