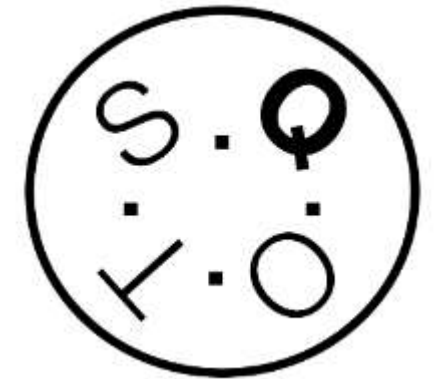


# How to measure Quality of Life in Clinical Obesity Care

The S.Q.O.T. Initiative

**Phillip J. Dijkhorst**, Claire E.E. de Vries, Caroline B. Terwee, Ignace M.C. Janssen, Ronald S.L. Liem, Bart A. van Wagenveld, Johan Ottosson, Bruno Halpern, Stuart W. Flint, Elisabeth F.C. van Rossum, Alend Saadi, Lisa West-Smith, Mary O'Kane, Jason C.G. Halford, Karen D. Coulman, Salman Al-Sabah, John B. Dixon, Wendy A. Brown, Ximena Ramos Salas, Sally Abbott, Alyssa J. Budin, Jennifer F. Holland, Lotte Poulsen, Richard Welbourn, Natasja Wijling, Laura Divine, Nadya Isack, Susie Birney, J.M. Bernadette Keenan, Theodore K. Kyle, Melanie Bahlke, Andrew Healing, Ian Patton, Valerie M. Montpellier



**[X] I have no potential conflict of interest to report**



# Introduction

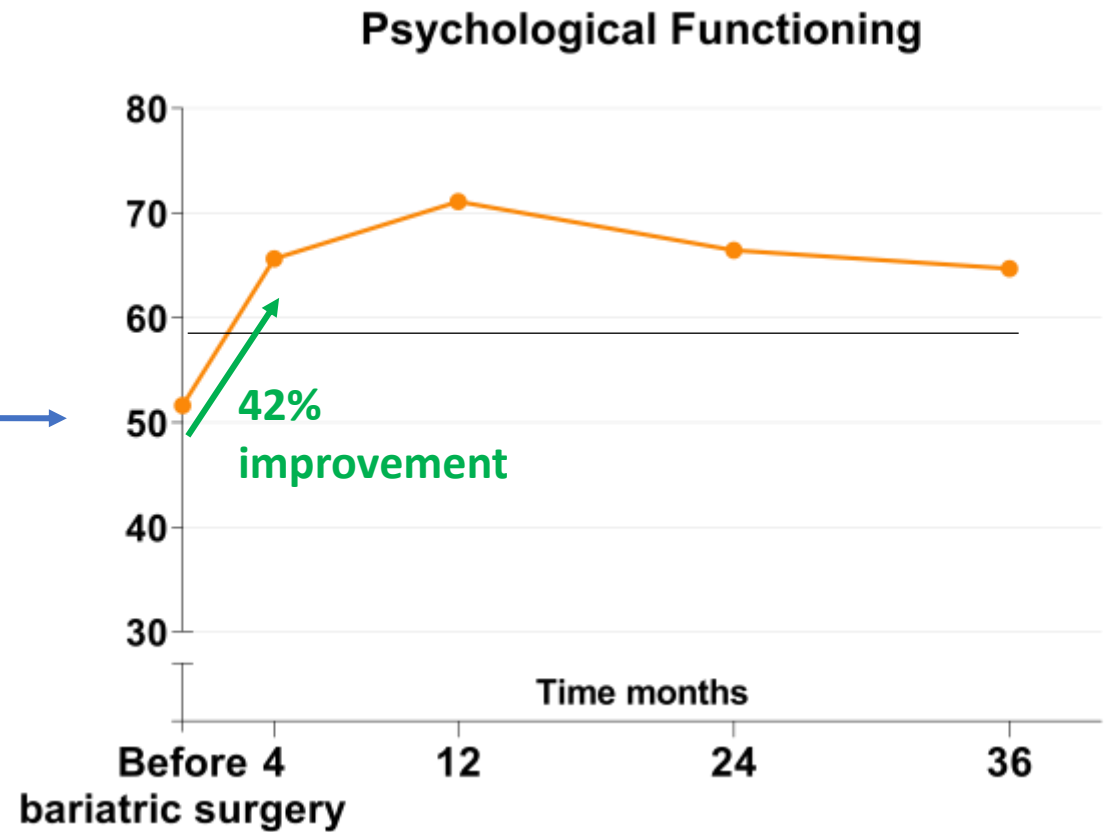


- Weight loss
- Remission of comorbidities
- Nutritional deficiencies
- Surgical complications

- Improved physical function
- Alleviation of psycho-social issues
- Improved body image
- Mitigation of daily challenges associated with obesity

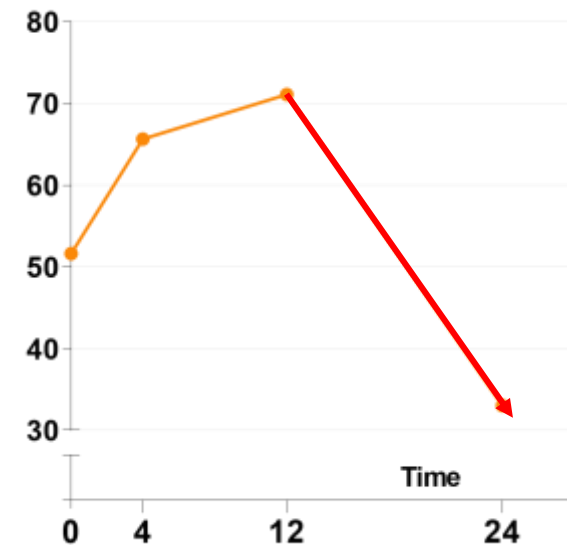


# Patient reported outcome measures



# Benefits of using PROMs

- ✓ Valuable insights into outcomes that truly matter to patients
- ✓ Improved patient-healthcare provider communication
- ✓ Support decision-making when additional interventions are necessary
- ✓ Support the comparative effectiveness of different interventions



# Challenges in quality of life measurement

- Measurement of irrelevant Patient Reported Outcome (PROs) domains
- The use of PROMs with poor measurement properties (reliability and validity)
- The wide variation in PROM use



Standardizing quality of life measures

# Standardizing Quality of Life Measures in Obesity Treatment (S.Q.O.T.)

## Goals:

- Standardization of Quality of Life measures in Obesity Treatment

## What to measure....

- Decide on the key components of QOL (PROs)

## How to measure...

- Selection of the preferred patient-reported outcome measure(s) (PROMs) to capture this information



| Domain                                   | PROM(s) available   | PROM(s) selected based on face validity                     | PROM(s) that were selected after the vote <sup>a</sup> |
|--|---|---|--|
| Self-esteem                              | IWQOL-Lite, IWQOL-Lite CT, PROS, WHO-QOL BREF   | IWQOL-Lite, IWQOL-Lite CT                                   | IWQOL-Lite   |
| Physical health/<br>functioning/symptoms | BAROS, BODY-Q, BOSS, BQL-Index, EQ-5D-5L, GIQLI, IWQOL-Lite, IWQOL-Lite CT, M-A QOL QII, OP-scale, PBOT, PROS, QOLOS, SF-36, TRIM, WHO-QOL BREF | BODY-Q, BQL-Index, IWQOL-Lite CT, SF-36                     | BODY-Q, IWQOL-Lite, SF-36,                             |
| Mental/psychological health              | BAROS, BODY-Q, BQL-Index, IWQOL-Lite CT, M-A QOL QII, SF-36, TRIM, WHO-QOL BREF   | BODY-Q, BQL-Index, IWQOL-Lite CT, SF-36                     | BODY-Q   |
| Social health                            | BAROS, BODY-Q, BOSS, BQL-Index, EQ-5D-5L, GIQLI, IWQOL-Lite, IWQOL-Lite CT, M-A QOL QII, OP-Scale, PBOT, PROS, QOLOS, SF-36, TRIM, WHO-QOL BREF | BODY-Q, BOSS, BQL-Index, GIQLI, IWQOL-Lite, OP-Scale, SF-36 | BODY-Q, IWQOL-Lite, OP-Scale                           |
| Stigma                                   | —   | —   | —  |
| Eating                                   | BODY-Q, BOSS, M-A QOL QII, QOLOS, TRIM  | BODY-Q, BOSS, QOLOS   | BODY-Q   |
| Body image                               | BODY-Q, QOLOS   | BODY-Q, QOLOS   | BODY-Q, QOLOS  |
| Excess skin                              | BODY-Q, QOLOS   | BODY-Q, QOLOS   | BODY-Q, QOLOS  |

.... >70%  
"Definitely include"

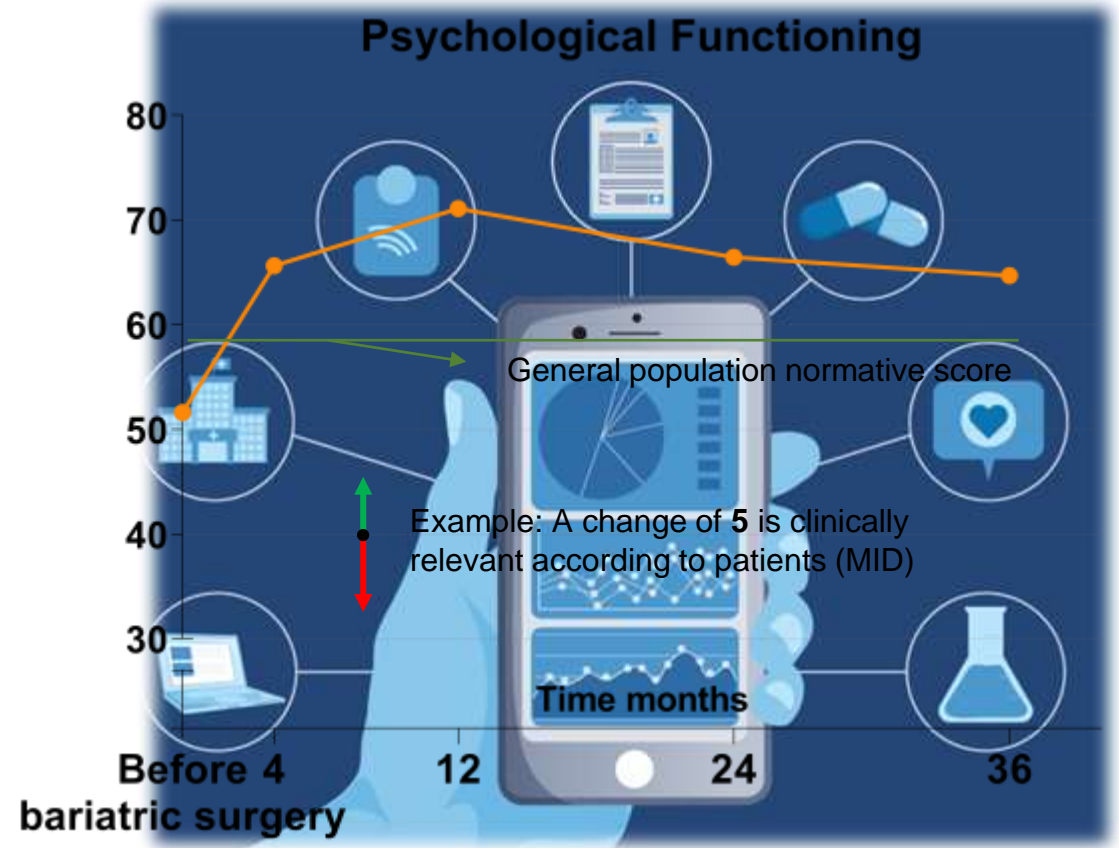
# Core set of PROMs for clinical obesity care

| <b>PRO</b>           | <b>Clinical practice</b> |
|----------------------|--------------------------|
| Self-esteem          | IWQOL-Lite               |
| Physical functioning | BODY-Q                   |
| Physical symptoms    | BODY-Q                   |
| Psychological health | BODY-Q                   |
| Social health        | BODY-Q                   |
| Stigma               | -                        |
| Eating               | BODY-Q                   |
| Body image           | BODY-Q                   |
| Excess Skin          | QOLOS                    |

# Examples of using PROMs in clinical practice

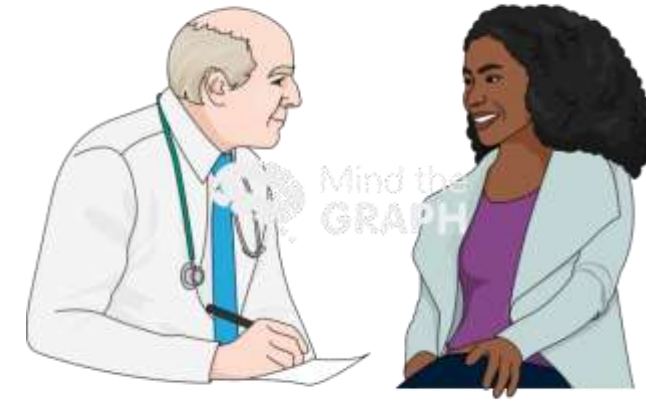


- Normative scores
- Minimal clinically important difference



# Conclusion

- To measure the outcomes that truly matter to patients, it is essential **to measure** quality of life in clinical obesity care and **to provide feedback** to the patient
- A core outcome set of PROs and PROMs for clinical practice is now available, incorporating the opinions of patients and experts



# Sponsors

- None of the members of the organizing committee (C.V., V.M., B.W., I.J., R.L. and P.D.) and none of the participants received payment for their participation.
- Travel expenses and the hotel overnight for the face-to-face meetings of the S.Q.O.T. were supported by Medtronic, Johnson & Johnson, Philips Vital Health, Novo Nordisk, Goodlife, Castor, Fitforme and Bart Torensma.

**Medtronic**

*Johnson & Johnson*



novo nordisk®

**goodlife**  
pharma

**fitforme**



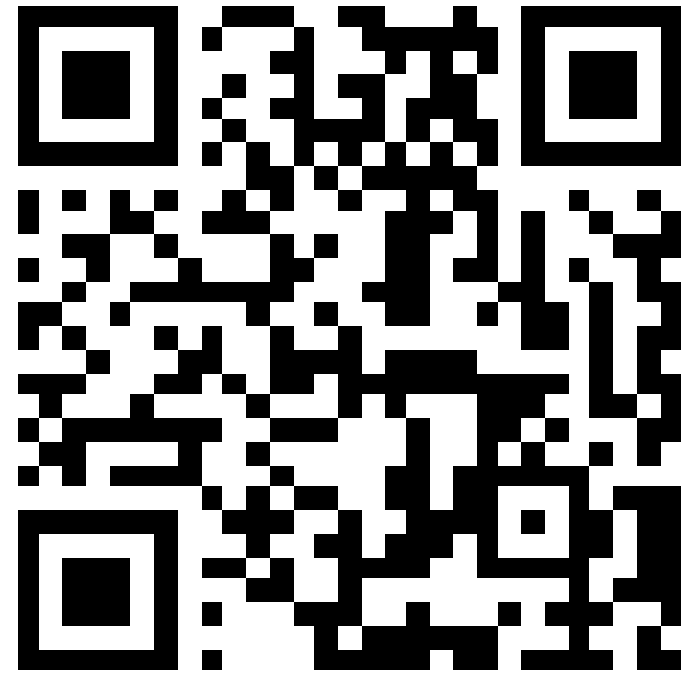
**NAPOLI**  
2023

# Questions?

<https://www.sqotinitiative.com/>

Twitter/Linkedin: S.Q.O.T. initiative

Sign up for our newsletter:



## Symptom Monitoring With Patient-Reported Outcomes During Routine Cancer Treatment: A Randomized Controlled Trial

Ethan Basch, Allison M. Deal, Mark G. Kris, Howard I. Scher, Clifford A. Hudis, Paul Sabbatini, Lauren Rogak, Antonia V. Bennett, Amylou C. Dueck, Thomas M. Atkinson, Joanne F. Chou, Dorothy Dulko, Laura Sit, Allison Barz, Paul Novotny, Michael Fruscione, Jeff A. Sloan, and Deborah Schrag

See accompanying editorial on page 527

Ethan Basch, Mark G. Kris, Howard I. Scher, Clifford A. Hudis, Paul Sabbatini, Lauren Rogak, Thomas M. Atkinson, Joanne F. Chou, Dorothy Dulko, Laura Sit, Michael Fruscione, and Deborah Schrag, Memorial Sloan Kettering Cancer Center, New York, NY; Ethan Basch, Allison M. Deal, and Antonia V. Bennett, University of North Carolina, Chapel Hill, NC; Amylou C. Dueck, Mayo Clinic, Scottsdale, AZ; Allison Barz, Children's Hospital of Philadelphia, Philadelphia, PA; Paul Novotny and Jeff A. Sloan, Mayo Clinic, Rochester, MN; and Deborah Schrag, Dana-Farber/Harvard Cancer Center, Boston, MA.

Published online ahead of print at [www.jco.org](http://www.jco.org) on December 7, 2015.

Supported by the National Cancer Institute and a grant from the Society of Memorial Sloan Kettering.

The National Cancer Institute and the Steps for Breath Fund of Memorial Sloan Kettering Cancer Center did not play any role in the design and conduct of the study; collection, management, analysis, and interpretation of the data; preparation, review, or approval of the manuscript; or decision to submit the manuscript for publication.

Authors' disclosures of potential conflicts of interest and author contributions are found at the end of this article.

### A B S T R A C T

#### Purpose

There is growing interest to enhance symptom monitoring during routine cancer care using patient-reported outcomes, but evidence of impact on clinical outcomes is limited.

#### Methods

We randomly assigned patients receiving routine outpatient chemotherapy for advanced solid tumors at Memorial Sloan Kettering Cancer Center to report 12 common symptoms via tablet computers or to receive usual care consisting of symptom monitoring at the discretion of clinicians. Those with home computers received weekly e-mail prompts to report between visits. Treating physicians received symptom printouts at visits, and nurses received e-mail alerts when participants reported severe or worsening symptoms. The primary outcome was change in health-related quality of life (HRQL) at 6 months compared with baseline, measured by the EuroQol EQ-5D Index. Secondary endpoints included emergency room (ER) visits, hospitalizations, and survival.

#### Results

Among 766 patients allocated, HRQL improved among more participants in the intervention group than usual care (34% v 18%) and worsened among fewer (38% v 53%;  $P < .001$ ). Overall, mean HRQL declined by less in the intervention group than usual care (1.4- v 7.1-point drop;  $P < .001$ ). Patients receiving intervention were less frequently admitted to the ER (34% v 41%;  $P = .02$ ) or hospitalized (45% v 49%;  $P = .08$ ) and remained on chemotherapy longer (mean, 8.2 v 6.3 months;  $P = .002$ ). Although 75% of the intervention group was alive at 1 year, 69% with usual care survived the year ( $P = .05$ ), with differences also seen in quality-adjusted survival (mean of 8.7 v 8.0 months;  $P = .004$ ). Benefits were greater for participants lacking prior computer experience. Most patients receiving intervention (63%) reported severe symptoms during the study. Nurses frequently initiated clinical actions in response to e-mail alerts.

#### Conclusion

Clinical benefits were associated with symptom self-reporting during cancer care.

*J Clin Oncol* 34:557-565. © 2015 by American Society of Clinical Oncology



## Psychological Functioning

