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Mechanistic modeling of turkey growth response to genotype and nutrition

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ABSTRACT: Along with the fast genetic improvement, nutritional and environmental effects on poultry growth performance have made it necessary to develop growth models that have the flexibility to adapt to different genotypes and growing conditions. A mechanistic simulation model of energy and nutrient utilization in growing turkeys is presented herein. The model consists of simulating the average homeorhetic and homeostatic regulations associated with the utilization of circulating glucose, fatty acid, AA, and acetyl-CoA for protein and lipid retention in carcass, viscera, and feathers in a turkey population. Homeorhesis plays a major role in the control of protein and lipid turnover for the definition of genetic potential and feed intake, whereas homeostasis adjusts growth rate through protein and lipid turnover rates and, therefore, BW gain and feed intake to the growing conditions. Also, homeostasis enables the maintenance of a dynamic balance state during all the growing period through the control of circulating nutrient concentration. The model was developed and calibrated with experimental data that described en-

ergy utilization in male and female growing turkeys. Then, the ability of the model to adapt to genotypes and to predict the average response of a turkey population to dietary energy was evaluated. Model calibration showed simulations of energy and nutrient utilization that fitted well with the experimental data because ME was satisfyingly partitioned into heat production and energy retention as protein and lipid, and nutrient intake accurately partitioned BW gain into carcass, viscera, and feathers. The evaluation of the model was also satisfactory because BW gain and feed-to-gain ratio were globally in accordance with the observations in different male and female genotypes, in spite of an overestimation of the feed-to-gain ratio during the first weeks of age. Model evaluation showed that the BW gain and feed intake response of growing turkeys to dietary energy was accurately predicted. The model can therefore be used in different growing conditions as it is capable of simulating the growth of different turkey genotypes fed under changing environmental and nutritional contexts.

Key words: energy, growth, model, nutrient utilization, regulation, turkey

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INTRODUCTION

The worldwide competitiveness of the meat poultry industry has made it necessary to develop new methods that relate nutrition to production cost and optimize the feeding strategies and the efficiency of nutrient utilization (Baldwin et al., 1980), according to input ingredients, production cost, and market price of poultry meat. Poultry growth models have been developed over the years to better relate BW gain to feed intake. Most

of these models are based on empirical equations that relate BW or BW gain to time (Buffington et al., 1973; Vitezica et al., 2010) or nutrient intake (Emmans, 1987; King, 2001). A limitation to the use of these models for commercial applications is that empirical equations define nutrient requirements specific to a particular genotype and are not adapted to different environmental conditions (Oviedo-Rondon and Waldroup, 2002). In turkeys, growth model flexibility is essential to account for the effect of sex, genotype, nutrition, and interindividual variability on commercial growth performances.

Metabolic approaches have been suggested as an alternative to simulate growth and nutrient requirements of some genotypes of poultry fed under different nutritional and environmental conditions (Firman, 1994).

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Although no such model seems to be yet available in the poultry literature, several growth models of nutrient metabolism have been developed in other species (Gerrits et al., 1997; Lovatto and Sauvant, 2003; Strathe et al., 2008). We hypothesized that models based on the mechanistic representation of the metabolic regulatory processes are a good means to account for the flexibility in the growth response of commercial turkeys to genotype, nutrition, and environment. The goal of this study was to develop a model capable of simulating the nutritional response of male and female turkey genotypes grown in a controlled environment or in experimental conditions. The study included the following objectives: 1) the development of a teleonomic and mechanistic model of male and female turkey growth; 2) the evaluation of the ability of the model to simulate different turkey genotypes; and 3) the evaluation of the ability of the model to predict the average growth response of a turkey population to nutritional inputs. The ultimate goal of this project is to use the model in different genetic, nutritional, and environmental contexts to adapt nutrient recommendations and better promote the competitiveness and sustainability of turkey production.

MATERIALS AND METHODS

Animal Care and Use committee approval was not obtained for this study because all data used were from previously published studies (i.e., Rivera-Torres et al., 2010b; 2011a,b).

Model Description

General Structure. Our dynamic mechanistic model simulates the average response of male and female commercial turkeys to genotype and nutrition by simulating the metabolic regulations of energy and nutrient utilization from hatching to slaughter age. The turkey is considered as a teleonomic system, organized and regulated to grow and reproduce for the sustainability of its own species. The model differentiates the regulatory processes from the biological response. Two subsystems constitute the model and continuously interact at different levels of organization (Sauvant, 1992): 1) the decisional system defines the homeorhetic (HR) and homeostatic (HS) regulations that drive growth; and 2) the operational system simulates the composition of the whole turkey body as a response to the decisional system.

The decisional system is associated with HR and HS regulations that control the partitioning of nutrient intake among retention, dissipation, and excretion. The HR regulations define the genetic potential, whereas the HS regulations define the actual growth trajectory as the adaptive response of the genetic potential to growing conditions. The growing conditions refer to the nutritional and the ambient constraints that may result in a growth response that deviates from the genetic

potential of the turkey. The nutritional constraints correspond to both the quantity and the nature of nutrient intake. In this model, the ambient constraints are assumed to neither restrict feed intake nor generate any specific response to a stress (i.e., heat stress, or physiological or immunological stress), so the turkey is capable of ingesting the desired amount of feed that allows it to reach its growth trajectory (Emmans, 1987). Feed intake is calculated in the decisional system from the HR and HS regulations that define desired growth. In the operational system, feed intake results in energy and nutrient utilization for the accomplishment of daily BW gain as a result of protein and lipid retention.

The operational system describes the transformation flows of dietary macronutrients intake into intermediary metabolites for protein and lipid retention (Figure 1). The macronutrients are partitioned into carbohydrates, triglycerides, and AA. The indigestible fraction of these macronutrients is eliminated in the excreta (i.e., feces and urine), whereas the digestible fraction enters the circulating glucose (GI), AA, and fatty acid (FA) compartments. The circulating GI compartment represents the major source of energy available in plasma. Each of the circulating AA and FA compartments represents a pivot for N and lipid utilization, respectively. The utilization of GI, AA, and FA results in ATP producing and ATP consuming reactions, of which acetyl-CoA (Ay) is an intermediary metabolite. The Ay is preferred over other circulating metabolites because it is a common intermediate in the utilization of circulating GI, AA, and FA compartments, and it can easily define the ATP production from the different transactions represented in the model. Protein retention corresponds to the difference between the anabolic and the catabolic flows with the circulating AA compartment in carcass, viscera, and feathers (i.e., body component b). Protein retention defines ash and water retention in the body tissues (Emmans, 1989; Eits et al., 2002), and the protein, ash, water, and lipid compartments all constitute the whole body. Similar to protein retention, lipid retention consumes ATP and results from the difference between lipid anabolism and catabolism in carcass and viscera, with the interaction of circulating GI and FA compartments. The partitioning of protein and lipid retention among carcass, viscera, and feathers enables to account for the differences in the growth rates among these body components (Hurwitz et al., 1991). Finally, the operational system is constituted of 11 body compartments associated with the 4 circulating compartments.

Homeorhetic Regulations. The HR regulations constitute “pull” driving forces that define the genetic potential as a dynamic balance state controlled by both protein and lipid retention in the body compartments of the operational system. Protein and lipid retention result from anabolic and catabolic flows (g/d), whereas ash and water retention are defined by allometric relations with protein retention (Emmans, 1989; Gous et al., 1999; Eits et al., 2003). The anabolic and catabolic

Figure 1. Schematic representation of the operational system of the turkey growth model. The dietary nutrients ingested are triglycerides (TG), AA, and carbohydrates (CH). The indigestible fraction is excreted (E) with NH_3 , whereas the digestible fraction is an input to the circulating fatty acids (FA), glucose (Gl), AA, and acetyl-CoA (Ay) compartments in the plasma. Protein and lipid metabolism result in the accretion of protein (P_b), lipid (L_b), ash (A_b), and water (W_b) in the body compartment of component (b) represented by carcass, viscera, and feathers. No lipid compartment is represented in the feathers. Dotted curved lines represent the effect of P_b on the inflows to A_i and W_i .

flows of protein and lipid are controlled by fractional synthesis and degradation rates (**FSR** and **FDR**, respectively; % per day). The daily quantity of protein or lipid retained (R_{Mb} , g/d) is defined as a function of the mass of protein or lipid in the body component b (M_b , g) at time t:

$$R_{Mb}(t) = [\text{FSR}_{Mb}(t) - \text{FDR}_{Mb}(t)]M_b(t).$$

The FSR and FDR were shown to decrease in the muscle of broilers (Figure 2) and turkeys (Kang et al., 1985a), and in the adipose tissue of broilers during growth (Lemarchal et al., 1988; Foglia et al., 1994). To account for the effect of age on these fractional turnover rates, both the FSR and FDR are defined by a declining exponential function of age (Danfaer, 1991; Lovatto and Sauvant, 2003):

$$\text{FSR}_{Mb}(t) = k_s + A_{iMb} \exp(-A_{eMb}t);$$

$$\text{FDR}_{Mb}(t) = k_s + C_{iMb} \exp(-C_{eMb}t),$$

where the k_s , A_{iMb} , C_{iMb} , A_{eMb} , and C_{eMb} parameters constitute inputs to the model. At the adult state (i.e., zero retention), both the FSR and FDR reach the common decreased value k_s . The A_{iMb} (or C_{iMb} , % per day) corresponds to the difference between the initial FSR (or FDR) and k_s . The A_{eMb} (or C_{eMb} , % per day) refers to the rate of change of M_b . Differentiating the FSR and FDR of carcass and viscera protein enables distinguishing the retention rates in these 2 body compo-

nents during growth (Hurwitz et al., 1991). The FSR and FDR of carcass protein are assumed to be less than that in the viscera (Lovatto and Sauvant, 2003). The FSR and FDR of lipid are assumed similar between carcass and viscera because little information is available on the change in lipid turnover with age and on the partitioning of adipose tissue between carcass and viscera. In feathers, protein retention only depends on its FSR because protein retention in feathers refers to an irreversible exportation from the AA pool (i.e., FDR equals zero).

Homeostatic Regulations. Depending on nutrition and management, protein and lipid retention, and thus daily BW gain, may deviate from the BW gain of the genetic potential. To control such deviations, HS regulations interact with HR regulations as “push” driving forces that limit the difference between the actual growth and the balance state (i.e., defined by the HR regulations) and the flows of nutrients among the compartments of the operational system.

The deviation of the growth rate from the balance state is defined by interday HS regulations. These regulations control the growth rate during the whole life of the individual by adapting the fractional turnover rates of protein and lipid to the growing conditions. Results from the literature demonstrated that nutritional levels (Tesseraud et al., 1992, 2001) and dietary macronutrient composition (Collin et al., 2003) may affect the FSR and FDR of protein and lipid in carcass. Also, lipid turnover changes with nutrition, depending on the amount of ME available in excess and depending on the physiological state, as demonstrated by Baldwin et al. (1980) in dairy cows. Thus, we assume 6 parameters related to the interday HS regulations may change according to growing conditions. These parameters refer to the FSR and FDR of protein in carcass (i.e., A_{iPc} ,

Figure 2. Fractional synthesis rate (FSR) of protein in the broiler with age. References mentioned in the legend are 1 = Jones et al. (1986), 2 = Kang et al. (1985b), 3 = Morgan et al. (1989), 4 = Muramatsu and Okumura (1985), 5 = Pym et al. (2004), 6 = Tesseraud et al. (1992), 7 = Tesseraud et al. (1996a), 8 = Tesseraud et al. (1996b), 9 = Tesseraud et al. (1996c), 10 = Tesseraud et al. (2000), 11 = Tesseraud et al. (2001), 12 = Tomas et al. (1991), and 13 = Urdaneta-Rincon and Leeson (2004).

A_{epc} , C_{ipc} , and C_{epc}) and to the FSR of lipid (i.e., A_{il} , and A_{el}). The fractional turnover rate of viscera protein is not affected by nutrition because changes in daily BW gain are assumed to be primarily due to changes in carcass gain (Veldkamp et al., 2005; Aimiwu and Lilburn, 2008). Also, the FDR of lipid is not affected by the interday HS regulations because it is independent of age and its value is small relative to the other fractional turnover rates (i.e., 1% per day). Although the water-to-protein ratio may be affected by nutrition in broilers (contrary to the ash-to-protein ratio; Eits et al., 2002), the interday HS regulations are assumed to have no effect on the allometric relation of water (and ash) with protein. As a consequence, the interday HS regulations are only due to changes in protein and lipid turnover.

In addition to the interday HS regulations, intraday HS regulations simulate the short-term adaptation of the flows of circulating nutrients to the amount and the nature of the ingredients ingested during the day. The intraday HS regulations generate a buffer for the partitioning of the circulating nutrients among the different compartments because they depend on the instantaneous concentration (i.e., the integration step is the minute) of the circulating nutrients (Gill et al., 1989). The intraday HS regulations are represented by exponential laws of response that depend on both the HR and the interday HS regulations, and on the relative difference between the concentration of the circulating nutrient i (in g/L) at time t [$C_i(t)$] and the balance concentration of this circulating nutrient (C_{oi}), defined as $\Delta C_i(t)$ (Lovatto and Sauvant, 2003). The flow resulting from the transformation of the circulating compartment X into Y is expressed in grams of Z (or in moles of Z when mentioned):

$$F_{X,Y/Z}(t) = f_{X,Y/Z}(t) \exp\{Ve[\Delta C_i(t)]\}, \text{ where}$$

$$\Delta C_i(t) = \{[C_i(t) - C_{oi}]/C_{oi}\}.$$

The $f_{X,Y/Z}(t)$ refers to the balance state defined by the HR and the interday HS regulations discussed before. The $F_{X,Y/Z}(t)$ flow equals $f_{X,Y/Z}(t)$ when $\Delta C_i(t)$ equals zero. Similar to a previous version of the model (Rivera-Torres et al., 2010a), the circulating nutrient i creates a return force for X when $\Delta C_i(t)$ differs from zero and ensures a fairly constant concentration of nutrient i by increasing or decreasing $F_{X,Y/Z}(t)$. Positive intraday HS regulations correspond to positive exponential values (Ve) and result in increasing $F_{X,Y/Z}(t)$ when $\Delta C_i(t)$ is greater than zero. On the contrary, negative intraday HS regulations refer to negative Ve values and induce a decrease in the $F_{X,Y/Z}(t)$ when $\Delta C_i(t)$ is greater than zero. Thus, the intraday HS regulations enable the maintenance of a state close to that defined by the HR and interday HS regulations.

Feed Intake. Feed intake results in the transformation of the digestible macronutrients into circulating

GI, AA, FA, or Ay compartments, which can be further used for protein and lipid retention in the body tissues. Feed intake is due to the HR and interday HS regulations that define the desired growth of each day. Because no nutrient is assumed to be limiting in the diet, feed intake regulation depends on dietary ME (Emmans, 1987). Feed intake is therefore calculated as the ME intake to dietary ME ratio. The ME intake [$MEI(t)$, in kJ/d] is driven by the HR and interday HS regulations and is defined as the sum of maintenance energy expenditure (ME_m) and the energy used for desired protein and lipid retention at time t (**REP** and **REL**, respectively), with k_p and k_f the energy efficiencies for REP and REL, respectively:

$$MEI(t) = ME_m(t) + 1/k_p \text{ REP}(t) + 1/k_f \text{ REL}(t).$$

The ME_m is commonly expressed per metabolic BW unit (usually per kilogram $BW^{0.75}$ or per kilogram $BW^{2/3}$ per day). In this study, the ME_m per metabolic BW is a linear function of BW (with a_{ME_m} and d_{ME_m} being, respectively, the intercept and the slope) and depends on sex (Rivera-Torres et al., 2011a):

$$ME_m(t) = [a_{ME_m} + d_{ME_m} BW(t)]BW(t)^{0.75}.$$

The macronutrient inputs are determined from feed intake and from the dietary CP, crude fat, and ME content. Ingested CP and crude fat are associated with AA and triglycerides, respectively, whereas starch, soluble sugars, and soluble fibers intake are assimilated to carbohydrates. The truly digestible fraction of the ingested macronutrients is absorbed by the gut and enters the circulating AA, FA, and GI compartments. All the N is provided by the digestible CP fraction, which is degraded into free AA that constitute inputs to the circulating AA compartment. The triglycerides are degraded into 3 FA and 0.5 GI. Fatty acids and GI constitute inputs to the circulating FA and GI compartments, respectively. The FA are defined as oleic acid equivalents because oleic acid is assumed to be the most concentrated FA in the adipose tissue of poultry, as demonstrated in lean and fat lines of broilers by Lemarchal et al. (1988). Along with the GI that is formed by degradation of triglycerides, the carbohydrates are ingested as GI equivalents to enter the circulating GI compartment. The remaining fraction of ME provided by the diet corresponds to the carbohydrates and is calculated from the energy content of GI and from the difference between ME intake and the energy content in the digestible CP and crude fat intake. Consequently, the macronutrient inputs, and thus feed intake, depend on ME intake and drive BW gain composition as related to protein and lipid turnover.

AA Utilization. The AA circulating compartment is a pivot for protein retention. The inputs to the AA circulating compartment are the digestible AA and the AA resulting from protein catabolism in carcass and

viscera. The dietary AA profile corresponds to the ideal protein ratio (Firman and Boling, 1998). The outputs to the AA circulating compartment are the anabolic flows of protein in carcass, viscera, and feathers; the basal ileal AA endogenous losses (**IEAA**); and the oxidative deamination (i.e., transamination).

Amino acid digestibility depends on the IEAA losses, which correspond to the excretion of protein from mucus, enzyme, and epithelium. To correct AA (or CP) digestibility from these losses, standardized ileal digestibility (**SID**) values are used. The average SID of AA is determined from the feed ingredient composition and from the CP content in the feed ingredients (Sauvant et al., 2004). Defining a SID has the advantage over the apparent digestibility in that it accounts for the effect of DM intake on the IEAA (Adedokun et al., 2007a). Although the IEAA losses may vary with CP and age, particularly during the first week of age (Adedokun et al., 2007b), the IEAA losses were defined as a constant function of DMI.

Oxidative deamination is due to inevitable oxidation losses and to the transformation of the excess of AA into a carbon chain for ATP production, and an amine group for further synthesis of essential AA or rejection as ammonia (or uric acid). The excess of AA corresponds to the difference between digestible CP intake, IEAA losses, and minimal AA oxidation. The carbon chain that is liberated after oxidative deamination, results in either Ay or Gl formation by intraday HS regulations. The oxidative deamination [AAd(t), in g/d] is positively related to $\Delta C_{AA}(t)$:

$$AAd(t) = k_{ox} AA(t) \exp\{Ve_{AAox}[\Delta C_{AA}(t)]\}.$$

The k_{ox} value refers to the minimal AA oxidation rate (in %), and AA(t) corresponds to the mass (in g) of the circulating AA compartment at time t. A deficiency in the concentration of circulating Gl results in increasing inputs of Gl from the circulating AA compartment, whereas the reverse is true in case of an increased concentration in the Gl compartment:

$$F_{AA,Gl/AA}(t) = AAd(t) \exp\{-Ve_{Gl}[\Delta C_{Gl}(t)]\}.$$

Based on the determination of the rate of oxidative deamination and of the quantity of AA transformed into Gl equivalents, the quantity of AA that results in Ay formation ($F_{AA,Ay/AA}$) is calculated:

$$F_{AA,Ay/AA}(t) = AAd(t) - F_{AA,Gl/AA}.$$

Gl and FA Utilization. Glucose and FA utilization enable lipid retention. The inputs to the circulating Gl compartment are the macronutrients absorbed as Gl equivalents, Gl originating from the catabolic flow of lipid into Gl and FA, and oxidative deamination flow of AA that are transformed into Gl equivalents. The outputs to the circulating Gl compartment are the Gl that

enters the lipid compartments, and Gl oxidation flow into Ay ($F_{Gl,Ay/Gl}$). Glucose oxidation depends on the intraday HS regulations defined by $\Delta C_{Gl}(t)$. Increasing Gl oxidation into Ay results from increasing circulating Gl concentration (i.e., Ve_{Glox} is greater than zero):

$$F_{Gl,Ay/Gl}(t) = k_{ox} Gl(t) \exp\{Ve_{Glox} [\Delta C_{Gl}(t)]\}.$$

The inputs to the FA circulating compartment are the ingested FA originating from triglycerides, the de novo lipogenesis of FA from Ay ($F_{Ay,FA/FA}$), and the FA formed by the degradation of lipid in carcass and viscera. Glucose is at the origin of the intraday HS regulations of the flows related to Ay compartment because FA synthesis is assumed to be primarily due to increased concentrations of circulating Gl:

$$F_{Ay,FA/FA}(t) = k_{ox} FA(t) \exp\{Ve_{Gl}[\Delta C_{Gl}(t)]\}.$$

The outputs to the circulating FA compartment are represented by the anabolic flows of lipid in body component b and by the oxidation flow of FA into Ay ($F_{FA,Ay/FA}$). The $F_{FA,Ay/FA}$ increases with increasing FA mass and circulating FA concentration:

$$F_{FA,Ay/FA}(t) = k_{ox} FA(t) \exp\{Ve_{FAox} [\Delta C_{FA}(t)]\}.$$

Thus, the circulating FA concentration is at the origin of FA oxidation, whereas the circulating Gl concentration regulates both Gl oxidation into Ay and de novo lipogenesis.

Energy Utilization. Metabolizable energy utilization corresponds to the partitioning of ME intake between heat production (**HP**) and energy retention as protein and lipid (kJ/d). The ME intake is determined by multiplication of the feed intake and the dietary ME value. Heat production refers to the ATP used during the endogenous reactions (i.e., ATP consuming reactions; Figure 1) and is calculated as the difference between ME intake and energy retention. Heat production is expected to equal the amount of energy produced during the exogenous reactions (i.e., ATP producing reactions; Figure 1), resulting in a zero energy balance.

The exogenous reactions are simulated mechanistically. They correspond to the oxidation of AA, Gl, and FA into Ay, to the Ay oxidation into CO_2 , and to the lipid catabolism into FA. The oxidation of Gl and AA into Ay (**OX_{Gl}** and **OX_{AA}**, respectively) is determined from the energy costs (i.e., as ATP equivalents) associated with the transformation of circulating Gl and AA into Ay, respectively. The **OX_{FA}** corresponds to the sum of the energy produced during FA oxidation into Ay and during lipid catabolism into FA and Gl. Finally, the **OX_{Ay}** is determined by difference between total HP and the sum of **OX_{Gl}**, **OX_{AA}**, and **OX_{FA}** to result in a zero energy balance between the endogenous and the exogenous reactions.

Model Development

Implementation. The model was developed with a software (Modelmaker version 3.0, Cherwell Scientific Publishing Inc., Palo Alto, CA). The model outputs were calculated on a daily basis using the Runge-Kutta method with a fixed integration step of 1 min (1,440 integration steps per day), which enabled accounting for the intraday HS regulations. Model integration started at hatching ($t = 0$). The mathematical statement of the model is presented in Table A1.

Parameters and Initial Values. The parameters were differentiated between user-defined, fitted, and constant parameters (Table 1). The user-defined parameters corresponded to the parameters that were adapted to every simulation. They referred to the dietary DM, ME, crude fat, and CP contents, and to the ages at feed transition. All the parameter values were first defined during calibration. The defined and constant parameters were not changed during evaluation, whereas some fitted parameters were adjusted during model evaluation. The constant parameters were taken from the literature and were assumed not to change between individuals of a same sex. They corresponded to stoichiometry parameters, digestibility values, the ATP produced during exogenous reactions, and the parameters related to the definition of ME intake (Table 2).

The initial values (i.e., day of hatching) of the composition in protein, lipid, ash, and water were taken from the serial slaughter analyses performed at 7 d of age (Rivera-Torres et al., 2011b). The initial values of the circulating compartments were determined from the balance concentration of the circulating nutrients and from the plasma volume of the turkey at hatching, assuming an initial BW of 60 g in both males and females.

Model Calibration

The calibration was performed to determine the values of the fitted parameters to validate the adequacy of the structure and functions of the model with observed data. To accomplish this goal, time series results from 3 experimental studies (Rivera-Torres et al., 2010b, 2011a, b) were used as observed values for calibration procedure.

Experimental Data

The model was calibrated on experimental data of male and female turkeys. For each of the 3 experimental studies used for calibration, the turkeys were grown under a controlled environment and fed standard diets; thus no environmental constraint was assumed to limit the expression of genetic potential. Two experimental studies consisted in energy and N balances that were carried out by indirect calorimetry on male medium-size turkeys (Rivera-Torres et al., 2010b) and males and females from a heavy strain (Rivera-Torres

et al., 2011a) to partition ME intake into HP and energy retention as protein and lipid. The third study corresponded to serial slaughter analyses performed in medium-size male turkeys (Rivera-Torres et al., 2011b) to determine the contribution of carcass, viscera, and feather in total protein, lipid, ash, and water retention. The user-defined parameters (i.e., age at feed transition, and dietary DM, ME, CP, and crude fat contents) were taken from Rivera-Torres et al. (2011a).

The experimental data from the serial slaughter analyses corresponded to the observed mass of carcass, viscera, and feather in male turkeys and to the observed mass of protein, ash, and water in the carcass with viscera and in the feathers, along with the lipid mass in carcass with viscera (g). The experimental data from the energy and N balances corresponded to the observed BW (g), feed intake (g/d), O_2 consumption (L/d), CO_2 production (L/d), respiratory quotient, ME intake (kJ/d), HP (kJ/d), AA ($N \times 6.25$) intake, excretion in droppings (g/d), and protein and lipid retention (g/d). In addition, the oxidation rates of AA (OX_P), carbohydrates (OX_{CHO} , as Gl equivalent), and fat (OX_F) into CO_2 were calculated as suggested by Chwalibog and Thorbek (1999) in laying hens, assuming a zero egg production; the OX_F , OX_{CHO} , and OX_P values calculated from the energy and N balances were used as observed values and compared with the simulated OX_F , OX_{CHO} , and OX_P values. The simulated OX_F , OX_{CHO} , and OX_P values were calculated from the simulated OX_{AA} , OX_{FA} , OX_{Gl} , and OX_{Ay} by attributing the OX_{Ay} to OX_{CHO} , OX_F , or OX_P , depending on the energetic contribution of OX_{AA} , OX_{FA} , and OX_{Gl} in the production of Ay:

$$OX_{CHO} = OX_{Gl} \times \left[1 + \frac{1}{(OX_{Gl} + OX_{FA} + OX_{AA})} OX_{Ay} \right];$$

$$OX_F = OX_{FA} \times \left[1 + \frac{1}{(OX_{Gl} + OX_{FA} + OX_{AA})} OX_{Ay} \right]; \text{ and}$$

$$OX_P = OX_{AA} \times \left[1 + \frac{1}{(OX_{Gl} + OX_{FA} + OX_{AA})} OX_{Ay} \right].$$

Calibration Procedure

Model calibration consisted in defining the parameter values of both male and female growth profiles as a basis for further model evaluation of the effect of genotype and nutrition on each sex. Model calibration was mostly performed manually. Although labor intensive, manual iterations seemed to be most adapted to compare the model outputs with the large number of experimental data because successive computer-based optimizations of 1 or few parameters did not result in the optimum of all the parameter estimates. In addi-

Table 1. Definition of the constant and fitted and user-defined parameters used in the turkey growth model

Item	Definition	Unit	Parameter		
			Constant	Fitted	User-defined
a_{ME_m}	Intercept of the linear regression of maintenance energy expenditure per metabolic BW unit	kJ/kg of BW ¹ per day	*		
AeL	Rate of change of lipid anabolism	%/d		*	
AePc	Rate of change of protein anabolism in carcass	%/d		*	
AePf	Rate of change of protein anabolism in feathers	%/d		*	
AePv	Rate of change of protein anabolism in viscera	%/d		*	
a_{IEAA}	Ileal AA endogenous losses	g/kg of DMI per day		*	
AiL	Difference between the initial fractional synthesis rate of lipid at time $t = 0$ and the adult state	%/d		*	
AiPb	Difference between the initial fractional synthesis rate of protein at time $t = 0$ and the adult state in body component b	%/d		*	
a_{PX}	Ash content relative to protein in X	%	*		
a_W	Parameter for the relation of water with protein in carcass and viscera ¹	%	*		
$Ay_{X,Y}$	Number of acetyl-CoA produced for the transformation of X into Y	mol	*		
b	Scaling parameter of BW		*		
B_V	Blood volume	% of BW	*		
b_W	Power value for the allometric relation of water with protein in carcass and viscera	—	*		
CeL	Rate of change of lipid catabolism	%/d		*	
CePb	Rate of change of protein catabolism in body component b	%/d		*	
CiL	Difference between the initial fractional degradation rate of lipid at time $t = 0$ and the adult state	%/d		*	
C_{iPb}	Difference between the initial fractional degradation rate of protein at time $t = 0$ and the adult state in body component b	%/d		*	
CoX_0	Balance concentration of X in plasma	g/mol	*		
d_{ME_m}	Slope of the regression of maintenance energy expenditure per metabolic BW unit in turkeys ¹	kJ/kg of BW/kg of BW ^{0.75} /d	*		
EX	Energy content of X	kJ/mol or kJ/g	*		
$FA_{L,FA}$	Number of moles of fatty acids produced for the transformation of lipid into fatty acids	mol	*		
feedage ²	Age at feed transition of feed i	%			*
feedCPi ²	CP content in feed i	%			*
feedCFi ²	Crude fat content in feed i	%			*
feedDM ²	DM content in feed i	%			*
feedMEi ²	ME content in feed i	%			*
$Gl_{X,Y}$	Number of moles of glucose produced for the transformation of X into Y	mol	*		
kf	Efficiency of energy utilization for lipid retention	%	*		
kox	Basal rate of oxidation of circulating glucose, fatty acid, and AA	%	*		
kp	Efficiency of energy utilization for protein retention	%	*		
ks	Basal fractional turnover rate	%/d	*		
MM_X	Molar mass of X	g/mol	*		
MV_X	Volume mass of X	g/L	*		
$NH3_{AA,NH2}$	Number of moles of NH ₂ produced for the transformation of AA in NH ₂	mol	*		
PoB	Amount of plasma in blood	% of blood volume	*		
SID_X	Standardized ileal digestibility of X	%	*		
$Ve_{X_{ox}}$	Exponential value for the oxidation of X for intraday homeostatic regulations	—		*	
Ve_{GI}	Exponential value of nutrient metabolism for the intraday homeostatic regulations driven by glucose	—		*	
Ve_{LT}	Exponential value of lipid turnover for intraday homeostatic regulations	—		*	
Ve_{PT}	Exponential value of protein turnover for intraday homeostatic regulations	—		*	
w_{Pf}	Water content in feather protein	%	*		

¹The value may differ depending on sex.²The number of parameters equals the number of feeds.

Table 2. Constant parameters defined in the turkey growth model

Constant parameter	Value ¹	References
a_{IEAA}	6.5	Adedokun et al. (2007b)
a_{MEem}	628	Rivera-Torres et al. (2011a)
a_{PCV}	17	Rivera-Torres et al. (2011b)
a_{Pf}	1.5	Rivera-Torres et al. (2011b)
$\text{ATP}_{\text{AA,Ay}}$	14	Gerrits et al. (1997)
$\text{ATP}_{\text{Ay,CO}_2}$	12	Gerrits et al. (1997)
$\text{ATP}_{\text{Ay,FA}}$	1.13	Gerrits et al. (1997)
$\text{ATP}_{\text{FA,Ay}}$	36	Gerrits et al. (1997)
$\text{ATP}_{\text{Gl,Ay}}$	14	Gerrits et al. (1997)
ATP_{La}	-10	Gerrits et al. (1997)
ATP_{Lc}	2.26	Gerrits et al. (1997)
ATP_{Pa}	-5	Gerrits et al. (1997)
ATP_{Pc}	-1	Gerrits et al. (1997)
a_{W}	290/341	Rivera-Torres et al. (2011b)/Emmans (1989)
$\text{Ay}_{\text{AA,Ay}}$	2.5	Gerrits et al. (1997)
$\text{Ay}_{\text{FA,Ay}}$	9	Gerrits et al. (1997)
$\text{Ay}_{\text{Gl,Ay}}$	2	Gerrits et al. (1997)
b	0.75	Rivera-Torres et al. (2010b)
b_{W}	1/0.90	Rivera-Torres et al. (2011b)/Emmans (1989)
B_{V}	7.2	McCartney (1952)
C_{AA0}	0.7	Geraert et al. (1987)
C_{FA0}	0.85	Sato et al. (2008)
C_{G10}	2	Geraert et al. (1987)
$\text{CO}_2_{\text{Gl,Ay}}$	2	van Milgen (2002)
$\text{CO}_2_{\text{Ay,CO}_2}$	2	van Milgen (2002)
d_{MEem}	-5/-13	Rivera-Torres et al. (2011a)
E_{ATP}	79.5	Campbell and Reece (2005)
E_{Gl}	15.7	Campbell and Reece (2005)
E_{L}	39.6	Rivera-Torres et al. (2010a)
E_{P}	23.7	Rivera-Torres et al. (2010a)
$\text{FA}_{\text{L,FA}}$	3	Gerrits et al. (1997)
$\text{Gl}_{\text{AA,Gl}}$	0.83	Gerrits et al. (1997)
$\text{Gl}_{\text{FA,Gl}}$	2.67	Gerrits et al. (1997)
$\text{Gl}_{\text{L,Gl}}$	0.5	Gerrits et al. (1997)
k_{f}	96	Rivera-Torres et al. (2011a)
k_{ox}	2	Gerrits et al. (1997)
k_{p}	65	Rivera-Torres et al. (2011a)
k_{s}	1	Lovatto and Sauvant (2003)
MM_{AA}	125	Campbell and Reece (2005)
MM_{CO_2}	48	Campbell and Reece (2005)
MM_{FA}	282	Campbell and Reece (2005)
MM_{Gl}	180	Campbell and Reece (2005)
MM_{N}	14	Campbell and Reece (2005)
MM_{NH_2}	16	Campbell and Reece (2005)
MM_{O_2}	32	Campbell and Reece (2005)
MM_{TG}	884	Campbell and Reece (2005)
MV_{CO_2}	1.87	Campbell and Reece (2005)
MV_{O_2}	1.35	Campbell and Reece (2005)
$\text{NH}_2_{\text{AA,NH}_2}$	1.4	Campbell and Reece (2005)
$\text{O}_{2\text{Gl,Ay}}$	2	van Milgen (2002)
$\text{O}_{2\text{Ay,CO}_2}$	2	van Milgen (2002)
PoB	55	Lovatto and Sauvant (2003)
SID_{AA}	90	Sauvant et al. (2004)
SID_{CF}	85	Honda et al. (2009)
w_{Pf}	12.7	Rivera-Torres et al. (2011b)

¹Parameter values differ between sexes (male/female).

tion, the experimental data had to be interpreted individually rather than being considered as a set of average observed values with SD (i.e., protein retention rate in males was that of both medium-size and heavy genotypes, whereas protein mass of carcass with viscera was only that of a medium-size genotype).

The values of the HR and interday HS regulation parameters were first defined on male and female growth profiles, whereas no intraday HS regulation was considered. The results from the meta-analysis of the FSR and FDR of protein in broilers were used as a basis for calibrating the parameters associated with protein

turnover. Similar to the FSR and FDR of lipid, the FSR of both feather and viscera protein were assumed independent of sex. The initial synthesis and degradation rates of protein were assumed to be 2- to 3-fold greater in viscera than in carcass, whereas the initial synthesis rate of protein in feathers was assumed to be less than that in carcass. Also, because of the earlier development of viscera relative to carcass (Hurwitz et al., 1991), the FSR and FDR of viscera protein were assumed to decrease faster than those of carcass protein.

On the basis of the estimates of the parameters associated with the HR and interday HS regulations, the intraday HS regulation parameters were calibrated. These parameters were associated with the metabolic regulations of circulating nutrient concentrations; thus, they were assumed to be independent of sex and genotype. The observed OX_{CHO} , OX_F , and OX_P values were used for manual calibration of the Ve_{Glox} , Ve_{FAox} , and Ve_{AAox} parameters, respectively. It was assumed that the main driving forces associated with intraday HS regulations corresponded to the oxidation of circulating nutrient (i.e., AA, Gl, and FA). Thus, the Ve_{Glox} , Ve_{AAox} , and Ve_{FAox} parameters were expected to have the greater values (i.e., greater than unity), whereas the Ve_{PT} and Ve_{LT} parameters were assumed to be less than unity. Finally, the Ve_{PT} and Ve_{LT} parameters were calibrated ultimately to maximize the goodness of fit of protein and lipid retention in both male and female growth profiles.

Sensitivity Analyses. The sensitivity analysis was performed on constant and fitted parameters to allow identifying the parameters that had a major influence on the model outputs. The analysis was performed on the parameters that regulate BW gain composition. Also, all the fitted parameters associated with the intraday HS regulations were analyzed because (contrary to the interday HS regulations) their values could not be calibrated or validated with biological observations. The sensitivity analysis evaluated the effect of a 5% (i.e., parameters less than unity) or 15% (i.e., parameters greater than unity) increase and decrease of the reference parameter values on major model outputs (e.g., protein retention, lipid retention, BW gain, and feed intake).

Statistical Analyses. The statistical analyses were performed to determine the accuracy of the model and to estimate the bias of simulation. The accuracy of the model defined how close the simulated values were to the observed values. The model was expected to be accurate from both a technical viewpoint (e.g., feed-to-gain ratio, BW gain) and a nutritional one (e.g., nutrient oxidation, CO_2 production). To observe the accuracy of the model at the different levels of organization of the turkey, the observed values (Y-variate) were plotted against the simulated values (X-variate) of the different outputs of the model. The simulated values were plotted on the X-variate because they corresponded to deterministic values, whereas the observed values were stochastic (i.e., replicate measurements). A linear

regression was performed to estimate the intercept and the slope (b value) of the linear regression of the Y-variate on the X-variate. Accurate simulations should result in an intercept and a regression slope close to zero and unity, respectively, and in a high coefficient of determination.

To evaluate the bias of simulation, the mean square prediction error (**MSPE**) was calculated. The MSPE was defined as follows (Tedeschi, 2006):

$$MSPE = (\bar{X} - \bar{Y})^2 + \delta_X^2 \times (1 - b)^2 + (1 - r^2) \times \delta_Y^2.$$

The first term of this equation $\left[(\bar{X} - \bar{Y})^2 \right]$ represented the mean bias of the simulated values and indicated an eventual over or underestimation of the observations. The second term $\left[\delta_X^2 \times (1 - b)^2 \right]$ corresponded to the bias of regression (i.e., deviation of the slope from unity), whereas the third term referred to the random errors unrelated to the error of simulation. The root MSPE was used so that it was expressed as the same unit as the observed values. Also, each of the 3 terms of the MSPE was expressed as a percentage of the total MSPE to interpret the source of error. Accurate simulations were expected to have a small MSPE (i.e., or root MSPE) and small mean bias and bias of regression.

Model Evaluation

Response to Genotype. Based on the average male and female growth profiles, simulations were performed on different genotypes of both sexes to evaluate the flexibility of the model to adapt to different genotypes. Similar to the male and female growth profiles defined during calibration, we assumed similar fractional turnover rates of protein in viscera and feathers among turkey individuals. Also, the FDR of lipid was assumed independent of genotype. The adjustment among genotypes of a same sex, therefore, included the Ai_{PC} , Ae_{PC} , Ci_{PC} , Ce_{PC} , Ai_L , and Ae_L parameters. The genotypes of the male (Nicholas 700, Aviagen Turkeys Ltd., Tattenhall, Cheshire, UK) and the female (Hybrid Converter, Hybrid Turkeys, Kitchener, Ontario, Canada) strains were used for evaluation. Feed characteristics, daily BW gain (g/d), feed intake (g/d), and feed conversion (g/g) expectations were those of the breeding company. The results from the evaluation were performed following the statistical analyses described in model calibration.

Response to Nutrition. The accuracy of the model to predict the response to changing energy or nutrient content was tested by comparing the model simulations with experimental data. The results from the measurements made on male turkeys from 4 to 16 wk of age (Veldkamp et al., 2005) were used for this evaluation. To use the model as a prediction tool of the response to nutrition, we first performed an adjustment of the growth profile to the daily BW gain, feed intake,

and feed-to-gain ratio measured in the control treatment [E100, with Lys content at 105% of NRC (1994) recommendations]. We assumed that all the turkeys were given a control diet that corresponded to 0 to 4 wk of age (NRC, 1994).

The model predictions consisted of comparing the predicted outputs with the observed daily BW gain, feed intake, and feed-to-gain ratio of the treatments, after modification of the user-defined parameters associated with the feed characteristics (i.e., ME levels). The treatments tested corresponded to 90 and 110% of the ME of the control treatment (i.e., treatments E90 and E110, respectively). Statistical analyses were performed as described in model calibration.

RESULTS

Model Calibration

Model Behavior. The results from the calibration agreed with the observed values from all 3 experimental studies. The modeling of energy and nutrient utilization allowed us to simulate the change in the retention rates of protein, lipid, ash, and water in carcass, viscera, and feathers of male and female turkeys during growth. As expected, carcass gain was greater than viscera gain. The growth rate of viscera was maximized at 6 wk of age in both male and female turkeys, and it then transiently stabilized after 11 wk of age when the carcass growth rate was maximized (Figure 3).

The calibration on both sexes was a means to define some differences in the outcomes of HR and HS regulations among males and females. Female turkeys had a decreased carcass gain but a greater viscera gain in comparison with males because of the greater mass of viscera protein relative to the mass of carcass protein.

Male turkeys deposited more protein and less lipid than females, resulting in a decreased feed-to-gain ratio, as indicated by the observed values. The greater protein retention and the reduced lipid retention in male turkeys were due to the greater FSR of protein and to the decreased FSR of lipid in comparison with females. Also, the FDR of carcass protein in females was less than that in males. In addition to the maintenance energy expenditure and the allometric relation of water with protein (i.e., d_{ME_m} and W_{PCV} were constant parameters), 6 fitted parameters related to the HR and interday HS regulations characterized the difference between the growth profiles of males and females (Table 3). These parameters corresponded to the FSR and FDR of carcass protein (i.e., Ai_{Pc} , Ae_{Pc} , Ci_{Pc} , and Ce_{Pc} , respectively) and the FSR of lipid (i.e., Ai_L and Ae_L) as defined during model calibration.

The fitted parameters associated with protein turnover in carcass, viscera, and feathers resulted in FSR and FDR estimates of carcass and viscera protein that were close to those measured by Kang et al. (1985a) in the breast muscle of turkeys after 2 wk of age (Figure 4). However, the simulated FSR of protein in carcass before 2 wk of age seemed to be less than those measured by the same authors in breast and thigh of turkeys. Also, the FDR of carcass determined by the model was similar to turkeys younger than 8 wk of age as reported by Kang et al. (1985a).

The intraday HS regulations ensured the maintenance of a constant concentration of circulating nutrients, primarily through the regulation of nutrient oxidation (i.e., Ve_{Glox} , Ve_{AAox} , Ve_{FAox}), and partly through the control of protein and lipid turnovers in carcass and viscera (i.e., Ve_{PT} and Ve_{LT}), as illustrated by the greater Ve_{Glox} , Ve_{AAox} , and Ve_{FAox} values in comparison with the Ve_{PT} and Ve_{LT} values (Table 3). The Ve_{Glox}

Figure 3. Carcass and viscera growth rates in male and female growing turkeys with age.

Figure 4. Fractional synthesis rate (FSR) of protein in carcass, viscera, and feathers with age in male growing turkeys (Kang et al., 1985a).

parameter was greater than Ve_{AAox} and Ve_{FAox} because the OX_{CHO} determined from the experimental data was greater than OX_P and OX_F . The Ve_{LT} parameter estimate was greater than Ve_{PT} because lipid turnover was greater than Ve_{PT} . Also, the energy provided in excess was primarily stored as lipid, whereas most of the AA in excess were deaminated and used as energy source. The adaptation of protein turnover to the circulating AA concentration intervened as a minor regulation be-

cause it was limited by the HR and interday HS regulations, which both defined protein retention rate on a long-term scale.

Accuracy and Bias of Model Outputs. The accuracy of the technical outputs of the model was illustrated by increased coefficients of determination for daily BW gain, feed intake, N utilization, and energy utilization in male and female turkeys. The MSPE were small in both males and females relative to the mean observed values. The MSPE were globally greater in males in comparison with females, averaging 25 and 10% of the mean observed values of males and females, respectively. The major simulation errors were associated with daily BW gain, lipid retention and mass, and feather mass components in male turkeys (Table 4). However, similar to the other simulated values, the MSPE were mostly due to unexplained variations, which means that the mean bias and regression bias remained low relative to the variability of the observed values. The unexplained variations exceeded 50% of the MSPE, except for simulated HP in females (Table 5), which showed a consistent underestimation (i.e., 55.0% of the MSPE was associated with mean bias). Similar observations were made when simulating HP in male turkeys.

In males, protein and lipid retention showed a consistent underestimation (i.e., increased mean bias relative to total MSPE), whereas these outputs were accurately

Table 3. Fitted parameters associated with protein and lipid turnover in male and female growing turkeys¹

	Value		
Parameter	Male	Female	Regulation ²
Fractional rate of change, %/d			
Ae _L	2.0	1.9	HR/HS _d
Ae _{Pc}	2.2	2.8	HR/HS _d
Ae _{Pf}	6.4		HR
Ae _{Pv}	5.3		HR
Ce _L	2.8		HR
Ce _{Pc}	2.3	4.0	HR/HS _d
Ce _{Pv}	6.3		HR
Difference in fractional turnover rate between the initial state and the adult state, %/d			
Ai _L	10.0	12.4	HR/HS _d
Ai _{Pc}	27.3	25.4	HR/HS _d
Ai _{Pf}	21.0		HR
Ai _{Pv}	55.0		HR
Ci _L	2.0		HR
Ci _{Pc}	18.0	17.0	HR/HS _d
Ci _{Pv}	50.0		HR
Intraday homeostatic regulations			
Ve _{AAox}	3.0		HS _m
Ve _{FAox}	7.0		HS _m
Ve _{Gl}	0.2		HS _m
Ve _{Glox}	10.0		HS _m
Ve _{LT}	0.1		HS _m
Ve _{PT}	0.03		HS _m

¹Values determined from the model calibration on the experimental values defined by Rivera-Torres et al. (2010b, 2011a,b).

²HR = homeorhetic; HS_d = interday homeostatic; and HS_m = intraday homeostatic.

Table 4. Mean observed values, coefficient of determination (R^2), root mean square prediction error (MSPE), and decomposition of MSPE of simulated outputs in male turkeys until 23 wk of age

Item	n	Mean observed value	R^2 , %	Root MSPE	Mean bias, % MSPE	Regression bias, % MSPE	Unexplained variation, % MSPE
Rivera-Torres et al. (2010b, 2011a)							
Daily BW gain, g/d	36	146	47.3	48	3.2	12.5	84.4
Feed intake, g/d	36	397	93.3	49	1.2	0.0	98.8
Feed-to-gain ratio, g/g	36	2.762	73.0	0.861	11.3	18.5	70.1
Protein retention, g/d	36	38.5	80.5	7.0	30.8	0.3	68.8
Lipid retention, g/d	36	29.1	76.0	12.2	30.6	11.7	57.8
N excretion, g/d	36	7.4	91.0	1.1	12.5	0.6	86.9
Heat production, kJ/d	36	3,506	97.1	358	23.8	0.2	76.0
CO ₂ production, L/d	36	163	96.9	16	0.7	7.4	91.9
Rivera-Torres et al. (2010b)							
Carcass mass, g	32	5,325	97.6	813	0.0	29.8	70.2
Viscera mass, g	32	580	89.7	144	3.4	9.7	86.9
Feather mass, g	32	188	86.7	64	15.5	21.0	63.5
Protein in carcass with viscera, g	32	1,262	96.8	223	3.4	31.6	65.0
Lipid in carcass with viscera, g	32	623	80.7	330	0.5	4.5	95.0
Water in carcass with viscera, g	32	3,708	97.3	598	1.1	33.4	65.5
Ash in carcass with viscera, g	32	217	96.6	38	1.2	31.4	67.4
Protein in feathers, g	32	166	86.3	57.1	16.9	20.3	62.8
Water in feathers, g	32	22.2	87.0	8.9	18.7	31.3	50.0
Ash in feathers, g	32	2.3	82.4	0.6	0.9	0.7	98.4

simulated in females (i.e., the root MSPE equaled 16 and 48 g/d in females and males, respectively). As a result of the underestimation of protein and lipid retention in males, BW gain was globally underestimated until 150 g/d. Thereafter, it was underestimated in heavy male turkeys, whereas it was overestimated in medium-size turkeys (Figure 5).

Sensitive Parameters. The sensitivity analysis showed that the parameters defining ME intake regulation (i.e., a_{MEm} , k_p , and k_f) and carcass protein turnover (i.e., A_{Pc} , C_{Pc} , A_{ePc} , and C_{ePc}) had a major effect on the model outputs (Table 6). As expected, increasing the a_{MEm} value by 15% resulted in increased ME intake (and thus feed intake) because of greater maintenance energy expenditure. On the contrary, increasing the k_p or k_f value resulted in a decrease of both ME and feed intake because of the decreased heat production associated with protein and lipid retention.

The parameters associated with carcass protein turnover affected both BW gain and feed intake (Table 6). Increasing the A_{Pc} parameter by 5% resulted in increased anabolic flows of protein in carcass (i.e., +89.1 g/d), and thus in increased BW gain (i.e., +60.9 g/d) and feed intake (i.e., +207.7 g/d). On the contrary, increasing the C_{Pc} parameter by 5% resulted in increased catabolic flow of carcass protein, and thus decreased protein retention, BW gain, and feed intake by 12.9, 57.3, and 89.6 g/d, respectively. Similarly, the A_{ePc} and C_{ePc} parameters strongly affected BW gain and feed intake by modifying the protein retention rate in carcass. In comparison with protein turnover, changes in the parameters associated with lipid turnover did not result in a major effect on lipid retention and feed intake.

The V_e parameters associated with the intraday HS regulations affected the growth response of the turkey to genotype and nutrition because these parameters were at the origin of the daily partitioning of the nutrients ingested among retention as protein and lipid, and dissipation as heat of AA, FA, and Gl origin. Increasing or decreasing the values of the V_e parameters by 15% mostly affected protein retention and thus BW gain and feed intake (Table 7) during all the observed growing period. More specifically, increasing the V_{eGlox} , V_{eAAox} , or V_{eFAox} parameter resulted in an increase in the corresponding oxidation rate (i.e., OX_{CHO} , OX_P , or

Figure 5. Observed and simulated BW gain (g/d) of male turkeys (Rivera-Torres et al., 2010b, 2011a). The straight regression line ($Y = X$) represents the expected simulated values for which observed values (Y) equal simulated values (X).

Table 5. Mean observed values, coefficient of determination (R^2), root mean square prediction error (MSPE), and decomposition of MSPE of simulated outputs in female turkeys from 0 to 23 wk of age (Rivera-Torres et al., 2011a)

Item	n	Mean observed value	R^2 , %	Root MSPE	Mean bias, % MSPE	Regression bias, % MSPE	Unexplained variation, % MSPE
Daily BW gain, g/d	19	147	92.5	16	3.2	12.0	84.7
Feed intake, g/d	19	368	98.5	23	0.9	25.8	73.3
Feed-to-gain ratio, g/g	19	2.461	78.1	0.479	12.6	10.5	76.9
Protein retention, g/d	19	28.4	96.6	2.3	0.4	24.6	75.0
Lipid retention, g/d	19	36.6	96.6	4.6	0.7	25.1	74.2
N excretion, g/d	19	7.3	93.6	0.9	3.2	16.1	80.7
Heat production, kJ/d	19	2,764	99.3	155	55.0	3.6	41.4
CO ₂ production, L/d	19	132	99.2	7.9	41.3	10.2	48.6

OX_F, respectively) at the expense of protein and lipid retention, and thus feed intake.

The parameters associated with AA oxidation (i.e., Ve_{AAox}) and protein turnover (i.e., Ve_{PT}) had the greater effects on the model outputs. An increase in the Ve_{AAox} parameter resulted in a decrease in daily BW gain, mostly because of a decreased protein retention, which was due to a greater oxidation of the AA in excess and a decreased amount of AA integrated in protein turnover (i.e., greater Ve_{AAox} -to- Ve_{PT} ratio). As a consequence of the decreased protein retention (and BW gain), feed intake was also decreased. An increase in the Ve_{PT} parameter was associated with greater protein retention due to an increase in protein synthesis and a decrease in protein degradation. Consequently, increasing the priority for protein turnover relative to nutrient oxidation resulted in a greater BW gain and feed intake.

Model Evaluation

Response to Genotype. The application of the model on different genotypes showed its ability to adapt to both male and female growth trajectories. The

growth trajectories of the male (Nicholas 700) and the female (Hybrid Converter) strains were globally in accordance with the observed daily BW gain, feed intake, and feed-to-gain ratio. The coefficients of determination were greater in the simulations performed in females in comparison with males (Table 8). In females, the root-MSPE of BW gain and feed intake were both less than those observed in males (i.e., 4 and 23 vs. 10 and 46 g/d, respectively). The greater error of simulation was observed for the feed-to-gain ratio because it cumulated the simulation errors of both BW gain and feed intake. In both sexes, the errors of simulation of BW gain were mostly due to unexplained variation; the regression bias represented more than 55% of the MSPE of feed intake and feed-to-gain ratio.

In the genotype of males (Nicholas 700), the regression bias was due to an overestimation of the feed-to-gain ratio at early ages and to an underestimation at later ages (i.e., after 11 wk of age; Figure 6). The error of simulation (i.e., difference between simulated and observed values) of the feed-to-gain ratio averaged 0.004 g/g between 1 to 22 wk of age. The error of simulation of the feed-to-gain ratio of the male genotype was particularly large before 5 wk of age, averaging 0.396 g/g.

Table 6. Sensitivity analysis of protein retention, lipid retention, BW gain, and feed intake (g/d) to the fitted parameters associated with HR regulations in male growing turkeys at 77 d of age

Fitted parameter	Change, %	Main transaction	Protein retention, g/d	Lipid retention, g/d	BW gain, g/d	Feed intake, g/d	Effect of main transaction
a_{ME}	-15	ME intake, kJ/d	-1.6	-2.9	-9.2	-57.0	-716
	+15		+1.3	+2.0	+7.4	+54.3	+682
k_p	-15	ME intake, kJ/d	+1.6	+1.9	+8.4	+43.0	+540
	+15		-1.3	-1.6	-7.1	-42.3	-405
k_f	-15	ME intake, kJ/d	+0.5	+0.9	+3.0	+20.9	+262
	+15		-0.4	-0.7	-2.4	-15.2	-191
Ae_{Pc}	-5	Carcass protein anabolism, g/d	+28.9	+2.5	+120.1	+142.7	+57.2
	+5		-17.7	-4.7	-76.7	-107.2	-37.2
AI_{Pc}	-5	Carcass protein anabolism, g/d	-21.2	-8.0	-94.5	-151.9	-50.9
	+5		+38.8	+3.1	+60.9	+207.7	+89.1
CI_{Pc}	-5	Carcass protein catabolism, g/d	+17.9	+2.2	+74.9	+101.7	+20.1
	+5		-12.9	-4.7	-57.3	-89.6	-15.8
CE_{Pc}	-5	Carcass protein catabolism, g/d	-11.2	-3.5	-49.0	-68.1	-8.1
	+5		+13.3	+1.7	+55.9	+69.5	+8.5

Table 7. Sensitivity analysis of protein retention, lipid retention, BW gain, and feed intake (in g/d) to the fitted parameters associated with intraday homeostatic regulations in male growing turkeys at 77 d of age

Fitted parameter	Change, %	Main transaction	Protein retention, g/d	Lipid retention, g/d	BW gain, g/d	Feed intake, g/d	Effect of main transaction, g/d
Ve _{AAox}	-15	AA oxidation	+13.3	+2.0	+56.3	+80.9	+2.2
	+15		-8.2	-2.9	-36.4	-58.2	-3.0
Ve _{FAox}	-15	Fatty acid oxidation	+0.1	+0.5	+1.1	+3.4	-0.3
	+15		-0.1	-1.3	-1.9	-5.9	+0.9
Ve _{Gl}	-15	Lipid turnover ¹	-0.3	-3.8	-5.2	-16.7	-3.2
	+15		+0.3	+0.6	+1.7	+5.2	+0.5
Ve _{Glox}	-15	Glucose oxidation	+0.3	+0.8	+1.9	+6.1	+1.8
	+15		-0.3	-3.1	-4.2	-13.5	-6.1
Ve _{LT}	-15	Lipid turnover ¹	-0.2	-1.6	-2.4	-0.2	-1.4
	+15		+0.1	+0.5	+1.0	+0.1	+0.4
Ve _{PT}	-15	Protein turnover ¹	-10.0	-3.7	-44.3	-71.4	-9.7
	+15		+12.4	+1.9	+52.2	+75.2	+12.0

¹The effect of main transaction was calculated from the synthesis and degradation rates of protein or lipid in carcass.

The daily BW gain in the genotype of females (Hybrid Converter) was alternately overestimated from 4 to 8 wk of age and after 16 wk of age (Figure 7). The average error of simulation of daily BW gain (i.e., difference between simulated and observed values) was 1.0 g/d and varied from -6 to +7 g/d from 0 to 20 wk of age.

Response to Nutrition. The prediction of the effect of different ME (and crude fat content) was in accordance with the observed daily BW gain, feed intake, and feed-to-gain ratio (Figure 8). As expected, increasing the dietary ME decreased feed intake and feed-to-gain ratio, whereas the reverse was true when dietary ME was decreased. In addition to the observations made by Veldkamp et al. (2005) during the study, the model showed that decreasing the ME to 90% of the control treatment resulted in reduced lipid retention, in spite of a daily BW gain close to that of the control treatment. The turkeys fed diets with reduced ME were therefore shown to be leaner than those fed ME at 100 and 110% of the control treatment. Although the feed-to-gain ratio was less in the turkeys fed ME at 110% of the control, these turkeys were less efficient in using nutrients for BW gain because they deposited more fat.

DISCUSSION

The model is capable of simulating the average growth response of a population of male and female turkeys to genetic and nutrition in a controlled (i.e., indirect calorimetry chambers) and experimental environment. Contrary to models based on empirical definitions of BW gain composition (Hurwitz et al., 1983; Emmans, 1989; King, 2001), the model was capable of simulating the differences in BW gain composition depending on feed ingredients, energy, and nutrient utilization. It was developed based on the hypothesis that differences in growth rate and feed-to-gain ratio are mainly due to the partitioning of energy retention between protein and lipid. In these conditions, no nutrient should be limiting in the diets.

The evaluation on both male and female genotypes showed the ability of the model to simulate the response of different genotypes of male and female turkeys to nutrition. The flexibility of the model was enabled by the mechanistic simulation of the interday and intraday HS regulations associated with energy and nutrient utilization. Both regulations were considered because the

Table 8. Mean observed values, coefficient of determination (R^2), root mean square prediction error (MSPE), and decomposition of MSPE of simulated outputs of 2 different sexes and genotypes of turkeys

Item ¹	n	Mean observed value	R^2	Root MSPE	Mean bias, % MSPE	Regression bias, % MSPE	Unexplained variation, % MSPE
Male							
BW gain, g/d	22	146	92.6	10	6.0	1.5	92.6
Feed-to-gain ratio, g/g	22	2.526	39.6	0.250	0.6	59.8	39.6
Feed intake, g/d	22	401	39.9	46	4.8	55.2	39.9
Female							
BW gain, g/d	21	88	98.5	4	6.4	3.3	90.3
Feed-to-gain ratio, g/g	21	2.699	97.7	0.388	0.6	71.7	27.8
Feed intake, g/d	21	236	98.7	23	2.3	66.9	30.8

¹Male = Nicholas 700 (Aviagen Turkeys Ltd., Tattenhall, Cheshire, UK), and female = Hybrid Converter (Hybrid Turkeys, Kitchener, Ontario, Canada).

Figure 6. Comparison between simulated and observed daily feed-to-gain ratio in a male genotype (Nicholas 700, Aviagen Turkeys Ltd., Tattenhall, Cheshire, UK) and evolution of the error of simulation (simulated – observed values, g/g) of daily feed-to-gain ratio (vertical bars) with BW from 1 to 21 wk of age. The relative error of simulation was calculated as the relative difference between the simulated and the observed values.

adaptive response of the turkey individual was assumed to be due to both external and endogenous rules. The interday HS regulations relied on the hypothesis that the growth trajectory results from complex interactions between genotype and ambient conditions (Bodin et al., 2010). Thus, it is not only homeorhesis that drives the growth trajectory. The adjustment of the 6 parameters of the interday HS regulations enabled the adaptation of the growth trajectory to different genotypes characterized by the observed data on BW gain, feed

intake, and feed-to-gain ratio during growth. Based on this fairly simple adjustment to BW gain, feed intake, and feed-to-gain ratio, the model may be fairly easy applicable to commercial facilities.

The energy and N balances measured experimentally (Rivera-Torres et al., 2010b, 2011a) and the serial slaughter analyses (Rivera-Torres et al., 2011b) provided datasets that enabled us to calibrate the parameters at different levels of organization of the teleonomic system (i.e., from feed intake to daily CO₂ production and ash retention in feathers). The energy and N balances provided quantitative information to calibrate the in-

Figure 7. Difference between simulated and observed daily BW gain in a female genotype of turkeys (Hybrid Converter, Hybrid Turkeys, Kitchener, Ontario, Canada) as a function of the observed values; the corresponding age is mentioned in the graph and junctions between points are made following increasing ages. The average error of simulation is 1.0 g/d.

Figure 8. Model simulation of the turkey response to different energy (E) content (90 vs. 110% of the control treatment); comparison between simulated and observed (Veldkamp et al., 2005) values. The E90 and E110 refer to an energy content of 90 and 110%, respectively, of the control treatment. The E90 and E110 observations are the values determined by Veldkamp et al. (2005).

traday HS regulations associated with nutrient utilization for retention or AA, Gl, and FA oxidation into CO₂, whereas the serial slaughter analyses enabled to define the contribution of carcass, viscera, and feathers in whole BW gain. Because of the lack of information on body composition of female turkeys, the results from the serial slaughter analyses were used as a basis for the definition of the growth dynamics of protein and lipid retention in carcass, viscera, and feathers, and the parameters associated with protein and lipid turnovers in viscera were assumed similar between sexes. However, more information on the evolution of the body composition of females during growth may be useful to validate this hypothesis.

Along with protein and lipid turnover in carcass and viscera, the differences in body composition between male and female turkeys were due to the allometric relation between water and protein. Gous et al. (1999) observed a statistically significant effect of sex and strain on the allometric relation of water with protein. In this study, we used the parameter estimates of a previous study (Rivera-Torres et al., 2011b) and those of Emmans (1989) to define the allometric relation of water with protein in male and female turkeys (i.e., 2.90 Prot and 3.41 Prot^{0.90}, respectively, where Prot equals the mass of carcass or viscera protein). The results of the simulations in females showed that the allometric relation of water with protein may be appropriate for females because both BW gain and protein and lipid retention were in accordance with observations. However, this observation was not true for male turkeys. Because water is the major constituent in daily BW gain, it is possible that the allometric relation of water with protein was the origin of the error in the simulation of BW gain at early and later ages because BW gain was globally overestimated, in spite of an accurate simulation of protein retention and an underestimation of lipid retention.

Contrary to other growth models (King, 2001; Eits et al., 2003), the version of the model described herein does not account for the effect of temperature and individual AA on the growth response. The effect of temperature is an important factor to consider in a model for simulating commercial conditions because heat stress is known to have a negative effect on feed intake. Also, Hurwitz and Bengal (1982) demonstrated that basal energy metabolism in turkeys can be defined as a function of ambient temperature. The model represents total AA as a single AA pool. Although such a development would increase the number of circulating compartments in the model, it would enable to propose AA recommendations while accounting for the changes in basal metabolism (i.e., basal protein turnover) and nutrient utilization (i.e., endogenous losses and AA oxidation) with genotype and nutrition.

The user should keep in mind that the model simulates average growth responses rather than individual ones. Indeed, the average growth profile of a population may differ from that of the average individual within

that same population (Pomar et al., 2003). The adaptability of the model to different genotypes and growing conditions, and its ability to simulate the effect of nutrition on growth performance, represent a step toward a better knowledge of turkey growth dynamics and net nutrient recommendations.

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APPENDIX

Table A1. Mathematical statement of the turkey growth model¹

Description	Equation	No.
Body compartment		
Carcass		
Protein (P _c), g		
Input	$F_{AA,Pc/Pc}(t) = k_s + A_{iPc} \exp(-A_{ePc} t) \exp\{Ve_{PT} [\Delta C_{AA}(t)]\} P_c(t)$	1.1
Output	$F_{Pc,AA/Pc}(t) = k_s + C_{iPc} \exp(-C_{ePc} t) \exp\{-Ve_{PT} [\Delta C_{AA}(t)]\} P_c(t)$	1.2
Differential equation	$dP_c/dt = F_{AA,Pc/Pc}(t) - F_{Pc,AA/Pc}(t)$	1.3
Lipid (L _c), g		
Input	$F_{FA-GL/Lc/FA}(t) = k_s + A_{iLc} \exp(-A_{eLc} t) \exp\{Ve_{LT} [\Delta C_{FA}(t)]\} \exp\{Ve_{GL} [\Delta C_{FA}(t)]\} L_c(t)$	1.4
Output	$F_{Lc,FA-GL/Lc}(t) = k_s + C_{iLc} \exp(-C_{eLc} t) \exp\{-Ve_{LT} [\Delta C_{FA}(t)]\} \exp\{-Ve_{GL} [\Delta C_{FA}(t)]\} L_c(t)$	1.5
Differential equation	$dL_c/dt = F_{FA-GL/Lc/FA}(t) - F_{Lc,FA-GL/Lc}(t)$	1.6
Ash (A _c), g		
Input	$FA_c = a_{PCV} [F_{AA,Pc/Pc}(t) - F_{Pc,AA/Pc}(t)]$	1.7
Differential equation	$dA_c/dt = FA_c$	1.8
Water (W _c), g		
Input	$FW_c = a_W [F_{AA,Pc/Pc}(t) - F_{Pc,AA/Pc}(t)]$	1.9
Differential equation	$dW_c/dt = FW_c$	1.10
Viscera		
Protein (P _v), g		
Input	$F_{AA,Pv/Pv}(t) = k_s + A_{iPv} \exp(-A_{ePv} t) \exp\{Ve_{PT} [\Delta C_{AA}(t)]\} P_v(t)$	2.1
Output	$F_{Pv,AA/Pv}(t) = k_s + C_{iPv} \exp(-C_{ePv} t) \exp\{-Ve_{PT} [\Delta C_{AA}(t)]\} P_v(t)$	2.2
Differential equation	$dP_v/dt = F_{AA,Pv/Pv}(t) - F_{Pv,AA/Pv}(t)$	2.3
Lipid (L _v), g		
Input	$F_{FA-GL/Lv/FA}(t) = k_s + A_{iLv} \exp(-A_{eLv} t) \exp\{Ve_{LT} [\Delta C_{FA}(t)]\} \exp\{Ve_{GL} [\Delta C_{FA}(t)]\} L_v(t)$	2.4
Output	$F_{Lv,FA-GL/Lv}(t) = k_s + C_{iLv} \exp(-C_{eLv} t) \exp\{-Ve_{LT} [\Delta C_{FA}(t)]\} \exp\{-Ve_{GL} [\Delta C_{FA}(t)]\} L_v(t)$	2.5
Differential equation	$dL_v/dt = F_{FA-GL/Lv/FA}(t) - F_{Lv,FA-GL/Lv}(t)$	2.6
Ash (A _v), g		
Input	$FA_v = a_{PCV} [F_{AA,Pv/Pv}(t) - F_{Pv,AA/Pv}(t)]$	2.7
Differential equation	$dA_v/dt = FA_v$	2.8
Water (W _v), g		
Input	$FW_v = a_W [F_{AA,Pv/Pv}(t) - F_{Pv,AA/Pv}(t)]$	2.9
Differential equation	$dW_v/dt = FW_v$	2.10
Feathers		
Protein (P _f), g		
Input	$F_{AA,Pf/Pf}(t) = k_s + A_{iPf} \exp(-A_{ePf} t) P_f(t)$	3.1
Differential equation	$dP_f/dt = F_{AA,Pf/Pf}(t)$	3.2
Ash (A _f), g		
Input	$FA_f = a_{Pf} F_{AA,Pf/Pf}(t)$	3.3
Differential equation	$dA_f/dt = FA_f$	3.4
Water (W _f), g		
Input	$FW_f = w_{Pf} F_{AA,Pf/Pf}(t)$	3.5
Differential equation	$dW_f/dt = FW_f$	3.6
Summative equations		
Carcass (C), g	$C(t) = A_c(t) + W_c(t) + P_c(t) + L_c(t)$	3.7
Viscera (V), g	$C(t) = A_v(t) + W_v(t) + P_v(t) + L_v(t)$	3.8
Feathers (F), g	$F(t) = A_f(t) + W_f(t) + P_f(t)$	3.9
BW, g	$BW(t) = A_c(t) + W_c(t) + P_c(t) + L_c(t) + A_v(t) + W_v(t) + P_v(t) + L_v(t) + A_f(t) + W_f(t) + P_f(t)$	3.10
Circulating compartments		
AA, g		
Concentration	$C_{AA}(t) = AA(t)/[Bv \text{ PoB } BW(t)]$	4.1
Inputs	$F_{Pc,AA/AA}(t) + F_{Pv,AA/AA}(t) + \text{FeedCPI } FI(t) \text{ SID}_{AA}$	4.2
Outputs	$F_{AA,Pc/AA}(t) + F_{AA,Pv/AA}(t) + F_{AA,Pf/AA}(t) + F_{AA,GI/AA}(t) + F_{AA,Ay/AA}(t) + IEAA(t)$	4.3
Differential equation	$F_{AA,GI/AA} = k_{ox} AA(t) \exp\{Ve_{AAox} [\Delta C_{AA}(t)]\} \exp\{-Ve_{GI} [\Delta C_{GI}(t)]\}$	4.4
Auxiliary equations	$F_{AA,Ay/AA}(t) = k_{ox} AA(t) \exp\{Ve_{AAox} [\Delta C_{AA}(t)]\} - F_{AA,GI/AA}$	4.5
	$IEAA(t) = a_{IEAA} DM_{intake}/1000$	4.6
	$FI(t) = MEI(t)/\text{feedMEI}$	4.7
	$MEI(t) = [a_{MEI} + d_{MEI} BW(t)] BW(t)^{0.75} + 1/k_p \text{ REP}(t) + 1/k_f \text{ REL}(t)$	4.8
	$\text{REP}(t) = [F_{AA,Pc/Pc}(t) - F_{Pc,AA/Pc}(t) + F_{AA,Pv/Pv}(t) - F_{Pv,AA/Pv}(t) + F_{AA,Pf/Pf}(t)] E_p$	4.9
	$\text{REL}(t) = [F_{FA-GL/Lc/Lc}(t) - F_{Lc,FA-GL/Lc}(t) + F_{FA-GL/Lv/Lv}(t) - F_{Lv,FA-GL/Lv}(t)] E_L$	4.10
	$DM_{intake}(t) = FI(t) \text{ feedDM}$	4.11
Glucose (Gl), g		
Concentration	$C_{Gl}(t) = Gl(t)/[Bv \text{ PoB } BW(t)]$	5.1
Inputs	$F_{AA,GI/Gl}(t) + F_{Lc,GI/Gl}(t) + F_{Lv,GI/Gl}(t) + Gl_{intake}$	5.2

Continued

Table A1 (Continued). Mathematical statement of the turkey growth model¹

Description	Equation	No.
Outputs	$F_{\text{Gl,Lc/Gl}}(t) + F_{\text{Gl,Lv/Gl}}(t) + F_{\text{Gl,Ay/Gl}}(t)$	5.3
Differential equation	$d\text{Gl}/dt = F_{\text{AA,Gl/Gl}}(t) + F_{\text{Lc,Gl/Gl}}(t) + F_{\text{Lv,Gl/Gl}}(t) + \text{Gl}_{\text{intake}} - F_{\text{Gl,Lc/Gl}}(t) - F_{\text{Gl,Lv/Gl}}(t) - F_{\text{Gl,Ay/Gl}}(t)$	5.4
Auxiliary equations	$\text{Gl}_{\text{intake}} = [\text{MEI}(t) - \text{FeedCPi FI}(t) \text{SID}_{\text{AA}} E_{\text{P}} - \text{FeedCFi FI}(t) \text{SID}_{\text{CF}} E_{\text{L}}]/E_{\text{Gl}}$	5.5
	$F_{\text{Gl,Ay/Gl}}(t) = k_{\text{ox}} \text{Gl}(t) \exp\{\text{Ve}_{\text{Glox}} [\Delta C_{\text{Gl}}(t)]\}$	5.6
Fatty acid (FA), g		
Concentration	$C_{\text{FA}}(t) = \text{FA}(t)/[\text{Bv PoB BW}(t)]$	6.1
Inputs	$F_{\text{Ay,FA/FA}}(t) + F_{\text{Lc,FA/FA}}(t) + F_{\text{Lv,FA/FA}}(t) + \text{CF}_{\text{intake}}$	6.2
Outputs	$F_{\text{FA,Ay/FA}}(t) + F_{\text{FA,Lc/FA}}(t) + F_{\text{FA,Lv/FA}}(t)$	6.3
Differential equation	$d\text{FA}/dt = F_{\text{Ay,FA/FA}}(t) + F_{\text{Lc,FA/FA}}(t) + F_{\text{Lv,FA/FA}}(t) + \text{CF}_{\text{intake}} - F_{\text{FA,Ay/FA}}(t) - F_{\text{FA,Lc/FA}}(t) - F_{\text{FA,Lv/FA}}(t)$	6.4
	$\text{CF}_{\text{intake}}(t) = \text{FeedCFi FI}(t)$	6.5
Auxiliary equation	$F_{\text{FA,Ay/FA}}(t) = k_{\text{ox}} \text{FA}(t) \exp\{\text{Ve}_{\text{FAox}} [\Delta C_{\text{FA}}(t)]\}$	6.6
	$F_{\text{Ay,FA/FA}}(t) = k_{\text{ox}} \text{FA}(t) \exp\{\text{Ve}_{\text{Gl}} [\Delta C_{\text{Gl}}(t)]\}$	6.7
	$F_{\text{FA,Lc/FA}}(t) = k_{\text{ox}} \text{FA}(t) \exp\{\text{Ve}_{\text{Gl}} [\Delta C_{\text{Gl}}(t)]\}$	6.8
Acetyl-coA (Ay), mol		
Inputs	$F_{\text{AA,Ay/Ay}} + F_{\text{Gl,Ay/Ay}}(t) + F_{\text{FA,Ay/Ay}}(t)$	6.9
Output	$F_{\text{Ay,FA/Ay}}(t) = k_{\text{ox}} \text{FA}(t) \exp\{\text{Ve}_{\text{Gl}} [\Delta C_{\text{Gl}}(t)]\}$	6.10
Differential equation	$d\text{Ay}/dt = F_{\text{AA,Ay/Ay}} + F_{\text{Gl,Ay/Ay}}(t) + F_{\text{FA,Ay/Ay}}(t) - F_{\text{Ay,FA/Ay}}(t)$	6.11

¹Refer to Tables 1, 2, and 3 for explanation of the notation and parameter values.

References

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