The Agreement between Measured and Predicted Resting Energy Expenditure in Patients with Pancreatic Cancer: A Pilot Study

Judith Bauer^{1,2}, Marina M Reeves², Sandra Capra³

¹The Wesley Research Institute. Brisbane. Australia. ²Centre for Health Research, Queensland University of Technology. Brisbane, Australia. ³School of Health Sciences, University of Newcastle. Callaghan, Australia

ABSTRACT

Objective To compare measured resting energy expenditure to resting energy expenditure predicted from eight published prediction equations in a sample of patients with pancreatic cancer.

Design Cross-sectional study.

Setting Ambulatory patients of a tertiary private hospital.

Participants Eight patients with pancreatic cancer (5 males, 3 females; age: 62.0±5.2 years; BMI: 24.4±3.2 kg/m²; weight loss: 12.1±6.0%; mean±SD).

Methods Resting energy expenditure was measured using indirect calorimetry and predicted from eight published prediction methods (Harris-Benedict with no injury factor, Harris-Benedict with 1.3 injury factor, Schofield, Owen, Mifflin, Cunningham, and Wang equations and the 20 kcal/kg ratio). Body composition was assessed by deuterium oxide dilution technique. Statistical analysis was performed by using the method of Bland and Altman, and the Student's t-test.

Results The Harris-Benedict equations with an injury factor of 1.3 resulted in a significantly higher mean predicted resting energy expenditure compared to measured resting energy expenditure, while there was

no significant difference between mean measured and predicted resting energy expenditure and the other 7 methods. At an individual level, the limits of agreement are wide for all equations. The best combination of low bias and narrowest limits of agreement was observed in the prediction of resting energy expenditure from the Wang equation (based on fat free mass) and the Harris-Benedict equation (based on weight and height).

Conclusion At a group level, there is agreement between mean measured and predicted resting energy expenditure with the exception of the Harris-Benedict equation with an injury factor of 1.3. The results of this pilot study suggest that, for an individual, the limits of agreement are wide, and clinically important differences in resting energy expenditure would be obtained. Clinicians need to be aware of the limitations of the use of resting energy expenditure prediction equations for individuals.

INTRODUCTION

The assessment of energy requirements is necessary for the development of individualised nutrition support regimens and recommendations. Measurement of energy expenditure is the most accurate method for determining energy requirements [1, 2, 3]. In the clinical setting, however, measurement of

energy expenditure is impractical, time-consuming and expensive. As such, prediction equations using easily measurable variables are commonly used to predict the basal metabolic rate (BMR) or resting energy expenditure (REE), which account for approximately 60-80% of total energy expenditure [4, 5, 6].

The development of cachexia is common in people with solid tumours such as pancreatic, lung, gastric and colorectal cancer. Cancer cachexia is associated with metabolic alterations that are likely to affect energy expenditure [7, 8, 9]. Patients with cancer cachexia are often malnourished: therefore. the provision of adequate nutrition support is essential for maintaining or improving nutritional status. Underfeeding and possible further weight loss in malnourished patients can result in an increase in complications and increased length of stay [10]. Overfeeding is also associated with complications such as hyperglycaemia, hyperlipidemia, hepatic dysfunction and respiratory distress [11].

Studies comparing mean measured REE and predicted REE by Harris-Benedict equations [12] in patients with pancreatic cancer have shown inconsistent results [13, 14, 15]. These studies have all compared, measured and predicted REE at the group level, whereas individual predictive accuracy is important in

the clinical setting. The method described by Bland and Altman is the appropriate statistical analysis for assessing agreement between two measurement methods [16].

Although no prediction equations have specifically been developed for people with cancer, a number have been developed in healthy populations. Comparison of measured REE with estimates of REE using these equations prediction would determine whether they are appropriate for use in people with cancer, whether alone or in combination with a constant stress factor. The combination of an injury factor of 1.3 with the Harris-Benedict equations commonly is recommended for patients with cancer [17, 18, 19]. Only the Harris-Benedict equations previously been compared have measurements of REE in cancer patients.

The aim of this study was to compare predicted REE from a number of published prediction equations to the measured REE of a sample of patients with pancreatic cancer.

METHODS

Patients

Eight patients (5 males, 3 females; age 62.0±5.2 years; body mass index 24.4±3.2 kg/m²; weight loss in past six months

Table 1. Characteristics of eight patients with pancreatic cancer.

Patient	Gender	Age	Replication	Weight	BMI	FFM
Number		(years)		(kg)	(kg/m^2)	(kg)
1	Male	55	1	94.4	28.8	60.1
			2	90.6	27.7	59.8
			3	91.8	28.0	60.8
			4	95.0	29.0	63.3
2	Female	70	1	40.2	17.0	34.4
3	Female	63	1	51.6	21.8	39.3
4	Male	65	1	79.0	23.6	64.0
5	Male	67	1	75.4	22.8	54.0
6	Male	59	1	79.2	24.4	52.8
			2	81.6	25.2	-
			3	82.1	25.3	57.4
7	Female	57	1	61.4	21.3	37.2
			2	61.6	21.3	41.9
8	Male	60	1	73.9	24.1	61.2
			2	77.4	25.3	65.7
Mean±SD		62.0±5.2		75.7±15.9	24.4±3.3	53.7±10.9

SD = standard deviation

12.1±6.0%; mean±SD) with pancreatic cancer receiving palliative treatment who had lost greater than 5% body weight in the previous months had their resting energy expenditure measured. Diagnosis was based histological and/or radiological (computerized axial tomography) or operative diagnosis. The disease was staged according to "Union Internationale Contre le Cancre (UICC)" criteria (stage II: local spread, 3 patients; stage IV: distant metastases, 5 patients). Four patients participated as part of a nutrition intervention trial and had more than one consecutive measurement at least 4 weeks apart. A total of 15 resting energy expenditure measurements were performed. Characteristics of patients who participated in the study are shown in Table 1.

Resting Energy Expenditure (REE)

Resting energy expenditure was measured by open circuit indirect calorimetry (Vmax 229, Sensormedics, Yorba Linda, CA, USA) using a mouthpiece and noseclip in a thermoneutral environment. Patients had fasted overnight (12 hours) and rested for 30 minutes prior to the measurement. Measurements were taken between 7:00 and 9:00 am. Gas analysers were calibrated prior to each measurement against standard mixed reference gases. The mass flow sensor was also calibrated prior to each measurement. Measurements carried out for at least 30 minutes. The steady state period was determined by the computer software and was defined as a 5-minute period during which average minute oxygen consumption (VO₂) changed by less than 10% and the respiratory quotient (RQ) changed by less than 5%. REE was determined from VO₂ and VCO₂ using the abbreviated Weir equation [20].

Body Composition

Weight was measured to the nearest 0.1 kg (Tanita Inc., Tokyo, Japan, Model 300GS) and height measured to the nearest 0.1 cm with a stadiometer (Harpenden, Holtain Ltd,

Crosswell, Dyfed, UK). Total body water was measured using the stable isotope deuterium in the form of water (²H₂O). Patients provided a urine sample and then drank a 10% solution of ²H₂O based on their body weight (0.5 g/kg body weight). The amount of deuterium consumed was weighed to 2 decimal places. A second urine sample was provided 5 h later. The enrichment of the pre-dose urine sample, post-dose urine sample, local tap water and the dose given were measured using isotope radio mass spectrometry (Hydra, Europa Scientific, Crewe, UK). The deuterium oxide dilution space was divided by a factor of 1.04 to calculate total body water [21]. Fat free mass (FFM) was estimated from total body water assuming a constant hydration factor of 0.73.

Prediction Equations

Four prediction equations using combination of easily measurable variables (weight, height, age) and two prediction equations based on a measure of FFM were examined (Table 2). The ratio method commonly recommended nutrition bv societies and clinical nutrition textbooks [22, 23, 24] was also included in the analysis. Although a value of 30-35 kcal/kg is commonly recommended for cancer patients to predict total energy expenditure, for the purpose of this study, a value of 20 kcal/kg was used as the estimate was compared to resting energy expenditure. The equations by Ireton-Jones et al. [25] were not included in these analyses as they predict total daily energy expenditure and are therefore not comparable to measured resting energy expenditure. The units of energy calculated from the equations varied. All predicted REE values were calculated as per equation then converted to kilojoules.

ETHICS

The Ethics Committee of the hospital approved the study and informed written consent was obtained from all participants.

Table 2. Prediction equations for estimating energy requirements.

Equation	Subset of subjects	Formula
Harris and Benedict [12]	Males Females	BMR (kJ/day) = $(57.5 \times \mathbf{W}) + (20.9 \times \mathbf{H}) - (28.3 \times \mathbf{A}) + 278$ BMR (kJ/day) = $(40.0 \times \mathbf{W}) + (7.7 \times \mathbf{H}) - (19.6 \times \mathbf{A}) + 2,741$
Schofield [40]	Males, 30-60 years Females, 30-60 years Males, over 60 years Females, over 60 years	BMR (MJ/day) = $(0.048 \times W) + 3.653$ BMR (MJ/day) = $(0.034 \times W) + 3.538$ BMR (MJ/day) = $(0.049 \times W) + 2.459$ BMR (MJ/day) = $(0.038 \times W) + 2.755$
Owen et al. [36, 37]	Males Females	RMR (kcal/day) = $(10.2 \text{ x W}) + 875$ RMR (kcal/day) = $(7.18 \text{ x W}) + 795$
Mifflin et al. [32]	Males Females	REE (kcal/day) = $(10 \times \mathbf{W}) + (6.25 \times \mathbf{H}) - (5 \times \mathbf{A}) + 5$ REE (kcal/day) = $(10 \times \mathbf{W}) + (6.25 \times \mathbf{H}) - (5 \times \mathbf{A}) - 161$
Cunningham [41]	All subjects	REE (kcal/day) = $(21.6 \text{ x } \text{FFM}) + 370$
Wang et al. [42]	All subjects	REE (kcal/day) = $(21.5 \text{ x } \text{FFM}) + 407$
20 kcal/kg ratio [24]	All subjects	REE (kcal/day) = $\mathbf{W} \times 20$

BMR: basal metabolic rate; RMR: resting metabolic rate; REE: resting energy expenditure; W: body weight (kg); H: height (cm); A: age (years); FFM: fat free mass (kg); kJ: kilojoules; MJ: megajoules; kcal: kilocalories

STATISTICS

Statistical analysis was carried out using SPSS for Windows (Version 11.0.1, 2001, SPSS Inc., Chicago, USA). Mean, standard deviation (SD), and frequencies were the descriptive statistics used to present patient characteristics. The paired Student's t-test was used to assess individual differences between measured REE and predicted REE by each equation. The approach of Bland and Altman was used to compare the agreement between measured REE and REE estimated from the prediction equations [16]. This analysis allowed for the calculation of bias (mean of the individual differences between estimates) and the limits of agreement (± 2 SD from the mean bias). Correlation analysis was used to examine the association between the mean of the measured and predicted REE, and the differences between the two methods. Statistical significance was reported at the conventional P<0.05 level (two-tailed). The interpretation of the differences was not based solely on statistical testing but also on clinically important differences in energy expenditure which was defined a priori to be greater than 400 kJ/day (equivalent to greater than $\pm 3-6\%$) for this sample of patients [26].

RESULTS

Mean measured REE was 6,557±1,131 kJ/day $(1,568\pm270 \text{ kcal/day})$ or $123.0\pm14.5 \text{ kJ/kg}$ FFM (29.4±3.5 kcal/kg FFM). Table 3 shows mean predicted REE, the bias (mean difference between predicted REE and measured REE) and limits of agreement (±2 standard deviations of the bias) for the prediction of REE for each equation relative to the measured REE. There was no significant difference between mean measured and predicted REE from the Harris-Benedict (no injury factor), Schofield, Owen, Mifflin, Cunningham, and Wang equations and the 20 kcal/kg ratio method. The Harris-Benedict equations with an injury factor of 1.3 resulted in a significantly higher (P<0.001) mean predicted REE as compared to measured REE.

There was a significant correlation between measured REE and REE as predicted by Harris-Benedict equations alone, Harris-Benedict equations with an injury factor of 1.3, Schofield equations, Owen equations, Mifflin equations, Cunningham equation, Wang equation and 20 kcal/kg ratio method; however, this is a measure of the strength of the relationship - and not a measure of the

Table 3. Predicted resting energy expenditure (REE), bias and limits of agreement for predicted REE relative to

measured REE (6,557±1,131 kJ/day) in patients with pancreatic cancer.

Equation	Predicted REE	Bias Limits of		Student's t-test*		Correlation**	
	(mean±SD; kJ/day)	(kJ/day)	agreement (±2SD; kJ/day)	t value	P value	r value	P value
Harris and Benedict ^a [12]	6,579±1,170	22	1,717	0.097^{b}	0.924	0.722	0.002
Harris and Benedict x 1.3 [12]	8,552±1,521	1,995	2,105	7.342^{b}	< 0.001	0.722	0.722
Schofield [40]	6,825±1,285	268	2,134	0.974^{b}	0.347	0.617	0.014
Owen et al. [36, 37]	6,617±1,095	60	1,574	0.297^{b}	0.771	0.750	0.001
Mifflin et al. [32]	6,329±1,241	-228	1,810	-0.975 ^b	0.346	0.713	0.003
Cunningham [41]	6,404±982	-117	1,280	-0.686 ^c	0.505	0.836	< 0.001
Wang et al. [42]	6,537±977	15	1,280	0.088 $^{\rm c}$	0.931	0.836	< 0.001
20 kcal/kg ratio [24]	6,330±1,332	-227	1,889	-0.932 ^b	0.367	0.717	0.003

SD: standard deviation

level of agreement - between the two methods [16].

The smallest bias was observed with the Wang equation (15 kJ) followed by the Harris-Benedict equations alone (22 kJ) and Owen equations (60 kJ). The 20 kcal/kg ratio method, Mifflin and Cunningham equations tended to underestimate REE values, while the Schofield equations and Harris-Benedict equations with an injury factor of 1.3 tended to overestimate REE values. The bias for all prediction methods was consistent across the range of measurements of REE as the mean of the measured REE, and predicted REE was not significantly correlated with the difference between measured and predicted REE for all prediction equations.

Individual data indicates that the limits of agreement for each of the prediction equations are wide, with the narrowest limit evident with the Cunningham and Wang equations (1,280 kJ for both) and the widest limits with the Schofield equations (2,134 kJ) and Harris-Benedict equations with an injury factor of 1.3 (2,105 kJ).

Nine (60%) measurements of REE were within $\pm 10\%$ of REE predicted by the Harris-

Benedict equations, three measurements (20%) were classified as hypermetabolic (REE greater than 110% of the Harris-Benedict-predicted value) and three measurements (20%) were classified as hypometabolic (REE less than 90% of the Harris-Benedict-predicted value).

DISCUSSION

The aim of this study was to compare the agreement between predicted REE from a number of published equations to measured REE in patients with pancreatic cancer. For good agreement, it is expected that bias will be close to zero, that 95% of the differences will lie between the limits of agreement and that there is no clear evidence of a relationship between difference and mean of measured and predicted REE. Based on these assumptions, mean predicted REE from the equations developed by Harris-Benedict, Schofield. Owen, Mifflin, Wang, Cunningham and the 20 kcal/kg ratio method agree with mean measured REE at a group level. The Harris-Benedict equation with an injury factor of 1.3 for cancer is not suitable

^a no injury factor

^b Evaluated on 15 cases (14 degrees of freedom)

^c 14 available cases only (13 degrees of freedom): FFM missing for 1 case

^{*} Comparison between predicted and measured REE

^{**} Correlation bias against mean of predicted and measured REE

at the group level due to the large overestimation.

To consider how well the methods are likely to agree for an individual, the limits of agreement are assessed and, provided they are not clinically important, the methods can be used interchangeably. In this study the limits of agreement are wide (greater than 400 kJ) for all equations. For an individual with cancer cachexia, predicted REE commonly used prediction equations cannot be used interchangeably with measured REE, as there would be clinically important differences in the REE results obtained. As an example, the Wang et al. equations may predict REE as much as 1,295 kJ above or 1,265 kJ less than measured REE for an individual. while the Harris-Benedict equations with an injury factor of 1.3 may predict REE as much as 4,100 kJ above or 110 kJ less than measured REE.

Reeves and Capra [27], in their survey of more than 300 clinical dietitians in Australia, determined that 67.4% of respondents used the Schofield equations and 25.9% used the Harris-Benedict equations for prediction of energy requirements for a cancer case study. The majority added an injury factor for cancer to their estimate (median injury factor of 1.3) [27]. In the present study, the addition of the injury factor of 1.3 recommended for cancer to the Harris-Benedict equations significantly overestimated energy requirement. The use of injury factors has recently been questioned by Reeves and Capra [26]. This study indicates that the automatic application of an injury factor for patients with pancreatic cancer is not appropriate, as only 20% of measurements were greater than 110% of predicted REE from the Harris-Benedict equations. Such gross overestimation of energy requirements could lead to negative complications associated with overfeeding [11].

A wide range of metabolism from hypometabolic to hypermetabolic has been observed in cancer patients [14, 28, 29, 30]. The metabolic response to cancer is highly variable as even within the same type of cancer, REE is increased in some patients and not in others [31]. Dempsey *et al.* [14] found

that patients with pancreatic cancer were predominantly hypometabolic Falconer et al. [15] and Barber et al. [13] found elevated REE in patients with pancreatic cancer. In this study, the mean REE predicted from Harris-Benedict was 100.8±14.1% although 20% of measurements were classified as hypometabolic and 20% hypermetabolic. Several investigators have shown that the Harris-Benedict equations overestimate energy expenditure in the healthy population by 5-15% [32, 33, 34, 35, 36, 37]. Therefore, the number of patients true with elevated **REE** mav underestimated

In clinical practice, measurement of energy expenditure is impractical, time-consuming and expensive. In spite of the limitations of prediction equations for use in individuals, clinicians require an estimate of REE. In this study the best combination of low bias and narrow limits of agreement was observed in the prediction of REE from the Wang equation based on FFM. Measurement of FFM using the deuterium dilution technique is impractical in the clinical setting, and other methods such as bioelectrical impedance analysis have been shown to be inaccurate at prediction of FFM at the individual level in patients with cancer [38]. Therefore, most clinicians rely on REE prediction equations that incorporate easily measurable variables. such as body weight and height. Of these, the Harris-Benedict equations gave the lowest bias and narrowest limits of agreement, followed by the Owen equations. Therefore if both weight and height were available, the Harris-Benedict equations would recommended and if only weight is available, the Owen equations. As the limits of agreement for both equations are wide. clinicians should be aware of the limitations of the equations for individuals and that monitoring of patient outcomes is essential. A limitation of this study was that more than

A limitation of this study was that more than one REE measurement was performed on four of the study patients who were participating in a nutrition intervention trial. As the aim of the study was to compare agreement between REE methods, repeated measurements have been included in the analyses as there were significant changes in weight and body composition between repeated measurements for all four patients. Hence changes in measured REE due to changes in body composition should also be reflected in predictions of REE. An additional limitation relates to the assumption that the hydration of lean tissue is constant at 73%. In pancreatic cancer, development of oedema or ascites is a feature of the end stage of the disease. Changes in fluid distribution such as oedema or ascites may lead to inaccurate body composition determinations [39]. Although no patients with visible oedema or ascites were included in the study, it is possible that some patients had subclinical fluid retention. Finally, the variation in the level of metabolism observed might be due to differences in disease stage or other factors associated with the disease. The sample size in this study was too small to investigate this further. From a clinical practice perspective however, this study has been informative in identifying the potential for overestimation of REE with the automatic application of injury factors with prediction equations.

CONCLUSION

At a *group* level, the equations developed by Harris-Benedict, Schofield, Owen, Mifflin, Wang, Cunningham and the 20 kcal/kg ratio method are suitable for predicting the REE of pancreatic cancer patients. However, these results suggest that for an *individual*, the limits of agreement are wide for all prediction equations and clinically important differences in REE would be obtained. Clinicians need to be aware of the limitations of the use of REE prediction equations for estimating individual REE in pancreatic cancer patients. Monitoring of patient outcomes is the most effective method of determining whether patients are receiving adequate nutrition support.

Received September 1st, 2003 - Accepted December 5th, 2003

Keywords Cachexia; Calorimetry, Indirect; Nutritional Requirement; Metabolism; Pancreatic Neoplasms; Pilot Projects; Protein-Energy Malnutrition

Abbreviations BMR: basal metabolic rate; FFM: fat free mass; REE: resting energy expenditure; RQ: respiratory quotient; VCO₂ carbon dioxide consumption; VO₂ oxygen consumption

Acknowledgment The study was sponsored by The Wesley Research Institute

Correspondence

Judith Bauer
The Wesley Research Institute
PO Box 499
Toowong, Queensland
Australia 4066
Phone: +61-7-3232.7918

Phone: +61-7-3232.7918 Fax: +61-7-3371.6834

E-mail address: judithba@wesley.com.au

References

- 1. Soares MJ, Sheela ML, Kurpad AV, Kulkarni RN, Shetty PS. The influence of different methods on basal metabolic rate measurements in human subjects. Am J Clin Nutr 1989; 50:731-6. [PMID 2508459]
- 2. Jequier E, Schutz Y. Long-term measurements of energy expenditure in humans using a respiration chamber. Am J Clin Nutr 1983; 38:989-98. [PMID 6650455]
- 3. Flancbaum L, Choban PS, Sambucco S, Verducci J, Burge JC. Comparison of indirect calorimetry, the Fick method, and prediction equations in estimating the energy requirements of critically ill patients. Am J Clin Nutr 1999; 69:461-6. [PMID 10075331]
- 4. Shetty PS, Henry CJ, Black AE, Prentice AM. Energy requirements of adults: an update on basal metabolic rates (BMRs) and physical activity levels (PALs). Eur J Clin Nutr 1996; 50:S11-23. [PMID 8641254]
- 5. Ravussin E, Bogardus C. Relationship of genetics, age, and physical fitness to daily energy expenditure and fuel utilization. Am J Clin Nutr 1989; 49:968-75. [PMID 2655422]
- 6. Carpenter WH, Poehlman ET, O'Connell M, Goran MI. Influence of body composition and resting metabolic rate on variation in total energy expenditure:

- a meta-analysis. Am J Clin Nutr 1995; 61:4-10. [PMID 7825536]
- 7. Tisdale MJ. Metabolic abnormalities in cachexia and anorexia. Nutrition 2000; 16:1013-4. [PMID 11054609]
- 8. Nelson KA, Walsh D, Sheehan FA. The cancer anorexia-cachexia syndrome. J Clin Oncol 1994; 12:213-25. [PMID 8270980]
- 9. Fearon KC, Barber MD, Moses AG. The cancer cachexia syndrome. Surg Oncol Clin N Am 2001; 10:109-26. [PMID 11406454]
- 10. Ottery FD. Definition of standardized nutritional assessment and interventional pathways in oncology. Nutrition 1996; 12:S15-9. [PMID 8850213]
- 11. Klein CJ, Stanek GS, Wiles CE, 3rd. Overfeeding macronutrients to critically ill adults: metabolic complications. J Am Diet Assoc 1998; 98:795-806. [PMID 9664922]
- 12. Harris JA, Benedict FG. A Biometric Study of Basal Metabolism in Man. Washington, DC, USA: Carnegie Institution of Washington, 1919.
- 13. Barber MD, McMillan DC, Preston T, Ross JA, Fearon KC. Metabolic response to feeding in weight-losing pancreatic cancer patients and its modulation by a fish-oil-enriched nutritional supplement. Clin Sci 2000; 98:389-99. [PMID 10731472]
- 14. Dempsey DT, Feurer ID, Knox LS, Crosby LO, Buzby GP, Mullen JL. Energy expenditure in malnourished gastrointestinal cancer patients. Cancer 1984; 53:1265-73. [PMID 6692317]
- 15. Falconer JS, Fearon KC, Plester CE, Ross JA, Carter DC. Cytokines, the acute-phase response, and resting energy expenditure in cachectic patients with pancreatic cancer. Ann Surg 1994; 219:325-31. [PMID 7512810]
- 16. Bland JM, Altman DG. Statistical methods for assessing agreement between two methods of clinical measurement. Lancet 1986;1: 307-10. [PMID 2868172]
- 17. Lysen LK. Quick Reference to Clinical Dietetics. Gaithersburg, MD, USA: Aspen Publishers Inc., 1997.
- 18. Alpers D, Stenson W, Bier D. Manual of Nutritional Therapeutics. 4th edition. Philadelphia, PA, USA: Lippincott Williams & Wilkins, 2002.
- 19. Dietitians' Pocket Book. Perth, Australia: School of Public Health, Curtin University of Technology, 1999.
- 20. Weir JB. New methods for calculating metabolic rate with special reference to protein metabolism. J Physiol 1949; 109:1-9.
- 21. Schoeller DA, van Santen E, Peterson DW, Dietz W, Jaspan J, Klein PD. Total body water measurement

- in humans with 18O and 2H labeled water. Am J Clin Nutr 1980; 33:2686-93. [PMID 6776801]
- 22. Bloch AS. Cancer. In: Matarese LE, Gottschlich MM, eds. Contemporary Nutrition Support Practice: A Clinical Guide. Philadelphia, PA, USA: W.B.Saunders Company, 1998.
- 23. Bloch AS. Oncology. In: Lysen LK, ed. Quick Reference to Clinical Dietetics. Gaithersburg, MD, USA: Aspen Publishers Inc., 1997.
- 24. ASPEN Board of Directors and the Clinical Guidelines Task Force. Guidelines for the use of parenteral and enteral nutrition in adult and pediatric patients. JPEN. J Parenter Enteral Nutr 2002; 26:1-138. [PMID 11841046]
- 25. Ireton-Jones CS, Turner WW Jr, Liepa GU, Baxter CR. Equations for the estimation of energy expenditures in patients with burns with special reference to ventilatory status. J Burn Care Rehabil 1992; 13:330-3. [PMID 1618877]
- 26. Reeves MM, Capra S. Predicting energy requirements in the clinical setting: are current methods evidence based? Nutr Rev 2003; 61:143-151. [PMID 12795449]
- 27. Reeves MM, Capra S. Variation in the application of methods used for predicting energy requirements in acutely ill adult patients: a survey of practice. Eur J Clin Nutr 2003; 57:1530-5. [PMID 14647217]
- 28. Knox LS, Crosby LO, Feurer ID, Buzby GP, Miller CL, Mullen JL. Energy expenditure in malnourished cancer patients. Ann Surg 1983; 197:152-62. [PMID 6824369]
- 29. Bosaeus I, Daneryd P, Svanberg E, Lundholm K. Dietary intake and resting energy expenditure in relation to weight loss in unselected cancer patients. Int J Cancer 2001; 93:380-3. [PMID 11433403]
- 30. Hyltander A, Drott C, Korner U, Sandstrom R, Lundholm K. Elevated energy expenditure in cancer patients with solid tumours. Eur J Cancer 1991; 27:9-15. [PMID 1826450]
- 31. Fredrix EW, Soeters PB, Rouflart MJ, von Meyenfeldt MF, Saris WH. Resting energy expenditure in patients with newly detected gastric and colorectal cancers. Am J Clin Nutr 1991; 53:1318-22. [PMID 2021140]
- 32. Mifflin MD, St Jeor ST, Hill LA, Scott BJ, Daugherty SA, Koh YO. A new predictive equation for resting energy expenditure in healthy individuals. Am J Clin Nutr 1990; 51:241-7. [PMID 2305711]
- 33. Foster GD, Wadden TA, Mullen JL, Stunkard AJ, Wang J, Feurer ID, et al. Resting energy expenditure, body composition, and excess weight in the obese. Metabolism 1988; 37:467-72. [PMID 3367789]

- 34. Daly JM, Heymsfield SB, Head CA, Harvey LP, Nixon DW, Katzeff H, et al. Human energy requirements: overestimation by widely used prediction equation. Am J Clin Nutr 1985; 42:1170-4. [PMID 4072952]
- 35. Case KO, Brahler CJ, Heiss C. Resting energy expenditures in Asian women measured by indirect calorimetry are lower than expenditures calculated from prediction equations. J Am Diet Assoc 1997; 97:1288-92. [PMID 9366867]
- 36. Owen OE, Holup JL, D'Alessio DA, Craig ES, Polansky M, Smalley KJ, et al. A reappraisal of the caloric requirements of men. Am J Clin Nutr 1987; 46:875-85. [PMID 3687821]
- 37. Owen OE, Kavle E, Owen RS, Polansky M, Caprio S, Mozzoli MA, et al. A reappraisal of caloric requirements in healthy women. Am J Clin Nutr 1986; 44:1-19. [PMID 3728346]
- 38. Isenring E, Bauer J, Capra S, Davies PSW. Evaluation of foot-to-foot bioelectrical impedance

- analysis for the prediction of total body water in oncology outpatients. Eur J Clin Nutr 2004; 58:46-51. [PMID 14679366]
- 39. Kushner RF, Gudivaka R, Schoeller DA. Clinical characteristics influencing bioelectrical impedance analysis measurements. Am J Clin Nutr 1996; 64:423S-7S. [PMID 8780358]
- 40. Schofield WN. Predicting basal metabolic rate, new standards and review of previous work. Hum Nutr Clin Nutr 1985; 39:5-41. [PMID 4044297]
- 41. Cunningham JJ. Body composition as a determinant of energy expenditure: a synthetic review and a proposed general prediction equation. Am J Clin Nutr 1991; 54:963-9. [PMID 1957828]
- 42. Wang Z, Heshka S, Gallagher D, Boozer CN, Kotler DP, Heymsfield SB. Resting energy expenditure-fat-free mass relationship: new insights provided by body composition modeling. Am J Physiol Endocrinol Metab 2000; 279:E539-45. [PMID 10950820]