# **PALLIATIVE & SUPPORTIVE CARE**

**VOLUME 7, 2009** 

Author and Title Index Volume 7 Contents



## **Palliative & Supportive Care**

#### **Editor-in-Chief**

William Breitbart, M.D. Memorial Sloan-Kettering Cancer Center Department of Psychiatry & Behavioral Sciences 641 Lexington Avenue New York, NY 10022 Phone: (646) 888-0020 Fax: (212) 888-2356 E-mail: breitbaw@mskcc.org

#### **Managing Editor**

Donna Cassetta, M.F.A. Memorial Sloan-Kettering Cancer Center Department of Psychiatry & Behavioral Sciences 641 Lexington Avenue New York, NY 10022 Phone: (646) 888-0020 Fax: (212) 888-2356 E-mail: cassettd@mskcc.org

#### **Co-Editors**

Harvey Max Chochinov, M.D., Ph.D., F.R.C.P.C. University of Manitoba CancerCare Department of Psychiatry **McDermot** Avenue Winnipeg, Manitoba RSE 0V9, Canada Phone: (204) 787-4933 Fax: (204) 787-4937 E-mail: harvey.chochinov@cancercare.mb.ca

Walter Baile, M.D., MD Anderson Cancer Center Sandra L. Bertman, Ph.D., Mount Ida College

Susan Block, M.D., Dana-Farber Cancer Institute

University Robert Brescia, Calvary Hospital

Gian Domenico Borasio, M.D., Ludwig Maximilians

Eduardo Bruera, M.D., MD Anderson Cancer Center

Nathan I. Cherny, M.B.B.S., F.R.A.C.P., Shaare Zedek

Ira Byock, M.D., Robert Wood Johnson Foundation Edwin Cassem, M.D., Harvard Medical School

Albert Diefenbacher, M.D., Humboldt University

Kathleen Foley, M.D., Memorial Sloan-Kettering

Francisco Gil, Ph.D., Institut Català d'Oncologia

Paul Glare, M.D., Memorial Sloan-Kettering Cancer Center

Linda Ganzini, M.D., Oregon Health Sciences University

Sylvie Dolbeault, M.D., Institut Curie Derek Doyle, O.B.E., National Council for Hospice

David Cella, Ph.D., Northwestern University

Medical Center Nessa Coyle, R.N., Ph.D., N.P., Memorial Sloan-Kettering Cancer Center

and Specialist Palliative Care Services

Cancer Center

Simon Wein, M.D Palliative Care Unit **Davidoff Cancer Centre Rabin Medical Centre** 39 Jabotinski Street, Petach Tikvah Israel 49100 Phone: +972-3-9377216 Fax: +972-3-9240398

#### **Editorial Board**

- Luigi Grassi, M.D., University of Ferrara Irene Higginson, B.M., B.S., F.F.P.H.M., Ph.D., King's College London
- Jimmie Holland, M.D., Memorial Sloan-Kettering Cancer Center
- Paul Jacobsen, Ph.D., H. Lee Moffitt Cancer Center Stein Kaasa, M.D., Norwegian University of Science and Technology
- Virgilio Kasprzykowski, M.D., Instituto Portugues de Oncologia

Brian Kelly, M.D., Ph.D., University of Newcastle David Kissane, M.B., B.S., M.P.M., M.D., Memorial Sloan-Kettering Cancer Center

Uwe Koch, M.D., Ph.D., University Hospital at Hamburg-Eppendorf

Linda Kristjansen, Ph.D., Curtin University of Technology

Peter Lawlor, M.D., St. Francis Hospice Matthew Loscalzo, M.S.W., City of Hope Paola Luzzatto, PhD., A.T.R.-B.C., Art Therapy Italiana

Neil MacDonald, C.M., M.D., F.R.C.P.(C), F.R.C.P. (EDIN), McGill University Tatsuya Morita, M.E., D.O., Serei Hospice

Balfour Mount, M.D., McGill University Steven Passik, Ph.D., Memorial Sloan-Kettering Cancer Center

Russell Portenoy, M.D., Beth Israel Medical Center Christina M. Puchalski, M.D., George Washington Institute for Spirituality and Health

Barry Rosenfeld, Ph.D., Fordham University Carla Ripamonti, M.D., National Cancer Institute Tone Rustoen, R.N., Ph.D., Oslo University College John L. Shuster, Jr., M.D., University of Alabama

School of Medicine Daniel P. Sulhasy, O.F.M., M.D., Ph.D., University of Chicago

Frederic Stiefel, M.D., University of Lausanne James A. Tulsky, M.D., Durham VA Medical Center Yosuke Uchitomi, M.D., Ph.D., National Cancer Center Research Institute East

Frits S. A. M. van Dam, M.D., Ph.D., The Netherlands

Cancer Institute Maggie Watson, Ph.D., The Royal Marsden Hospital Roberto Wenk, M.D., Fundacion FEMEBA

Palliative & Supportive Care (ISSN: 1478-9515) is a quarterly international journal of palliative medicine that focuses on the psychiatric, psychosocial, spiritual, existential, ethical, and philosophical aspects of palliative care. The journal's aim is to serve as an educational resource for practitioners from a wide array of disciplines engaged in the delivery of care to those with life threatening illnesses along the entire continuum of care from diagnosis to the end of life.

Copyright © Cambridge University Press, 2009. All rights reserved. No part of this publication may be reproduced, in any form or by any means, electronic, photocopying, or otherwise, without permission in writing from Cambridge University Press. For further information see http://us.cambridge.org/formation/rights/ or http://www.cambridge.org/uk/information/rights/. Authorization to photocopy items for internal or personal use, or the internal or personal use of specific clients, is granted by Cambridge University Press, provided that the appropriate fee is paid directly to Copyright Clearance Center, 222 Rosewood Drive, Danvers, MA 01923, USA (Telephone: (508) 750-8400), stating the ISSN (1478-9515/09 \$20.00), the volume, and the first and last page numbers of each article copied. The copyright owner's consent does not include copying for general distribution, promotion, new works, or resale.

2010 Subscription Rates: Palliative & Supportive Care is published in March, June, September, and December by Cambridge University Press, 32 Avenue of the Americas, New York, NY 10013-2473, USA / Cambridge University Press, The Edinburgh Building, Shaffesbury Road, Cambridge CB2 8RU, UK. Institutions print and electronic: US \$444.00 in the USA, Canada, and Mexico; UK £255.00 + VAT elsewhere. Institutions electronic only: US \$369.00 in the USA, Canada, and Mexico; UK \$210.00 + VAT elsewhere. Institutions print only: US \$411.00 in the USA, Canada, and Mexico; UK £237.00 + VAT elsewhere. Individuals print and electronic only: US \$198.00 in the USA, Canada, and Mexico; UK £109.00 + VAT elsewhere. Individuals print only: US \$198.00 in the USA, Canada, and Mexico; UK £52.00 + VAT elsewhere. Subscribers in the USA, Canada, and Mexico should send their orders, with payment in US dollars or the equivalent value in Canadian dollars to: Cambridge University Press, Journals Customer Services Department, 100 Brook Hill Drive, West Nyack, NY 10994-2133, USA. Telephone: (845) 353-7500, Fax: (845) 353-4141. Orders may be phoned direct (toll free): (800) 872-7423. E-mail: subscriptions\_newyork@cambridge.org. Subscribers elsewhere should send their orders with payment in sterling to: Journals Customer Services Department Cambridge University Press, The Edinburgh Building, Shaftesbury Road, Cambridge, CB2 8RU, United Kingdom. Telephone: +44 (0) 122 332 5150. E-mail: journals@cambridge.org.

Abstracted and indexed in the following: CINAHL database; Cumulative Index to Nursing and Allied Health Liteature; MEDLINE

Change of address: Allow six weeks for all changes to become effective. All communication should include both old and new addresses (with postal codes) and should be accompanied by a mailing label from a recent issue.

Editorial Office: William Breitbart, Editor-in-Chief, Memorial Sloan-Kettering Cancer Center, Department of Psychiatry and Behavioral Sciences, 641 Lexington Avenue, New York, NY 10022, USA. Telephone: (646) 888-0020. Fax: (212) 888-2356. E-mail: breitbaw@mskcc.org.

Office of Publication: Cambridge University Press, 32 Avenue of the Americas, New York, NY 10013-2473, USA: Telephone: (212) 337-5000. Fax: (212) 337-5959.

Printed in the United States of America on acid-free paper.

Application to mail at periodicals rate is pending in New York, NY, and additional mailing offices. Postmaster: send address changes in the USA and Canada to Palliative & Supportive Care, Subscription Department, Cambridge University Press, 100 Brook Hill Drive, West Nyack, NY 10994-2133.

# **Title and Author Index**

- A call for simple and rational palliative medicine psychotropic prescribing guidelines: Response to Barnhorst et al. (Palliative & Supportive Care, Volume 6, Number 1, 2008), Scott Irwin, 3
- A feasibility study of a two-session home-based cognitive behavioral therapy-insomnia intervention for bereaved family caregivers, Patricia A. Carter, Sabrina Q. Mikan, and Cherie Simpson, 197

Abboud, Sarah, 339

- Activity and meaning making in the everyday lives of people with advanced cancer, Karen la Cour, Helle Johannessen, and Staffan Josephsson, 469
- Adjusting to pancreatic cancer: Perspectives from first-degree relatives, Kerri Petrin, Deborah J. Bowen, Catherine M. Alfano, and Robin Bennett, 281
- Adolescents' experiences of a parent's serious illness and death, Lena Dehlin and Lena Mårtensson, 13 Aging, Bonnie Raingruber, 383

Aging, bonnie Kamgruber

- Ahles, Tim A., 75
- Ahn, Sung-Hee, 349
- Akechi, Tatsuo, 229
- Alexander, Stewart C., 000
- Alfano, Catherine M., 281
- Allan, Diane E., 435
- Ando, Michiyo, 349
- André, Beate, 57
- Applying feminist, multicultural, and social justice theory to diverse women who function as caregivers in end-of-life and palliative home care, Christopher J. MacKinnon, 501

Arnaert, Antonia, 357

Arnold, Robert, 143

Associations with worry about dying and hopelessness in ambulatory ovarian cancer patients, Eileen Huh Shinn, Cindy L. Carmack Taylor, Kelly Kilgore, Alan Valentine, Diane C. Bodurka, John Kavanagh, Anil Sood, Yisheng Li, and Karen Basen-Engquist, 299

Åström, Sture, 87 Astrow, Alan, 315 Attitudes of Quebec doctors toward sedation at the end of life: An exploratory study, Danielle Blondeau, Serge Dumont, Louis Roy, and Isabelle Martineau, 331

Bakitas, Marie, 75 Balan, Stefan, 75 Barclay, Stephen I. G., 289 Barnett, Kathleen N., 75 Basen-Engquist, Karen, 299 Bates, Ursula, 323 Bennett, Robin, 281 Blondeau, Danielle, 331 Blooming, Bonnie Raingruber, 381 Bodurka, Diane C., 299 Boettger, Soenke, 50 Bowen, Deborah J., 281 Breitbart, William, 50, 139, 269 Brokaw, Frances C., 75 Brown, Richard F., 171 Brufsky, Adam, 143 Bylund, Carma L., 171 Byock, Ira R., 40, 75

Can short hospice enrollment be long enough? Comparing the perspectives of hospice professionals and family caregivers, Deborah P. Waldrop and Elaine S. Rinfrette, 37

Cancer patients' reluctance to discuss psychological distress with their physicians was not associated with underrecognition of depression by physicians: A preliminary study, Toru Okuyama, Chiharu Endo, Takashi Seto, Masashi Kato, Nobuhiko Seki, Tatsuo Akechi, Toshiaki A. Furukawa, Kenji Eguchi, and Takashi Hosaka, 229

Caring for a person in advanced illness and suffering from breathlessness at home: Threats and resources, Marjolein H. Gysels and Irene J. Higginson, 153

"Caring as if it were my family": Health care aides' perspectives about expert care of the dying

- Carlson, Linda E., 49
- Carlsson, Maria E., 207
- Carmack Taylor, Cindy L., 299
- Carter, Patricia A., 197
- Challenges of illness in metastatic breast cancer: A low-income African American perspective, Margaret Quinn Rosenzweig, Theresa Wiehagen, Adam Brufsky, and Robert Arnold, 143
- Chan, Lisa S., 235 Changes, Bonnie Raingruber, 531
- Childhood abuse uncovered in a palliative care audit,
- Catriona Macpherson, 481
- Clawson, Lora, 315
- Complications, Paul Rousseau, 379
- Conducting family meetings in palliative care: Themes, techniques, and preliminary evaluation of a communication skills module, Jennifer A. Gueguen, Carma L. Bylund, Richard F. Brown, Tomer T. Levin, and David W. Kissane, 171 Cooper, Dan, 365 Copel, Laure, 27
- Cypress Forests, Bonnie Raingruber, 385
- Dehlin, Lena, 13
- Delirium superimposed on dementia versus delirium in the absence of dementia: Phenomenologic differences, Soenke Boettger, Steven Passik, and William Breitbart, 495
- Development and validation of the Family Decision-Making Self-Efficacy Scale, Marie T. Nolan, Mark T. Hughes, Joan Kub, Peter B. Terry, Alan Astrow, Richard E. Thompson, Lora Clawson, Kenneth Texeira, and Daniel P. Sulmasy, 315
- Dimassi, Hani, 339
- Dooley, Barbara, 323
- Dossey, Barbara M., 405
- Doyle Lyons, Kathleen, 75
- Duggleby, Wendy, 365
- Dumont, Isabelle, 163
- Dumont, Serge, 331
- Effect of music therapy on oncologic staff bystanders: A substantive grounded theory, Clare O'Callaghan and Lucanne Magill, 218 Eguchi, Kenji, 229 Endo, Chiharu, 229 Expertise, Bonnie Raingruber, 527
- Farquhar, Morag C., 289 Fatigue in relatives of palliative patients, Maria E. Carlsson, 207 Flahault, Cecile 27

Fricchione, Gregory L., 213 Funk, Laura M., 435 Furukawa, Toshiaki A., 229 Garland, Sheila N., 49 Gaudron, Cecile, 253 George, Linda K., 40 Gold, Michelle, 181 Grande, Gunn E., 289 Granet, Roger, 259, 261 Greaves, Judi, 245 Greenstein, Mindy, 271 Gueguen, Jennifer A., 171 Gysels, Marjolein H., 153 Håvard Loge, Jon, 459 Hagen, Neil A., 415 Hagino, Masashi, 307 Halifax, Joan, 404 Hamel-Bissell, Brenda P., 65 Hegel, Mark T., 75 Heller, Karen S., 405 Hendrickson, Karrie, 109 Higginson, Irene J., 153 Holland, Jimmie, 271 Hope in palliative care: An integrative review, Jari Kylmä, Wendy Duggleby, Dan Cooper, and Gustaf Molander, 365 Hosaka, Takashi, 229 Hughes, Mark T., 315 Huijer, Huda Abu-Saad, 339 Hull, Jay G., 75

- Humor and death: A qualitative study of *The New Yorker* cartoons (1986–2006), Marianne Matzo and David Miller, 487
- Ide, Saburo, 349
- Illuminating the path: What literature can teach doctors about death and dying, Andrea Killick, 521
- Impact of a contemplative end-of-life training program: Being with dying, Cynda Hylton Rushton, Deborah E. Sellers, Karen S. Heller, Beverly Spring, Barbara M. Dossey, and Joan Halifax, 405
- Implementation of computerized technology in a palliative care unit, Beate André, Gerd I. Ringdal, Jon H. Loge, Toril Rannestad, and Stein Kaasa, 57
- Individual brief art therapy can be helpful for women with breast cancer: A randomized controlled clinical study, Karin Egberg Thyme, Eva C. Sundin, Britt Wiberg, Inger Öster, Sture Åström, and Jack Lindh, 87
- International comparison study on the primary concerns of, terminally ill cancer patients in shortterm life review interviews among Japanese,

Koreans, and Americans, Michiyo Ando, Tatsuya Morita, Sung-Hee Ahn, Felicia Marquez-Wong, and Saburo Ide, 349 Irwin, Scott 3 Is House, M.D., always right?, Simon Wein, 1 Iwamitsu, Yumi, 307 Jerking Awake, Bonnie Raingruber, 429 Johannessen, Helle, 496 Josephsson, Staffan, 496 Kaasa, Stein, 459, 57 Kato, Masashi, 229 Kaufman, Hannah, 517 Kavanagh, John, 299 Kilgore, Kelly, 299 Killick, Andrea, 521 King, Nigel, 7 Kissane, David W., 171 Kissane, David, 163 Klaasen, Kathleen, 449 Komesaroff, Paul, 181 Kub, Joan, 315 Kylmä, Jari, 365 la Cour, Karen, 47 Laurence, Valerie, 27 Leavings, Rita Sherman, 257 Lee, Susan, 245 Levin, Tomer T., 171 Li, Yisheng, 299 Li, Zhongze, 75 Lightening up before death, A. D. (Sandy) Macleod, 513 Lindh, Jack, 87 Loge, Jon H., 459 Mårtensson, Lena, 13 MacKinnon, Christopher J., 501 Macleod, A. D. (Sandy), 513 Macpherson, Catriona, 48 Magill, Lucanne, 97, 218 Marquez-Wong, Felicia, 349 Marr, Hubert, 49 Martin, Anne, 271 Martineau, Isabelle, 331 Masaru, Kuranami, 307 Matzo, Marianne, 487 McCahill, Laurence, 65 McClement, Susan, 449 McKinstry, Elizabeth, 75 Mehta, Anita, 235 Mikan, Sabrina Q., 197 Miller, David, 487 Miyaoka, Hitoshi, 307 Molander, Gustaf, 365

Montel, Sebastien, 27 Morbidity, mortality, and parental grief: A review of the literature on the relationship between the death of a child and the subsequent health of parents, Karrie Hendrickson, 109 Morita, Tatsuya, 349 Mullane, Mary, 323 Mysak, Marlene, 415 Nelson, Christian, 271 Nolan, Marie T., 315 O'Callaghan, Clare, 218 O'Connor, Margaret, 245 Okazaki, Shigemi, 307 Okuyama, Toru, 229 Oldervoll, Line, 459 One hundred meters, Simon Wein, 391 Öster, Inger, 87 Pacquement, Helene, 27 Paddison, Johanna S., 213 Palliative care: A need for a family systems approach, Anita Mehta, S. Robin Cohen, and Lisa S. Chan, 235 Palliative family caregivers' accounts of health care experiences: The importance of "security", Laura M. Funk, Diane E. Allan, and Kelli I. Stajduhar, 435 Paltiel, Hanne, 459 Passik, Steven, 495 Patients' views on decision making in advanced cancer, Jennifer Philip, Michelle Gold, Max Schwarz, and Paul Komesaroff, 181 Penel, Nicolas, 253 Personal reflections on love, death and suffering: My narrative of the final days of my husband's life, Claire Turenne Sjolander, 121 Perspectives on palliative care in Lebanon: Knowledge, attitudes, and practices of medical and nursing specialties, Huda Abu-Saad Huijer, Hani Dimassi, and Sarah Abboud, 339 Petrin, Kerri, 281 Philip, Jennifer, 181 Pirl, William F., 213 Place of death of adolescents and young adults with cancer: First study in a French population, Sebastien Montel, Valerie Laurence, Laure Copel, Helene Pacquement, and Cecile Flahault, 27 Poppito, Shannon, 271 Providing care and sharing expertise: Reflections of nurse-specialists in palliative home care, Antonia Arnaert and Megan Wainwright, 357

Quality of life measures (EORTC QLQ-C30 and SF-36) as predictors of survival in palliative colorectal and lung cancer patients, Gunn E. Grande, Morag C. Farquhar, Stephen I. G. Barclay, and Christopher J. Todd, 289

Raingruber, Bonnie, 531, 529, 527, 381 383, 385 Rannestad, Toril, 57

Reappraisal in the eighth life cycle stage:A theoretical psychoeducational intervention in elderly patients with cancer, Jimmie Holland, Shannon Poppito, Christian Nelson, Talia Weiss, Mindy Greenstein, Anne Martin, Phoutthasone Thirakul, and Andrew Roth, 271

Recruitment and retention of palliative cancer patients and their partners participating in a longitudinal evaluation of a psychosocial retreat program, Sheila N. Garland, Linda E. Carlson, Hubert Marr, and Steve Simpson, 49

Reich, Michel, 253

Rinfrette, Elaine S., 37

Ringdal, Gerd I., 57

Robin Cohen, S., 235

Rodriguez, Alison, 7

Rosenzweig, Margaret Quinn, 143

Roth, Andrew, 271

Rousseau, Paul, 379

Roy, Louis, 331 Rushton, Cynda Hylton, 405

Schwarz, Max, 181

Seki, Nobuhiko, 229

Sellers, Deborah E., 405

Seriously ill patients' discussions of preparation and life completion: An intervention to assist with transition at the end of life, Karen E. Steinhauser, Stewart C. Alexander, Ira R. Byock, Linda K. George, and James A. Tulsky, 393
Seto, Takashi, 229
Seville, Janette L., 75
Sherman, Rita, 257
Shimoinaba, Kaori, 245
Shinn, Eileen Huh, 299
Simpson, Cherie, 197
Simpson, Steve, 49
Sinclair, Shane, 415
Sjolander, Claire Turenne, 121

Solvoll, Edel, 459

Sood, Anil, 299 Spring, Beverly, 405

Charles in the second s

Staff grief and support systems for Japanese health care professionals working in palliative care, Kaori Shimoinaba, Margaret O'Connor, Susan

Lee, and Judi Greaves, 245

Stajduhar, Kelli I., 435

Steinhauser, Karen E., 393

Stories, Roger Granet, 259
Strength through adversity: Bereaved cancer carers' accounts of rewards and personal growth from caring, W. K. Tim Wong and Jane Ussher, 187
Sulmasy, Daniel P., 315
Sundin, Eva C., 87
Sutherland, Nisha, 423

Suzuki, Shimako, 307

Techniques for framing questions in conducting family meetings in palliative care, Isabelle Dumont and David Kissane, 163

Temel, Jennifer S., 213

Terry, Peter B., 315

Texeira, Kenneth, 315

The Chair, Roger Granet, 261

The Congress of Vienna, William Breitbart, 269

"The healthy me appears": Palliative cancer patients' experiences of participation in a physical group exercise program, Hanne Paltiel, Edel Solvoll, Jon Håvard Loge, Stein Kaasa, and Line Oldervoll, 459

The lived experience of parenting a child with a life-limiting condition: A focus on the mental health realm, Alison Rodriguez and Nigel King, 7

The meaning of being in transition to end-of-life care for female partners of spouses with cancer, Nisha Sutherland, 423

The patient lived experience for surgical treatment of colorectal liver metastases: A phenomenological study, Laurence McCahill and Brenda P. Hamel-Bissell, 65

The project ENABLE II randomized controlled trial to improve palliative care for rural patients with advanced cancer: Baseline findings, methodological challenges, and solutions, Marie Bakitas, Kathleen Doyle Lyons, Mark T. Hegel, Stefan Balan, Kathleen N. Barnett, Frances C. Brokaw, Ira R. Byock, Jay G. Hull, Zhongze Li, Elizabeth McKinstry, Janette L. Seville, and Tim A. Ahles, 75

The psychological responses of outpatient breast cancer patients before and during first medical consultation, Shigemi Okazaki, Yumi Iwamitsu, Kuranami Masaru, Keiko Todoroki, Shimako Suzuki, Kenji Yamamoto, Masashi Hagino, Masahiko Watanabe, and Hitoshi Miyaoka, 307

The spiritual domain of palliative care: Who should be "spiritual care professionals"?, William Breitbart, 139

The spiritual meaning of pre-loss music therapy to bereaved caregivers of advanced cancer patients, Lucanne Magill, 97 Thirakul, Phoutthasone, 271
This won't hurt a bit: The ethics of promising pain relief, Hannah Kaufman, 517
Thompson, Richard E., 315
Thyme, Karin Egberg, 87
Tiernan, Eoin, 323
Tim Wong, W. K., 187
Todd, Christopher J., 289
Todoroki, Keiko, 307
Tulsky, James A., 40

Using the differential from complete blood counts as a biomarker of fatigue in advanced non-small-cell lung cancer: An exploratory analysis, Johanna S. Paddison, Jennifer S. Temel, Gregory L. Fricchione, and William F. Pirl, 213 Ussher, Jane, 187

Valentine, Alan, 299

Validation of the Demoralization Scale in an Irish advanced cancer sample, Mary Mullane, Barbara Dooley, Eoin Tiernan, and Ursula Bates, 323 Wainwright, Megan, 357 Waldrop, Deborah P., 37 Watanabe, Masahiko, 307 Wein, Simon, 1, 391 Weiss, Talia, 271 What are the core elements of oncology spiritual care programs?, Shane Sinclair, Marlene Mysak, and Neil A. Hagen, 415 When cancerophobia and denial lead to death, Michel Reich, Cecile Gaudron, and Nicolas Penel, 253 Wiberg, Britt, 87 Wiehagen, Theresa, 143 Wowchuk, Suzanne, 449

Yamamoto, Kenji, 307

# **PALLIATIVE & SUPPORTIVE CARE**

March 2009

Number 1

		CONTENTS
		From the Editor
Simon Wein	1	Is House, M.D., always right?
		Letter to the Editor
Scott Irwin	3	A call for simple and rational palliative medicine psychotropic prescribing guidelines: Response to Barnhorst et al. (Palliative & Supportive Care, Volume 6, Number 1, 2008)
		Original Articles
Alison Rodriguez and Nigel King	7	The lived experience of parenting a child with a life- limiting condition: A focus on the mental health realm
Lena Dehlin and Lena Mårtensson	13	Adolescents' experiences of a parent's serious illness and death
Sebastien Montel, Valerie Laurence, Laure Copel, Helene Pacquement, and Cecile Flahault	27	Place of death of adolescents and young adults with cancer: First study in a French population
DEBORAH P. WALDROP AND ELAINE S. Rinfrette	37	Can short hospice enrollment be long enough? Comparing the perspectives of hospice professionals and family caregivers
Sheila N. Garland, Linda E. Carlson, Hubert Marr, and Steve Simpson	49	Recruitment and retention of palliative cancer patients and their partners participating in a longitudinal evaluation of a psychosocial retreat program
Beate André, Gerd I. Ringdal, Jon H. Loge, Toril Rannestad, and Stein Kaasa	57	Implementation of computerized technology in a palliative care unit
LAURENCE MCCAHILL AND BRENDA P. HAMEL-BISSELL	65	The patient lived experience for surgical treatment of colorectal liver metastases: A phenomenological study
Marie Bakitas, Kathleen Doyle Lyons, Mark T. Hegel, Stefan Balan, Kathleen N. Barnett, Frances C. Brokaw, Ira R. Byock, Jay G. Hull, Zhongze Li, Elizabeth McKinstry, Janette L. Seville, and Tim A. Ahles	75	The project ENABLE II randomized controlled trial to improve palliative care for rural patients with advanced cancer: Baseline findings,methodological challenges, and solutions
Karin Egberg Thyme, Eva C. Sundin, Britt Wiberg, Inger Öster, Sture	87	Individual brief art therapy can be helpful for women with breast cancer: A randomized controlled clinical study

ÅSTRÖM, AND JACK LINDH

Volume 7

Lucanne Magill	97	The spiritual meaning of pre-loss music therapy to bereaved caregivers of advanced cancer patients
		Review Article
KARRIE HENDRICKSON	109	Morbidity, mortality, and parental grief: A review of the literature on the relationship between the death of a child and the subsequent health of parents
		Essay/Personal Reflections
Claire Turenne Sjolander	121	Personal reflections on love, death and suffering: My narrative of the final days of my husband's life
	133	Calendar of Events
	133 135	

Volume 7

# June 2009

# Number 2

## **CONTENTS**

## **From the Editor**

William Breitbart	139	The spiritual domain of palliative care: Who should be "spiritual care professionals"?
		Original Articles
Margaret Quinn Rosenzweig, Theresa Wiehagen, Adam Brufsky, and Robert Arnold	143	Challenges of illness in metastatic breast cancer: A low-income African American perspective
Marjolein H. Gysels and Irene J. Higginson	153	Caring for a person in advanced illness and suffering from breathlessness at home: Threats and resources
ISABELLE DUMONT AND DAVID KISSANE	163	Techniques for framing questions in conducting family meetings in palliative care
Jennifer A. Gueguen, Carma L. Bylund, Richard F. Brown, Tomer T. Levin, and David W. Kissane	171	Conducting family meetings in palliative care: Themes, techniques, and preliminary evaluation of a communication skills module
Jennifer Philip, Michelle Gold, Max Schwarz, and Paul Komesaroff	181	Patients' views on decision making in advanced cancer
W.K. TIM WONG AND JANE USSHER	187	Strength through adversity: Bereaved cancer carers' accounts of rewards and personal growth from caring
PATRICIA A. CARTER, SABRINA Q. MIKAN, AND CHERIE SIMPSON	197	A feasibility study of a two-session home-based cognitive behavioral therapy–insomnia intervention for bereaved family caregivers
Maria E. Carlsson	207	Fatigue in relatives of palliative patients
Johanna S. Paddison, Jennifer S. Temel, Gregory L. Fricchione, and William F. Pirl	213	Using the differential from complete blood counts as a biomarker of fatigue in advanced non-small-cell lung cancer: An exploratory analysis
1017/51472051500001052 Published online by Combridge Unive	rcity Brocc	

https://doi.org/10.1017/S1478951509991052 Published online by Cambridge University Press

Clare O'Callaghan and Lucanne Magill	218	Effect of music therapy on oncologic staff bystanders: A substantive grounded theory
Toru Okuyama, Chiharu Endo, Takashi Seto, Masashi Kato, Nobuhiko Seki, Tatsuo Akechi, Toshiaki A. Furukawa, Kenji Eguchi, and Takashi Hosaka	229	Cancer patients' reluctance to discuss psychological distress with their physicians was not associated with underrecognition of depression by physicians: A preliminary study
		Review Articles
Anita Mehta, S. Robin Cohen, and Lisa S. Chan	235	Palliative care: A need for a family systems approach
Kaori Shimoinaba, Margaret O'Connor, Susan Lee, and Judi Greaves	245	Staff grief and support systems for Japanese health care professionals working in palliative care
		Case Report
Michel Reich, Cecile Gaudron, and Nicolas Penel	253	When cancerophobia and denial lead to death
		Poetry/Fiction
Rita Sherman	257	Leavings
Roger Granet	259	Stories
Roger Granet	261	The Chair
	263	Calendar of Events
	265	Instructions for Contributors

τ7	1			
1/1	$\mathbf{n}$	11	m	. /
- V 1		u		

## September 2009

Number 3

### CONTENTS

cancer

#### **From the Editor**

WILLIAM BREITBART **269** The Congress of Vienna

#### **Original Articles**

first-degree relatives

Reappraisal in the eighth life cycle stage: A theoretical

psychoeducational intervention in elderly patients with

Adjusting to pancreatic cancer: Perspectives from

- JIMMIE HOLLAND, SHANNON POPPITO, CHRISTIAN NELSON, TALIA WEISS, MINDY GREENSTEIN, ANNE MARTIN, PHOUTTHASONE THIRAKUL, AND ANDREW ROTH
  - Kerri Petrin, Deborah J. Bowen, 281 Catherine M. Alfano, and Robin Bennett
- GUNN E. GRANDE, MORAG C. FARQUHAR,<br/>STEPHEN I. G. BARCLAY, AND<br/>CHRISTOPHER J. TODD289Quality of life measures (EORTC QLQ-C30 and SF-36) as<br/>predictors of survival in palliative colorectal and lung<br/>cancer patients

EILEEN HUH SHINN, CINDY L. CARMACK	299
TAYLOR, KELLY KILGORE, ALAN	
VALENTINE, DIANE C. BODURKA, JOHN	
KAVANAGH, ANIL SOOD, YISHENG LI, AND	
KAREN BASEN-ENGQUIST	

307

315

Shigemi Okazaki, Yumi Iwamitsu, Kuranami Masaru, Keiko Todoroki, Shimako Suzuki, Kenji Yamamoto, Masashi Hagino, Masahiko Watanabe, and Hitoshi Miyaoka

MARIE T. NOLAN, MARK T. HUGHES, JOAN KUB, PETER B. TERRY, ALAN ASTROW, RICHARD E. THOMPSON, LORA CLAWSON, KENNETH TEXEIRA, AND DANIEL P. SULMASY

- Mary Mullane, Barbara Dooley, Eoin 323 Tiernan, and Ursula Bates
  - DANIELLE BLONDEAU, SERGE DUMONT, 331 LOUIS ROY, AND ISABELLE MARTINEAU
- Huda Abu-Saad Huijer, Hani Dimassi, 339 and Sarah Abboud
- Michiyo Ando, Tatsuya Morita, Sung-Hee Ahn, Felicia Marquez-Wong and Saburo Ide

JARI KYLMÄ, WENDY DUGGLEBY, DAN

COOPER, AND GUSTAF MOLANDER

- Associations with worry about dying and hopelessness in ambulatory ovarian cancer patients
- The psychological responses of outpatient breast cancer patients before and during first medical consultation
- Development and validation of the Family Decision-Making Self-Efficacy Scale
- Validation of the Demoralization Scale in an Irish advanced cancer sample

Attitudes of Quebec doctors toward sedation at the end of life: An exploratory study

- Perspectives on palliative care in Lebanon: Knowledge, attitudes, and practices of medical and nursing specialties
- International comparison study on the primary concerns of, terminally ill cancer patients in short-term life review interviews among Japanese, Koreans, and Americans

#### **Review Articles**

- ANTONIA ARNAERT AND MEGAN WAINWRIGHT 357 Providing care and sharing expertise: Reflections of nursespecialists in palliative home care
  - **365** Hope in palliative care: An integrative review

#### **Essay/Personal Reflections**

PAUL ROUSSEAU 379 Complications

# Poetry/Fiction

BONNIE RAINGRUBER381BloomingBONNIE RAINGRUBER383AgingBONNIE RAINGRUBER385Cypress Forests387Instructions for Contributors

### Number 4

#### **CONTENTS**

#### **From the Editor**

One hundred meters

SIMON WEIN 391

405

#### **Original Articles**

Being with dying

- KAREN E. STEINHAUSER, STEWART C.<br/>ALEXANDER, IRA R. BYOCK, LINDA K.<br/>GEORGE, AND JAMES A. TULSKY**393**Seriously ill patients' discussions of preparation and life<br/>completion: An intervention to assist with transition at the<br/>end of life
- Cynda Hylton Rushton, Deborah E. Sellers, Karen S. Heller, Beverly Spring, Barbara M. Dossey, and Joan Halifax
- Shane Sinclair, Marlene Mysak, and 415 Neil A. Hagen
  - NISHA SUTHERLAND 423
- Laura M. Funk, Diane E. Allan, and 435 Kelli I. Stajduhar
  - SUSAN MCCLEMENT, SUZANNE 449 WOWCHUK, AND KATHLEEN KLAASEN
- HANNE PALTIEL, EDEL SOLVOLL, JON HÅVARD LOGE, STEIN KAASA, AND LINE OLDERVOLL
- KAREN LA COUR, HELLE JOHANNESSEN, 469 AND STAFFAN JOSEPHSSON
  - CATRIONA MACPHERSON 481
  - MARIANNE MATZO AND DAVID MILLER 487
- SOENKE BOETTGER, STEVEN PASSIK, AND 495 WILLIAM BREITBART
  - Christopher J. MacKinnon 501

What are the core elements of oncology spiritual care programs?

Impact of a contemplative end-of-life training program:

- **423** The meaning of being in transition to end-of-life care for female partners of spouses with cancer
  - Palliative family caregivers' accounts of health care experiences: The importance of "security"
  - "Caring as if it were my family": Health care aides' perspectives about expert care of the dying resident in a personal care home
  - "The healthy me appears": Palliative cancer patients' experiences of participation in a physical group exercise program
  - Activity and meaning making in the everyday lives of people with advanced cancer
  - Childhood abuse uncovered in a palliative care audit
  - Humor and death: A qualitative study of *The New Yorker* cartoons (1986–2006)
  - Delirium superimposed on dementia versus delirium in the absence of dementia: Phenomenologic differences

#### **Review Article**

Applying feminist, multicultural, and social justice theory to diverse women who function as caregivers in end-of-life and palliative home care

#### **Case Report**

A. D. (SANDY) MACLEOD 513 Lightening up before death

#### **Essay/Personal Reflections**

Hannah Kaufman 517

This won't hurt a bit: The ethics of promising pain relief

ANDREA KILLICK	<b>521</b>	Illuminating the path: What literature can teach doctors
		about death and dying

## **Poetry/Fiction**

Bonnie Raingruber	527	Expertise
Bonnie Raingruber	529	Jerking Awake
Bonnie Raingruber	<b>531</b> Changes	
	533	Erratum
	537	Title and Author Index
	543	Content to Volume 7
	549	Instructions for Contributors

# **Instructions for Contributors**

AIMS AND SCOPE. Palliative & Supportive Care publishes papers on psychiatric disorders and their management in the palliative care setting, including but not limited to depression, delirium, anxiety, and post-traumatic stress disorder and bereavement. Evolving constructs of particular relevance to the interface of psychiatry/psychology and palliative medicine, such as demoralization, meaning, dignity, hopelessness, will to live, suffering, and developmental growth at the end of life are a major focus of this journal. Research focusing on these issues, including epidemiology, diagnostic screening, assessment, management, and intervention drug and psychotherapy trials, are also addressed. In addition, the journal provides a forum for the discussion of psychosocial and sociocultural matters such as caregiver burden, health care provider burnout, counseling interventions, the impact of psychosocial factors related to intervention of pain and physical symptom control, and communication issues. The journal features full-length research reports and review articles as well as short communications.

**ORIGINALITY AND COPYRIGHT.** To be considered for publication in *Palliative & Supportive Care* a manuscript cannot have been published previously, nor can it be under review for publication elsewhere. Papers with multiple authors are reviewed with the assumption that all authors have approved the submitted manuscript and concur in its submission to *Palliative & Supportive Care*. A Transfer of Copyright Agreement must be executed before an article can be published. Government authors whose articles were created in the course of their employment must so certify in lieu of copyright transfer. Authors are responsible for obtaining written permission from the copyright owners to reprint any previously published material included in their article.

**MANUSCRIPT SUBMISSION AND REVIEW.** An original and three high quality photocopies with the Copyright Transfer Agreement should be submitted to:

Donna Cassetta Managing Editor, Palliative & Supportive Care Memorial Sloan Kettering Cancer Center Department of Psychiatry and Behavioral Sciences, Box 421 1242 Second Avenue New York, NY 10021 Phone: 212-639-3907 Fax: 212-717-3763

#### Contributors may choose to submit their manuscript electronically by sending an email to palliative@mskcc.org.

Each manuscript will normally be reviewed by at least two referees with relevant scientific experience. Authors may suggest appropriate reviewers, but final selection of referees will be made by the Editor. Reviewers are asked to evaluate manuscripts for their scientific merit and clarity of presentation.

**MANUSCRIPT PREPARATION AND STYLE.** Manuscripts must be in English and typed double-spaced on one side only of 8 1/2 x 11" or A4 size good quality paper. Allow margins of at least 1" (20 mm); use a 5-space paragraph indent; do not hyphenate words at the end of lines and do not justify right margins. Minor corrections to the manuscript may be typed or neatly printed in ink; retyping is required for significant changes. Numbers should be spelled out when they occur at the beginning of a sentence; use Arabic numerals elsewhere. Abbreviations should be used sparingly and nonstandard abbreviations should be defined at their first occurrence. Metric system (SI) units should be used. Manuscripts that do not conform to the style of **Palliative & Supportive Care** will be returned without review. Authors of accepted manuscripts will be requested to provide the final text both as hard copy and on diskette. The diskette should be formatted in the latest version of MS Word for Macintosh or IBM compatible computers.

**MANUSCRIPT ELEMENTS AND ORDER.** Unless there are obvious and compelling reasons for variation (e.g. review articles, short communications), manuscripts should be organized as follows:

**Title page.** This is page 1. The title should be concise, informative, and free of abbreviations, chemical formulae, technical jargon, and esoteric terms. This page should include (a) the article's full title, (b) names and affiliations of all authors, (c) the name, mailing address, email address, and telephone number of the corresponding author, (d) the address for reprint requests if different from that of the corresponding author, (e) a short title of 50 characters or less, and (f) a list of the number of manuscript pages, number of tables, and number of figures.

**Abstract and keywords page.** This is page 2 and should include (a) the article's full title, (b) an abstract of no more than 300 words, and (c) up to 5 keywords or phrases that reflect the content and major thrust of the article. The abstract should give a succinct account of the objective, methods, results, and significance of the research.

**Introduction.** This section begins on page 3 and should clearly state the objective of the research in the context of previous work bearing directly on the subject. An extensive review of the literature is not usually appropriate.

**Methods.** This section should be brief but provide sufficient information to permit others to replicate the study. Pertinent details of species, apparatus and equipment, procedures and experimental design should be described.

All experiments involving human subjects must be conducted in accordance with principles embodied in the Declaration of Helsinki (Code of Ethics of the World Medical Association). Experiments involving animal subjects must conform to the principles regarding the care and use of animals adopted by the American Physiological Society and the Society for Neuroscience. The editor may refuse papers that provide insufficient evidence of adherence to these principles.

**Results.** The results should be presented clearly and concisely, using figures and tables to summarize or illustrate the important findings. Quantitative observations are often more effectively displayed in graphs than in tables.

**Discussion.** The discussion should summarize the major findings and explain their significance in terms of the study's objectives and relationship to previous, relevant work. This section should present compact, clearly developed arguments rather than wide-ranging speculation or uncritical collation of earlier reports.

**Acknowledgments.** Use a separate page to recognize the contributions of individuals and supporting institutions.

**References.** In the text, references should be cited as follows:

as shown by Cella and Tusky (1990) (Bloom et al., 2000) (Cella & Tusky, 1990; Bloom et al., 2000)

The alphabetical list of references begins a new page, and must be typed double-spaced. Each in-text citation must have a corresponding reference and vice versa. List works by different authors who are cited within the same parentheses in chronological order, beginning with the earlier work. Journal titles should not be abbreviated. Only published articles and articles in press should appear in this list. Responsibility for the accuracy of references cited lies with the authors. Brief examples:

#### Journal article

Cella, D.F. & Tusky, D.S. (1990). Measuring quality of life today: Methodological aspects. *Oncology*, *4*, 29–38.

#### Book

Tompar-Tiu, A. & Sustento-Seneriches, J. (1995). Depression and Other Mental Health Issues: The Filipino American Experience. San Francisco: Jossey-Bass.

#### Chapter in an edited book

Karnofsky, D.A. & Burchenal, J.H. (1949). The clinical evaluation of chemotherapeutic agents in cancer. In *Evaluation of Chemotherapeutic Agents in Cancer*, Macleod, C.M. (ed.), pp. 191–205. New York: Columbia University Press. For more than one work by the same author(s) published in the same year, use (Jones, 1986*a*, 1986*b*) in text and likewise in the reference section.

**Tables.** Tables should be numbered consecutively with Arabic numerals and each should be typed double-spaced on a separate sheet. All tables are to be grouped together after the references. A short explanatory title and column headings should make the table intelligible without reference to the text. All tables must be cited and their approximate positions indicated in the text.

Figures and legends. The number of figures should be the minimum necessary to make the essential points of the paper. Figures should be supplied no larger than 8 imes 10''(approx.  $200 \times 250$  mm) and must be cameraready. Photographs for halftone reproduction must be on white glossy paper. Figures should be composed to occupy a single column (8.3 cm) or two columns (17 cm) after reduction. Diagrams and illustrations must have a professional appearance and be typed or drawn with sharp, black lettering to permit reduction. To assure legibility, letters, numbers, and symbols on figures should have a minimum height of 1 mm when reduced. Photomicrographs must include a calibration bar; if symbols are used on micrographs, they must contrast sufficiently with the background to be clearly visible when printed. Photocopies of micrographs are not acceptable for review purposes.

Artwork should normally be in black and white; if authors have color figures, the publisher will provide a price quotation for the additional production costs. All figures must be identified on the back with the short title of the paper, figure number, and figure orientation (top or bottom). Preferably, figures should be mounted on heavy sheets of the same size as the manuscript. Four complete sets of figures should be carefully packaged in protective envelopes, one to accompany each copy of the manuscript. Each figure must be cited and its approximate position clearly indicated within the text.

Figures must be numbered consecutively with Arabic numerals and be accompanied by a descriptive caption typed double-spaced on a separate sheet. The captions, collected at the end of the manuscript, should concisely describe the figure and identify any symbols and/or calibration bars.

**COPYEDITING AND PAGE PROOFS.** The publisher reserves the right to copyedit manuscripts to conform to the style of *Palliative & Supportive Care*. The corresponding author will receive page proofs for final proofreading. No rewriting of the final accepted manuscript is permitted at the proof stage, and substantial changes may be charged to the authors. Page proofs can be distributed as PDF files by email, and authors are encouraged to choose this option.

# **Palliative & Supportive Care**

#### **Editor-in-Chief**

William Breithart, M.D. Memorial Sloan-Kettering Cancer Center Department of Psychiatry & Behavioral Sciences 641 Lexington Avenue New York, NY 10022 Phone: (646) 888-0020 Fax: (212) 888-2356 E-mail: breitbaw@mskcc.org

#### **Co-Editors**

Harvey Max Chochinov, M.D., Ph.D., F.R.C.P.C. University of Manitoba CancerCare Department of Psychiatry McDermot Avenue Winnipeg, Manitoba R3E 0V9, Canada Phone: (204) 787-4933 Fax: (204) 787-4937 E-mail: harvey.chochinov@cancercare.mb.ca Simon Wein, M.D Palliative Care Unit Davidoff Cancer Centre Rabin Medical Centre 39 Jabotinski Street, Petach Tikvah Israel 49100 Phone: +972-3-9377216 Fax: +972-3-9240398

#### **Managing Editor**

Donna Cassetta, M.F.A. Memorial Sloan-Kettering Cancer Center Department of Psychiatry & Behavioral Sciences 641 Lexington Avenue New York, NY 10022 Phone: (646) 888-0020 Fax: (212) 888-2356 E-mail: cassettd@mskcc.org

#### **Editorial Board**

- Walter Baile, M.D., MD Anderson Cancer Center
- Sandra L. Bertman, Ph.D., Mount Ida College
- Susan Block, M.D., Dana-Farber Cancer Institute Gian Domenico Borasio, M.D., Ludwig Maximilians
- University
- Robert Brescia, Calvary Hospital
- Eduardo Bruera, M.D., MD Anderson Cancer Center
- Ira Byock, M.D., MD Anderson Cancer Center
- Edwin Cassem, M.D., Harvard Medical School
- David Cella, Ph.D., Northwestern University
- Nathan I. Cherny, M.B.B.S., F.R.A.C.P., Shaare Zedek Medical Center
- Nessa Coyle, R.N., M.S., F.A.A.N., Memorial Sloan-Kettering Cancer Center
- Albert Diefenbacher, M.D., Humboldt University
- Sylvie Dolbeault, M.D., Institut Curie
- Derek Doyle, O.B.E., National Council for Hospice and Specialist Palliative Care Services
- Kathleen Foley, M.D., Memorial Sloan-Kettering Cancer Center
- Linda Ganzini, M.D., Oregon Health Sciences University
- Francisco Gil, Ph.D., Institut Català d'Oncologia
- Paul Glare, M.D., Memorial Sloan-Kettering Cancer Center
- Luigi Grassi, M.D., University of Ferrara
- Irene Higginson, B.M., B.S., F.F.P.H.M., Ph.D., King's College London
- Jimmie Holland, M.D., Memorial Sloan-Kettering Cancer Center
- Paul Jacobsen, Ph.D., H. Lee Moffitt Cancer Center
- Stein Kaasa, M.D., Norwegian University of Science and Technology
- Virgilio Kasprzykowski, M.D., Instituto Portugues de Oncologia
- Brian Kelly, M.D., Ph.D., University of Newcastle

- David Kissane, MB., BS., MPM., MD., Memorial Sloan-Kettering Cancer Center
- Uwe Koch, M.D., Ph.D., University Hospital at Hamburg-Eppendorf
- Linda Kristjansen, Ph.D., Curtin University of Technology
- Peter Lawlor, M.D., St. Francis Hospice
- Matthew Loscalzo, M.S.W., City of Hope
- Paola Luzzatto, PhD., A.T.R.-B.C., Art Therapy Italiana
- Neil MacDonald, C.M., M.D., F.R.C.P.(C), F.R.C.P. (EDIN), McGill University
- Tatsuya Morita, M.E., D.O., Serei Hospice
- Balfour Mount, M.D., McGill University
- Steven Passik, Ph.D., Memorial Sloan-Kettering Caner Center
- Russell Portenoy, M.D., Beth Israel Medical Center
- Christina M. Puchalski, M.D., George Washington Institute for Spirituality and Health
- Barry Rosenfeld, Ph.D., Fordham University
- Carla Ripamonti, M.D., National Cancer Institute
- Tone Rustoen, R.N., Ph.D., Oslo University College
- John L. Shuster, Jr., M.D., University of Alabama School of Medicine
- Daniel P. Sulmasy, O.F.M., M.D., Ph.D., University of Chicago
- Frederic Stiefel, M.D., University of Lausanne
- James A. Tulsky, M.D., Durham VA Medical Center
- Yosuke Uchitomi, M.D., Ph.D., National Cancer Center Research Institute East
- Frits S. A. M. van Dam, Ph.D., The Netherlands Cancer Institute
- Maggie Watson, Ph.D., The Royal Marsden Hospital
- Roberto Wenk, M.D., Fundacion FEMEBA

# CAMBRIDGE



# CAMBRIDGE LIBRARY COLLECTION Books of enduring scholarly value



In the spirit of Cambridge University Press's continued commitment to innovation and enterprise, the Cambridge Library Collection will launch in July 2009, making an initial group of 475 out-of-print books available again. Another 1,000 will follow by the end of 2009 and 3,000 titles a year will be added as the program continues.

Drawing from the world-renowned collections in the Cambridge University Library, these books are carefully selected by experts in each subject area because of their scholarly importance or as landmarks in the history of their academic discipline.

Cambridge University Press is using state-ofthe art scanning machines to capture the contents of each book. The files are processed to give a consistently clear, crisp image, and the books finished to the high-quality standard for which the press is recognized around the world. The latest print-on-demand technology ensures that the books will remain available indefinitely.

The Cambridge Library Collection will revive books of enduring scholarly value across a wide range of disciplines in the humanities, social sciences, science and technology.

www.cambridge.org/us

CAMBRIDGE UNIVERSITY PRESS

 $1584 \cdot 2009$ 

425 YEARS OF CAMBRIDGE PRINTING AND PUBLISHING

https://doi.org/10.1017/S1478951509991052 Published online by Cambridge University Press

## **Case Report**

A.D. (SANDY) MACLEOD	513	Lightening up before death	
		Essay/Personal Reflections	
HANNAH KAUFMAN	517	This won't hurt a bit: The ethics of promising pain relief	
ANDREA KILLICK	521	Illuminating the path: What literature can teach doctors about death and dying	
		Poetry/Fiction	
BONNIE RAINGRUBER	527	Expertise	
BONNIE RAINGRUBER	529	Jerking Awake	
BONNIE RAINGRUBER	531	Changes	
	533	Erratum	
	537	Title and Author Index	
	543	Content to Volume 7	
	549	Instructions for Contributors	

# **PALLIATIVE & SUPPORTIVE CARE**

Volume 7

December 2009

Number 4

Seriously ill patients' discussions of preparation and life

Impact of a contemplative end-of-life training program:

completion: An intervention to assist with transition at the

What are the core elements of oncology spiritual care programs?

The meaning of being in transition to end-of-life care for

Palliative family caregivers' accounts of health care

"Caring as if it were my family": Health care aides' perspectives about expert care of the dying resident in a

"The healthy me appears": Palliative cancer patients'

Childhood abuse uncovered in a palliative care audit

absence of dementia: Phenomenologic differences

Humor and death: A qualitative study of The New Yorker

Delirium superimposed on dementia versus delirium in the

experiences of participation in a physical group exercise

Activity and meaning making in the everyday lives of people

female partners of spouses with cancer

experiences: The importance of "security"

CONTENTS

end of life

Being with dying

personal care home

with advanced cancer

cartoons (1986-2006)

**Review Article** 

program

### From the Editor

SIMON WEIN 391 One hundred meters

### **Original Articles**

- KAREN E. STEINHAUSER, 393 STEWART C. ALEXANDER, IRA R. BYOCK, LINDA K. GEORGE, AND JAMES A. TULSKY
  - CYNDA HYLTON RUSHTON, 405 DEBORAH E. SELLERS, KAREN S. HELLER, BEVERLY SPRING, BARBARA M. DOSSEY, AND JOAN HALIFAX
    - SHANE SINCLAIR, 415 MARLENE MYSAK, AND NEIL A. HAGEN
      - NISHA SUTHERLAND 423
- LAURA M. FUNK, DIANE E. ALLAN, 435 AND KELLI I. STAJDUHAR
  - SUSAN MCCLEMENT, 449 SUZANNE WOWCHUK, AND KATHLEEN KLAASEN
- HANNE PALTIEL, EDEL SOLVOLL, 459 JON HÅVARD LOGE, STEIN KAASA, AND LINE OLDERVOLL
  - KAREN LA COUR, 469 HELLE JOHANNESSEN, AND STAFFAN JOSEPHSSON
    - CATRIONA MACPHERSON 481
      - MARIANNE MATZO AND 487 DAVID MILLER

SOENKE BOETTGER, 495 STEVEN PASSIK, AND WILLIAM BREITBART

CHRISTOPHER J. MACKINNON 501

Applying feminist, multicultural, and social justice theory to diverse women who function as caregivers in end-of-life and palliative home care

continued in inside back cover

Cambridge Journals Online For further information about this journal please go to the journal website at: journals.cambridge.org/pax