

Estimation of the Total Efficiency of Metabolizable Energy Utilization for Maintenance and Growth by Cattle in Tropical Conditions

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ABSTRACT - Data of 320 animals were obtained from eight comparative slaughter studies performed under tropical conditions and used to estimate the total efficiency of utilization of the metabolizable energy intake (MEI), which varied from 77 to 419 kcal kg^{-0.75} d⁻¹. The provided data also contained direct measures of the recovered energy (RE), which allowed calculating the heat production (HE) by difference. The RE was regressed on MEI and deviations from linearity were evaluated by using the *F*-test. The respective estimates of the fasting heat production and the intercept and the slope that composes the relationship between RE and MEI were 73 kcal kg^{-0.75} d⁻¹, -42 kcal kg^{-0.75} d⁻¹ and 0.37. Hence, the total efficiency was estimated by dividing the net energy for maintenance and growth by the metabolizable energy intake. The estimated total efficiency of the ME utilization and analogous estimates based on the beef cattle NRC model were employed in an additional study to evaluate their predictive powers in terms of the mean square deviations for both temperate and tropical conditions. The two approaches presented similar predictive powers but the proposed one had a 22% lower mean squared deviation even with its more simplified structure.

Key Words: bioenergetics, growth, net energy, ruminants

Estimação da Eficiência Total de Utilização da Energia Metabolizável para Manutenção e Crescimento por Bovinos em Condições Tropicais

RESUMO - Foram utilizadas 320 informações obtidas a partir de 8 estudos de abate comparativo conduzidos em condições tropicais para se estimar a eficiência total de utilização da energia metabolizável consumida, a qual variou de 77 a 419 kcal kg^{-0.75} d⁻¹. Os dados também continham informações sobre a energia retida (RE), o que permitiu o cálculo da produção de calor por diferença. As estimativas da produção de calor em jejum e dos coeficientes linear e angular da regressão entre RE e MEI foram respectivamente, 73 kcal kg^{-0.75} d⁻¹, -42 kcal kg^{-0.75} d⁻¹ e 0,37. Em seguida, a eficiência total foi estimada dividindo-se a energia líquida para manutenção e produção pelo consumo de energia metabolizável. A eficiência total de utilização da MEI obtida e estimativas análogas baseadas no modelo do NRC gado de corte foram empregadas em um estudo adicional para avaliar as previsões da energia líquida total para manutenção e crescimento em condições de clima temperado e tropical. Os dois modelos apresentaram poder de predição semelhante, mas o quadrado do desvio médio do método proposto foi 22% menor, mesmo com uma estrutura mais simplificada.

Palavras-chave: bioenergética, crescimento, energia líquida, ruminantes

Introduction

The transformation of food energy into products of animal origin, as in any other energy transformation system, is not devoid of losses since there is an efficiency by which the food energy is used for maintenance and production. Brody (1945) and Kleiber (1975) categorized two types of energy

efficiency in the production systems: gross or total efficiency and net or partial efficiency. The former is obtained by dividing the energy recovered in the animal product by the total intake of a determined food energy category (gross, digestible or metabolizable), while the latter is the recovered energy divided by the subtraction between the food energy intake and its amount used for maintenance purposes.

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Among the losses observed in the energy transformation processes, there are pronounced ones that can not be neglected: the contribution of the heat of combustion from both the partially digestible and indigestible residues recovered in feces; the energy losses associated to gaseous products of digestion and urinary excretion residues; and the heat increment due to digestion, absorption and uptake of nutrients (NRC, 1981, 1996; Baldwin, 1995). Subtracting these losses from the total energy intake one could obtain the net energy of the food, i.e., the fraction that will be available for both the maintenance and productive functions.

The observation that animals present different performances by consuming rations with the same contents in total digestible nutrients (TDN) led to the development of the net energy systems, allowing the prediction of these differences in performance because energy losses associated to heat increments are considered and are directly related to the fiber content of the ration (Blaxter, 1966; Van Soest, 1994).

The net energy system is based on comparative slaughter experiments to determine the energy requirements of beef cattle and the net energy value of the food. The heat energy (HE) is calculated by the difference between the metabolizable energy intake (MEI) and the energy retained by the animal body (RE). According to the current terminology recommended by the NRC (1981), the heat energy could be further fractionated into the energy costs due to fasting metabolism (H_eE) and to maintenance and productive heat increments (H_iE).

The maintenance and retained net energy contents of the food are both defined according to the following expressions (NRC, 1981, 1996):

$$[NE_m] = H_eE/I_m, \quad NE_m = H_eE \quad (1)$$

$$[NE_r] = RE/(I - I_m), \quad NE_r = RE \quad (2)$$

The terms $[NE_m]$ and $[NE_r]$, expressed in kcal kg^{-1} of dry matter, correspond to the food net energies concentrations of maintenance and production, respectively; the terms H_eE and RE are expressed in $\text{kcal kg}^{-0.75} \text{d}^{-1}$; I_m ($\text{kg kg}^{-0.75} \text{d}^{-1}$) is the food intake necessary to meet the maintenance functions; and I ($\text{kg kg}^{-0.75} \text{d}^{-1}$) represents the voluntary dry matter intake under ad libitum conditions.

The relationship among the metabolizable energy, the net energy for maintenance (NE_m), the retained energy ($RE = NE_r$), and the heat increments associated

to these processes is classically described by the expression below (NRC, 1981; NRC, 1996):

$$ME = NE_m + NE_r + H_iE \quad (3)$$

where ME represents the animal requirement for metabolizable energy and all terms are expressed in $\text{kcal kg}^{-0.75} \text{d}^{-1}$.

In AFRC (1993) the square brackets involving the variables are used to denote energy concentration (energy units per unit of mass), while the variables without this notation represent the total energy category required by the animal (energy units per unit of time). In the present study, the same notation was used. Nevertheless, in the NRC (1981; 1996) the notation used to describe both the energy requirements ($\text{kcal kg}^{-0.75} \text{d}^{-1}$) and the food energy values (kcal kg^{-1}) were the same and this may cause misunderstanding. An example is the improper sum between $[NE_r]$ and $[NE_m]$, since the replacement of the terms in Eq. (3) by its concentration counterparts of Eq. (1) and (2) will result in the following paradox:

$$H_eE/I_m + RE/(I - I_m) + H_iE/I \neq ME/I. \quad (4)$$

The goals of this study were the estimation of the total efficiency of metabolizable energy utilization for maintenance and growth by cattle raised under tropical conditions and the evaluation of the predictive power of the statistical approach performed.

Materials and Methods

Experimental data

The data used in this study were obtained from the appendices of five Doctoral thesis (Teixeira, 1984; Paulino, 1996; Ferreira, 1997; Signoretti, 1998; Vêras, 2000) and two Master of Science dissertations (Salvador, 1980; Freitas, 1995) presented to the Animal Science Department of the Federal University of Viçosa and one Doctor of Philosophy thesis presented to the Faculty of the Graduate School of Cornell University (Tedeschi, 2001). In these studies, the comparative slaughter technique was used to estimate the net energy requirements for maintenance and growth, based on both the metabolizable energy intake and the retained energy. The studies above provided 325 observations from which 5 were considered outliers and discarded after residual analysis (Draper & Smith, 1966).

Data were composed by animals belonging to Nellore ($n_1 = 132$), Gir ($n_2 = 12$), 1/4 Fleckvieh×5/16 Angus×7/16 Nellore ($n_3 = 11$), F_1 Holstein×Nellore ($n_4 = 12$), Guzerat ($n_5 = 12$), F_1 Holstein×Zebu ($n_6 = 42$), 3/4 Holstein×Zebu ($n_7 = 10$), Tabapuan ($n_8 = 10$), 5/8 Holstein×Zebu ($n_9 = 9$), F_1 Nellore

×Simmental ($n_{10} = 24$), and Holstein ($n_{11} = 46$) breed and crossbreed types. All animals were male and 134 of the total were castrated. The experiments were carried out for five months on average, with the animals initial age varying from 2 to 42 months old. Information about the animals is presented in Table 1.

Table 1 - Description of the estimation set of data

Breed ¹	n	Initial age, days	Gender	MEI kcal kg ^{-0.75} d ⁻¹		Reference
				Minimum	Maximum	
H-Z	34	1260	Steer	77	250	Salvador (1980)
1/2 H-Z	8	480	Steer	148	292	Teixeira (1984)
3/4 H-Z	10	480	Steer	129	257	Teixeira (1984)
5/8 H-Z	9	480	Steer	145	254	Teixeira (1984)
H	10	480	Steer	143	236	Teixeira (1984)
N	10	480	Steer	138	218	Teixeira (1984)
N	10	720	Bull	100	203	Freitas (1995)
F-A-N	11	720	Bull	103	266	Freitas (1995)
H-N	12	720	Bull	114	280	Freitas (1995)
Gir	12	720	Bull	129	325	Paulino (1996)
Guz	12	720	Bull	140	321	Paulino (1996)
N	10	720	Bull	147	419	Paulino (1996)
T	10	720	Bull	143	332	Paulino (1996)
N-S	24	480	Bull	231	342	Ferreira (1997)
H	36	60	Bull	235	396	Signoretto (1998)
N	25	600	Bull	202	296	Véras (2000)
N	53	675	Steer	136	261	Tedeschi (2001)
N	12	675	Bull	145	221	Tedeschi (2001)
N	12	315	Bull	142	216	Tedeschi (2001)

¹Breed: H = Holstein, N = Nellore, T = Tabapuan, Guz = Guzerat, H-Z = Holstein-Zebu, F-A-N = Fleckvieh-Angus-Nellore, H-N = Holstein-Nellore, N-S = Nellore-Simmental.

The nonlinear form of the model employed by Lofgreen and Garret (1968) was used to describe the heat production as a function of the metabolizable energy intake:

$$HE_i = \alpha \cdot \exp(\beta \cdot MEI_i) + e_i \quad (5)$$

where HE_i and MEI_i represented the heat energy and the metabolizable energy intake in the i^{th} animal, both expressed in kcal kg^{-0.75} d⁻¹; α is the parameter that allowed estimating the fasting heat production and corresponds to the net energy requirement for maintenance (NE_m , kcal kg^{-0.75} d⁻¹); β is defined as the specific transformation coefficient of the consumed metabolizable energy into heat (d kg^{0.75} Kcal⁻¹), equivalent to the slope of the original linear model used by Lofgreen & Garret (1968) multiplied by $\ln 10$; and the term e_i used to describe the experimental error.

The parameters estimates of the nonlinear model described by Eq. (5) were obtained according to the Marquardt's compromise by means of a nonlinear least squares estimation procedure. Following the first fitting, outliers were excluded from the original data when the absolute values of their studentized residuals were greater than three (Draper & Smith, 1966). Then, the parameters were again estimated by using the new data set without the outliers.

The metabolizable energy intake for maintenance (MEI_m) was established at the energy equilibrium, where assumptions made were the completely conversion of MEI into heat and at this point RE should be equated to 0. Afterwards MEI_m was estimated by an iterative procedure in which increasing values of MEI (within the range of the observed data) were substituted in the equation:

$$HE = \hat{\alpha} \cdot \exp(\hat{\beta} \cdot MEI) \quad (6)$$

until the ratio HE/MEI approached 1.000.

The validity of the linearity assumption for the relationship between energy retention and the metabolizable energy intake, widely discussed in the literature (Blaxter, 1966; NRC, 1981; Garret & Johnson, 1983; Baldwin, 1995), was verified by using *F*-tests based on the sum of squares of first, second and third degrees models (Searle, 1971). Since linearity assumptions were not violated, the estimated MEI_m was used as a component of the parametric restriction in the following model:

$$\begin{cases} RE_i = \tilde{a} + \tilde{a} \cdot MEI_i + e_i \\ \mathbf{P}'\tilde{\mathbf{e}} = 0 \end{cases} \quad (7)$$

where RE_i corresponds to the energy retained by the i^{th} animal; γ and δ represent both the intercept and the slope of the linear regression of the RE_i over MEI_i ; and e_i is the experimental error under the usual assumptions of normally and independently distributions with mean 0 and variance σ^2 . \mathbf{P}' is the transpose of the restriction vector applied to the parameter vector $\tilde{\mathbf{e}}$, such that $\mathbf{P}' = [1 \quad MEI_m]$ and $\tilde{\mathbf{e}}' = [\gamma \quad \delta]$. The solution of the restricted model was obtained by the ordinary least squares estimation procedure (Searle, 1971) and illustrated by its normal equations in matrix terms as follows:

$$\begin{bmatrix} \hat{\gamma} \\ \hat{\delta} \\ \dots \\ \hat{\lambda} \end{bmatrix} = \begin{bmatrix} n & \sum_{i=1}^n MEI_i & \dots & 1 \\ \sum_{i=1}^n MEI_i & \sum_{i=1}^n MEI_i^2 & \dots & MEI_m \\ \dots & \dots & \dots & \dots \\ 1 & MEI_m & \dots & 0 \end{bmatrix}^{-1} \begin{bmatrix} \sum_{i=1}^n RE_i \\ \sum_{i=1}^n MEI_i RE_i \\ \dots \\ 0 \end{bmatrix} \quad (8)$$

where λ is a Lagrange multiplier. The logic of this approach was the avoidance of different MEI_m estimates after relating both the retained energy and the heat production on MEI.

Energetic efficiency

The total efficiency of metabolizable energy utilization (k_i) was individually estimated for those

animals belonging to the evaluation set of data by using the following expression:

$$\hat{k}_i = (NE_m + RE_i)/MEI_i = \hat{\alpha}/MEI_i + \hat{\gamma}/MEI_i + \hat{\delta} \quad (9)$$

where $\hat{\alpha}$, $\hat{\gamma}$ and $\hat{\delta}$ were estimated by fitting Eq. (5) and (7) to the experimental data.

The comparison of predictions and observations to assess model performance (COPAMP, according to P. L. Mitchell, personal communication) was done by using 134 paired observations (MEI and RE) obtained from the appendices of four Master of Science dissertations (Teixeira, 1975; Piekarski, 1983; Galvão, 1991; Paulino, 2002) and one Doctoral thesis (Freitas, 2004) presented to the Animal Science Department of the Federal University of Viçosa (Table 2), and 65 paired means (each one calculated from 6 to 8 observations) published by Lofgreen & Garret (1968). These data sets were independent from the experimental one used for parameters estimation, a common criterion for empirical validation (Mertens, 1976; Mitchell & Sheehy, 1997).

The fasting heat production of the evaluation set of data was ad hoc estimated after fitting Eq. (5) and added to the observed RE of each animal to obtain the related total net energy required for maintenance and growth (NE_{mg}). The observed MEI_i of this group were used as inputs to Eq. (9) and allowed predicting the NE_{mg} by using parameters estimates ($\hat{\alpha}$, $\hat{\gamma}$ and $\hat{\delta}$) after fitting Eq. (5) and (7) to the set of experimental data. The same procedure was done by using the average values published by Lofgreen & Garret (1968). The NE_{mg} estimates were also obtained by simulating the NRC (1996) model with the evaluation set of data as an input. The predictive powers of these approaches were compared by means of residual analysis (Draper & Smith, 1966; Mitchell & Sheehy, 1997) and by the decomposition of the mean square deviations as suggested by Kobayashi & Salam (2000).

Results and Discussion

Net energy for maintenance

The procedure used to fit the model described by Eq. (5) to the experimental data set reached the established convergence criteria for the Marquardt's

Table 2 - Description of the evaluation set of data

Breed ¹	n	Initial age, days	Gender	MEI kcal kg ^{-0.75} d ⁻¹		Reference
				Minimum	Maximum	
1/2 H-Z	19	1080	Steer	140	218	Teixeira (1975)
H-Z	16	900	Steer	134	434	Piekarski (1983)
N-M	12	60	Bull	249	342	Galvão (1991)
N-L	12	60	Bull	273	325	Galvão (1991)
N	12	60	Bull	227	319	Galvão (1991)
Zebu	15	720	Steer	153	307	Paulino (2002)
N-A	12	300	Bull	231	310	Freitas (2004)
N	12	300	Bull	199	274	Freitas (2004)
N-B	12	300	Bull	220	298	Freitas (2004)
N-S	12	300	Bull	193	267	Freitas (2004)

¹ Breed: N = Nellore, H-Z = Holstein-Zebu, N-M = Nellore-Marchigiana, N-L = Nellore-Limousin, N-A = Nellore-Aberdeen Angus, N-B = Nellore-Brown Swiss, N-S = Nellore-Simmental.

algorithm with three iterations. The examination of the studentized residuals plotted against the metabolizable energy intake allowed identifying five outliers. After their elimination from the original set of data, a new fitting procedure was done which resulted in the estimates presented in Table 3.

The plot of the residuals (Figure 1b) exhibited an undesirable curvature nearest the upper and lower values of the MEI. However, the good adjustment of the function's line to the observed data (Figure 1a), the asymptotic confidence intervals (Table 3) and the small coefficient of variation (10^0 order of magnitude, %) of the parameters estimates were indicative of a satisfactory quality of fit.

The fasting heat production estimated in this study (Table 3) was slightly lower than $77 \text{ kcal kg}^{-0.75} \text{ d}^{-1}$ found by Lofgreen & Garret (1968) and Tedeschi et al. (2002) for animals belonging, respectively, to *Bos taurus* and *Bos indicus* species. The estimate $1.7 \cdot 10^{-3} \text{ d kg}^{0.75} \text{ kcal}^{-1}$ obtained by Lofgreen & Garret (1968) with the linearized form of the model did not fall within the asymptotic confidence interval estimated for

parameter β in the present study (Table 3). More closer with the presented results were the estimates obtained by Ferrell & Jenkins (1998) for animals of diverse genotypes (74.9 for α and $3.7 \cdot 10^{-3}$ for β). A point estimate alone, however, is not satisfactory. Random errors are presented in all measurements, and no mathematical model could be used to account all facets of a physical situation. Therefore, it is virtually impossible to obtain the true values of the parameters. Nor point estimates calculated from different data samples will be equal, even if the samples were obtained under similar conditions. Thus it is necessary to augment the point estimates with some information on its variability, which is provided by the standard deviations and confidence intervals associated to the parameters (Bard, 1974).

The fasting heat production estimated (Table 3) would corroborate the hypothesis that the *Bos indicus* breeds and crossbreeds have lower net energy requirements for maintenance due to their lower genetic production potential and better adaptation to unfavorable environmental conditions (NRC, 1996). In this study, the majority of the information used to

Table 3 - Point and interval estimates of the parameters obtained by fitting the model that describes the heat production as a function of the metabolizable energy intake (Eq. (5))

Parameters	Estimates	SE ^a	Asymptotic confidence interval at the probability level of 0.95	
			Lower limit	Upper limit
α^*	73	2	70	76
β^{**}	3.9×10^{-3}	8.0×10^{-5}	3.7×10^{-3}	4.0×10^{-3}

* kcal kg^{-0.75} d⁻¹.

** d kg^{0.75} kcal⁻¹.

^a Asymptotic standard error.

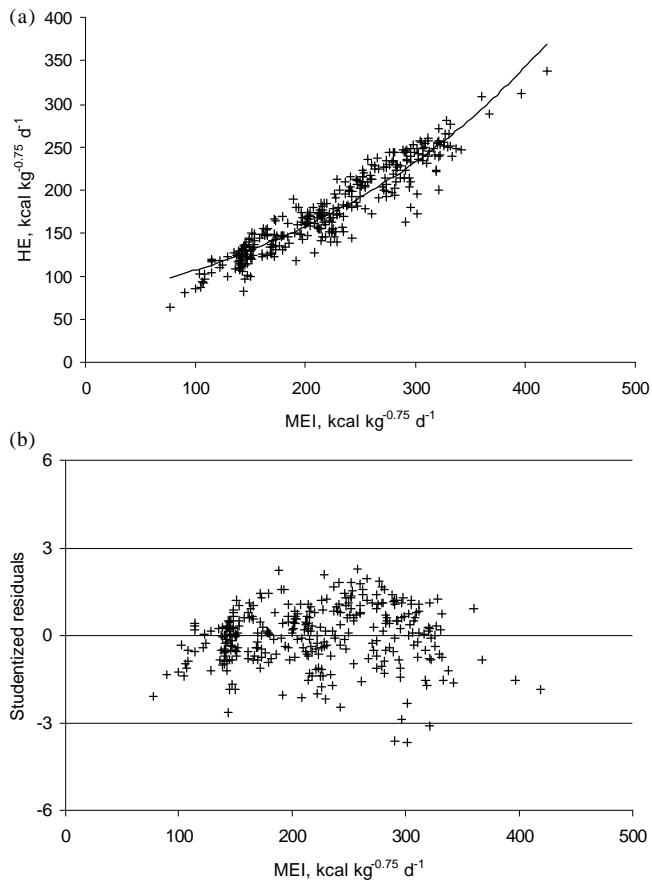


Figure 1 - Relationship of heat production (HE) to metabolizable energy intake (MEI) (a). Estimates of the studentized residuals related to heat production plotted against the metabolizable energy intake (b).

fit Eq (5) was obtained from *Bos indicus* breeds and crossbreeds (n=274). Nevertheless, this subject is open to question since other studies have not found differences between *Bos indicus* and *Bos taurus* in relation to the net energy requirements for maintenance (Ferrell & Jenkins, 1998; Tedeschi et al., 2002). Brody (1945) and Baldwin (1995) suggested that there are factors such as the body mass, age, sex, and

physiological stage affecting estimates for parameter α . Kleiber (1975) emphasized that a larger interval for the animal masses causes a greater influence over the estimate of this parameter, thus reducing the impact of other factors. The modeling process is based on assumptions and aggregations about the components of the real-world systems (Mertens, 1976; 1993). This theoretical reduction is thus an intrinsic characteristic of the modeling process and as mentioned earlier some aspects of the real system must be disregarded for mathematical treatment and simplicity (Bard, 1974; Mertens, 1976; Kobayashi & Salam, 2000).

The assertion about the existence of differences among the mentioned estimates for parameter α could not be valid, since we found a high asymptotic correlation between the estimates for parameters α and β after the fitting procedure. This may be a problem intrinsically related to model characteristics and fitting procedures that are beyond the scope of the present study.

The simulation of the model under extreme conditions provides an evaluation tool for either its applicability or its lack of generality (Mertens, 1976). If a continuous increase of the metabolizable energy intake is simulated, the Eq. (5) with its exponential behavior predicts a heat production that could lead the animal to a theoretical overheating. However, no signs of either an inflection point or an asymptotic phase were identified after a visual appraisal of the plotted relationship between HE and MEI (Fig 1a), which still justifies the application of the model described by Eq. (5).

The net energy requirement for maintenance was considered constant in the approach presented. This assumption has been made for the feed evaluation and nutritional requirement systems in current use for ruminants (NRC, 1981; 1996). However, Milligan & Summers (1986) and Baldwin (1995) asserted that the heat increment above the maintenance level is

Table 4 - Variation range of the studied variables

Variables	Minimum observed	Maximum observed	Number of observations
MEI ¹	77	419	320
HE ²	63	339	320
RE ³	-6	129	320
EBW ⁴	102	486	320

¹ Metabolizable energy intake (kcal kg^{-0.75} d⁻¹).

² Heat energy (kcal kg^{-0.75} d⁻¹).

³ Retained energy (kcal kg^{-0.75} d⁻¹).

⁴ Empty body weight (kg).

attributed to the heat increments of the productive functions and to support energy expending processes that are not part of the productive related pathways. In fact, the latter processes are further accentuated by intake above energy balance, i.e., in line with the plane of nutrition.

The relationship between the recovered energy and the metabolizable energy intake was presumed linear and checked by using the F -test (Searle, 1971). After fitting the unrestricted models of first, second and third degrees, it was observed that the quadratic ($\hat{F}=1.81$; $P=0.1794$) and cubic effects ($\hat{F}<1$) were not significant and the linear model (Eq. (7)) adopted to describe this relationship.

Estimated the value $112 \text{ kcal kg}^{-0.75} \text{ d}^{-1}$ for MEI_m , the restricted linear model described by Eq. (7) was then fitted to obtain the point estimates of the parameters and their respective confidence intervals (Table 5, Figure 2).

The fitted equation accounted for only 49% of the variation in the RE. We are conscious about the effects of other variables over RE, among them are sex (Ferrell & Jenkins, 1985), breed (Ferrell & Jenkins, 1998) and stage of growth (Geay, 1984, Williams & Jenkins, 2003; Tedeschi et al., 2004). Geay (1984) argued that the diet ME utilization efficiency for growth is inversely related to the proportion of energy retained as protein in the animal body. The data used in his study were mean values obtained from literature, which in turn probably arose from large data sets. The objective of this study, however, was to estimate the diet $[\text{NE}_{\text{mg}}]$ and the size of the data set used hamper accurate estimates of the above-mentioned effects. Our strategy was the same employed by Ferrell & Jenkins (1998) who studied the diet ME utilization by diverse genotypes of cattle by pooling data of all genotypes together to obtain “a more robust equation”,

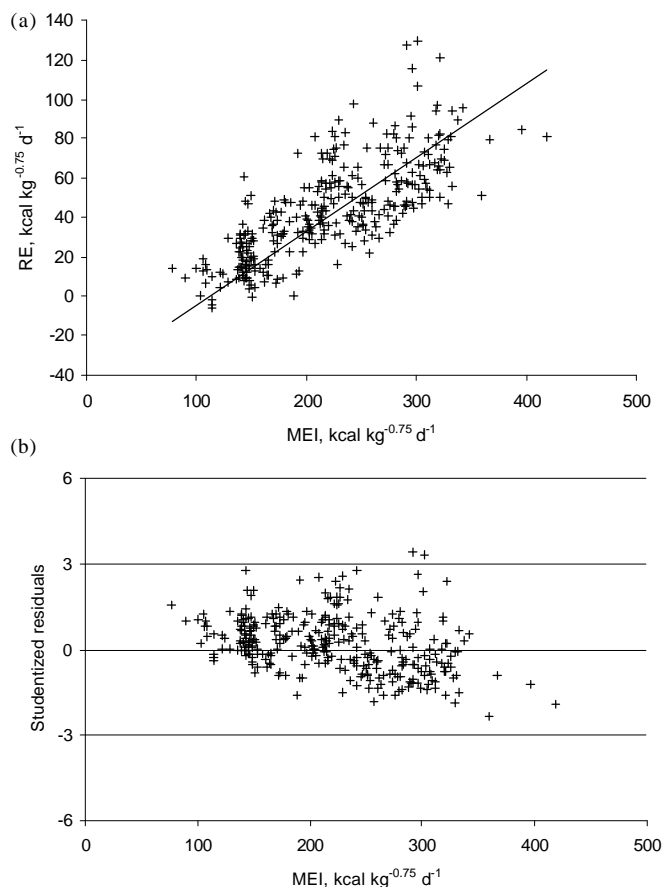


Figure 2 - (a) Relationship between recovered energy (RE) and metabolizable energy intake (MEI). (b) Studentized residuals plotted against the metabolizable energy intake.

despite statistical differences detected among breeds. Nevertheless, this matter will be considered in further developments of the approach just described.

The net energy available for maintenance and growth (NE_{mg}) of the evaluation data set was predicted from the observed MEI by using a simple rearrangement of Eq. (9):

Table 5 - Point and interval estimates of the parameters obtained by fitting the model that describes the retained energy as a function of the metabolizable energy intake (Eq. (8))

Parameters	Estimates	SE ^a	Confidence interval at the probability level of 0.95	
			Lower limit	Upper limit
γ^*	-42	1	-44	-40
δ^{**}	3.7×10^{-1}	1.0×10^{-2}	3.6×10^{-1}	3.9×10^{-1}

* $\text{kcal kg}^{-0.75} \text{ d}^{-1}$.

** Unitless parameter that represents the asymptotic partial efficiency of the metabolizable energy utilization.

^a Standard error.

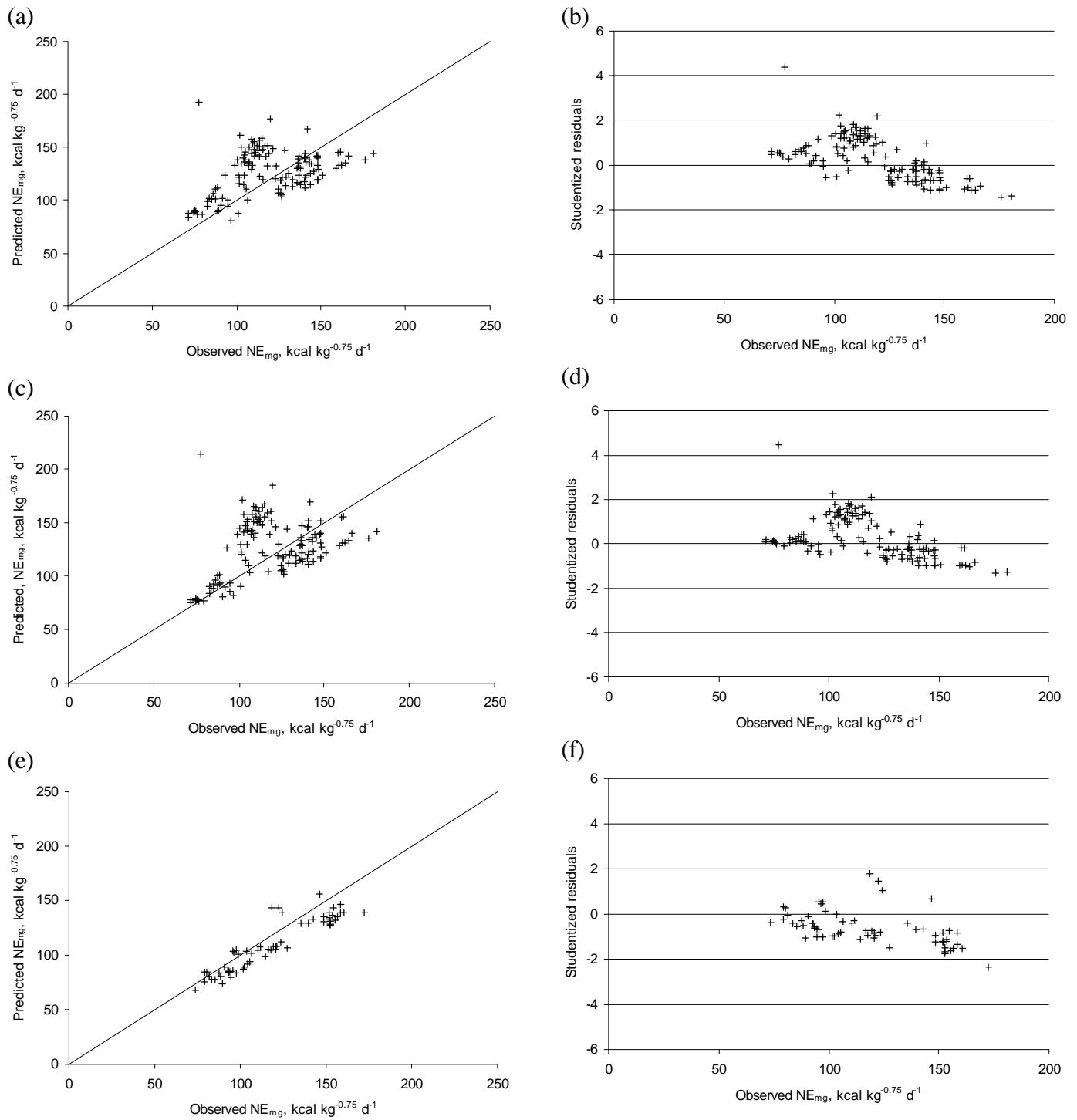


Figure 3 - (a) NE_{mg} predicted by using Eq. (10) and (c) NE_{mg} predicted by using the NRC (1996) equations either plotted against observed values obtained under tropical conditions; (e) NE_{mg} predicted by using Eq. (10) against observed values obtained by Lofgreen & Garret (1968); the residuals (predicted minus observed) of each scatter graph (a, c and e) were also plotted against the observed values (b, d and f).

$$NE_{mg_i} = MEI_i \cdot \hat{k}_i \quad (10)$$

where subscript i denotes the i^{th} animal of the evaluation set of data.

The concentration of the ration metabolizable

energy, [ME], the empty body weight (EBW), and the dry matter intake (DMI) of the evaluation data were also used as inputs in the NRC (1996) equations to estimate the NE_{mg} . This procedure was not done with the data published by Lofgreen & Garret (1968),

Table 6 - Mean squared deviations and its components in a comparison between the current approach and the NRC based approach

Components	Current approach	NRC based approach
Squared bias	67	72
Squared difference between standard deviations	11	4
Lack of correlation weighted by the standard deviation	611	803
Mean squared variation	622	808
Mean squared deviation	688	879

because the DMI were not available, hence they were only used in the evaluation of the estimates obtained with Eq. (10). The latter observations provided an evaluation of the model performance under temperate climate conditions (Figure 3e,f).

The evaluation of the performances of both models allowed identifying its similar predictive powers and systematic errors (Figure 3), but the presented approach had 22, 7, 24 and 23% respectively smaller mean squared deviation (MSD), squared bias (SB), lack of correlation weighed by the standard deviation (LCS) and mean squared variation (MSV) than the NRC model. The latter had a 59% lower squared difference between standard deviations (SDSD) in relation to our estimates (Table 6, Figure 4). According to Kobayashi & Salam (2000), when comparing predicted and observed values, the lower the value of MSD, the closer the simulation is to the measurement, a bigger MSV indicates that the model failed to simulate the variability of the measurement around the mean, a larger SDSD indicates that the model failed to simulate the magnitude of fluctuation among the n measurements and, a greater LCS means that the model failed to simulate the pattern of the fluctuation across the n measurements.

According to the analysis presented, the ration content of net energy for maintenance and growth estimated on the basis of the total efficiency of metabolizable energy utilization can be treated on an additive fashion:

$$k = NE_m / ([ME] \cdot I) + RE / ([ME] \cdot I) \quad (11)$$

Since

$$[ME] \cdot I = MEI \quad (12)$$

and by dividing the right-hand side of Eq. (10) by the dry matter intake (I), and substituting k by the expression described in Eq. (11) yield:

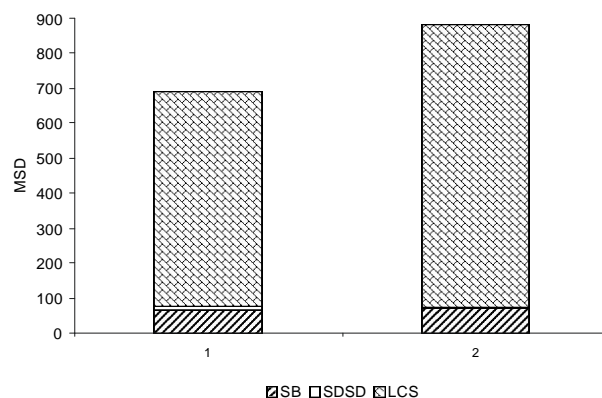


Figure 4 - Mean squared deviation (MSD) and its components, squared bias (SB), squared difference between standard deviations (SDSD), and lack of correlation weighted by the standard deviation (LCS) in a comparison between the presented approach (1) and the NRC model (2).

$$MEI \cdot k / I = [ME] \cdot k = [ME] \cdot \{NE_m / ([ME] \cdot I) + RE / ([ME] \cdot I)\} \quad (13)$$

and then,

$$(NE_m + RE) / I = NE_{mg} / I \quad (14)$$

Nutritionally, the terms of the equation related to the second law of thermodynamics are described as follows (Brody, 1945; Baldwin, 1995):

$$\Delta G = \Delta H - T\Delta S \quad (15)$$

where ΔG corresponds to the amount of free energy of the food available for productive functions and maintenance processes, $[NE_{mg}]$; ΔH represents the food energy available for oxidation by tissues, and corresponds to the food metabolizable energy concentration, $[ME]$; and the component $T\Delta S$ is the

heat increment due to the processes of digestion, absorption and uptake of nutrients, $[H_iE]$. Therefore,

$$NE_{mg}/I = ME/I - H_iE/I \quad (16)$$

by substituting NE_{mg} by the linear combination between NE_m and RE in the Eq. (14), rearranging the components of Eq. (16) and applying the notation used to describe the food energy in terms of concentration (AFRC, 1993), it is demonstrated that:

$$[ME] = [NE_m] + [RE] + [H_iE] \quad (17)$$

or

$$ME/I = H_eE/I + RE/I + H_iE/I \quad (18)$$

now obeying, differently from the left hand side of Eq. (4), the additive nature established in the first and second laws of thermodynamics.

Conclusions

The use of the function of the total efficiency of the diet metabolizable energy utilization yields reasonable estimates of the available net energy for maintenance and growth for both tropical and temperate conditions. Despite the similar predictive powers between the presented method and the NRC (1996) equations, the former has a lower mean squared deviation even with its more simplified structure.

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In Memoriam

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