

Modelling livestock

Julien Arino

April 2023

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2023-04-17

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Yes, why, really, must I tell you about this ?

I model the spread of infectious diseases and, less frequently, more general interactions between species (mathematical ecology)

I have even modelled how filaments grow (assemble) in cells

But "modelling livestock" ? Nope, never

So let's take a (for the most part uninformed) dive in models of livestock

Modelling livestock

└ Why?

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I model the spread of infectious diseases and, less frequently, more general interactions between species (mathematical ecology)

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So let's take a (for the most part uninformed) dive in models of livestock

Digging through the literature, there's actually quite a lot and some of it is quite fun!

This will be a completely non-exhaustive and random review of some models I have found **excluding** anything infectious disease related

Wait..

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This will be a completely non-exhaustive and random review of some models I have found **excluding** anything infectious disease related

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└ Wait..

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└ Outline

Outline

Models for cattle

Models for other herds

Models for management

Conclusion

Models for cattle

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Models for management

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Models for cattle

Models for individuals

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Conclusion

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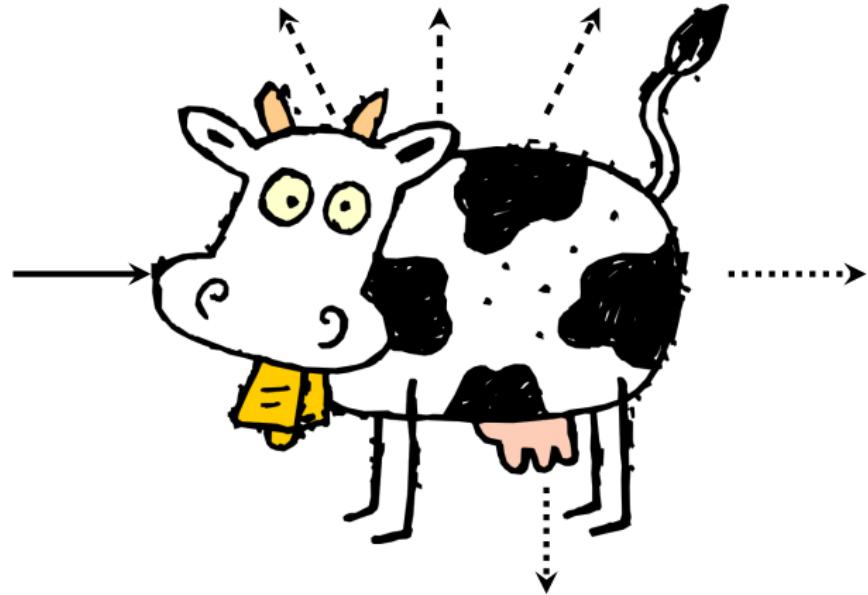
Models for cattle
Models for individuals
Models for herds

Models for other herds

Models for management

Conclusion

At the individual level



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 └ Models for individuals
 └ At the individual level



MODELLING ANIMAL SYSTEMS PAPER

Development of a dynamic mathematical model for investigating mammary gland metabolism in lactating cows

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| Abbreviation | Meaning |
|--------------|---|
| Aa | Amino acids |
| AaE | Essential amino acids |
| AaN | Non-essential amino acids |
| Ac | Acetate |
| Acm | Mitochondrial acetyl-CoA |
| Adp | Adenosine-diphosphate |
| AP | Mammary tissue actively perfused |
| Atp | Adenosine-triphosphate |
| Bhb | β -hydroxybutyrate |
| Cit | Citrate |
| eAa | Amino acids in the arterial plasma |
| eAc | Acetate in the arterial plasma |
| eBhb | β -hydroxybutyrate in the arterial plasma |
| eGlc | Glucose in the arterial plasma |
| eFa | FAs in the arterial plasma |
| Fa | FAs |
| F6p | Fructose-6-phosphate |
| Fbp | Fructose-1,6-biphosphate |
| Glc | Glucose |
| G6p | Glucose-6-phosphate |
| Gcl | Glycerol |
| Lac | Lactose |
| Oa | Oxaloacetate |
| Pga | Phospho-glyceraldehyde |
| Ptm | Milk protein synthesized in the gland |
| Pyr | Pyruvate |
| R5p | Ribulose-5-phosphate |
| Succ | Succinate |
| Tca | Tricarboxylic acids |
| Tgm | Triacylglycerol in milk |

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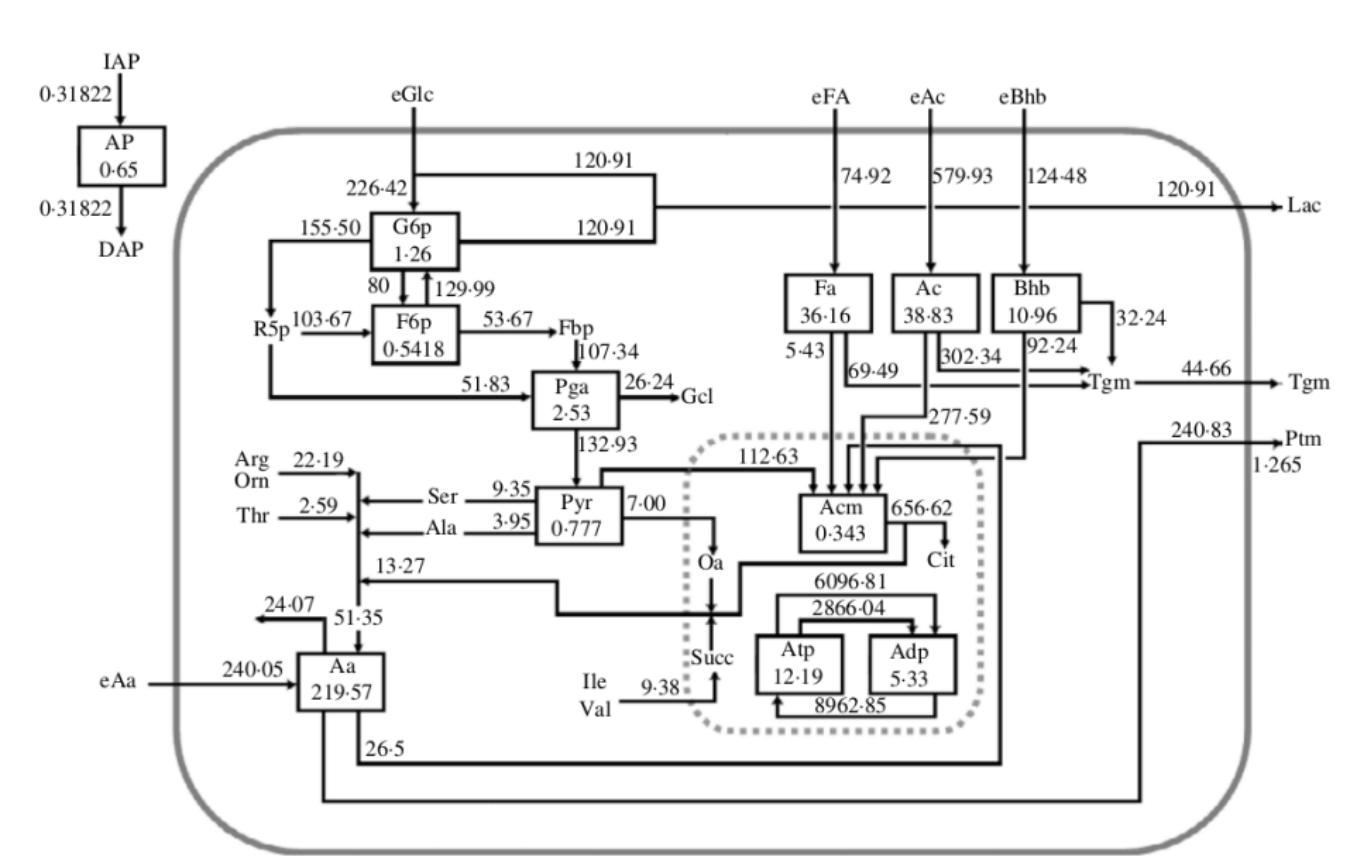


Fig. 1. Reference steady state diagram. Each individual compartment is represented through a box. Symbols are described in Table 2. AP is the volume of tissue actively perfused by blood. Fluxes are mmol/h except for IAP and DAP. Numbers in the boxes indicate the initial condition size of the compartment (mmol).

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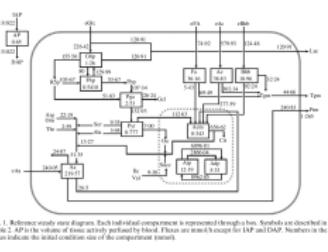


Fig. 1. Reference steady state diagram. Each individual compartment is represented through a box. Symbols are described in Table 2. AP is the volume of tissue actively perfused by blood. Fluxes are mmol/h except for IAP and DAP. Numbers in the boxes indicate the initial condition size of the compartment (mmol).

| Notation | Meaning | Units |
|--------------|---|-----------|
| c_A | Intracellular concentration of A | mmol/l |
| ce_A | Arterial concentration of A | mmol/l |
| $^o ce_A$ | Reference arterial concentration of A | mmol/l |
| $INSH$ | Insonaemia | Units |
| J_A | Inhibition constant for the inflow of A | mmol/l |
| $J_{C,AB}$ | Inhibition constant for transaction A→B with respect to C | mmol/l |
| K_{InA} | Affinity constant for the inflow of A | mmol/l |
| K_{OutA} | Affinity constant for the outflow of A | mmol/l |
| $K_{A,B}$ | Affinity constant for A in the transaction A→B | mmol/l |
| $K_{C,AB}$ | Affinity constant for transaction A→B with respect to C | mmol/l |
| $P_{B,A}$ | Rate of synthesis of B in the transaction A→B | mmol/h |
| PPS | Potential milk protein yield | mmol/h |
| $R_{A,B}$ | Requirement of A in the synthesis of B | mmol/mmol |
| R_i | Content of i th AA in milk protein synthesized in the gland | mmol/mmol |
| RPS | Real milk protein yield | mmol/h |
| $U_{A,B}$ | Rate of utilization of A in the transaction A→B | mmol/h |
| $^o U_{A,B}$ | Reference rate of utilization of A in the transaction A→B | mmol/h |
| UO_i | Uptake:output ratio for the i th AA | mmol/mmol |
| UW | Udder weight | kg |
| $v_{ceA,A}$ | Net uptake rate of A | mmol/h |
| $V_{A,B}$ | V_{max} for the transaction A→B | mmol/h |
| V_{InA} | V_{max} for the inflow of A | μmol/l |
| V_{OutA} | V_{max} for the outflow of A | μmol/l |
| $Y_{A,B}$ | Rate of yield of B in the transaction A→B | mmol/mmol |

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| $P_{B,A}$ | Rate of synthesis of B in the transaction A→B | mmol/h |
| PPS | Potential milk protein yield | mmol/h |
| R | Requirement of A in the synthesis of B | mmol/mmol |
| RPS | Real milk protein yield | mmol/h |
| R_i | Content of i th AA in milk protein synthesized in the gland | mmol/mmol |
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Table 4. MM parameters of the equations describing uptakes of nutrients

| Equation no. | V_{InA} ($\mu\text{mol/l}$) | K_{InA} (mm) | K_{InB} (mm) | J_A (mm) | V_{outA} ($\mu\text{mol/l}$) | K_{outA} (mm) |
|--------------|---|--------------------------|--------------------------|---------------|--|---------------------------|
| (2.2): Arg | 135.610 | 0.095 | — | 12.158 | 22.600 | 12.158 |
| (2.2): His | 49.060 | 0.053 | — | 12.158 | 8.200 | 12.158 |
| (2.2): Ile | 164.920 | 0.101 | — | 12.158 | 27.490 | 12.158 |
| (2.2): Leu | 232.0 | 0.065 | — | 12.158 | 62.500 | 12.158 |
| (2.2): Lys | 282.120 | 0.125 | — | 12.158 | 47.020 | 12.158 |
| (2.2): Met | 50.990 | 0.025 | — | 12.158 | 8.500 | 12.158 |
| (2.2): Phe | 85.490 | 0.039 | — | 12.158 | 14.250 | 12.158 |
| (2.2): Thr | 127.710 | 0.080 | — | 12.158 | 21.290 | 12.158 |
| (2.2): Trp | 18.990 | 0.047 | — | 12.158 | 3.170 | 12.158 |
| (2.2): Val | 215.330 | 0.147 | — | 12.158 | 35.890 | 12.158 |
| (2.9): Ala | 67.370 | 0.189 | — | 12.158 | 11.230 | 12.158 |
| (2.9): Asp | 10.540 | 0.004 | — | 12.158 | 1.0 | 12.158 |
| (2.9): Asn | 127.230 | 0.129 | — | 12.158 | 21.210 | 12.158 |
| (2.9): Cys | 36.340 | 0.045 | — | 12.158 | 6.060 | 12.158 |
| (2.9): Glu | 159.480 | 0.058 | — | 12.158 | 15.190 | 12.158 |
| (2.9): Gln | 134.270 | 0.221 | — | 12.158 | 22.380 | 12.158 |
| (2.9): Gly | 30.550 | 0.231 | — | 12.158 | 5.090 | 12.158 |
| (2.9): Pro | 34.020 | 0.110 | — | 12.158 | 5.671 | 12.158 |
| (2.9): Ser | 95.230 | 0.084 | — | 12.158 | 15.870 | 12.158 |
| (2.9): Tyr | 77.300 | 0.045 | — | 12.158 | 12.880 | 12.158 |
| (2.9): Orn | 133.690 | 0.073 | — | 12.158 | 22.280 | 12.158 |
| (3.1): Ac* | 3.600 | 2.150 | — | — | 0.450 | 0.253 |
| (5.1): Bhb* | 1.250 | 0.750 | — | — | 0.550 | 0.420 |
| (6.1): Fa*† | 1.100 | 0.280 | 0.100 | — | 0.150 | 0.309 |
| (7.1): Glc*‡ | 2.250 | 3.640 | — | — | 1.250 | 0.081 |

* For Ac, Bhb, Fa and Glc parameters V_{InA} and V_{outA} are expressed in mmol/l.† Equation (6.1), K_{InA} and K_{InB} are K_{InNefA} and K_{InTg} respectively.‡ Equation (7.1), parameters V_{InA} , K_{InA} , V_{outA} and K_{outA} are $V_{\text{eGlc}, \text{G}6\text{p}}$, $K_{\text{eGlc}, \text{G}6\text{p}}$, $V_{\text{G}6\text{p}, \text{eGlc}}$ and $K_{\text{G}6\text{p}, \text{eGlc}}$, respectively.

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| Equation no. | F_{InA} ($\mu\text{mol/l}$) | K_{InA} (mm) | K_{InB} (mm) | J_A (mm) | F_{outA} ($\mu\text{mol/l}$) | K_{outA} (mm) |
|--------------|---|--------------------------|--------------------------|---------------|--|---------------------------|
| G.2.1: Arg | 170.000 | 0.000 | — | 12.158 | 40.0 | 21.200 |
| G.2.2: His | 100.000 | 0.003 | — | 12.158 | 21.400 | 21.200 |
| G.2.3: Ile | 160.020 | 0.101 | — | 12.158 | 50.0 | 21.200 |
| G.2.4: Leu | 200.020 | 0.058 | — | 12.158 | 80.0 | 21.200 |
| G.2.5: Lys | 240.120 | 0.121 | — | 12.158 | 47.830 | 21.200 |
| G.2.6: Met | 50.000 | 0.023 | — | 12.158 | 10.0 | 21.200 |
| G.2.7: Phe | 85.490 | 0.039 | — | 12.158 | 14.280 | 21.200 |
| G.2.8: Thr | 127.710 | 0.080 | — | 12.158 | 21.750 | 21.200 |
| G.2.9: Trp | 18.990 | 0.047 | — | 12.158 | 3.170 | 21.200 |
| G.2.10: Val | 215.330 | 0.147 | — | 12.158 | 35.890 | 21.200 |
| G.2.11: Ala | 67.370 | 0.189 | — | 12.158 | 11.230 | 21.200 |
| G.2.12: Asp | 10.540 | 0.004 | — | 12.158 | 1.0 | 21.200 |
| G.2.13: Asn | 127.230 | 0.129 | — | 12.158 | 21.210 | 21.200 |
| G.2.14: Cys | 36.340 | 0.045 | — | 12.158 | 6.060 | 21.200 |
| G.2.15: Glu | 159.480 | 0.058 | — | 12.158 | 15.190 | 21.200 |
| G.2.16: Gln | 134.270 | 0.221 | — | 12.158 | 22.380 | 21.200 |
| G.2.17: Gly | 30.550 | 0.231 | — | 12.158 | 5.090 | 21.200 |
| G.2.18: Pro | 34.020 | 0.110 | — | 12.158 | 5.671 | 21.200 |
| G.2.19: Ser | 95.230 | 0.084 | — | 12.158 | 15.870 | 21.200 |
| G.2.20: Tyr | 77.300 | 0.045 | — | 12.158 | 12.880 | 21.200 |
| G.2.21: Orn | 133.690 | 0.073 | — | 12.158 | 22.280 | 21.200 |
| G.3.1: Ac* | 3.600 | 2.150 | — | — | 0.450 | 0.253 |
| G.5.1: Bhb* | 1.250 | 0.750 | — | — | 0.550 | 0.420 |
| G.6.1: Fa*† | 1.100 | 0.280 | 0.100 | — | 0.150 | 0.309 |
| G.7.1: Glc*‡ | 2.250 | 3.640 | — | — | 1.250 | 0.081 |

* For Ac, Bhb, Fa and Glc parameters V_{InA} and V_{outA} are expressed in mmol/l.† Equations (6.1), K_{InA} and K_{InB} are K_{InNefA} and K_{InTg} respectively.‡ Equation (7.1), parameters V_{InA} , K_{InA} , V_{outA} and K_{outA} are $V_{\text{eGlc}, \text{G}6\text{p}}$, $K_{\text{eGlc}, \text{G}6\text{p}}$, $V_{\text{G}6\text{p}, \text{eGlc}}$ and $K_{\text{G}6\text{p}, \text{eGlc}}$, respectively.

APPENDIX

MATHEMATICAL STATEMENT OF THE MODEL

Mammary plasma flow

$$dAP/dt = IAP - DAP \quad (1.0)$$

– *Inputs:*

$$IAP = PPS / \text{MIN}\{RPS, PPS\} \text{ INSH}^{0.20} 0.350 / (1 + 0.035 / (1 - AP)) \quad (1.1)$$

– *Outputs:*

$$DAP = cAtp / cAdp 0.1522 / (1 + 0.065 / AP) \quad (1.2)$$

$$MPF = MS^{\circ} 0.9974^{(\text{DIM} - 133)} UW753AP \quad (1.3)$$

AA compartment (Aa)

$$dAa/dt = v_{eAa, Aa} + P_{AaN, AaE} - U_{Aa, Deg} - U_{Aa, Mtb} - U_{Aa, Ptm} \quad (2.0)$$

– *Inputs:*

$$v_{eAa, Aa} = \sum v_{eAaEi, AaEi} + \sum v_{eAaNi, AaNi} \quad (2.1)$$

Uptake of essential AAs

$$\begin{aligned} v_{eAaEi, AaEi} &= V_{InAai} MPF (\text{ceAaE}_i / \text{ceAaE}_i)^{0.2} / (1 + K_{InAai} (\text{oRPS}_i / ePS)^{\varepsilon} / \text{ceAaE}_i + (\text{cAa} / J_{Aa})^{3.5}) \\ &\quad - V_{OutAai} MPF / (1 + K_{OutAa} / \text{cAa}) \end{aligned} \quad (2.2)$$

$$\begin{aligned} \text{oRPS}_{His} &= \{V_{InHis} MPF (\text{ceHis} / \text{ceHis})^{0.2} / (1 + K_{InHis} / \text{ceHis} + (\text{cAa} / J_{Aa})^{3.5}) \\ &\quad - V_{OutHis} MPF / (1 + K_{OutAa} / \text{cAa})\} 1.1 / (UO_{His} R_{His}) \end{aligned} \quad (2.3)$$

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| MATHEMATICAL STATEMENT OF THE MODEL | |
| <i>Mammary plasma flow</i> | $dAP/dt = IAP - DAP$ (1.0) |
| – <i>Inputs:</i> | $IAP = PPS / \text{MIN}\{RPS, PPS\} \text{ INSH}^{0.20} 0.350 / (1 + 0.035 / (1 - AP))$ (1.1) |
| – <i>Outputs:</i> | $DAP = cAtp / cAdp 0.1522 / (1 + 0.065 / AP)$ (1.2) |
| | $MPF = MS^{\circ} 0.9974^{(\text{DIM} - 133)} UW753AP$ (1.3) |
| <i>AA compartment (Aa)</i> | |
| | $dAa/dt = v_{eAa, Aa} + P_{AaN, AaE} - U_{Aa, Deg} - U_{Aa, Mtb} - U_{Aa, Ptm}$ (2.0) |
| <i>Update of essential AAs</i> | |
| | $v_{eAaEi, AaEi} = V_{InAai} MPF (\text{ceAaE}_i / \text{ceAaE}_i)^{0.2} / (1 + K_{InAai} (\text{oRPS}_i / ePS)^{\varepsilon} / \text{ceAaE}_i + (\text{cAa} / J_{Aa})^{3.5})$ (2.1) |
| | $- V_{OutAai} MPF / (1 + K_{OutAa} / \text{cAa})$ (2.2) |
| | $\text{oRPS}_{His} = \{V_{InHis} MPF (\text{ceHis} / \text{ceHis})^{0.2} / (1 + K_{InHis} / \text{ceHis} + (\text{cAa} / J_{Aa})^{3.5})$ (2.3) |
| | $- V_{OutHis} MPF / (1 + K_{OutAa} / \text{cAa})\} 1.1 / (UO_{His} R_{His})$ (2.3) |

$$\text{oRPS}_{\text{Met}} = \{V_{\text{InMet}} \text{MPF} (\text{ceMet}/\text{ceMet})^{0.2} / (1 + K_{\text{InMet}}/\text{ceMet} + (\text{cAa}/J_{\text{Aa}})^{3.5}) - V_{\text{OutMet}} \text{MPF} / (1 + K_{\text{OutAa}}/\text{cAa})\} \cdot 1 \cdot 1 / (\text{UO}_{\text{Met}} R_{\text{Met}}) \quad (2.4)$$

$$\text{oRPS}_{\text{Phe}} = \{V_{\text{InPhe}} \text{MPF} (\text{cePhe}/\text{cePhe})^{0.2} / (1 + K_{\text{InPhe}}/\text{cePhe} + (\text{cAa}/J_{\text{Aa}})^{3.5}) - V_{\text{OutPhe}} \text{MPF} / (1 + K_{\text{OutAa}}/\text{cAa})\} \cdot 1 \cdot 1 / (\text{UO}_{\text{Phe}} R_{\text{Phe}}) \quad (2.5)$$

$$\text{oRPS}_{\text{Leu}} = \{V_{\text{InLeu}} \text{MPF} (\text{ceLeu}/\text{ceLeu})^{0.2} / (1 + K_{\text{InLeu}}/\text{ceLeu} + (\text{cAa}/J_{\text{Aa}})^{3.5}) - V_{\text{OutLeu}} \text{MPF} / (1 + K_{\text{OutAa}}/\text{cAa})\} / (\text{UO}_{\text{Leu}} R_{\text{Leu}}) \quad (2.6)$$

$$\text{oRPS}_{\text{AaO}} = \text{MIN}\{\text{oRPS}_{\text{His}}, \text{oRPS}_{\text{Met}}, \text{oRPS}_{\text{Phe}}\} \quad (2.7)$$

$$\text{ePS} = \text{PPS INSH}^{0.050} \quad (2.8)$$

Uptake of non-essential AAs

$$v_{\text{eAaNi}, \text{AaNi}} = V_{\text{InAai}} \text{MPF} (\text{ceAaNi}/\text{ceAaNi})^{0.2} / (1 + K_{\text{InAai}}/\text{ceAaNi} + (\text{cAa}/J_{\text{Aa}})^{3.5}) - V_{\text{OutAai}} \text{MPF} / (1 + K_{\text{OutAai}}/\text{cAa}) \quad (2.9)$$

Synthesis of non-essential AAs

$$P_{\text{AaN}, \text{AaE}} = U_{\text{Pyr}, \text{Ser}} + U_{\text{Pyr}, \text{Ala}} + P_{\text{GlAs}, \text{AaE}} + P_{\text{Pro}, \text{Arg}} + P_{\text{Gly}, \text{Thr}} \quad (2.10)$$

$$U_{\text{Pyr}, \text{Ser}} = (\text{RPS } R_{\text{Ser}} - v_{\text{eSer}, \text{Ser}} + \text{RPS } R_{\text{Gly}} - v_{\text{eGly}, \text{Gly}}) * V_{\text{Pyr}, \text{Ser}} / (1 + K_{\text{Pyr}, \text{Ser}}/\text{cPyr}) \quad (2.11)$$

$$U_{\text{Pyr}, \text{Ala}} = (\text{RPS } R_{\text{Ala}} - v_{\text{eAla}, \text{Ala}}) V_{\text{Pyr}, \text{Ala}} / (1 + K_{\text{Pyr}, \text{Ala}}/\text{cPyr}) \quad (2.12)$$

$$P_{\text{GlAs}, \text{AaE}} = 0.019717(U_{\text{Pyr}, \text{Oa}} + U_{\text{AcM}, \text{Cit}} + U_{\text{Val}, \text{SucTca}} + U_{\text{Ile}, \text{SucTca}}) \quad (2.13)$$

$$U_{\text{Val}, \text{SucTca}} = v_{\text{eVal}, \text{Val}} - \text{RPS } R_{\text{Val}} \quad (2.14)$$

$$U_{\text{Ile}, \text{SucTca}} = v_{\text{eIle}, \text{Ile}} - \text{RPS } R_{\text{Ile}} \quad (2.15)$$

$$P_{\text{Pro}, \text{Arg}} = \text{RPS } R_{\text{Pro}} - v_{\text{ePro}, \text{Pro}} \quad (2.16)$$

$$P_{\text{Gly}, \text{Thr}} = \text{RPS } R_{\text{Gly}} - v_{\text{eGly}, \text{Gly}} + \text{RPS } R_{\text{Ser}} - U_{\text{Pyr}, \text{Ser}} - v_{\text{eSer}, \text{Ser}} \quad (2.17)$$

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$$\begin{aligned} dP_{\text{Met}} &= \{V_{\text{InMet}} \text{MPF} (\text{ceMet}/\text{ceMet})^{0.2} / (1 + K_{\text{InMet}}/\text{ceMet} + (\text{cAa}/J_{\text{Aa}})^{3.5}) \\ &\quad - V_{\text{OutMet}} \text{MPF} / (1 + K_{\text{OutAa}}/\text{cAa})\} \cdot 1 \cdot 1 / (\text{UO}_{\text{Met}} R_{\text{Met}}) \quad (2.4) \\ dP_{\text{Phe}} &= \{V_{\text{InPhe}} \text{MPF} (\text{cePhe}/\text{cePhe})^{0.2} / (1 + K_{\text{InPhe}}/\text{cePhe} + (\text{cAa}/J_{\text{Aa}})^{3.5}) \\ &\quad - V_{\text{OutPhe}} \text{MPF} / (1 + K_{\text{OutAa}}/\text{cAa})\} \cdot 1 \cdot 1 / (\text{UO}_{\text{Phe}} R_{\text{Phe}}) \quad (2.5) \\ dP_{\text{Leu}} &= \{V_{\text{InLeu}} \text{MPF} (\text{ceLeu}/\text{ceLeu})^{0.2} / (1 + K_{\text{InLeu}}/\text{ceLeu} + (\text{cAa}/J_{\text{Aa}})^{3.5}) \\ &\quad - V_{\text{OutLeu}} \text{MPF} / (1 + K_{\text{OutAa}}/\text{cAa})\} / (\text{UO}_{\text{Leu}} R_{\text{Leu}}) \quad (2.6) \\ dP_{\text{AaO}} &= \text{MIN}\{\text{oRPS}_{\text{His}}, \text{oRPS}_{\text{Met}}, \text{oRPS}_{\text{Phe}}\} \quad (2.7) \\ \text{ePS} &= \text{PPS INSH}^{0.050} \quad (2.8) \end{aligned}$$

Uptake of non-essential AAs

$$v_{\text{eAaNi}, \text{AaNi}} = V_{\text{InAai}} \text{MPF} (\text{ceAaNi}/\text{ceAaNi})^{0.2} / (1 + K_{\text{InAai}}/\text{ceAaNi} + (\text{cAa}/J_{\text{Aa}})^{3.5}) - V_{\text{OutAai}} \text{MPF} / (1 + K_{\text{OutAai}}/\text{cAa}) \quad (2.9)$$

Synthesis of non-essential AAs

$$\begin{aligned} P_{\text{AaN}, \text{AaE}} &= U_{\text{Pyr}, \text{Ser}} + U_{\text{Pyr}, \text{Ala}} + P_{\text{GlAs}, \text{AaE}} + P_{\text{Pro}, \text{Arg}} + P_{\text{Gly}, \text{Thr}} \quad (2.10) \\ U_{\text{Pyr}, \text{Ser}} &= (\text{RPS } R_{\text{Ser}} - v_{\text{eSer}, \text{Ser}} + \text{RPS } R_{\text{Gly}} - v_{\text{eGly}, \text{Gly}}) * V_{\text{Pyr}, \text{Ser}} / (1 + K_{\text{Pyr}, \text{Ser}}/\text{cPyr}) \quad (2.11) \\ U_{\text{Pyr}, \text{Ala}} &= (\text{RPS } R_{\text{Ala}} - v_{\text{eAla}, \text{Ala}}) V_{\text{Pyr}, \text{Ala}} / (1 + K_{\text{Pyr}, \text{Ala}}/\text{cPyr}) \quad (2.12) \\ P_{\text{GlAs}, \text{AaE}} &= 0.019717(U_{\text{Pyr}, \text{Oa}} + U_{\text{AcM}, \text{Cit}} + U_{\text{Val}, \text{SucTca}} + U_{\text{Ile}, \text{SucTca}}) \quad (2.13) \\ U_{\text{Val}, \text{SucTca}} &= v_{\text{eVal}, \text{Val}} - \text{RPS } R_{\text{Val}} \quad (2.14) \\ U_{\text{Ile}, \text{SucTca}} &= v_{\text{eIle}, \text{Ile}} - \text{RPS } R_{\text{Ile}} \quad (2.15) \\ P_{\text{Pro}, \text{Arg}} &= \text{RPS } R_{\text{Pro}} - v_{\text{ePro}, \text{Pro}} \quad (2.16) \\ P_{\text{Gly}, \text{Thr}} &= \text{RPS } R_{\text{Gly}} - v_{\text{eGly}, \text{Gly}} + \text{RPS } R_{\text{Ser}} - U_{\text{Pyr}, \text{Ser}} - v_{\text{eSer}, \text{Ser}} \quad (2.17) \end{aligned}$$

$$\begin{aligned} U_{\text{Aa}, \text{Deg}} &= V_{\text{Aa}, \text{Deg}} / (1 + (\text{INSH}/5)^{0.30}) (1 + (K_{\text{Aa}, \text{Deg}}/\text{cAa})^{2.50}) & (2.18) \\ U_{\text{Aa}, \text{Mtb}} &= (v_{\text{eArg}, \text{Arg}} - \text{RPS } R_{\text{Arg}} + v_{\text{eOrn}, \text{Orn}}) V_{\text{Aa}, \text{Mtb}} / (1 + (K_{\text{Aa}, \text{Mtb}}/\text{cAa})^{2.50}) & (2.19) \\ U_{\text{Aa}, \text{Ptm}} &= \text{RPS } V_{\text{Aa}, \text{Ptm}} / (1 + K_{\text{Aa}, \text{Ptm}}/\text{cAa}) & (2.20) \\ \text{RPS} &= P_{\text{Ptm}, \text{Aa}} \ Y_{\text{Atp}, \text{Ptm}} & (2.21) \\ P_{\text{Ptm}, \text{Aa}} &= \text{MIN}\{1.09 v_{\text{eHis}, \text{His}}/R_{\text{His}}; 1.09 v_{\text{eMet}, \text{Met}}/R_{\text{Met}}; 1.09 v_{\text{ePhe}, \text{Phe}}/R_{\text{Phe}}\} & (2.22) \\ Y_{\text{Atp}, \text{Ptm}} &= \text{MIN}\{1.000; V_{\text{Atp}, \text{Ptm}} / (1 + K_{\text{Atp}, \text{Ptm}}/\text{cAtp})\} & (2.23) \\ dAc/dt &= v_{\text{eAc}, \text{Ac}} - U_{\text{Ac}, \text{Acm}} - U_{\text{Ac}, \text{Tgm}} & (3.0) \\ v_{\text{eAc}, \text{Ac}} &= V_{\text{InAc}} \text{MPF} / (1 + K_{\text{InAc}}/\text{cAc}) - V_{\text{OutAc}} \text{MPF} / (1 + K_{\text{OutAc}}/\text{cAc}) & (3.1) \end{aligned}$$

- Outputs:

$$U_{\text{Aa}, \text{Deg}} = V_{\text{Aa}, \text{Deg}} / ((1 + (\text{INSH}/5)^{0.30})(1 + (K_{\text{Aa}, \text{Deg}}/\text{cAa})^{2.50})) \quad (2.18)$$

- Inputs:

$$U_{\text{Aa}, \text{Deg}} = V_{\text{Aa}, \text{Deg}} / (1 + (\text{INSH}/5)^{0.30}) (1 + (K_{\text{Aa}, \text{Deg}}/\text{cAa})^{2.50}) \quad (2.18)$$

$$U_{\text{Aa}, \text{Mtb}} = (v_{\text{eArg}, \text{Arg}} - \text{RPS } R_{\text{Arg}} + v_{\text{eOrn}, \text{Orn}}) V_{\text{Aa}, \text{Mtb}} / (1 + (K_{\text{Aa}, \text{Mtb}}/\text{cAa})^{2.50}) \quad (2.19)$$

$$\text{RPS} = P_{\text{Ptm}, \text{Aa}} \ Y_{\text{Atp}, \text{Ptm}} \quad (2.20)$$

$$P_{\text{Ptm}, \text{Aa}} = \text{MIN}\{1.09 v_{\text{eHis}, \text{His}}/R_{\text{His}}; 1.09 v_{\text{eMet}, \text{Met}}/R_{\text{Met}}; 1.09 v_{\text{ePhe}, \text{Phe}}/R_{\text{Phe}}\} \quad (2.21)$$

$$Y_{\text{Atp}, \text{Ptm}} = \text{MIN}\{1.000; V_{\text{Atp}, \text{Ptm}} / (1 + K_{\text{Atp}, \text{Ptm}}/\text{cAtp})\} \quad (2.22)$$

$$dAc/dt = v_{\text{eAc}, \text{Ac}} - U_{\text{Ac}, \text{Acm}} - U_{\text{Ac}, \text{Tgm}} \quad (3.0)$$

$$v_{\text{eAc}, \text{Ac}} = V_{\text{InAc}} \text{MPF} / (1 + K_{\text{InAc}}/\text{cAc}) - V_{\text{OutAc}} \text{MPF} / (1 + K_{\text{OutAc}}/\text{cAc}) \quad (3.1)$$

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- Outputs:

$$U_{\text{Aa}, \text{Deg}} = V_{\text{Aa}, \text{Deg}} / ((1 + (\text{INSH}/5)^{0.30})(1 + (K_{\text{Aa}, \text{Deg}}/\text{cAa})^{2.50})) \quad (2.18)$$

$$U_{\text{Aa}, \text{Mtb}} = (v_{\text{eArg}, \text{Arg}} - \text{RPS } R_{\text{Arg}} + v_{\text{eOrn}, \text{Orn}}) V_{\text{Aa}, \text{Mtb}} / (1 + (K_{\text{Aa}, \text{Mtb}}/\text{cAa})^{2.50}) \quad (2.19)$$

$$U_{\text{Aa}, \text{Ptm}} = \text{RPS } V_{\text{Aa}, \text{Ptm}} / (1 + K_{\text{Aa}, \text{Ptm}}/\text{cAa}) \quad (2.20)$$

$$\text{RPS} = P_{\text{Ptm}, \text{Aa}} \ Y_{\text{Atp}, \text{Ptm}} \quad (2.21)$$

$$P_{\text{Ptm}, \text{Aa}} = \text{MIN}\{1.09 v_{\text{eHis}, \text{His}}/R_{\text{His}}; 1.09 v_{\text{eMet}, \text{Met}}/R_{\text{Met}}; 1.09 v_{\text{ePhe}, \text{Phe}}/R_{\text{Phe}}\} \quad (2.22)$$

$$Y_{\text{Atp}, \text{Ptm}} = \text{MIN}\{1.000; V_{\text{Atp}, \text{Ptm}} / (1 + K_{\text{Atp}, \text{Ptm}}/\text{cAtp})\} \quad (2.23)$$

Acetate compartment (Ac)

$$dAc/dt = v_{\text{eAc}, \text{Ac}} - U_{\text{Ac}, \text{Acm}} - U_{\text{Ac}, \text{Tgm}} \quad (3.0)$$

- Inputs:

$$v_{\text{eAc}, \text{Ac}} = V_{\text{InAc}} \text{MPF} / (1 + K_{\text{InAc}}/\text{cAc}) - V_{\text{OutAc}} \text{MPF} / (1 + K_{\text{OutAc}}/\text{cAc}) \quad (3.1)$$



Original research article

A mathematical model to describe the diurnal pattern of enteric methane emissions from non-lactating dairy cows post-feeding

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2.1. The model

The parameters for the model development are summarized in Table 1. Methane is emitted during the metabolism of methanogens that use hydrogen as an energy source, and this hydrogen is produced mainly during fermentation of degradable substrate by microorganisms in the rumen (Wang et al., 2013a). Methane emission rate (dV/dt , g/h) is assumed to be proportional to methanogen mass (M_r , g), activity of methanogens and degradable substrate (S_r , g) in the rumen, and is expressed as:

$$\frac{dV}{dt} = \alpha \beta_M M_r S_r, \quad (1)$$

where α is a proportionality constant [/(h·g)], β_M is the activity of methanogens linking the methane production and methanogen mass (g/g).

The substrate in the rumen was separated into two components: newly ingested and the residue, representing potential nutrient sources from the current and previous feeding, respectively. The total enteric methane produced associated with these feed fractions was a combination of that produced from use of residual (basal) substrate (V_1) and newly ingested (V_2) feed in the rumen.

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that changes in methane emissions are a response to substrate supply and activity of the methanogens, while methanogen mass (M_r) was assumed fixed for an individual animal on a particular ration. The rate of enteric methane emission, thus, can be expressed as follows:

$$\frac{dV}{dt} = \frac{dV_1}{dt} + \frac{dV_2}{dt}, \quad (2)$$

$$\frac{dV_1}{dt} = \alpha_1 \beta_{M1} M_r S_{rr}, \quad (2a)$$

$$\frac{dV_2}{dt} = \alpha_2 \beta_{M2} M_r S_{Ir}, \quad (2b)$$

where α_1 and α_2 are proportionality constants $[/(h \cdot g)]$ for basal V_1 and feeding V_2 , respectively; β_{M1} is the activity of methanogens to generate basal V_1 ; S_{rr} is the amount of degradable substrate in the residue of rumen before feeding (g); β_{M2} is the activity of methanogens to generate feeding V_2 ; S_{Ir} is the amount of degradable substrate in the rumen from the newly ingested feed (g).

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$$\frac{dV_2}{dt} = \alpha_2 \beta_{M2} M_r S_{Ir}. \quad (2b)$$

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| Term | Unit | Explanation |
|--------------|--------|---|
| V | g | Volume of enteric methane emission |
| V_1 | g | Volume of enteric methane emission generated by the residual substrate in the rumen |
| V_2 | g | Volume of enteric methane emission generated by the newly ingested feed |
| dV/dt | g/h | Rate of enteric methane emission |
| dV_1/dt | g/h | Rate of enteric methane emission for basal V_1 |
| dV_2/dt | g/h | Rate of enteric methane emission for feeding V_2 |
| α | /(h·g) | Proportionality constant |
| α_1 | /(h·g) | Proportionality constant for basal V_1 |
| α_2 | /(h·g) | Proportionality constant for feeding V_2 |
| β_M | — | Activity of methanogens |
| β_{M1} | — | Activity of methanogens to generate basal V_1 |
| β_{M2} | — | Activity of methanogens to generate feeding V_2 |
| S_r | g | Degrable Substrate in the rumen |
| S_{rr} | g | Degrable substrate in the residue in the rumen before feeding |
| S_{lr} | g | Degrable substrate in the rumen from the newly ingested feed |
| S_l | g | Degrable substrate from newly ingested feed |
| S_{le} | g | Degrable substrate from newly ingested feed which outflow from rumen |
| M_r | g | Methanogens in the rumen |
| k_p | /h | Ruminal passage rates |
| S_T | g | Potential degradable substrate in the newly ingested feed |
| VF_2 | g | Final asymptotic accumulated enteric methane emissions for feeding V_2 |
| γ | g/h | Shape parameter |
| d | — | Shape parameter |
| a | g | Shape parameter |

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| Term | Unit | Explanation |
|--------------|--------|---|
| V | g | Volume of enteric methane emission |
| V_1 | g | Volume of enteric methane emission generated by the residual substrate in the rumen |
| V_2 | g | Volume of enteric methane emission generated by the newly ingested feed |
| dV/dt | g/h | Rate of enteric methane emission |
| dV_1/dt | g/h | Rate of enteric methane emission for basal V_1 |
| dV_2/dt | g/h | Rate of enteric methane emission for feeding V_2 |
| α | /(h·g) | Proportionality constant |
| α_1 | /(h·g) | Proportionality constant for basal V_1 |
| α_2 | /(h·g) | Proportionality constant for feeding V_2 |
| β_M | — | Activity of methanogens |
| β_{M1} | — | Activity of methanogens to generate basal V_1 |
| β_{M2} | — | Activity of methanogens to generate feeding V_2 |
| S_r | g | Degrable Substrate in the rumen |
| S_{rr} | g | Degrable substrate in the residue in the rumen before feeding |
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| a | g | Shape parameter |

| Item | Mean | SD | Median | Minimum | Maximum |
|--|-------|-------|--------|---------|---------|
| BW, kg | 222 | 110 | 215 | 98 | 420 |
| DMI, kg/d | 4.45 | 1.38 | 4.44 | 2.66 | 7.35 |
| DMI _a :DMI _m ratio | 0.957 | 0.085 | 0.987 | 0.854 | 1.040 |
| Concentrate, kg/d | 2.96 | 0.82 | 3.22 | 1.61 | 4.02 |
| Rice straw, kg/d | 1.48 | 0.703 | 1.26 | 0.83 | 3.39 |
| Concentrate proportion in the diet, % | 66.8 | 7.55 | 68.2 | 53.9 | 78.8 |
| NDFI, kg/d | 2.30 | 0.781 | 2.20 | 1.37 | 4.14 |
| ADFI, kg/d | 1.18 | 0.426 | 1.09 | 0.70 | 2.23 |
| CPI, kg/d | 0.553 | 0.159 | 0.571 | 0.319 | 0.827 |
| GEI, MJ/d | 72.3 | 22.3 | 72.3 | 43.2 | 119 |
| Methane, g/d | 88.3 | 38.0 | 82.2 | 42.6 | 170 |
| Methane, % of GEI | 6.59 | 1.00 | 6.44 | 5.11 | 8.04 |

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Summary of variables for non-lactating dairy cows ($n = 16$).

| Item | Mean | Median | Minimum | Maximum | SD |
|--|-------|--------|---------|---------|-------|
| BW, kg | 222 | 215 | 98 | 420 | 110 |
| DMI, kg/d | 4.45 | 4.44 | 2.66 | 7.35 | 1.38 |
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BW = body weight; DMI = dry matter intake; DMI_m = DMI for morning feeding from 0600 to 1600 h; DMI_a = DMI for afternoon feeding from 1600 to 0600 h; NDFI = neutral detergent fibre intake; ADFI = acid detergent fiber intake; CPI = crude protein intake; GEI = gross energy intake; SD = standard deviation.

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|--|-------|-------|--------|---------|---------|
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2.2. Animal and housing

The use of the animals and the experimental procedure were approved by the Animal Care Committee, Institute of Subtropical Agriculture. The experiment was conducted at a local farm in the Wang-Cheng County of Hunan Province, China. Sixteen non-lactating Chinese Holstein dairy cows with a wide range of BW ([Table 2](#)) were assigned to the air-flow controlled chamber for enteric methane emission measurement.

Cows were housed in a tie-stall dairy barn, and were accustomed to restricted movement. Both gaseous exchange and feed intake were individually determined when the cow was placed in the respiration chamber. Cows were allocated to the single respiration chamber for two consecutive days in a staggered manner. The data presented are averaged from the two days of chamber. The experiment lasted from early Feb. 2012 to late Apr. 2013.

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The use of the animals and the experimental procedure were approved by the Animal Care Committee, Institute of Subtropical Agriculture. The experiment was conducted at a local farm in the Wang-Cheng County of Hunan Province, China. Sixteen non-lactating Chinese Holstein dairy cows with a wide range of BW ([Table 2](#)) were assigned to the air-flow controlled chamber for enteric methane emission measurement.

Cows were housed in a tie-stall dairy barn, and were accustomed to restricted movement. Both gaseous exchange and feed intake were individually determined when the cow was placed in the respiration chamber. Cows were allocated to the single respiration chamber for two consecutive days in a staggered manner. The data presented are averaged from the two days of chamber. The experiment lasted from early Feb. 2012 to late Apr. 2013.

2.3. Diet and feeding

The diet consisted of concentrate and roughage (rice straw). The concentrate contained maize, soybean meal, cottonseed meal and corn distiller's dried grains and maize with solubles, purchased from Agribrands Purina Feed mill Co., Ltd. The chemical composition of the concentrate was 950 g DM/kg and 155 g of CP, 415 g of neutral detergent fibre (NDF) and 157 g of acid detergent fibre (ADF) per kg of DM. The chemical composition for the rice straw was (on a DM basis) 975 g/kg DM, 63 g/kg CP, 760 g/kg NDF and 466 g/kg ADF.

The allowances of concentrate and roughage were decided by the farmer, based on experience and according to the live weight of individual cows (each around 1% of live weight). As a result, the amount of concentrate supplied was different for each animal ([Table 2](#)). The concentrate and roughage were placed in two separate feeding troughs, with the concentrate provided first. All animals had ad libitum access to water. The restricted supply of concentrate was divided into two portions for the morning and afternoon feeds (0600 and 1605 h) while the rice straw was provided in slight excess for both periods. Orts were collected twice daily before the new feed was provided. The characteristics of feed intake for all animals are shown in [Table 2](#).

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2.3. Diet and feeding

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One simple respiration chamber was built for the measurement of methane emissions from cows. Briefly, the chamber was made of galvanized steel plate with internal dimensions of 3 m length × 2 m width × 2 m height. The chamber had one front and one rear door

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2.4. Measurement of methane emissions

One simple respiration chamber was built for the measurement of methane emissions from cows. Briefly, the chamber was made of galvanized steel plate with internal dimensions of 3 m length × 2 m width × 2 m height. The chamber had one front and one rear door

fitted with internal rubber seals. The cow was restrained in the chamber with access to a feed bin and a drinking water container. A fresh air inlet was located at the top left of the chamber, and air inlets were piped from an intake vent, located 15 m from the chamber. The outlet consisted of two round polyethylene pipes (outside diameter, 50 mm) fixed to the left and right insides of the chamber, and each pipe comprised of 50 intake holes equally distributed around the entire circumference of the duct. These two ducts were piped through the right side of chamber via a 50 mm outside diameter polyethylene pipe. The outlet was connected via a 50 mm air filter, to a gas flow meter, followed by the pump. Airflow (150 to 190 m^3/h) under negative pressure was controlled by the pump. The chamber was fitted with four internal ventilation fans for efficient mixing of exhaled gases and incoming air. The outlet pipe from the chamber was connected to a plastic buffer box (50 cm length \times 50 cm width \times 50 cm height) for gas sampling.

The outlet gas was sampled from the box every 15 min during 0600 to 2200 h, at 2300, 2400 h, next day 0200 and 0530 h. A 50-mL syringe was used for sampling, and then injected into a vacuum tube for methane determination by gas chromatography (Agilent 7890A, Agilent Inc., Palo Alto, CA).

The cows were placed in the chamber at 0600 h. The cows were fed after entering the chamber at 0600 h, and the chamber was opened once a day at 1605 h for 5 min to deliver diet. The first sample of outlet gas was collected after the cows had been shut in the chamber for 10 min. Three inlet gas samples were collect at 0600, 1200 and 1700 h, and their mean value used to represent the methane concentration of the inflowing air.

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7890A, Agilent Inc., Palo Alto, CA)

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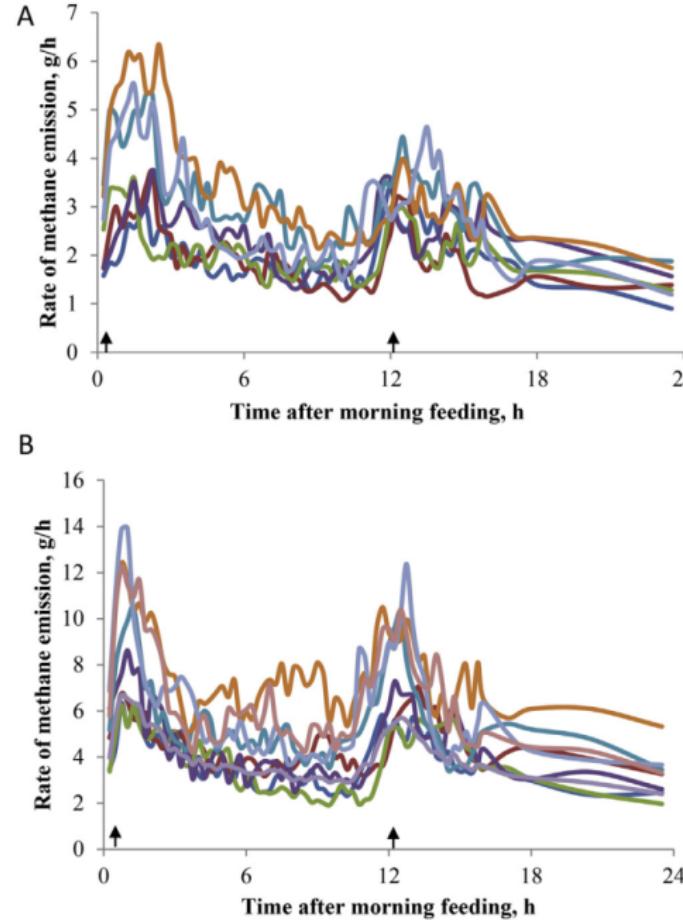
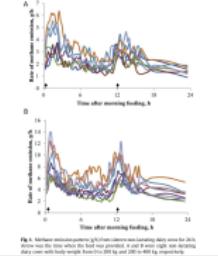
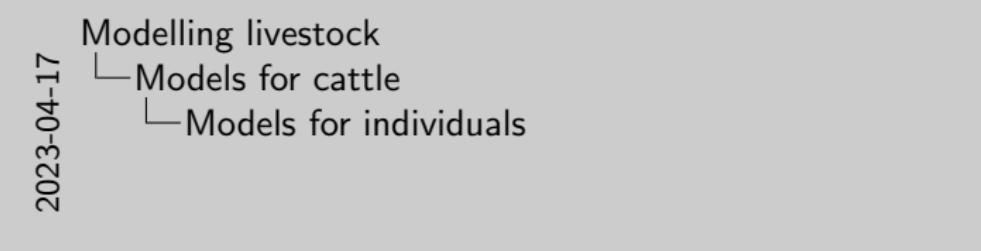
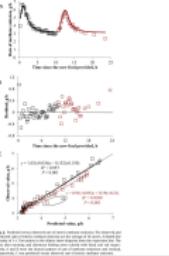


Fig. 1. Methane emission pattern (g/h) from sixteen non-lactating dairy cows for 24 h. Arrow was the time when the feed was provided. A and B were eight non-lactating dairy cows with body weight from 0 to 200 kg and 200 to 400 kg, respectively.





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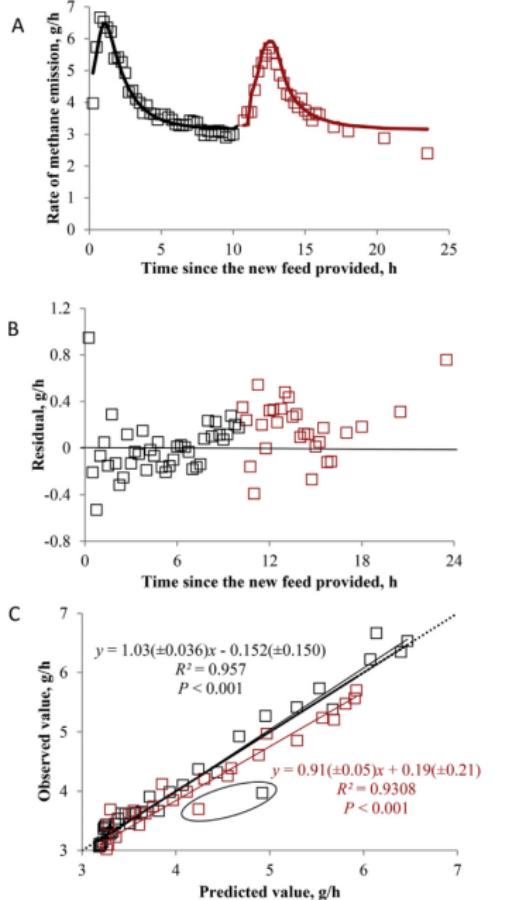


Fig. 2. Predicted versus observed rate of enteric methane emission. The observed and predicted rates of enteric methane emission are the average of 16 curves. A dotted line is unity of 1:1. Two points in the ellipse show disparity from the regression line. The data after morning and afternoon feeding were colored with black and red respectively. A and B were the diurnal pattern of rate of methane emissions and residual, respectively; C was predicted versus observed rate of enteric methane emission.

Development of mathematical models to predict volume and nutrient composition of fresh manure from lactating Holstein cows

J. A. D. Ranga Niroshan Appuhamy^{A,E}, L. E. Moraes^A, C. Wagner-Riddle^B, D. P. Casper^C, J. France^D and E. Kebreab^A

^ADepartment of Animal Science, University of California, Davis, CA 95616, USA.

^BSchool of Environmental Sciences, University of Guelph, Guelph, ON, N1G 2W1, Canada.

^CDepartment of Dairy Science, South Dakota State University, Brookings, SD 57007, USA.

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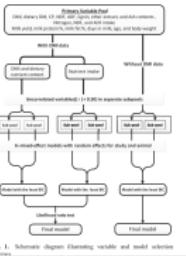
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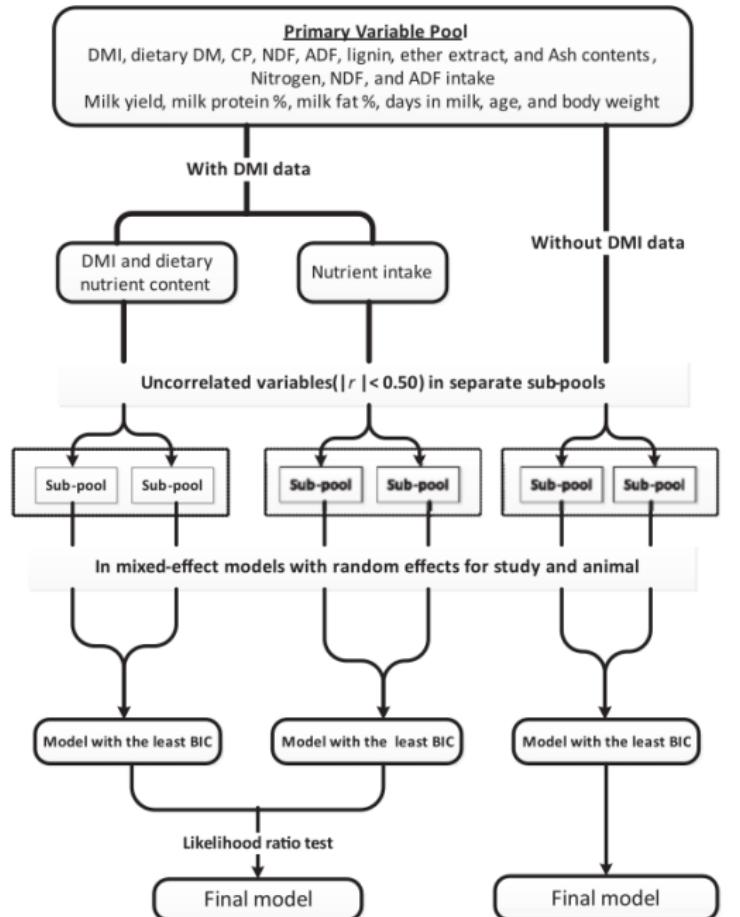


Fig. 1. Schematic diagram illustrating variable and model selection schemes.

Table 5. Prediction equations (standard errors of parameters in parentheses), maximum variance inflation factor (max_VIF) and the determinant of the correlation matrix ($\det\{R\}$) of the selected variables

F_{DM} = faecal dry matter, F_C = faecal carbon, F_{NDF} = faecal neutral detergent fibre (NDF), F_{ADF} = faecal acid detergent fibre (ADF), F_N = faecal nitrogen, F_{Water} = faecal water, U_E = total urine, U_C = urine carbon, and U_N = urine nitrogen outputs (all in kg/day). DMI = dry matter intake (kg/day), CP, ADF, NDF and LIG = dietary crude protein, ADF, NDF and lignin content, respectively (% of DM). Milk = milk yield (kg/day), mPr = milk protein percentage, DIM = days in milk, BW = bodyweight (kg/cow), Age = age of the cows (years)

| Equation | Variables and parameter estimates \pm standard error | max_VIF | $\det\{R\}$ |
|--------------------|---|---------|-------------|
| <i>With DMI</i> | | | |
| (1) | $F_{DM} = -0.576 \pm 0.222 + (0.370 \pm 0.006 \times DMI) + (-0.075 \pm 0.010 \times CP) + (0.059 \pm 0.006 \times ADF)$ | 1.02 | 0.90 |
| (2) | $F_C = (0.169 \pm 0.003 \times DMI) + (-0.034 \pm 0.004 \times CP) + (0.027 \pm 0.003 \times ADF) + (-0.075 \pm 0.019 \times mPr)$ | 1.04 | 0.87 |
| (3) | $F_{NDF} = -0.864 \pm 0.172 + (0.217 \pm 0.004 \times DMI) + (0.035 \pm 0.003 \times NDF) + (-0.039 \pm 0.007 \times CP)$ | 1.01 | 0.86 |
| (4) | $F_{ADF} = -1.272 \pm 0.084 + (0.125 \pm 0.003 \times DMI) + (0.061 \pm 0.003 \times ADF)$ | 1.00 | 0.99 |
| (5) | $F_N = -0.0368 \pm 0.007 + (0.0096 \pm 0.000 \times DMI) + (0.0022 \pm 0.000 \times CP) + (0.0034 \pm 0.001 \times lignin)$ + (-0.000043 \pm 0.000010 $\times BW$) | 1.09 | 0.80 |
| (6) | $F_{Water} = (1.987 \pm 0.034 \times DMI) + (0.348 \pm 0.032 \times ADF) + (-0.412 \pm 0.052 \times CP) + (-0.074 \pm 0.009 \times DM)$ + (-0.0057 \pm 0.0012 $\times DIM$) | 1.16 | 0.80 |
| (7) | $U_E = -7.742 \pm 2.367 + (0.388 \pm 0.055 \times DMI) + (0.726 \pm 0.096 \times CP) + (2.066 \pm 0.421 \times mPr)$ | 1.05 | 0.94 |
| (8) | $U_C = -0.1601 \pm 0.0169 + (0.0082 \pm 0.0005 \times DMI) + (0.0107 \pm 0.0008 \times CP) + (0.00013 \pm 0.00002 \times BW)$ | 1.13 | 0.84 |
| (9) | $U_N = -0.2837 \pm 0.0135 + (0.0068 \pm 0.0004 \times DMI) + (0.0155 \pm 0.0006 \times CP) + (0.00013 \pm 0.00001 \times DIM)$ + (0.000092 \pm 0.000017 $\times BW$) | 1.34 | 0.66 |
| <i>Without DMI</i> | | | |
| (10) | $F_{DM} = 0.846 \pm 0.469 + (0.098 \pm 0.004 \times Milk) + (-0.097 \pm 0.021 \times CP) + (0.080 \pm 0.012 \times ADF)$ + (0.0038 \pm 0.0005 $\times BW$) | 1.04 | 0.80 |
| (11) | $F_C = 0.468 \pm 0.232 + (0.046 \pm 0.002 \times Milk) + (-0.047 \pm 0.010 \times CP) + (0.037 \pm 0.006 \times ADF)$ + (0.0016 \pm 0.0002 $\times BW$) | 1.04 | 0.80 |
| (12) | $F_{NDF} = (0.056 \pm 0.003 \times Milk) + (-0.059 \pm 0.010 \times CP) + (0.0435 \pm 0.0042 \times NDF) + (0.0023 \pm 0.0003 \times BW)$ | 1.00 | 0.77 |
| (13) | $F_{ADF} = -0.973 \pm 0.152 + (0.0325 \pm 0.0016 \times Milk) + (0.0675 \pm 0.0043 \times ADF) + (0.0014 \pm 0.0002 \times BW)$ | 1.00 | 0.98 |
| (14) | $F_N = (0.00245 \pm 0.00011 \times Milk) + (0.00643 \pm 0.00082 \times LIG) + (0.000094 \pm 0.000009 \times BW)$ | 1.00 | 0.99 |
| (15) | $F_{Water} = (0.559 \pm 0.025 \times Milk) + (0.521 \pm 0.060 \times ADF) + (0.569 \pm 0.100 \times CP) + (0.024 \pm 0.003 \times BW)$ + (-0.033 \pm 0.012 $\times Age$) | 1.17 | 0.66 |
| (16) | $U_E = -0.644 \pm 0.226 + (0.778 \pm 0.099 \times CP) + (1.520 \pm 0.426 \times mPr)$ | 1.00 | 0.99 |
| (17) | $U_C = -0.1167 \pm 0.0201 + (0.0013 \pm 0.0002 \times Milk) + (0.0106 \pm 0.0009 \times CP) + (0.00024 \pm 0.00002 \times BW)$ | 1.03 | 0.87 |
| (18) | $U_N = -0.2578 \pm 0.0183 + (0.0152 \pm 0.0007 \times CP) + (0.0132 \pm 0.0031 \times mPr) + (0.00021 \pm 0.00002 \times BW)$ | 1.01 | 0.98 |

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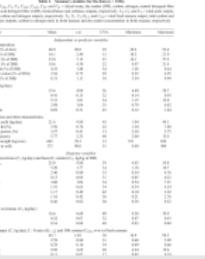
Table 1. Summary statistics for the data ($n = 1106$)
 F_{Water} , F_{DM} , F_C , F_N , F_{NDF} , F_{ADF} , F_{HC} and F_{CL} = faecal water, dry matter (DM), carbon, nitrogen, neutral detergent fibre (NDF), acid detergent fibre (ADF), hemicellulose and cellulose outputs, respectively. U_E , U_C and U_N = total urine output, urinary carbon and nitrogen outputs, respectively. T_E , T_C , T_N , $R_{C:N}$ and C_{DM} = total fresh manure output, total carbon and nitrogen outputs, carbon to nitrogen ratio in fresh manure and dry matter concentration in fresh manure, respectively

| Variable | Mean | s.d. | CV% | Minimum | Maximum |
|---|------|------|-----|---------|---------|
| <i>Independent or predictor variables</i> | | | | | |
| Diet composition | | | | | |
| DM (% of diet) | 68.0 | 20.0 | 29 | 30.2 | 93.8 |
| CP (% of DM) | 16.1 | 2.40 | 15 | 10.3 | 21.9 |
| NDF (% of DM) | 33.8 | 7.13 | 21 | 16.1 | 57.2 |
| ADF (% of DM) | 19.6 | 4.30 | 22 | 8.97 | 31.4 |
| Lignin (% of DM) | 4.33 | 1.48 | 34 | 1.26 | 8.44 |
| Ether extract (% of DM) | 2.56 | 0.75 | 29 | 0.52 | 4.95 |
| Ash (% of DM) | 6.31 | 1.11 | 18 | 3.54 | 9.99 |
| Intake (kg/day) | | | | | |
| DM | 15.6 | 4.08 | 26 | 6.40 | 28.7 |
| N | 0.41 | 0.13 | 32 | 0.14 | 0.93 |
| NDF | 5.31 | 1.81 | 34 | 1.15 | 12.0 |
| ADF | 3.08 | 1.06 | 35 | 0.70 | 6.82 |
| Lignin | 0.69 | 0.31 | 45 | 0.12 | 1.84 |
| Production and other characteristics | | | | | |
| Milk yield (kg/day) | 21.6 | 9.80 | 45 | 1.04 | 49.1 |
| Milk fat (%) | 3.50 | 0.76 | 22 | 1.30 | 7.60 |
| Milk protein (%) | 3.27 | 0.41 | 13 | 2.30 | 5.75 |
| Age (years) | 5.77 | 2.33 | 40 | 2.00 | 15.4 |
| Bodyweight (kg/cow) | 603 | 78.3 | 13 | 351 | 854 |
| Days in milk | 175 | 90.0 | 51 | 0.00 | 488 |
| <i>Response variables</i> | | | | | |
| Faecal excretions (F_x , kg/day) and faecal C content (C_x , kg/kg of DM) | | | | | |
| F_{Water} | 25.0 | 9.80 | 39 | 4.03 | 59.8 |
| F_{DM} | 5.20 | 1.77 | 34 | 1.18 | 10.7 |
| F_C | 2.40 | 0.80 | 33 | 0.54 | 4.76 |
| F_N | 0.13 | 0.04 | 31 | 0.05 | 0.25 |
| F_{NDF} | 3.06 | 1.04 | 34 | 0.54 | 7.21 |
| F_{ADF} | 1.91 | 0.65 | 34 | 0.34 | 4.24 |
| F_{HC} | 1.17 | 0.49 | 42 | 0.10 | 3.22 |
| F_{CL} | 1.16 | 0.42 | 36 | 0.21 | 2.76 |
| C_C | 0.46 | 0.02 | 04 | 0.38 | 0.52 |
| Urine excretions (U_x , kg/day) | | | | | |
| U_E | 16.6 | 6.60 | 40 | 4.38 | 34.9 |
| U_C | 0.22 | 0.07 | 32 | 0.07 | 0.43 |
| U_N | 0.16 | 0.08 | 48 | 0.03 | 0.40 |
| Total output (T_x , kg/day), C : N ratio ($R_{C:N}$), and DM content (C_{DM} , w/w) of fresh manure | | | | | |
| T_E | 46.7 | 14.0 | 30 | 16.9 | 98.5 |
| T_C | 2.59 | 0.80 | 31 | 0.68 | 5.09 |
| T_N | 0.29 | 0.10 | 34 | 0.09 | 0.66 |
| $R_{C:N}$ | 9.50 | 2.69 | 28 | 4.24 | 19.6 |
| C_{DM} | 0.11 | 0.02 | 17 | 0.05 | 0.19 |

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Models for individuals

Models for herds

Models for other herds

Models for management

Conclusion

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Models for cattle
 Models for individuals
 Models for herds

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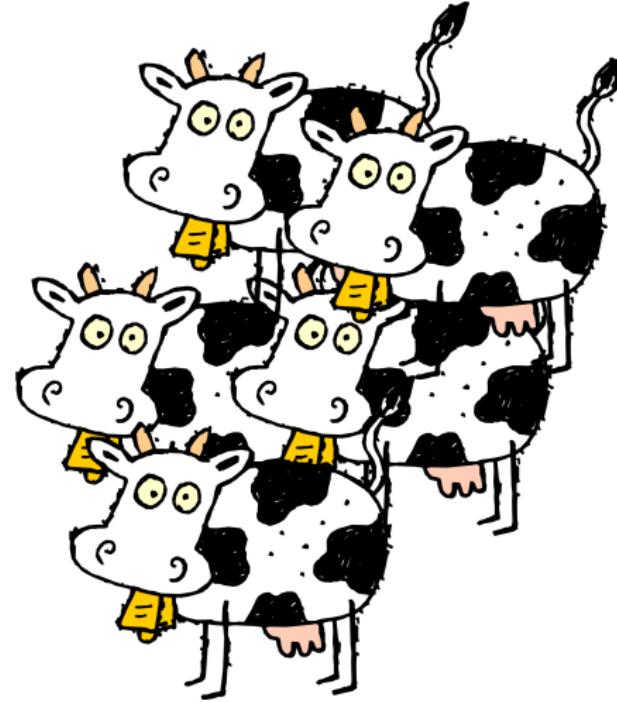
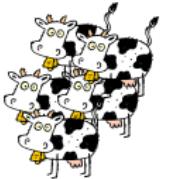
Conclusion

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At the herd level



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A mathematical model for the dynamics and synchronization of cows

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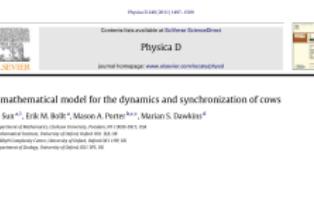
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We model the biological status of a single cow by

$$w = (x, y; \theta) \in [0, 1] \times [0, 1] \times \Theta. \quad (1)$$

The real variables x and y represent, respectively, the extent of desire to eat and lie down of the cow, and

$$\theta \in \Theta = \{\mathcal{E}, \mathcal{R}, \mathcal{S}\} \quad (2)$$

is a discrete variable that represents the state of the cow (see the equations below for descriptions of the states). Throughout this paper, we will refer to θ as a *symbolic variable* or a *state variable*. One can think of the symbolic variable θ as describing a switch that triggers different time-evolution rules for the other two variables x and y .

We model the dynamics of a single cow in different states using

$$(\mathcal{E}) \text{ Eating state: } \begin{cases} \dot{x} = -\alpha_2 x, \\ \dot{y} = \beta_1 y. \end{cases} \quad (3)$$

$$(\mathcal{R}) \text{ Resting state: } \begin{cases} \dot{x} = \alpha_1 x, \\ \dot{y} = -\beta_2 y. \end{cases} \quad (4)$$

$$(\mathcal{S}) \text{ Standing state: } \begin{cases} \dot{x} = \alpha_1 x, \\ \dot{y} = \beta_1 y, \end{cases} \quad (5)$$

where the calligraphic letters inside parentheses indicate the corresponding values of θ . For biological reasons, the parameters α_1 , α_2 , β_1 , and β_2 must all be positive real numbers. They can be interpreted as follows:

$$\begin{cases} \alpha_1 : \text{rate of increase of hunger,} \\ \alpha_2 : \text{decay rate of hunger,} \\ \beta_1 : \text{rate of increase of desire to lie down,} \\ \beta_2 : \text{decay rate of desire to lie down.} \end{cases}$$

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We model the biological state of a single cow by
 $w = (x, y; \theta) \in [0, 1] \times [0, 1] \times \Theta$ (1)
The real variables x and y represent, respectively, the extent of desire to eat and lie down of the cow, and
 $\theta \in \Theta \subseteq [1, 8, 4]$ (2)
 θ is a discrete variable that represents the state of the cow (see the equations below for descriptions of the states). Throughout this paper, we will refer to θ as a *symbolic variable* or a *state variable*. One can think of the symbolic variable θ as describing a switch that triggers different time-evolution rules for the other two variables x and y .
We model the dynamics of a single cow in different states using:
(E) Eating state: $\begin{cases} \dot{x} = -\alpha_2 x, \\ \dot{y} = \beta_1 y. \end{cases}$ (3)
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(S) Standing state: $\begin{cases} \dot{x} = \alpha_1 x, \\ \dot{y} = \beta_1 y. \end{cases}$ (5)
where the calligraphic letters inside parentheses indicate the corresponding values of θ . For biological reasons, the parameters $\alpha_1, \alpha_2, \beta_1, \beta_2$ must all be positive real numbers. They can be interpreted as follows:
 α_1 : rate of increase of hunger,
 α_2 : decay rate of hunger,
 β_1 : rate of increase of desire to lie down,
 β_2 : decay rate of desire to lie down.

The dynamics within each state does not fully specify the equations governing a single cow. To close the bovine equations, we also need switching conditions that determine how the state variable θ changes. We illustrate these switching conditions in Fig. 1 and describe them in terms of equations as follows:

$$\theta \rightarrow \begin{cases} \mathcal{E} & \text{if } \theta \in \{\mathcal{R}, \mathcal{S}\} \text{ and } x = 1, \\ \mathcal{R} & \text{if } \theta \in \{\mathcal{E}, \mathcal{S}\} \text{ and } x < 1, y = 1, \\ \mathcal{S} & \text{if } \theta \in \{\mathcal{E}, \mathcal{R}\} \text{ and } x < 1, y = \delta \text{ (or } x = \delta, y < 1). \end{cases} \quad (6)$$

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2.2. Switching conditions

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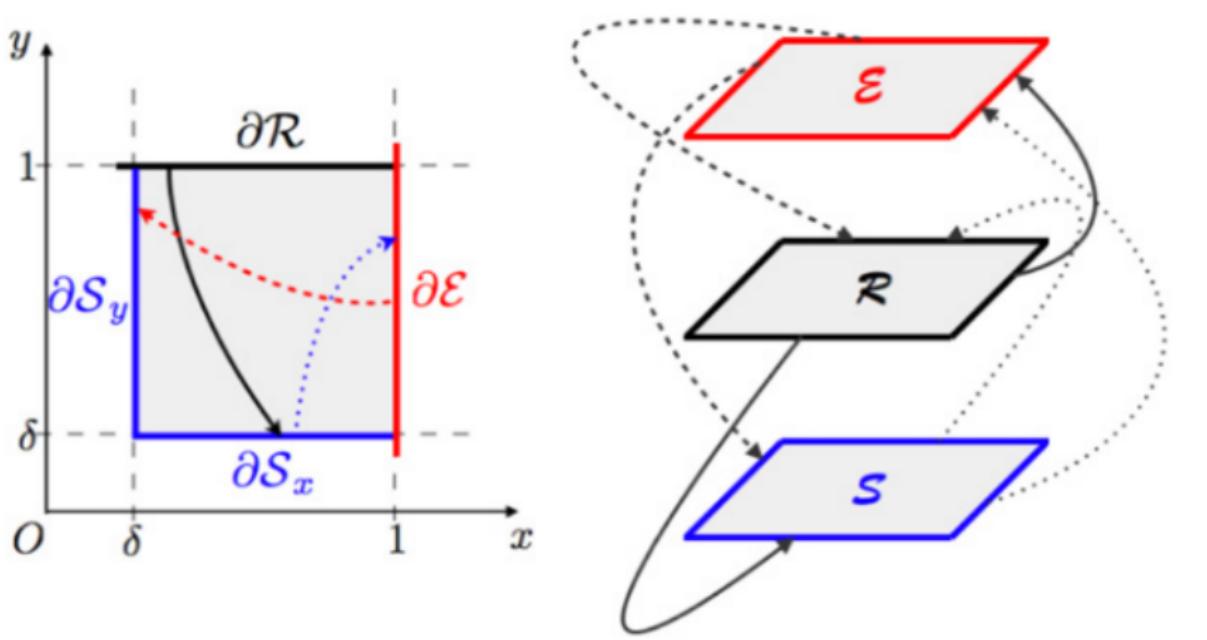
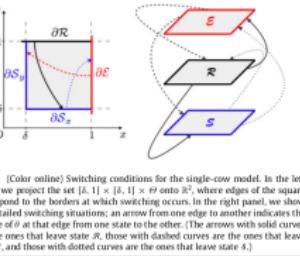


Fig. 1. (Color online) Switching conditions for the single-cow model. In the left panel, we project the set $[\delta, 1] \times [\delta, 1] \times \Theta$ onto \mathbb{R}^2 , where edges of the square correspond to the borders at which switching occurs. In the right panel, we show the detailed switching situations; an arrow from one edge to another indicates the change of θ at that edge from one state to the other. (The arrows with solid curves are the ones that leave state \mathcal{R} , those with dashed curves are the ones that leave state \mathcal{E} , and those with dotted curves are the ones that leave state \mathcal{S} .)

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4.1. The coupling scheme

There are numerous possible ways to model the coupling between cows. We have chosen one based on the hypothesis that a cow feels hungrier when it notices other cows eating and feels a greater desire to lie down when it notices other cows lying down. (We briefly discuss other possibilities in Section 5.) This provides a coupling that does not have a spatial component, in contrast to the agent-based approach of Ref. [30]. We therefore assume implicitly that space is unlimited, so we are considering cows to be in a field rather than in a pen. We suppose that the herd consists of n cows and use i to represent the i th cow in the herd. This yields herd equations given by

$$\begin{cases} \dot{x}_i = \left[\alpha^{(i)}(\theta_i) + \frac{\sigma_x}{k_i} \sum_{j=1}^n a_{ij} \chi_{\mathcal{E}}(\theta_j) \right] x_i, \\ \dot{y}_i = \left[\beta^{(i)}(\theta_i) + \frac{\sigma_y}{k_i} \sum_{j=1}^n a_{ij} \chi_{\mathcal{R}}(\theta_j) \right] y_i, \end{cases} \quad (26)$$

with the switching condition given by Eq. (6) for each individual cow. The summation terms in both equations give the coupling terms of this system. The matrix $A = [a_{ij}]_{n \times n}$ is a time-dependent adjacency matrix that represents the network of cows. Its components are given by

$$a_{ij}(t) = \begin{cases} 1 & \text{if the } i\text{th cow perceives the} \\ & \text{ } j\text{th cow at time } t, \\ 0 & \text{if the } i\text{th cow does not perceive the} \\ & \text{ } j\text{th cow at time } t. \end{cases} \quad (27)$$

Additionally, $k_i = \sum_{j=1}^n A_{ij}$ is the degree of node i (i.e., the number of cows to which it is connected), and the coupling strengths σ_x and σ_y are non-negative (and usually positive) real numbers. This is designed to emphasize that animal interaction strengths consider proximity to neighboring animals.

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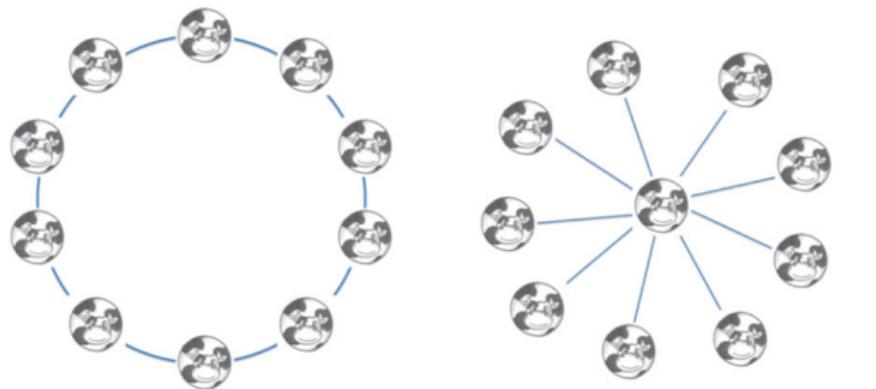


Fig. 8. (Color online) Example network architectures for coupled cows: (left) circular lattice with 10 nodes and (right) star graph with 10 nodes. (The spherical cow image was created for this paper by Yulian Ng and is used with her permission.)

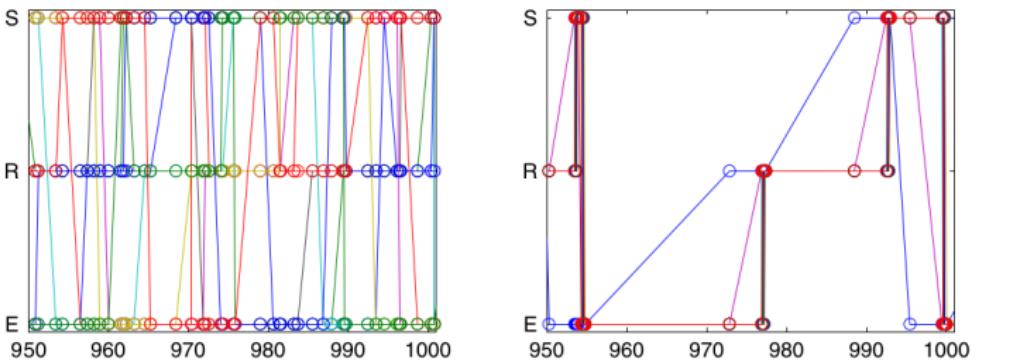
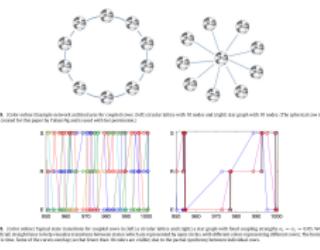


Fig. 9. (Color online) Typical state transitions for coupled cows in (left) a circular lattice and (right) a star graph with fixed coupling strengths $\sigma_x = \sigma_y = 0.05$. We plot (artificial) straight lines to help visualize transitions between states (which are represented by open circles, with different colors representing different cows). The horizontal axis is time. Some of the curves overlap (so that fewer than 10 colors are visible) due to the partial synchrony between individual cows.

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Model for the spatial pattern formed by a small herd in grazing cattle

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increase. In other words, the area reaches an equilibrium, and attraction activities (desire to be in a group) and repulsion activities (maintenance of individual space) operating among individuals are well-balanced in the herd although the area they occupy is elastic within fences. Because the area they occupy indicates the strength of unity or closeness among individuals in the herd, the analysis of the area should provide basic information for managing a cattle herd in paddocks. In the present study, only the distance between the far-left and far-right individuals in a small herd grazed in an experimental strip-wise pasture was observed instead of the area the cattle herd occupied. This distance is referred to as ‘troop length’ thereafter.

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Suppose that there is a straight line with length of θ , and n individuals are located independently and randomly at points whose distances from the origin are x_1, x_2, \dots, x_n according to the following rectangular distribution:

$$\begin{aligned}f(x) dx &= dx/\theta \quad \text{for } 0 \leq x \leq \theta \\f(x) dx &= 0 \quad \text{elsewhere.}\end{aligned}\tag{1}$$

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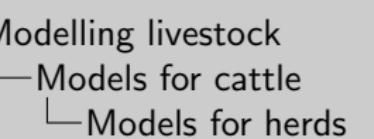
$$f(x) dx = 0 \quad \text{elsewhere.} \tag{1}$$

2.1. Probability density function of the troop length in the case of random patterns

We assume that the n independent individuals are located randomly on the line segment with a length of θ , and let y be the ‘troop length’. The probability density function for the troop length was derived using a sampling theory of order statistics (e.g. Wilks, 1962), in the following form:

$$f(y) \, dy = n(n-1)y^{n-2}(\theta-y)/\theta^n \, dy \text{ for } 0 \leq y \leq \theta \\ f(y) \, dy = 0 \text{ elsewhere.} \quad (2)$$

The expected distance, μ , and the variance, σ^2 , for y are expressed by the following equations, respectively: $\mu = (n-1)\theta/(n+1)$ and $\sigma^2 = 2(n-1)\theta^2/\{(n+2)(n+1)^2\}$.



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2.2. Deterministic model describing the changes in troop length

We express the troop length at time t by y , and assume that y changes during an infinitesimal period of time, dt , according to the following relationship (Fig. 1):

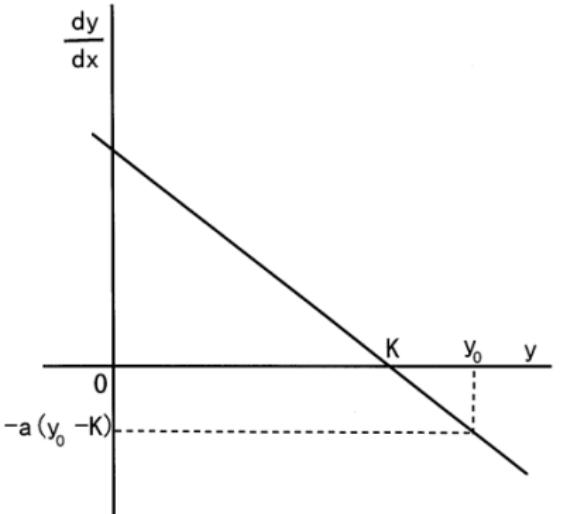


Fig. 1. Deterministic model representing the relationship between the troop length, y , and the change in an infinitesimal time period, dy/dt .

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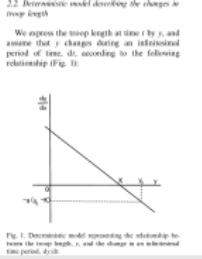


Fig. 1. Deterministic model representing the relationship between the troop length, y , and the change in an infinitesimal time period, dy/dt .

$$\frac{dy}{dt} = -a(y - K), \quad a \geq 0, \quad K \geq 0, \quad (3)$$

where a indicates the ‘convergence rate’ of troop length to the ‘equilibrium’ troop length K . Eq. (3) indicates that: (1) troop length, y , decreases during the following dt if $y > K$ (attraction); (2) y increases during the following dt if $y < K$ (repulsion); and (3) y does not change during the following dt if $y = K$ (equilibrium). Eq. (3) implies that the changes in the troop length with time occur solely based on attraction and repulsion operating among individuals within the herd. When we assume that the troop length at time t_0 is y_0 , the following solution is obtained from Eq. (3):

$$y = K - (K - y_0) e^{-at}. \quad (4)$$

For $t \rightarrow \infty$, the troop length, y , in Eq. (4) approaches K in monotone. It is empirically evident that this deterministic model does not fit to the actual behavior of a cattle herd because it is unlikely that the troop length of cattle converges to a constant, K , with time, without fluctuations. Actual troop length may fluctuate around K as the example shown below (Fig. 3). A stochastic model modifying Eq. (3) to describe the actual fluctuations is proposed in the following sections.

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2.3. Stochastic model describing changes in troop length

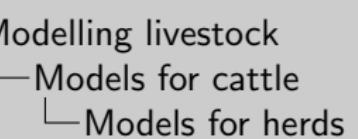
We assume that dy/dt follows: (1) the attractive and repulsive activities operating between individuals; and (2) a random movement or involuntary activity referred to as white noise in physics, ε , in the changes of the troop length at time t . Then, we have

$$dy/dt = -a(y - K) + \varepsilon. \quad (5)$$

Let us assume that the probability that the troop length, y , occurs between Y and $Y + \Delta Y$ at t , where ΔY denotes an infinitesimal length, is expressed by $g(y, t) dy$:

$$\text{Prob}\{Y \leq y(t) \leq Y + \Delta Y\} = g(y, t) dy,$$

where g denotes a probability density function of y at t .



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Then, by applying the Kolmogorov diffusion equation (e.g. Bharucha-Reid, 1960) to Eq. (5), we obtain the following equation:

$$\frac{\partial g(y, t)}{\partial t} = \frac{-\partial}{\partial y}[-a(y - K)g(y, t)] + \frac{1}{2} \frac{\partial^2}{\partial y^2}[\sigma^2 g(y, t)], \quad (6)$$

where σ^2 denotes a constant relating to the intensity of the involuntary activity.

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In a pasture in which a given cattle herd is grazed for a long period of time, the interrelationships between the herd members and the troop length of the herd are likely to be stable. We do not need to solve Eq. (6) directly, because of this stability, and we obtain a probability density function of y , only by putting $\partial g(y, t)/\partial t = 0$. Then, we have:

$$a(y - K)g(y) + \frac{1}{2} \frac{d}{dy} [\sigma^2 g(y)] = 0, \quad (7)$$

where $g(y)$ is independent of t .

From Eq. (7), we obtain the probability density function, $g(y)$, for troop length, y , as follows:

$$g(y) = \exp \left[-\frac{a}{\sigma^2}(y - K)^2 \right] / R,$$

where

$$R = \int_0^\theta \exp \left[-\frac{a}{\sigma^2}(y - k)^2 \right] dy, \quad a \geq 0, \quad 0 \leq K \leq \theta \quad (8)$$

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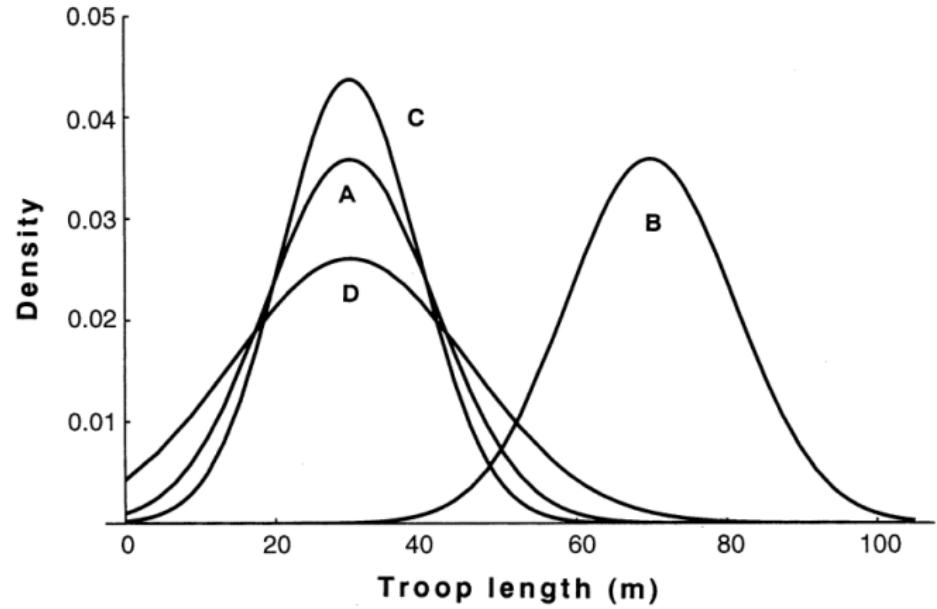


Fig. 2. Examples for various values of three parameters in Eq. (8) ($\theta = 100$). (A) $K = 30$, $a = 1.0$, $\sigma^2 = 250$; (B) $K = 70$, $a = 1.0$, $\sigma^2 = 250$; (C) $K = 30$, $a = 1.5$, $\sigma^2 = 250$; and (D) $K = 30$, $a = 1.0$, $\sigma^2 = 500$.

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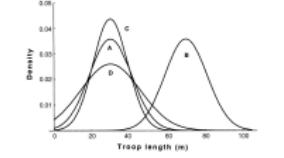


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3.1. Materials and methods

A pasture 88 m × 6 m in size was designed, to observe easily the cattle activities and to facilitate theoretical considerations under the experimental conditions, at the National Grassland Research Institute at Nishinasuno, Tochigi, Japan. The main plant species in the pasture were orchard grass, tall fescue and white clover. A grazing experiment was carried out using six Holstein heifers aged 1–2 years with a body weight ranging from 200 to 300 kg in 1979. The width of the pasture, 6 m, was sufficient for three or more cows to walk side by side. The positions of each of the six cows were visually observed and recorded every 20 min. The observation was started at 07:30 h on 5 June, and continued for ≈3 days except during the night when the positions could not be observed visually.

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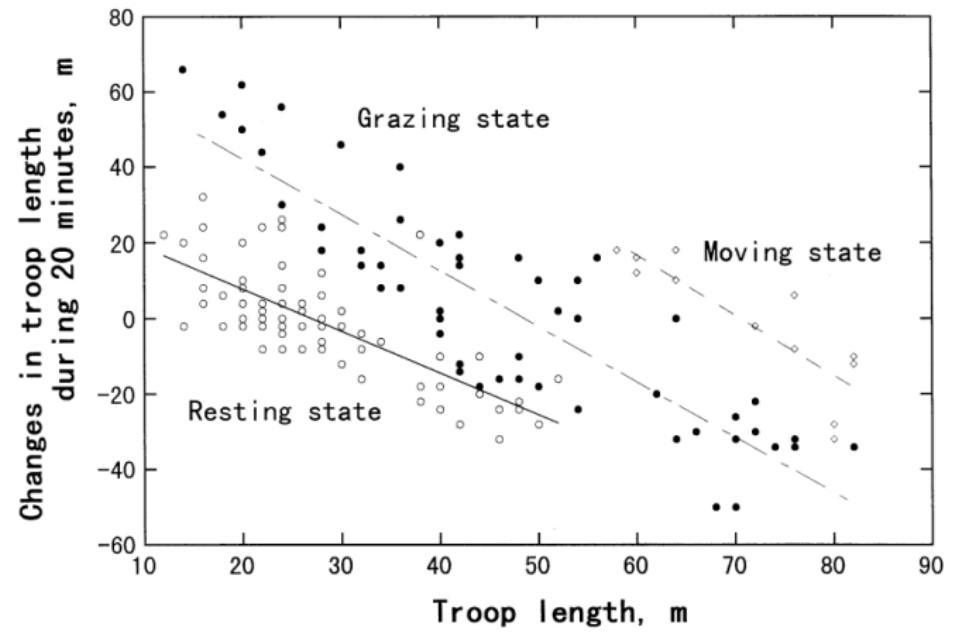


Fig. 3. Relationships, obtained in a field experiment, between the troop length at time t and change during successive 20 min intervals. Symbols ●, ✕ and ○ denote the resting, feeding and moving states, respectively.



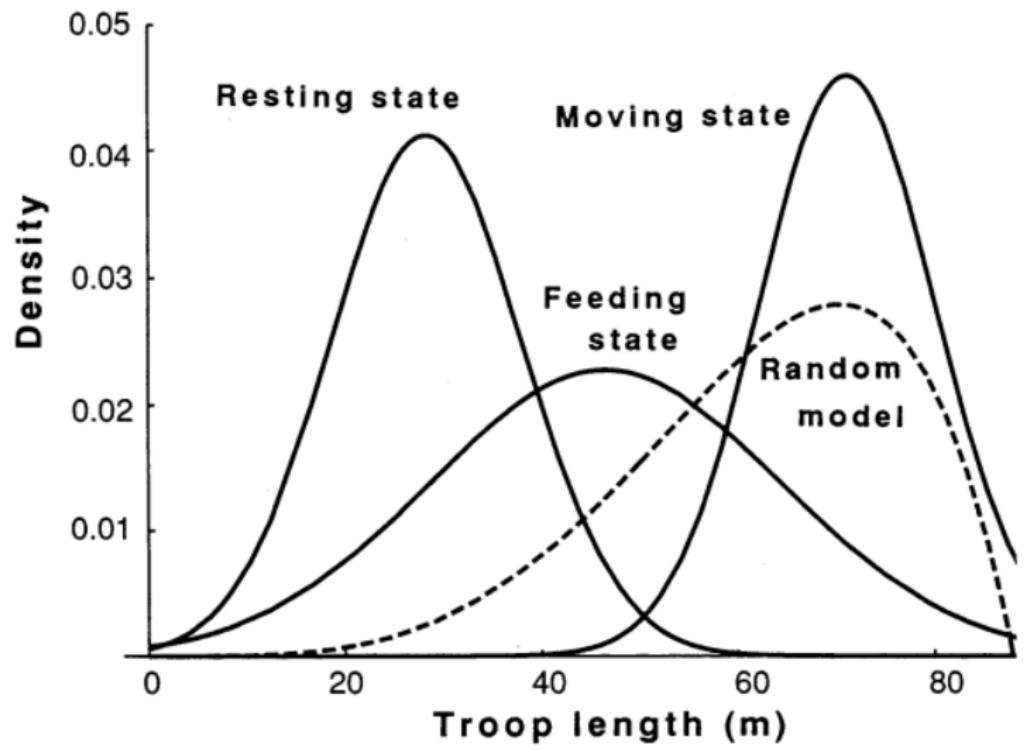


Fig. 4. Probability density functions, obtained for the experimental data, for the resting, feeding and moving states (Eq. (8)) and for the hypothetical random spatial pattern (Eq. 2). The parameter values used in the calculations are listed in Table 1.



4

Published 1989

Measurements of the Plant-Animal Interface in Grazing Research¹

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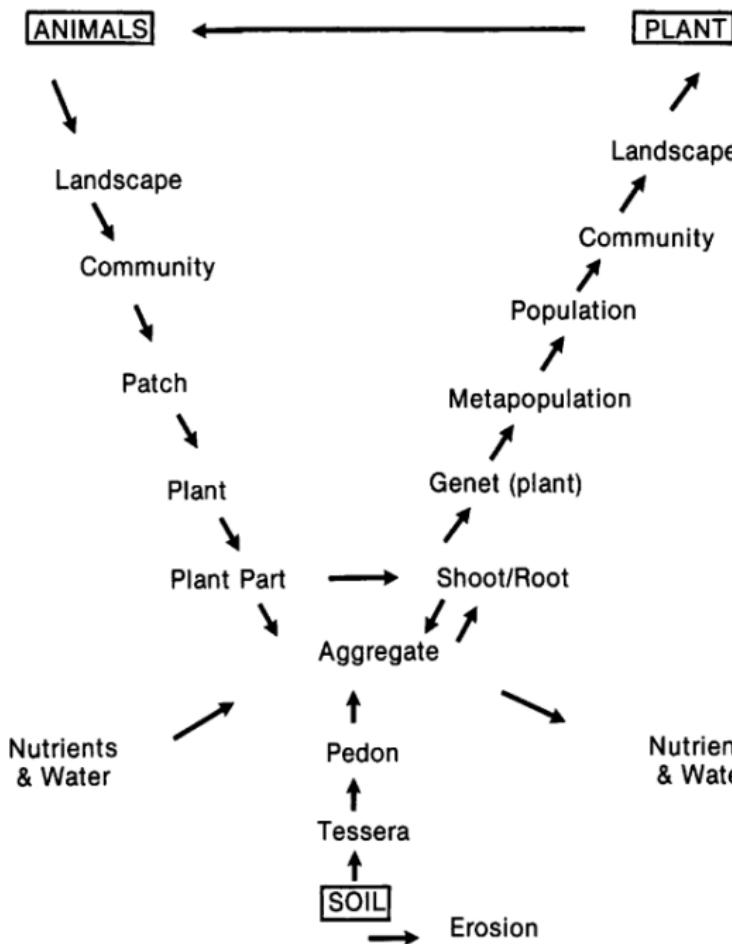
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The hierarchical context of the plant-animal interface.

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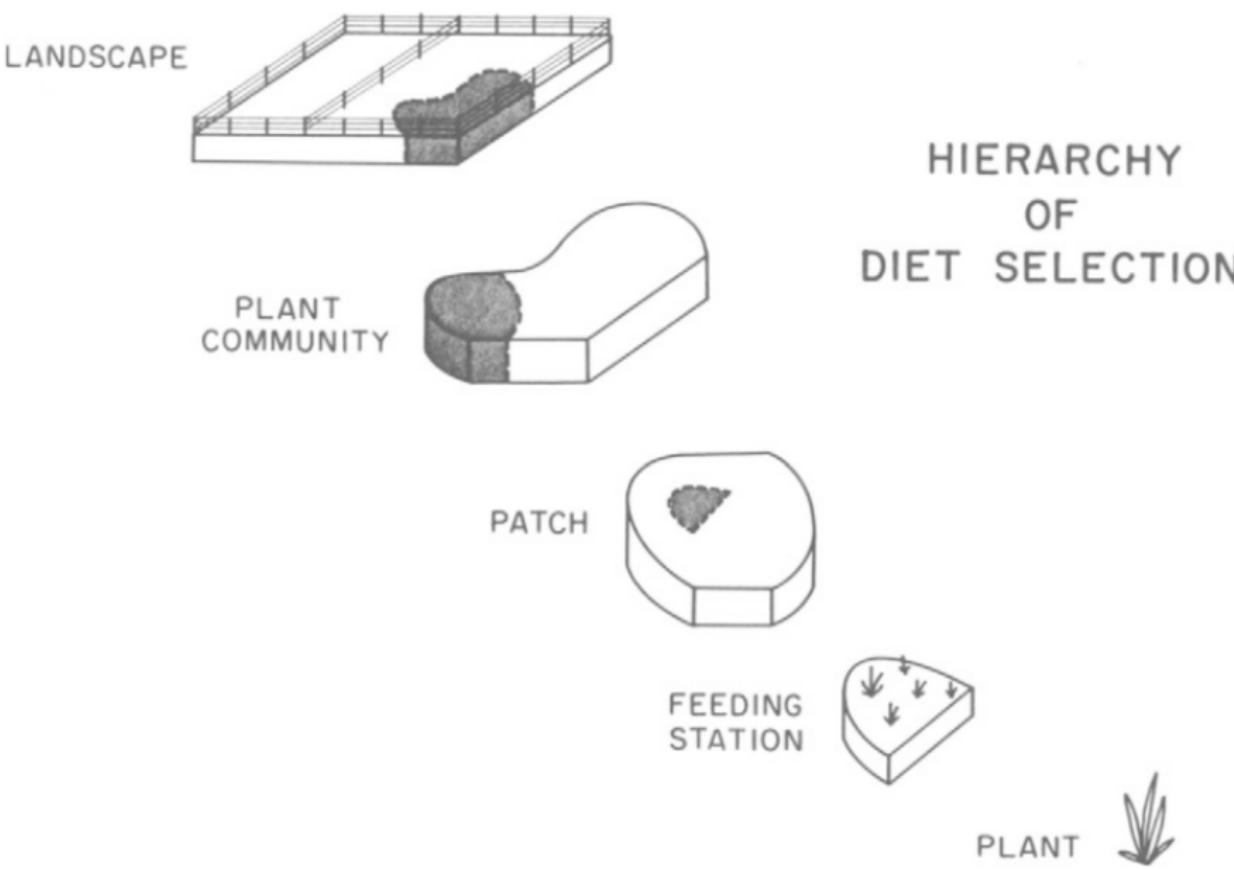
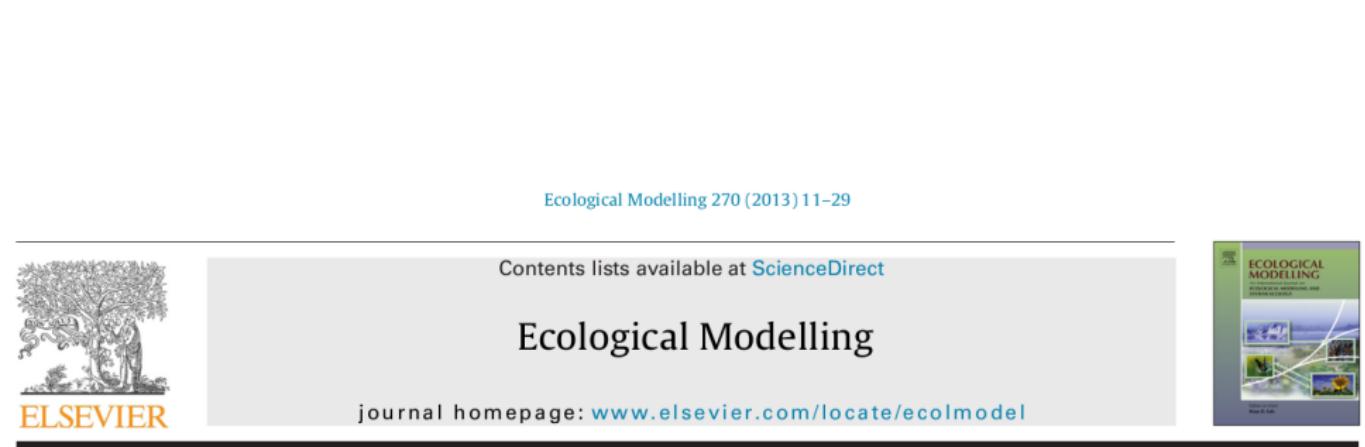


Fig. 4-2. The hierarchical context of diet selection as it descends from the landscape to the individual plant.



A model of diurnal grazing patterns and herbage intake of a dairy cow,
MINDY: Model description

Pablo Gregorini^{a,*}, Pierre C. Beukes^a, Alvaro J. Romera^a, Gil Levy^a, Mark D. Hanigan^b

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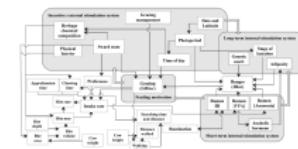


Fig. 1. Schematic representation of MINDY: a mechanistic and dynamic model to simulate diurnal patterns of herbage intake and grazing behavior of a grazing dairy cow. White boxes with solid lines represent true pools (hard components) of the model, white dashed with dashed lines represent soft components of the model, solid arrows represent modifiers. Grey boxes (functional components) and arrows represent the motivational system of feeding behavior adapted from Jensen and Toates (1993), Hughes and Duncan (1988) and Smith (1996).

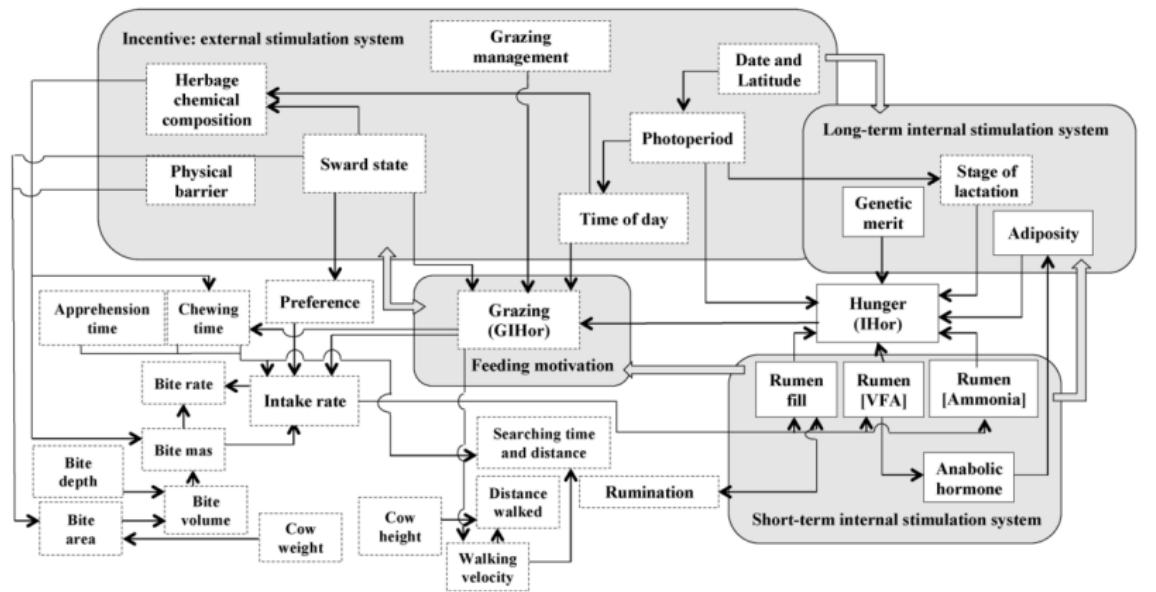


Fig. 1. Schematic representation of MINDY: a mechanistic and dynamic model to simulate diurnal patterns of herbage intake and grazing behavior of a grazing dairy cow. White boxes with solid lines represent true pools (hard components) of the model, white dashed with dashed lines represent soft components of the model, solid arrows represent modifiers. Grey boxes (functional components) and arrows represent the motivational system of feeding behavior adapted from Jensen and Toates (1993), Hughes and Duncan (1988) and Smith (1996).

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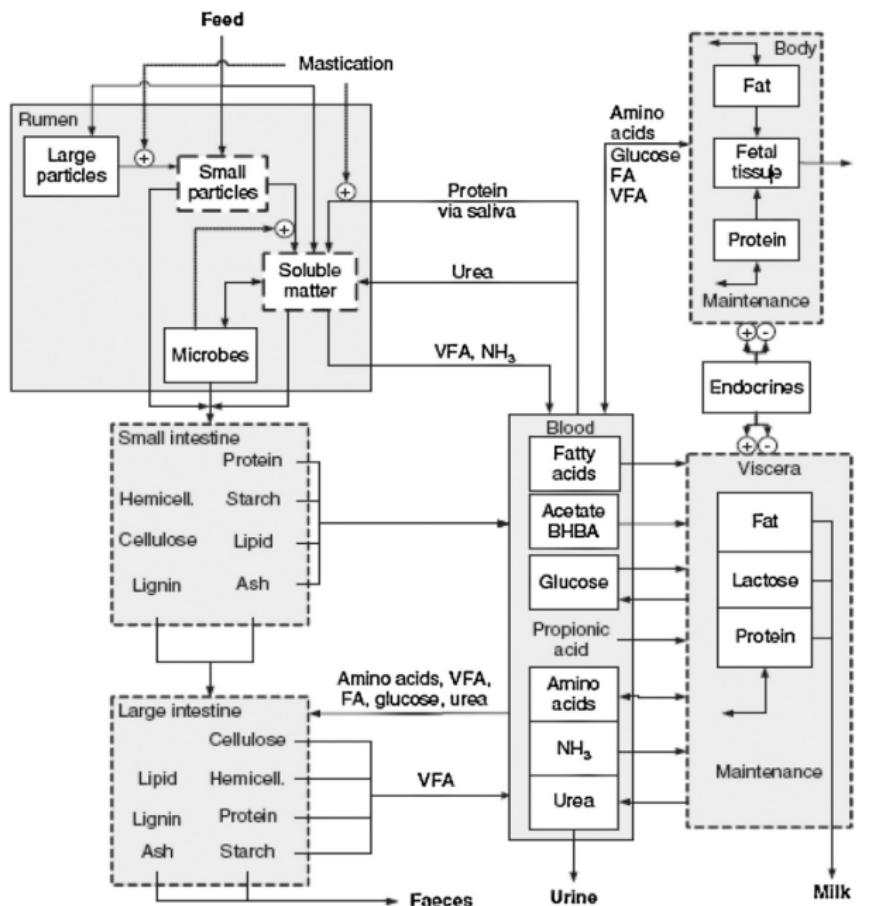
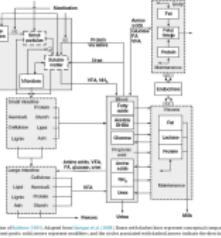
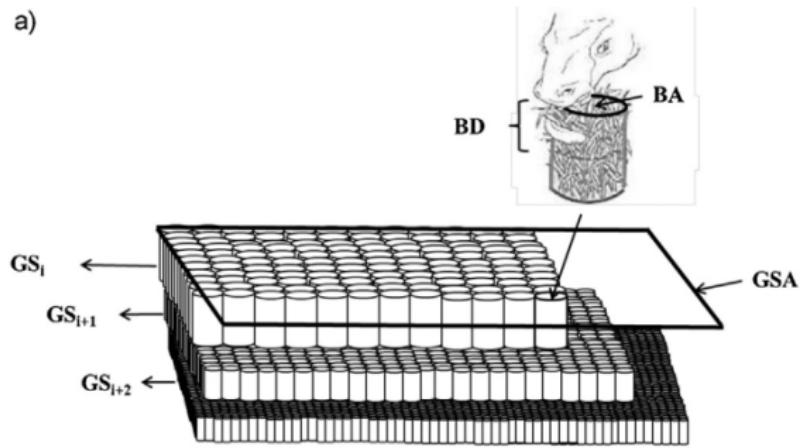


Fig. 2. A simplified schematic representation of Baldwin (1995). Adapted from Hanigan et al. (2008). Boxes with dashes lines represent conceptual compartments as defined in the model; boxes with solid lines represent pools; solid arrows represent modifiers; and the circles associated with dashed arrows indicate the direction of the modifiers. FA, fatty acids; VFA, volatile fatty acids.

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a)



b)

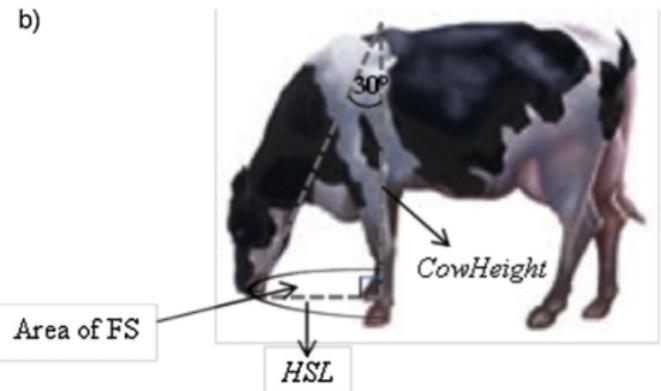
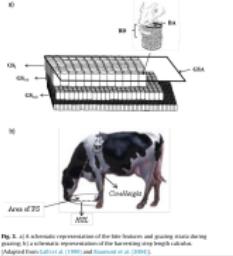


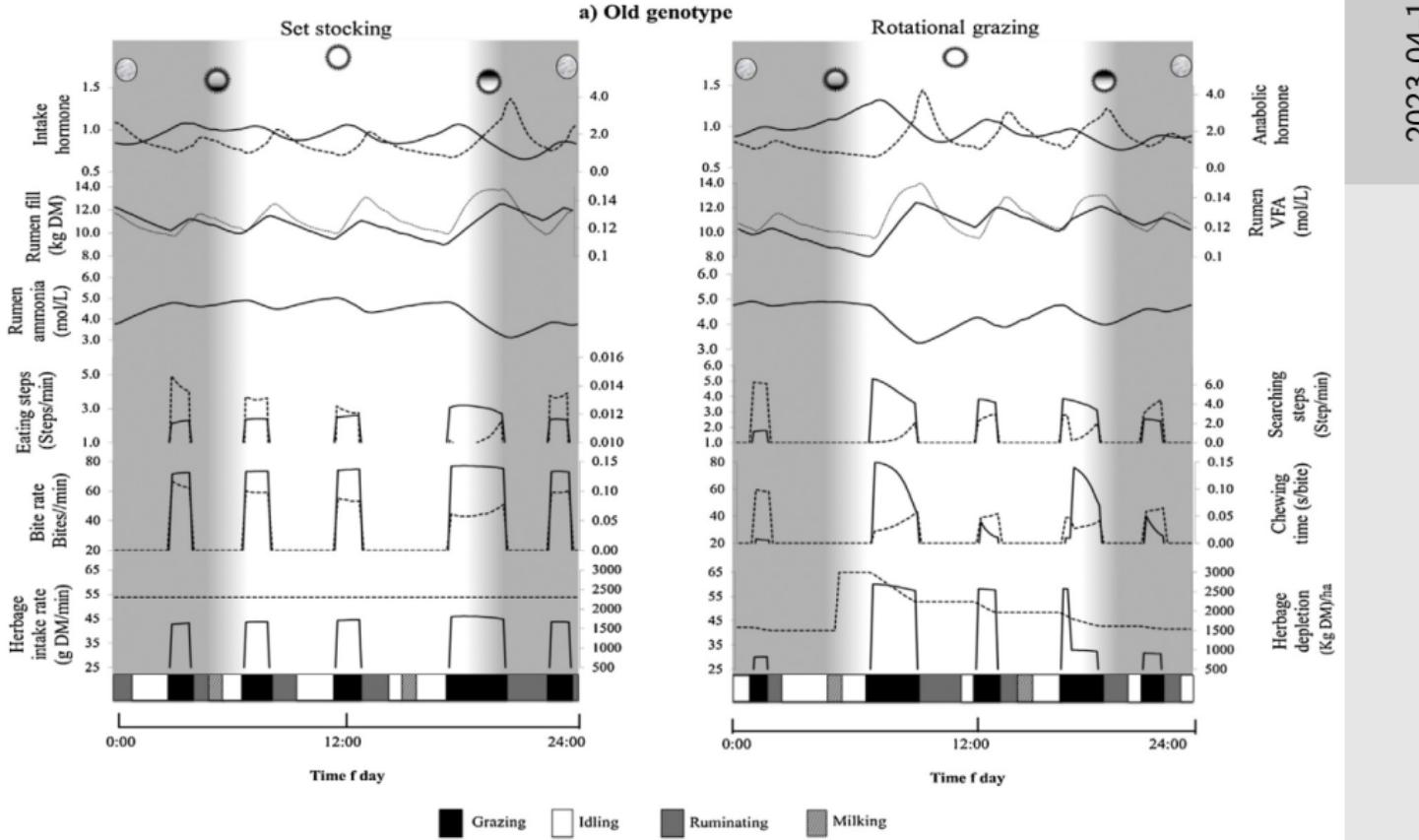
Fig. 3. a) A schematic representation of the bite features and grazing strata during grazing; b) a schematic representation of the harvesting step length calculus.
(Adapted from Galli et al. (1999) and Baumont et al. (2004)).

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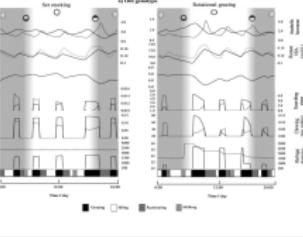




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| Symbol | Definition | Value/Unit |
|------------------------------------|--|---------------------|
| <i>ACHT</i> | Actual chewing time | Days |
| <i>ActHrR</i> | Actual herbage intake rate of the grazing stratum <i>i</i> | kg/min |
| <i>AHM</i> | Available herbage mass modulator | Unitless |
| <i>Ahor</i> | Anabolic hormone | Unitless |
| <i>Am</i> | Ruminal ammonia concentration | mmol/L |
| <i>AmCor</i> | Ruminal ammonia correction factor | Unitless |
| <i>BA</i> | Bite area of the grazing stratum <i>i</i> | m ² |
| <i>BCS</i> | Body condition score | PointS |
| <i>BCS_{target}</i> | Body condition score target | PointS |
| <i>BD</i> | Bite depth of the grazing stratum <i>i</i> | mm |
| <i>BM_i</i> | Bite mass of the grazing stratum <i>i</i> | kg |
| <i>BR</i> | Bite rate | Bites/day |
| <i>Chewingfactor</i> | Motivation to chew | Unitless |
| <i>CowHeight</i> | Animals height to the shoulder | m |
| <i>CR</i> | Consumption rate area of grazing stratum <i>i</i> | m ² /day |
| <i>CurrentStratum</i> | Upper stratum from the pair strata currently being grazed | mmol/L |
| <i>CurrentStratum₋₁</i> | Lower stratum from the pair strata currently being grazed | mmol/L |
| <i>C_{VFA}</i> | Ruminal concentration of volatile fatty acids | mmol/L |
| <i>DA</i> | Dental arcade | m |
| <i>Dailydistancewalked</i> | Daily distance walked | m |
| <i>DaylengthP1</i> | Daylength excluding twilight hours for lactation module | Unitless |
| <i>Daylight</i> | Value representing light intensity | Days |
| <i>DayTwelvethP1</i> | Length of the day including twilight hours | Days |
| <i>DayTwelvethP2</i> | Daylength including twilight hours | Proportion |
| <i>DD</i> | Depth of soil depth | m |
| <i>DWH</i> | Distance walked while harvesting | m |
| <i>eDMI</i> | Shape factor | Unitless |
| <i>ETH_i</i> | Extended tiller height of the grazing stratum <i>i</i> | m |
| <i>ETH_{hi}</i> | Initial extended tiller height | m |
| <i>ETH_{hi}in</i> | Physical barrier under what cows are not allowed to or are not capable to graze | m |
| <i>F_{adjustment}</i> | Adjustment factor to the herbage chemical composition | Unitless |
| <i>FdRat</i> | Herbage intake rate | kg/day |
| <i>FSR</i> | Number of feeding stations per unit of time | FS/day |
| <i>GACurrentStratum</i> | Area harvested at the upper grazing stratum from the pair of grazing strata being grazed at the time | m ² |
| <i>GACurrentStratum - 1</i> | Area harvested at the lower grazing stratum from the pair of grazing strata being grazed at the time | m ² |
| <i>GA_i</i> | Rates of changes in CSA due to herbage consumption in grazing stratum <i>i</i> | m ² /day |
| <i>GBhor</i> | Motivation to graze | Unitless |
| <i>GrazingSw</i> | Switch to turn on and off grazing | Unitless |
| <i>HDMI</i> | Herbage dry matter intake | kg/day |
| <i>HGR</i> | Herbage growth rate | m/day |
| <i>H</i> | Median point height of each grazing stratum <i>i</i> | m |
| <i>HighChewingMot</i> | Consume more | 1, unitless, 31 |
| <i>HM</i> | Herbage mass | kg/m ² |
| <i>HM_{available}</i> | Sum of the herbage mass remaining in each grazing stratum | kg |
| <i>HM_{unavail}</i> | Unavailable herbage mass | kg |
| <i>HSA</i> | Length of a step while harvesting | m |
| <i>HM_{pre}</i> | Pre-grazing herbage mass | kg |
| <i>HM_{post}</i> | Post-grazing available herbage mass | kg |
| <i>IHor</i> | Hunger hormone | Unitless |
| <i>IHorCor</i> | Scalar | 1.0, unitless |
| <i>IHorDeg</i> | Intake hormone degradation | Unitless |
| <i>IHorRange</i> | Constant, Range of <i>IHor</i> | 0.02, unitless |
| <i>IHorSyn</i> | Intake hormone synthesis | Unitless |
| <i>iHor</i> | Initial <i>iHor</i> | 1.0, unitless |
| <i>k_{Adt}</i> | Scalar | 1.0, unitless |
| <i>k_{Am}</i> | Scalar | 0.75, unitless |
| <i>k_{Chewfactor}</i> | Scalar | Unitless |
| <i>k_{DMI}</i> | Function adjusting for adiposity and genetic potential | Unitless |
| <i>k_{IFor}</i> | Constant | 0.001, unitless |
| <i>k_{IFor}</i> | Scalar (it scales <i>FdRat</i> with genetic potential) | 0.1, unitless |
| <i>k_{MassCells}</i> | Rate of particle breakdown while ruminating | Unitless |
| <i>k_{RP}</i> | Scalar to correct PIR(<i>j</i>) - 1)*PIR(<i>i</i>) to achieve a proper curve shape | Unitless |
| <i>k_{RPf}</i> | Scalar | 8.5, unitless |
| <i>k_{RPst}</i> | Scalar | 0.11, unitless |
| <i>k_{RS}</i> | Scalar | Unitless |
| <i>LogDMI</i> | Intake lag function | Unitless |
| <i>LateFeeding</i> | Adjusting variable to reduce <i>MinDHr</i> | Days |
| <i>LM</i> | Linear masses of lamina | kg/m |
| <i>LM_i</i> | Linear mass index of each grazing stratum <i>i</i> | Unitless |
| <i>LowChewingMot</i> | Constant | 0.15, unitless |
| <i>LP</i> | Large particle size pool in the rumen | kg |
| <i>MamCellPart</i> | Number of milk secrete cells in the udder | Unitless |
| <i>MBD_i</i> | Mean bulk density of the grazing stratum <i>i</i> | kg/m ³ |
| <i>MBD_{sword}</i> | Mean bulk density of the sword | kg/m ³ |
| <i>MeanLM</i> | Mean linear mass of the tiller | kg/m |
| <i>MeanSWheight</i> | Half of the ETH _{hi} | m |

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Table A.1 (Continued)

| Symbol | Definition | Value/Unit |
|----------------------------------|--|---------------------|
| minimunGSA | Area threshold at which hrazing strating has '0' preference | m ² |
| MinLPRumntn | Minimum LP size required to initiate a ruination bout | kg |
| MSH | Momentary speed of harvesting | m/d |
| NightMealInter | Length of the last meal of the day | 0.1, unitless |
| NightMealTime | Interval of the last meal of the day and the next meal during the night | 0.044, unitless |
| Nstrata | Number of sward canopy accessible grazing strata | |
| Nutrient _{adjustment} | Adjustment factor to the herbage nutrients | Unitless |
| PCHT | Potential chewing time | Days |
| PIR _{Current stratum} | Potential intake rate in the upper stratum from the pair strata currently being grazed | kg/day |
| PIR _{Current stratum+1} | Potential intake rate in the lower stratum from the pair strata currently being grazed | kg/day |
| PIR _i | Potential herbage dry matter intake rate of the grazing stratum <i>i</i> | kg/day |
| PREF _{CurrentStratum} | Partial preference for the upper stratum from the pair strata currently being grazed | Unitless |
| PREF _{inter} | Constant to affect the intercept of the curve of partial preference for current currently being grazed | Unitless |
| pSTI | Momentary average proportion of time searching | Unitless |
| PT | Prehension time | Days |
| Rest | Resting (idling) time | Days |
| RumDM | Ruminal dry matter load | kg DM |
| Rumntn | Rumination time | Days |
| SDI | Distance walked while searching | m |
| SGR | Sward growth rate | m ² /d |
| SM | Linear masses of sheath | kg/m |
| SSpeedS | Momentary average speed while searching | m/day |
| SSR | Searching step rate | Searching steps/day |
| StartlHor | Constant, trigger point for starting a grazing bout | Unitless |
| STI | Searching time | Days |
| StoplHor | Constant, trigger point for ending a meal | Unitless |
| T | Time of day | Days |
| TA | Total area offered | m ² |
| TB _i | Time per bite at the grazing stratum <i>i</i> | Days |
| Vm _{lHorSyn} | Maximum velocity of <i>lHor</i> synthesis | Unitless |
| xaHor | Scalar | 1.0, unitless |
| xAm | Scalar | 1.12, unitless |
| xFdRatLag | Scalar (it rescales Roseler et al. (1997) lag function) | 0.25, unitless |
| xRumDM | Scalar | 1.0, unitless |
| xVFA | Scalar | 10.0, unitless |

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| Table A.1 (Continued) | |
|-----------------------|-------------|
| Variables | Definitions |



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Bioresource Technology 87 (2003) 113–124

Mathematical modeling of non-ideal mixing continuous flow reactors for anaerobic digestion of cattle manure

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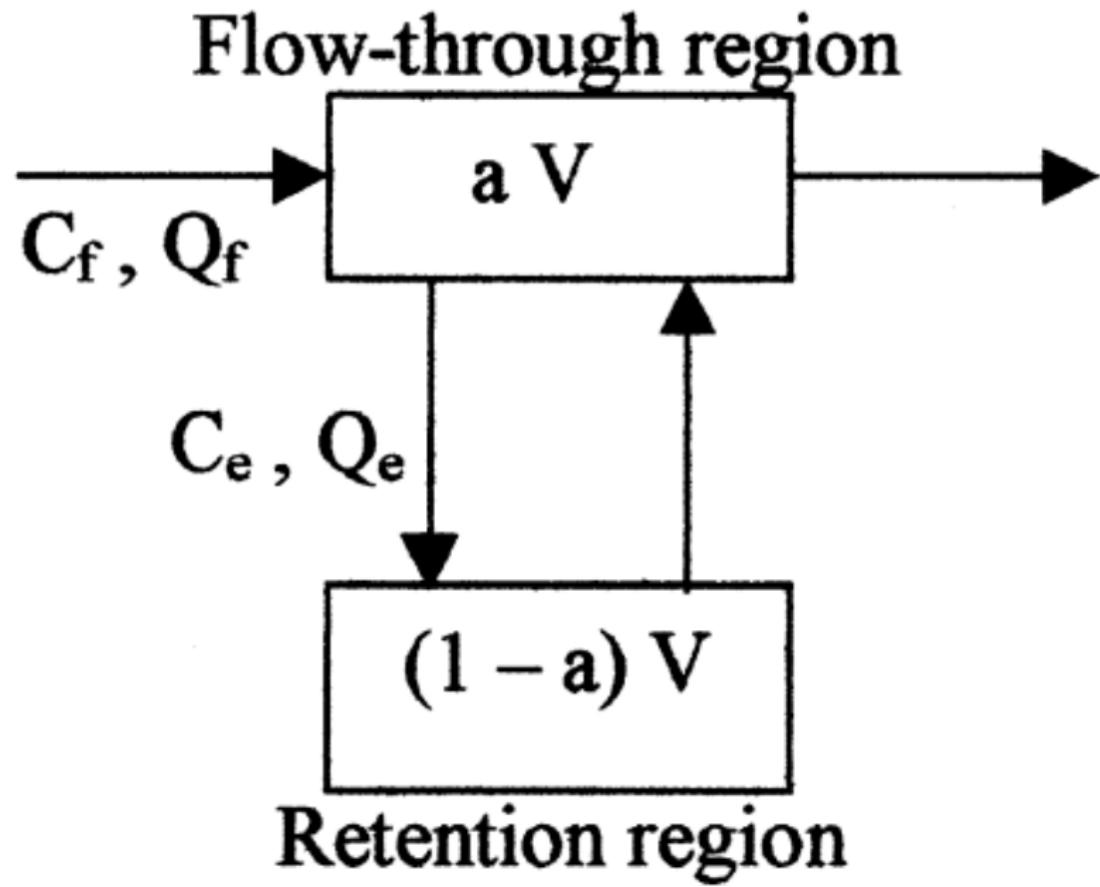
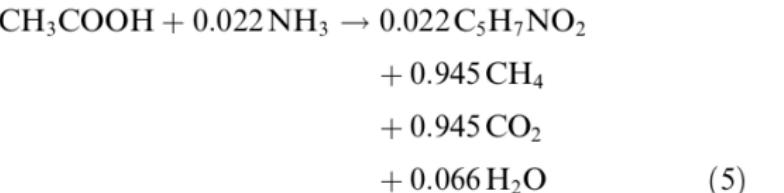
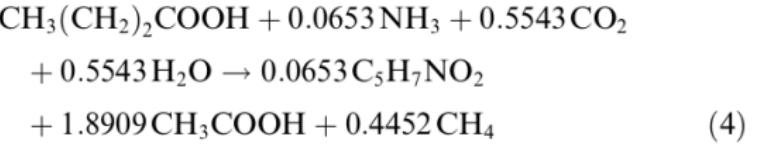
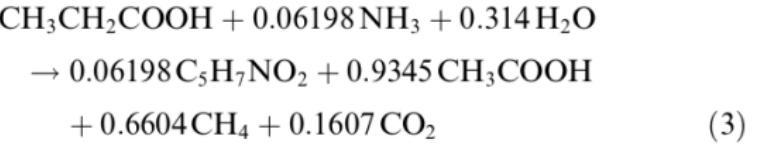
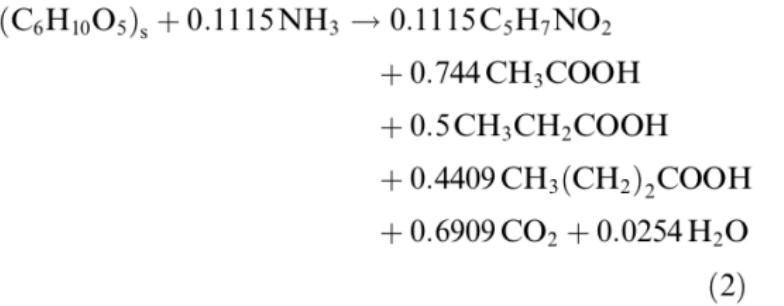
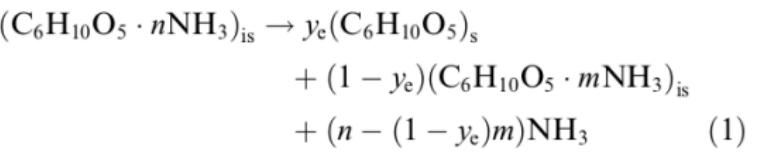


Fig. 1. Two-region mixing model.





Nomenclature

| | | | |
|-----------|--|-------------|--|
| a | mixing parameter | R | gas constant, atm l/mol K |
| b | mixing parameter | t | time, d |
| C | liquid concentration, g/l | T | temperature, K |
| $[CO_2]$ | free CO_2 in liquid concentration, mol/l | V_g | gas volume of reactor, l |
| f | individual bacterial fraction in initial total biomass | V_l | liquid volume of reactor, l |
| f_{pr} | mass conversion factor of propionate to acetate = 0.8108 | VFA | volatile fatty acids |
| f_{but} | mass conversion factor of butyrate to acetate = 0.6818 | X | microorganisms concentration, g/l |
| F_t | biogas transfer rate, mol/d | y_c | yield factor used in Eq. (10) |
| $F(pH)$ | pH function | α | flow-through region |
| H | Henry's constant, atm l/mol | β | retention region |
| HRT | hydraulic retention time | θ | HRT, d |
| SRT | sludge retention time | μ | specific growth rate, d^{-1} |
| k | hydrolysis rate constant, d^{-1} | μ_{max} | maximum specific growth rate, d^{-1} |
| K_0 | non-inhibited hydrolysis rate constant, d^{-1} | | |
| K_a | dissociation constant | | |
| k_d | bacterial decay rate constant, d^{-1} | | |
| K_i | inhibition constant, g/l | | |
| K_s | Monod saturation constant, g/l | | |
| m | feed constant used in Eq. (1) | | |
| n | feed constant used in Eq. (1) | | |
| N | gas transfer rate, g/d | | |
| $[NH_3]$ | free NH_3 in liquid concentration, mol/l | | |
| P | pressure, atm | | |
| pK_h | constant used in Eq. (16) | | |
| pK_l | constant used in Eq. (16) | | |
| Q | volumetric flow rate, d^{-1} | | |
| r_d | bacterial decay rate, g/l d | | |
| r_h | hydrolysis reaction rate, g/l d | | |
| r_s | substrate consumption rate, g/l d | | |
| r_x | bacterial growth rate, g/l d | | |

Subscripts

| | |
|-----|--|
| ac | acetate |
| am | ammonia |
| A | acidogenic bacteria |
| AB | butyric degrading acetogenic bacteria |
| AP | propionate degrading acetogenic bacteria |
| but | butyrate |
| c | carbon dioxide |
| e | exchange between zones |
| f | feed |
| i | component i |
| is | insoluble substrate |
| m | methane |
| M | methanogenic bacteria |
| pr | propionate |
| s | soluble substrate |
| t | total |
| w | water |

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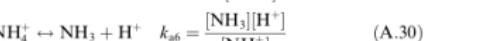
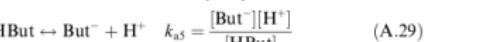
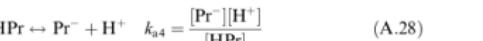
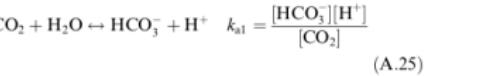
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| Nomenclature | |
|--------------|--|
| x | nitrate production, g/d |
| y | nitrate production, g/d |
| Z_{CO_2} | loss CO ₂ in liquid concentration, mol/l |
| Z_{M} | total bacterial fraction in mixed total biomass |
| z_{pr} | mass conversion factor of propionate to acetate = 0.8108 |
| z_{but} | mass conversion factor of butyrate to acetate = 0.6818 |
| z_{prt} | propionate transfer rate, mol/d |
| pH | pH of reactor, atm/mol |
| qH | reactor's retention, atm/mol |
| HRT | hydraulic retention time |
| SRT | sludge retention time |
| k_h | hydrolysis rate constant, d^{-1} |
| k_d | bacterial decay rate constant, d^{-1} |
| k_{dis} | dissolution rate constant, d^{-1} |
| K_s | saturation constant of hydrolysis reaction, g/l |
| pH_s | saturation pH of hydrolysis reaction, g/l |
| μ | bacterial growth rate, d^{-1} |
| μ_{max} | maximum bacterial growth rate, d^{-1} |
| pCO_2 | gas transfer rate, mol/d |
| pH_c | pressure, atm, used in Eq. (10) |
| pH_s | constant used in Eq. (10) |
| D | constant used in Eq. (10) |
| r_d | bacterial decay rate, g/l d |
| r_h | hydrolysis reaction rate, g/l d |
| r_s | substrate consumption rate, g/l d |
| r_x | bacterial growth rate, g/l d |
| x | gas constant, atm.l/mol.K |
| y | nitrate, g |
| Z | nitrate production, g/d |
| Z_{CO_2} | gas volume of reactor, l |
| Z_M | total bacterial fraction in mixed total biomass |
| z_{buty} | volatile fatty acids |
| z_{prop} | propionate concentration, g/l |
| z_{acet} | acetate concentration, g/l |
| z_{meth} | methane concentration, g/l |
| z_{metho} | methanogenic bacteria |
| z_{acid} | acidic bacteria |
| z_{buty} | butyric degrading acetogenic bacteria |
| z_{prop} | propionate degrading acetogenic bacteria |
| z_{acet} | acetate degrading acetogenic bacteria |
| z_{meth} | methane |
| z_{metho} | methanogenic bacteria |
| z_{acid} | acidic bacteria |
| z_{buty} | butyric degrading acetogenic bacteria |
| z_{prop} | propionate degrading acetogenic bacteria |
| z_{acet} | acetate degrading acetogenic bacteria |
| z_{meth} | methane |

A.4. Liquid phase equilibrium chemistry

Ionic dissociation equations



Ionic balance equations for both α and β liquid phases

$$\begin{aligned} [\text{H}^+] + [\text{NH}_4^+] &= [\text{OH}^-] + [\text{HCO}_3^-] + 2[\text{CO}_3^{2-}] \\ &\quad + [\text{Ac}^-] + [\text{Pr}^-] + [\text{But}^-] + [\text{A}^- \text{C}^+] \end{aligned} \quad (\text{A.32})$$

where

$$[\text{NH}_4^+] = \frac{C_{\text{am}}/17}{1 + k_{a6}/[\text{H}^+]} \quad (\text{A.33})$$

$$[\text{OH}^-] = k_w/[\text{H}^+] \quad (\text{A.34})$$

$$[\text{HCO}_3^-] = \frac{C_c/44}{1 + [\text{H}^+]/k_{a1} + k_{a2}/[\text{H}^+]} \quad (\text{A.35})$$

$$[\text{CO}_3^{2-}] = \frac{C_c/44}{1 + [\text{H}^+]/k_{a2} + [\text{H}^+]^2/k_{a1}k_{a2}} \quad (\text{A.36})$$

$$[\text{Ac}^-] = \frac{C_{\text{ac}}/60}{1 + [\text{H}^+]/k_{a3}} \quad (\text{A.37})$$

$$[\text{Pr}^-] = \frac{C_{\text{pr}}/74}{1 + [\text{H}^+]/k_{a4}} \quad (\text{A.38})$$

$$[\text{But}^-] = \frac{C_{\text{but}}/88}{1 + [\text{H}^+]/k_{a5}} \quad (\text{A.39})$$

Modelling livestock

- └ Models for cattle
- └ Models for herds

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| | |
|---|--|
| $\text{CH}_3 + \text{H}_2\text{O} \rightarrow \text{H}_2\text{O}_2 + \text{H}$ | $k_1 = \frac{[\text{H}_2\text{O}_2][\text{H}]}{[\text{CH}_3]} \quad (\text{A.36})$ |
| $\text{H}_2\text{O}_2 \rightarrow \text{O} + \text{H}_2\text{O}$ | $k_2 = \frac{[\text{O}][\text{H}_2\text{O}]}{[\text{H}_2\text{O}_2]} \quad (\text{A.37})$ |
| $\text{O} + \text{H} \rightarrow \text{OH}$ | $k_3 = \frac{[\text{OH}]}{[\text{O}][\text{H}]} \quad (\text{A.38})$ |
| $\text{OH} + \text{H} \rightarrow \text{H}_2\text{O}$ | $k_4 = \frac{[\text{H}_2\text{O}]}{[\text{OH}][\text{H}]} \quad (\text{A.39})$ |
| $\text{NO}_2 + \text{H}_2\text{O} \rightarrow \text{HNO}_3 + \text{HNO}$ | $k_5 = \frac{[\text{HNO}_3][\text{HNO}]}{[\text{NO}_2][\text{H}_2\text{O}]} \quad (\text{A.40})$ |
| $\text{HNO}_3 + \text{H} \rightarrow \text{NO} + \text{H}_2\text{O}$ | $k_6 = \frac{[\text{NO}]}{[\text{HNO}_3][\text{H}]} \quad (\text{A.41})$ |
| $\text{NO} + \text{H} \rightarrow \text{NO}_2 + \text{H}_2\text{O}$ | $k_7 = \frac{[\text{NO}_2]}{[\text{NO}][\text{H}]} \quad (\text{A.42})$ |
| $\text{NO}_2 + \text{NO} \rightarrow \text{N}_2\text{O}_4$ | $k_8 = \frac{[\text{N}_2\text{O}_4]}{[\text{NO}_2][\text{NO}]} \quad (\text{A.43})$ |
| $\text{N}_2\text{O}_4 + \text{H}_2\text{O} \rightarrow \text{HNO}_3 + \text{HNO}_2$ | $k_9 = \frac{[\text{HNO}_3][\text{HNO}_2]}{[\text{N}_2\text{O}_4][\text{H}_2\text{O}]} \quad (\text{A.44})$ |
| $\text{HNO}_2 + \text{H} \rightarrow \text{NO} + \text{H}_2\text{O}$ | $k_{10} = \frac{[\text{NO}]}{[\text{HNO}_2][\text{H}]} \quad (\text{A.45})$ |
| $\text{NO} + \text{H} \rightarrow \text{NO}_2 + \text{H}_2\text{O}$ | $k_{11} = \frac{[\text{NO}_2]}{[\text{NO}][\text{H}]} \quad (\text{A.46})$ |
| $\text{NO}_2 + \text{NO}_2 \rightarrow \text{N}_2\text{O}_4$ | $k_{12} = \frac{[\text{N}_2\text{O}_4]}{[\text{NO}_2]^2} \quad (\text{A.47})$ |
| $\text{N}_2\text{O}_4 + \text{H}_2\text{O} \rightarrow \text{HNO}_3 + \text{HNO}_2$ | $k_{13} = \frac{[\text{HNO}_3][\text{HNO}_2]}{[\text{N}_2\text{O}_4][\text{H}_2\text{O}]} \quad (\text{A.48})$ |
| $\text{HNO}_2 + \text{H} \rightarrow \text{NO} + \text{H}_2\text{O}$ | $k_{14} = \frac{[\text{NO}]}{[\text{HNO}_2][\text{H}]} \quad (\text{A.49})$ |
| $\text{NO} + \text{H} \rightarrow \text{NO}_2 + \text{H}_2\text{O}$ | $k_{15} = \frac{[\text{NO}_2]}{[\text{NO}][\text{H}]} \quad (\text{A.50})$ |

A.1. Liquid phase

Microbial biomass, X_i , $i = A, AP, AB, M$

$$\frac{dX_i^a}{dt} = \frac{X_{if} - X_i^a}{a\theta} + \frac{X_i^\beta - X_i^a}{a\theta/b} + (\mu_i^a - b_i)X_i^a \quad (A.1)$$

$$\frac{dX_i^\beta}{dt} = \frac{X_i^a - X_i^\beta}{(1-a)\theta/b} + (\mu_i^\beta - b_i)X_i^\beta \quad (A.2)$$

Insoluble substrate, C_{is}

$$\frac{dC_{is}^a}{dt} = \frac{C_{if} - C_{is}^a}{a\theta} + \frac{C_a^{\beta} - C_{is}^a}{a\theta/b} - k^a C_{is}^a \quad (A.3)$$

$$\frac{dC_{is}^{\beta}}{dt} = \frac{C_{is}^a - C_{is}^{\beta}}{(1-a)\theta/b} - k^{\beta} C_{is}^{\beta} \quad (A.4)$$

Soluble substrate, C_s

$$\begin{aligned} \frac{dC_s^a}{dt} &= \frac{C_{if} - C_s^a}{a\theta} + \frac{C_s^{\beta} - C_s^a}{a\theta/b} + \frac{162y_e}{162 + 17n} k^a C_{is}^a \\ &\quad - 12.858\mu_A^a X_A^a \end{aligned} \quad (A.5)$$

$$\begin{aligned} \frac{dC_s^{\beta}}{dt} &= \frac{C_s^a - C_s^{\beta}}{(1-a)\theta/b} + \frac{162y_e}{162 + 17n} k^{\beta} C_{is}^{\beta} \\ &\quad - 12.858\mu_A^{\beta} X_A^{\beta} \end{aligned} \quad (A.6)$$

Total acetate, C_{ac}

$$\begin{aligned} \frac{dC_{ac}^a}{dt} &= \frac{C_{acf} - C_{ac}^a}{a\theta} + \frac{C_{ac}^{\beta} - C_{ac}^a}{a\theta/b} + 3.54\mu_A^a X_A^a \\ &\quad + 8.006\mu_{AP}^a Y_{AP}^a + 15.366\mu_{AB}^a X_{AB}^a \\ &\quad - 24.135\mu_M^a Y_M^a \end{aligned} \quad (A.7)$$

$$\begin{aligned} \frac{dC_{ac}^{\beta}}{dt} &= \frac{C_{ac}^a - C_{ac}^{\beta}}{(1-a)\theta/b} + 3.54\mu_A^{\beta} Y_A^{\beta} + 8.006\mu_{AP}^{\beta} Y_{AP}^{\beta} \\ &\quad + 15.366\mu_{AB}^{\beta} X_{AB}^{\beta} - 24.135\mu_M^{\beta} Y_M^{\beta} \end{aligned} \quad (A.8)$$

Total propionate, C_{pr}

$$\begin{aligned} \frac{dC_{pr}^a}{dt} &= \frac{C_{ptf} - C_{pr}^a}{a\theta} + \frac{C_{pr}^{\beta} - C_{pr}^a}{a\theta/b} + 2.937\mu_A^a X_A^a \\ &\quad - 10.566\mu_{AP}^a X_{AP}^{\beta} \end{aligned} \quad (A.9)$$

$$\begin{aligned} \frac{dC_{pr}^{\beta}}{dt} &= \frac{C_{pr}^a - C_{pr}^{\beta}}{(1-a)\theta/b} + 2.937\mu_A^{\beta} X_A^{\beta} - 10.566\mu_{AP}^{\beta} X_{AP}^{\beta} \\ &\quad - 11.919\mu_{BP}^{\beta} X_{AB}^a \end{aligned} \quad (A.10)$$

Total butyrate, C_{but}

$$\begin{aligned} \frac{dC_{but}^a}{dt} &= \frac{C_{butf} - C_{but}^a}{a\theta} + \frac{C_{but}^{\beta} - C_{but}^a}{a\theta/b} + 3.079\mu_A^a X_A^a \\ &\quad - 11.919\mu_{BP}^a X_{AB}^a \end{aligned} \quad (A.11)$$

$$\begin{aligned} \frac{dC_{but}^{\beta}}{dt} &= \frac{C_{but}^a - C_{but}^{\beta}}{(1-a)\theta/b} + 3.079\mu_A^{\beta} X_A^{\beta} - 11.919\mu_{BP}^{\beta} X_{AB}^{\beta} \end{aligned} \quad (A.12)$$

Modelling livestock

- └ Models for cattle
- └ Models for herds

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| | |
|--|-------|
| Δ _s Lactate | 1.00 |
| Δ _s Acetate | -0.01 |
| Δ _s Propionate | -0.02 |
| Δ _s Butyrate | -0.03 |
| Δ _s NH_3N | -0.04 |
| Δ _s Ammonium | -0.05 |
| Δ _s NH_4^+ | -0.06 |
| Δ _s NH_3 | -0.07 |
| Δ _s NH_3N | -0.08 |
| Δ _s NH_4N | -0.09 |
| Δ _s NH_4NH_3 | -0.10 |
| Δ _s NH_3NH_4 | -0.11 |
| Δ _s NH_3NH_3 | -0.12 |
| Δ _s $\text{NH}_3\text{NH}_3\text{N}$ | -0.13 |
| Δ _s $\text{NH}_3\text{NH}_4\text{N}$ | -0.14 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_4$ | -0.15 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3$ | -0.16 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{N}$ | -0.17 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_4\text{N}$ | -0.18 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4$ | -0.19 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3$ | -0.20 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{N}$ | -0.21 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4\text{N}$ | -0.22 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4$ | -0.23 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3$ | -0.24 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{N}$ | -0.25 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4\text{N}$ | -0.26 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4$ | -0.27 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3$ | -0.28 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{N}$ | -0.29 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4\text{N}$ | -0.30 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_4$ | -0.31 |
| Δ _s $\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3\text{NH}_3$ | -0.32 |

$$\begin{aligned}
& \frac{dC_{\text{am}}}{dt} = \frac{C_{\text{am}} - C_{\text{am}}^{\beta}}{a\theta} + \frac{C_{\text{am}}^{\beta} - C_{\text{am}}^{\alpha}}{a\beta} \frac{17(n-m(1-y_e))}{162+17n} k^x C_{\text{u}}^{\alpha} \\
& - 0.15(\mu_A^{\alpha} X_A^{\alpha} + \mu_{\text{AP}}^{\alpha} X_{\text{AP}}^{\alpha} + \mu_{\text{AB}}^{\alpha} X_{\text{AB}}^{\alpha} + \mu_M^{\alpha} X_M^{\alpha}) \\
& \quad (\text{A.13}) \\
& \frac{dC_{\text{am}}^{\beta}}{dt} = \frac{C_{\text{am}}^{\alpha} - C_{\text{am}}^{\beta}}{(1-a)\theta/b} + \frac{17(n-m(1-y_e))}{162+17n} k^{\beta} C_{\text{u}}^{\beta} \\
& - 0.15(\mu_A^{\beta} X_A^{\beta} + \mu_{\text{AP}}^{\beta} X_{\text{AP}}^{\beta} + \mu_{\text{AB}}^{\beta} X_{\text{AB}}^{\beta} + \mu_M^{\beta} X_M^{\beta}) \\
& \quad (\text{A.14})
\end{aligned}$$

Total carbon dioxide in the liquid phase, C_c

$$\begin{aligned}
\frac{dC_c^{\alpha}}{dt} &= \frac{C_{\text{cf}} - C_c^{\alpha}}{a\theta} - \frac{C_c^{\beta} - C_c^{\alpha}}{a\theta/b} + 2.413\mu_A^{\alpha} X_A^{\alpha} \\
& + 1.01\mu_{\text{AP}}^{\alpha} X_{\text{AP}}^{\alpha} - 3.303\mu_{\text{AB}}^{\alpha} X_{\text{AB}}^{\alpha} \\
& + 16.726\mu_M^{\alpha} X_M^{\alpha} - \frac{N_c^{\alpha}}{aV_1} \\
& \quad (\text{A.15})
\end{aligned}$$

$$\begin{aligned}
\frac{dC_c^{\beta}}{dt} &= +\frac{C_c^{\alpha} - C_c^{\beta}}{(1-a)\theta/b} + 2.413\mu_A^{\beta} X_A^{\beta} \\
& + 1.01\mu_{\text{AP}}^{\beta} X_{\text{AP}}^{\beta} - 3.303\mu_{\text{AB}}^{\beta} X_{\text{AB}}^{\beta} \\
& + 16.726\mu_M^{\beta} X_M^{\beta} \\
& \quad (\text{A.16})
\end{aligned}$$

Methane in the liquid phase, C_m

$$\begin{aligned}
\frac{C_m^{\alpha}}{a\theta/b} + 1.509\mu_{\text{AP}}^{\alpha} X_{\text{AP}}^{\alpha} + 0.956\mu_{\text{AB}}^{\alpha} X_{\text{AB}}^{\alpha} \\
+ 6.082\mu_M^{\alpha} X_M^{\alpha} - \frac{N_m^{\alpha}}{aV_1} = 0 \\
& \quad (\text{A.17})
\end{aligned}$$

$$\begin{aligned}
\frac{dC_m^{\beta}}{dt} &= \frac{C_m^{\alpha} - C_m^{\beta}}{(1-a)\theta/b} + 1.509\mu_{\text{AP}}^{\beta} X_{\text{AP}}^{\beta} \\
& + 0.956\mu_{\text{AB}}^{\beta} X_{\text{AB}}^{\beta} + 6.082\mu_M^{\beta} X_M^{\beta} \\
& \quad (\text{A.18})
\end{aligned}$$

where

$$\theta = \frac{V_1}{Q_t} \quad (\text{A.19})$$

$$b = \frac{Q_e}{Q_t} \quad (\text{A.20})$$

A.2. Gas phase

Carbon dioxide in the gas phase, P_c

$$\frac{dP_c}{dt} = \frac{RT}{V_g} \left(\frac{N_c^{\alpha}}{44} - \frac{P_c}{P} F_i \right) \quad (\text{A.21})$$

Methane in the gas phase, P_m

$$\frac{dP_m}{dt} = \frac{RT}{V_g} \left(\frac{N_m^{\alpha}}{16} - \frac{P_m}{P} F_i \right) \quad (\text{A.22})$$

Total material balance in the gas phase, F_i

$$F_i = \frac{P}{P - P_u} \left(\frac{N_m^{\alpha}}{16} + \frac{N_c^{\alpha}}{44} \right) \quad (\text{A.23})$$

Modelling livestock

- └ Models for cattle
- └ Models for herds

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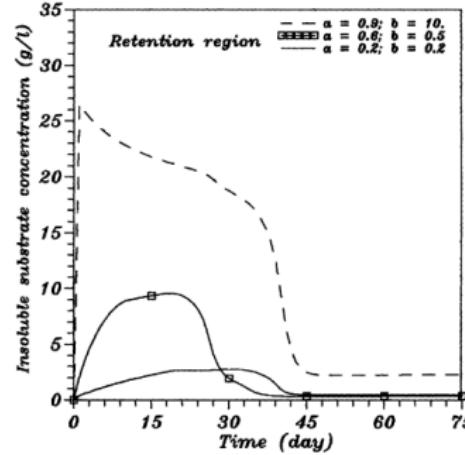
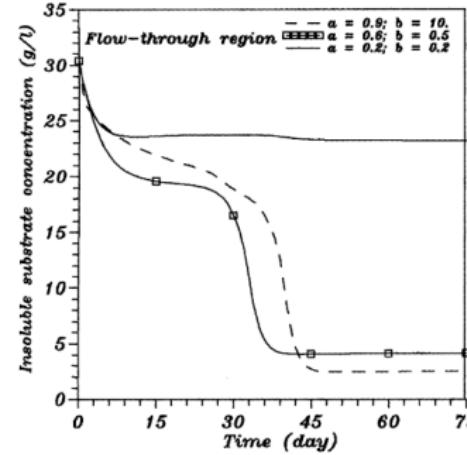


Fig. 2. Dynamic simulation of anaerobic digestion of cattle manure in a continuous flow reactor under HRT=15 days and different degrees of mixing for prediction of the insoluble substrate concentration in flow-through and retention regions.

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- └ Models for cattle
- └ Models for herds

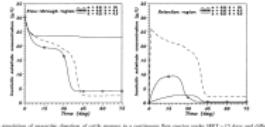


Fig. 2. Dynamic simulation of anaerobic digestion of cattle manure in a continuous flow reactor under HRT=15 days and different degrees of mixing for prediction of the insoluble substrate concentration in flow-through and retention regions.

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[Models for other herds](#)

[Models for management](#)

[Conclusion](#)

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Review article

Review of mathematical models for sow herd management

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Modelling livestock └ Models for other herds



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Review article

Review of mathematical models for sow herd management

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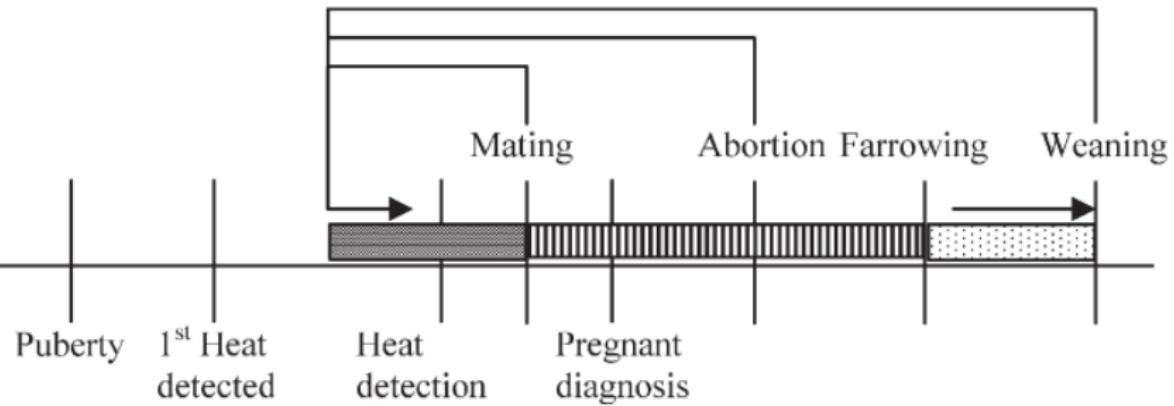
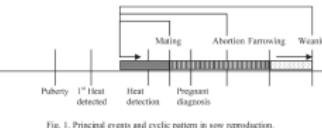


Fig. 1. Principal events and cyclic pattern in sow reproduction.

Main characteristics of sow herd models reviewed

| Authors | Year | Aspects | Model | Title |
|--------------------------|-------|----------|-------|--|
| Allen and Stewart | 1983 | R | S | A simulation model for a swine breeding unit producing feeder pigs |
| Tess et al. | 1983 | R, F, E | S | Simulation of genetic changes in life cycle efficiency of pork production I. A bioeconomic model |
| Dijkhuizen et al. | 1986 | RP, E | OP | Economic optimisation of culling strategies in swine breeding herds, using the "PORKCHOP computer program" |
| Marsh | 1986 | R, E | S | Economic decision making on health and management in livestock herds: examining complex problems through computer simulation |
| Pettigrew et al. | 1986 | R, E | S | Integration of factors affecting sow efficiency: a modelling approach |
| Signh | 1986 | R, E | S | Simulation of swine herd population dynamics |
| de Roo | 1987 | R, G, F | S | A stochastic model to study breeding schemes in a small pig population |
| Pomar et al. | 1991 | R, F | S | Computer simulation model of swine production systems: III. A dynamic herd simulation model including reproduction |
| Jalving et al. | 1992 | R, RP, E | S | Dynamic probabilistic modelling of reproduction and replacement management in sow herds. General aspects and model description |
| Huirne et al. | 1993 | R, RP, E | OP | An application of stochastic dynamic programming to support sow replacement decisions |
| Plà et al. | 1998 | R, RP, E | OP-S | A sow model for decision aid at farm level |
| Plà et al. | 2003 | R, E | S | A Markov decision sow model representing the productive lifespan of herd sows |
| Kristensen and Søllestad | 2004a | R, RP, E | OP | A sow replacement model using Bayesian updating in a three-level hierachic Markov process I. Biological model |
| | 2004b | | | A sow replacement model using Bayesian updating in a three-level hierachic Markov process II. Optimisation model |

R: reproduction, RP: replacement, E: economics, F: feeding, G: genetics, S: simulation, O: optimisation.

Modelling livestock └ Models for other herds

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| Main characteristics of sow herd models reviewed | | | |
|--|-------|----------|--|
| Authors | Year | Aspects | Model |
| Allen and Stewart | 1983 | R, C, E | S |
| Tess et al. | 1983 | R, F, S | Simulation of genetic changes in life cycle efficiency of pork production I. A bioeconomic model |
| Dijkhuizen et al. | 1986 | R, E | "PORKCHOP" computer program |
| Marsch | 1986 | R, E | Economic decision making on health and management in livestock herds: examining complex problems through computer simulation |
| de Roo | 1987 | R, G, F | A stochastic model to study breeding schemes in a small pig population |
| Pettigrew et al. | 1986 | R, E | A mechanistic model to study breeding solutions in a small pig population |
| Signh | 1986 | R, E | Simulation of swine herd population dynamics |
| Jalving et al. | 1992 | R, RP, E | Dynamic probabilistic modelling of reproduction and replacement management in sow herds. General aspects and model description |
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R: reproduction, RP: replacement, E: economics, C: feeding, G: genetics, S: simulation, O: optimisation.

Comparing non-linear mathematical models to describe growth of different animals

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Modelling livestock └ Models for other herds

| Model | Equation |
|-----------------|---|
| Brody | $W(t) = W_{\infty} \left[1 + \left[\left(\frac{W_0}{W_{\infty}} \right)^{\frac{1}{3}} - 1 \right] \exp(-kt) \right]^3 \quad (1)$ |
| von Bertalanffy | $W(t) = W_{\infty} \left[1 + \left[\left(\frac{W_0}{W_{\infty}} \right)^{\frac{1}{3}} - 1 \right] \exp(-kt) \right]^3 \quad (2)$ |
| Logistic | $W(t) = \frac{W_{\infty}}{1 + \left[\left(\frac{W_{\infty}}{W_0} \right)^{\frac{1}{3}} - 1 \right] \exp(-kt)} \quad (3)$ |
| Gompertz | $W(t) = W_{\infty} \exp \left[\ln \left(\frac{W_0}{W_{\infty}} \right) \exp(-kt) \right] \quad (4)$ |
| Richards | $W(t) = \frac{W_{\infty} \cdot W_0}{\left[W_0^m + (W_{\infty}^m - W_0^m) \exp(-kt) \right]^{\frac{1}{m}}} \quad (5)$ for $m \neq 0$ |

Modelling livestock

- Models for other herds

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| Model | Equation |
|-----------------|---|
| Brody | $W(t) = W_{\infty} \left[1 + \left[\left(\frac{W_0}{W_{\infty}} \right)^{\frac{1}{3}} - 1 \right] \exp(-kt) \right]^3 \quad (1)$ |
| von Bertalanffy | $W(t) = W_{\infty} \left[1 + \left[\left(\frac{W_0}{W_{\infty}} \right)^{\frac{1}{3}} - 1 \right] \exp(-kt) \right]^3 \quad (2)$ |
| Logistic | $W(t) = \frac{W_{\infty}}{1 + \left[\left(\frac{W_{\infty}}{W_0} \right)^{\frac{1}{3}} - 1 \right] \exp(-kt)} \quad (3)$ |
| Gompertz | $W(t) = W_{\infty} \exp \left[\ln \left(\frac{W_0}{W_{\infty}} \right) \exp(-kt) \right] \quad (4)$ |
| Richards | $W(t) = \frac{W_{\infty} \cdot W_0}{\left[W_0^m + (W_{\infty}^m - W_0^m) \exp(-kt) \right]^{\frac{1}{m}}} \quad (5)$ for $m \neq 0$ |

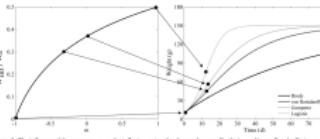


Figure 1. The influence of the parameter m on the inflection point of each growth curve: Brody ($m = -1$), von Bertalanffy ($m = -1/3$), Gompertz ($m = 0$), and Logistic ($m = 1$). The dots indicate the inflection points obtained for each growth model.

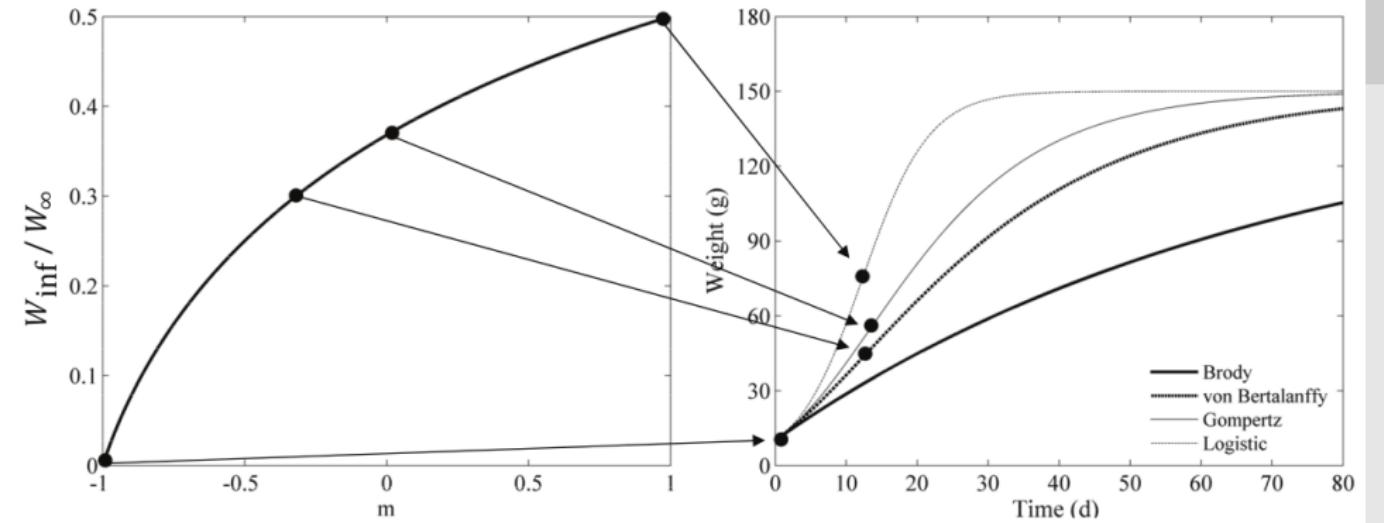


Figure 1. The influence of the parameter m on the inflection point of each growth curve: Brody ($m = -1$), von Bertalanffy ($m = -1/3$), Gompertz ($m = 0$), and Logistic ($m = 1$). The dots indicate the inflection points obtained for each growth model.

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Table 1. Data sets used in this study to evaluate five different growth models.

| Data set | Source |
|---|---|
| Holstein-Friesian bull ^a | Table 1. Means and standard deviations for body weight, growth rates and degree of maturity of Holstein-Friesian bulls from 6 months to 8 years of age. Calo, McDowell, Van Vleck, and Miller, 1973 |
| Nelore cow ^b | Figure 1. A) Estimation of weights based on age of Nelore females, observed and estimated by the models of Brody and von Bertalanffy. Silva, Alencar, Freitas, Packer, and Mourão, 2011 |
| Angus cow ^b | Figure 1. Growth curves of Lines A and K estimated with Brody model. Line least squares means for weight at fixed ages are used as reference for goodness of fit. Beltrán, Butts, Olson, and Koger, 1992 |
| Celta pig ^b (male and female) | Figure 2. Growth curve for males and females of the variety Barcina slaughtered at 14 months. Franco et al., 2011 |
| Karagouniko sheep ^b (male and female) | Figure 1. Growth curve and absolute growth rate for body weight of the Karagouniko male sheep: estimate growth curve; observed mean; estimated absolute growth rate. Goliomitis, Orfanos, Panopoulou, and Rogdakis, 2006 |
| Beetal goat ^a (male and female) | Figure 2. Growth curve and absolute growth rate for body weight of the Karagouniko female sheep: estimate growth curve; observed mean; estimated absolute growth rate. Waheed, Khan, Ali, and Sarwar, 2011 |
| New Zealand rabbit ^a | Table 1. Means (kg) and standard deviations (SD) of growth traits of Beetal goats. Curi, Nunes, and Curi, 1985 |
| Californian rabbit ^a | Table 2. Body weight of Norkfolk rabbit. Table 3. Body weight of Californian rabbit. |
| Norfolk rabbit ^a | Table 4. Body weight of New Zealand rabbit. Aggrey, 2002 |
| Athens-Canadian chicken ^a (male and female) | Table 1. Means and standard deviations for body weight at different ages in Athens-Canadian random-bred chickens. Nahashon, Aggrey, Adefope, Amenyenu, and Wright, 2006 |
| Guinea fowl ^a (male and female) | Table 2. Means and standard for body weight at different ages in a random-bred pearl guinea fowl population. |
| Japanese quail – white line ^a (male and female) | Sezer and Tarhan, 2005 |
| Japanese quail – brown line ^a (male and female) | Table 1. The results of statistical analyses for body weight of Japanese quail lines at different age (means \pm standard errors). |
| Japanese quail – wild line ^a (male and female) | |

^aExperimental data reported in the literature; ^bExperimental data taken from published figures by means of GetData Graph Digitizer 2.24.

| Table 1. Data sets used in this study to evaluate five different growth models. | |
|--|--|
| Brody | Calo, McDowell, Van Vleck, and Miller, 1973 |
| Holstein Friesian bull | Holstein Friesian cattle, number of days of age. |
| Moore cow ^a | Moore, 1961 |
| Angus cow ^b | Angus cattle, 1982 |
| Gilchrist pig ^b (male and female) | Gilchrist cattle, 1982 |
| Brown goat | Growth curve and absolute growth rate for body weight of Brown goat, Ingemarsson, 1996 |
| Black sheep | Growth curve and absolute growth rate for body weight of black sheep, Ingemarsson, 1996 |
| Merino sheep | Merino sheep, 1996 |
| Californian cattle ^a | Californian cattle, 1996 |
| Athens-Canadian chicken ^a | Athens-Canadian chicken, 1996 |
| Guinea fowl | Guinea fowl, 1996 |
| Japanese quail – white line ^a | Japanese quail – white line, 1996 |
| Japanese quail – brown line ^a | Japanese quail – brown line, 1996 |
| Japanese quail – wild line ^a | Japanese quail – wild line, 1996 |
| Beetal goat | Beetal goat, 1996 |
| Table 1. Means and standard deviations for body weight at different ages in Holstein Friesian bulls from 6 months to 8 years of age. | Table 1. Means and standard deviations for body weight at different ages in Holstein Friesian bulls from 6 months to 8 years of age. |
| Table 1. Males and females observed and estimated by the models of Brody and von Bertalanffy. | Table 1. Males and females observed and estimated by the models of Brody and von Bertalanffy. |
| Figure 1. A) Estimation of weights based on age of Nelore females, observed and estimated by the models of Brody and von Bertalanffy. | Figure 1. A) Estimation of weights based on age of Nelore females, observed and estimated by the models of Brody and von Bertalanffy. |
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| Figure 2. Growth curve for males and females of the variety Barcina slaughtered at 14 months. | Figure 2. Growth curve for males and females of the variety Barcina slaughtered at 14 months. |
| Figure 1. Growth curve and absolute growth rate for body weight of the Karagouniko male sheep: estimate growth curve; observed mean; estimated absolute growth rate. | Figure 1. Growth curve and absolute growth rate for body weight of the Karagouniko male sheep: estimate growth curve; observed mean; estimated absolute growth rate. |
| Figure 2. Growth curve and absolute growth rate for body weight of the Karagouniko female sheep: estimate growth curve; observed mean; estimated absolute growth rate. | Figure 2. Growth curve and absolute growth rate for body weight of the Karagouniko female sheep: estimate growth curve; observed mean; estimated absolute growth rate. |
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| Table 1. Means and standard deviations for body weight at different ages in Athens-Canadian random-bred chickens. | Table 1. Means and standard deviations for body weight at different ages in Athens-Canadian random-bred chickens. |
| Table 2. Means and standard for body weight at different ages in a random-bred pearl guinea fowl population. | Table 2. Means and standard for body weight at different ages in a random-bred pearl guinea fowl population. |
| Sezer and Tarhan, 2005 | Sezer and Tarhan, 2005 |
| Table 1. The results of statistical analyses for body weight of Japanese quail lines at different age (means \pm standard errors). | Table 1. The results of statistical analyses for body weight of Japanese quail lines at different age (means \pm standard errors). |
| Aggrey, 2002 | Aggrey, 2002 |
| Table 1. Means and standard deviations for body weight at different ages in Holstein Friesian bulls from 6 months to 8 years of age. | Table 1. Means and standard deviations for body weight at different ages in Holstein Friesian bulls from 6 months to 8 years of age. |
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| Figure 2. Growth curve and absolute growth rate for body weight of the Karagouniko female sheep: estimate growth curve; observed mean; estimated absolute growth rate. | Figure 2. Growth curve and absolute growth rate for body weight of the Karagouniko female sheep: estimate growth curve; observed mean; estimated absolute growth rate. |
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| Aggrey, 2002 | Aggrey, 2002 |

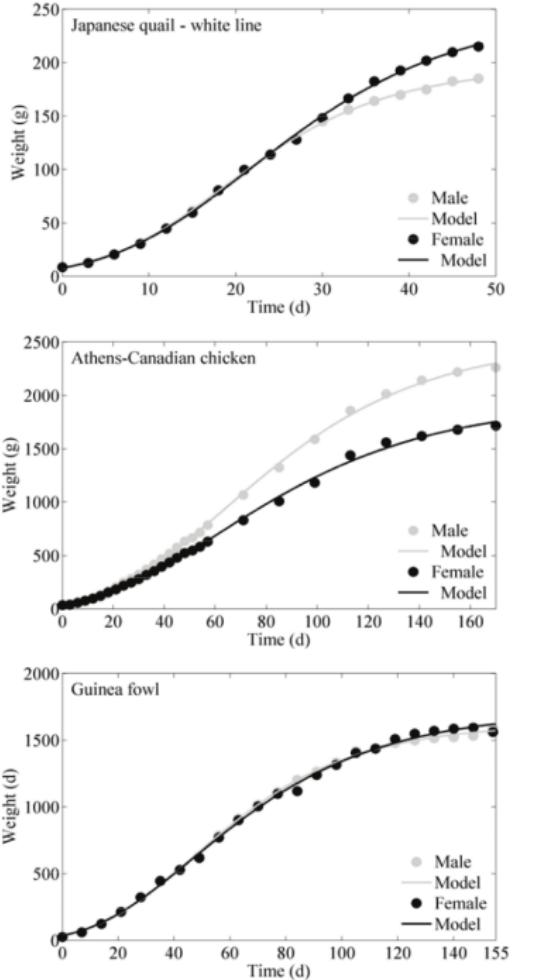


Figure 2. Birds growth kinetics fitted to the Richards' model.

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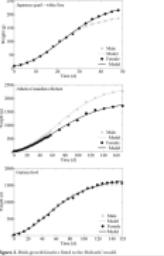


Table 6. Goodness of fit statistics obtained from the growth models applied to the experimental data set of mammals. Equations with the best goodness of fit are represented in bold.

| Animals | Growth models | | | | |
|------------------------|----------------|---------------|--------|---------------|--------|
| | Eq.(1) | Eq.(2) | Eq.(3) | Eq.(4) | Eq.(5) |
| Norfolk rabbit | | | | | |
| <i>R</i> ² | 0.9925 | 0.9991 | 0.9950 | 0.9990 | 0.9992 |
| RMSE | 106.8 | 37.30 | 87.57 | 39.84 | 36.83 |
| BIC | 75.373 | 58.922 | 72.266 | 59.953 | 59.444 |
| AIC _c | 79.322 | 62.870 | 76.214 | 63.901 | 65.500 |
| Californian rabbit | | | | | |
| <i>R</i> ² | 0.9882 | 0.9972 | 0.9943 | 0.9976 | 0.9976 |
| RMSE | 110.6 | 53.98 | 77.26 | 49.91 | 51.45 |
| BIC | 75.917 | 64.701 | 70.307 | 63.477 | 64.667 |
| AIC _c | 79.866 | 68.650 | 74.255 | 67.425 | 70.723 |
| New Zeland rabbit | | | | | |
| <i>R</i> ² | 0.9909 | 0.9985 | 0.9942 | 0.9984 | 0.9986 |
| RMSE | 103.3 | 42.07 | 82.27 | 43.15 | 41.88 |
| BIC | 74.851 | 60.803 | 71.289 | 61.201 | 61.450 |
| AIC _c | 78.799 | 64.752 | 75.238 | 65.150 | 67.506 |
| Holstein-Friesian Bull | | | | | |
| <i>R</i> ² | 0.9958 | 0.9988 | 0.9953 | 0.9986 | 0.9988 |
| RMSE | 17.54 | 9.57 | 18.61 | 10.02 | 9.60 |
| BIC | 67.549 | 53.886 | 68.892 | 54.910 | 54.872 |
| AIC _c | 70.395 | 55.641 | 71.738 | 57.756 | 59.117 |
| Nelore cow | | | | | |
| <i>R</i> ² | 0.9912 | 0.9832 | 0.9641 | 0.9781 | 0.9922 |
| RMSE | 14.84 | 20.55 | 30.01 | 23.41 | 14.81 |
| BIC | 29.856 | 33.243 | 37.193 | 34.603 | 30.294 |
| AIC _c | 35.6187 | 39.006 | 42.955 | 40.365 | 39.691 |
| Angus cow | | | | | |
| <i>R</i> ² | 0.9981 | 0.9921 | 0.9766 | 0.9878 | 0.9982 |
| RMSE | 8.992 | 18.26 | 31.47 | 22.68 | 9.955 |
| BIC | 14.188 | 18.497 | 21.804 | 19.813 | 14.778 |
| AIC _c | 25.653 | 29.961 | 33.268 | 31.277 | 39.397 |

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| Animals | Growth models | | | | |
|-----------------------|---------------|---------------|--------|---------------|--------|
| | Eq.(1) | Eq.(2) | Eq.(3) | Eq.(4) | Eq.(5) |
| New Zealand sheep | | | | | |
| <i>R</i> ² | 0.9992 | 0.9999 | 0.9998 | 0.9999 | 0.9992 |
| RMSE | 30.68 | 17.30 | 47.57 | 33.94 | 36.83 |
| BIC | 73.117 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |
| Merino sheep | | | | | |
| <i>R</i> ² | 0.9982 | 0.9992 | 0.9994 | 0.9996 | 0.9976 |
| RMSE | 44.94 | 24.70 | 54.30 | 23.47 | 44.83 |
| BIC | 73.147 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |
| Dorset sheep | | | | | |
| <i>R</i> ² | 0.9988 | 0.9999 | 0.9998 | 0.9999 | 0.9988 |
| RMSE | 30.68 | 17.30 | 47.57 | 33.94 | 36.83 |
| BIC | 73.117 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |
| Wiltshire Horn sheep | | | | | |
| <i>R</i> ² | 0.9988 | 0.9999 | 0.9998 | 0.9999 | 0.9988 |
| RMSE | 30.68 | 17.30 | 47.57 | 33.94 | 36.83 |
| BIC | 73.117 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |
| Lambing ewes | | | | | |
| <i>R</i> ² | 0.9988 | 0.9999 | 0.9998 | 0.9999 | 0.9988 |
| RMSE | 30.68 | 17.30 | 47.57 | 33.94 | 36.83 |
| BIC | 73.117 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |
| Lambing rams | | | | | |
| <i>R</i> ² | 0.9988 | 0.9999 | 0.9998 | 0.9999 | 0.9988 |
| RMSE | 30.68 | 17.30 | 47.57 | 33.94 | 36.83 |
| BIC | 73.117 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |
| Lambing ewes and rams | | | | | |
| <i>R</i> ² | 0.9988 | 0.9999 | 0.9998 | 0.9999 | 0.9988 |
| RMSE | 30.68 | 17.30 | 47.57 | 33.94 | 36.83 |
| BIC | 73.117 | 53.800 | 74.760 | 54.441 | 54.441 |
| AIC _c | 79.322 | 62.870 | 76.214 | 62.981 | 63.598 |

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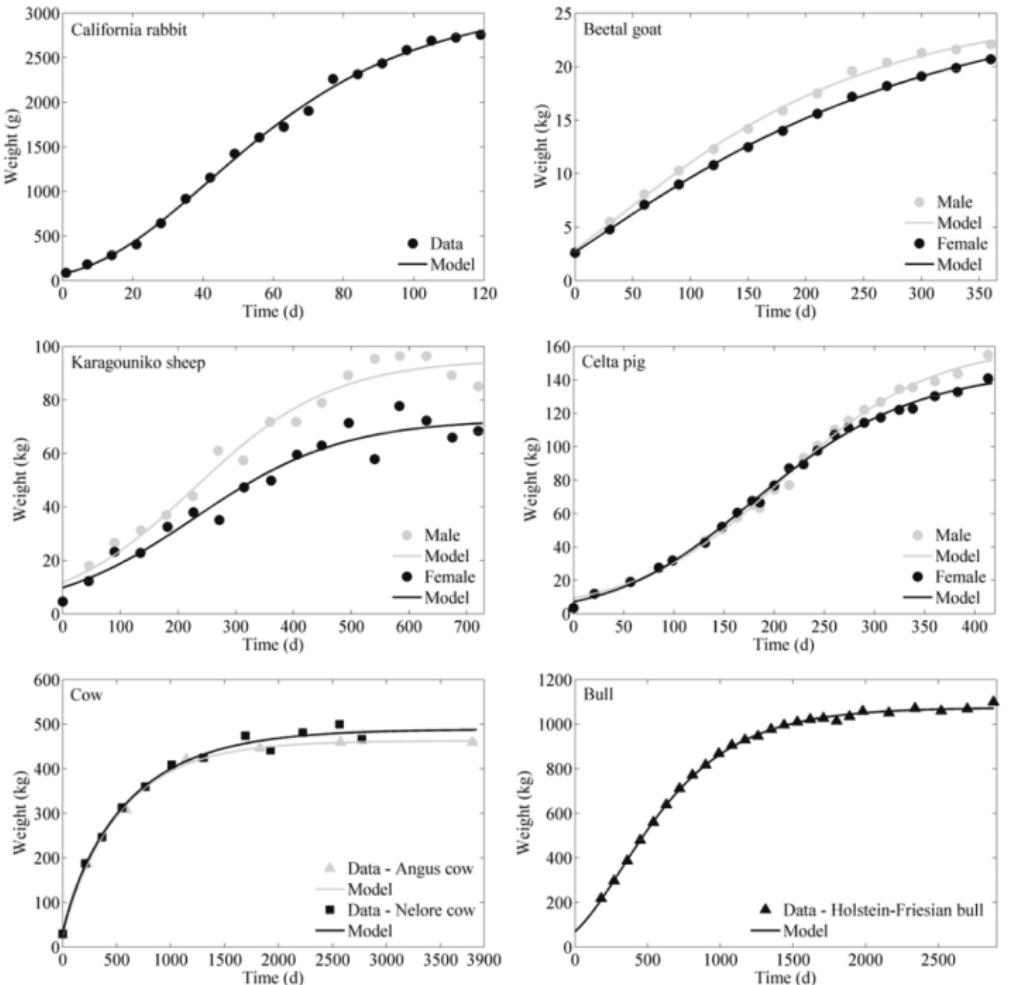
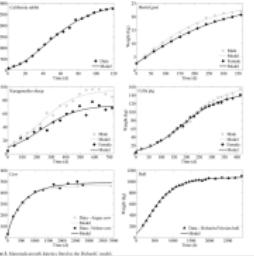


Figure 3. Mammals growth kinetics fitted to the Richards' model.

Comparison of five mathematical models that describe growth in tropically adapted dual-purpose breeds of chicken

Oludayo Michael Akinsola  ^a, Emmanuel Babafunso Sonaifya  ^b, Oladeji Bamidele  ^b, Waheed Akinola Hassan  ^c, Abdulmojeed Yakubu  ^d, Folasade Olubukola Ajayi  ^e, Uduak Ogundu  ^f, Olayinka Olubunmi Alabi  ^g and Oluwafunmilayo Ayoka Adebambo  ^h

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^aDepartment of Theriogenology and Production, University of Jos, Jos, Nigeria; ^bAfrican Chicken Genetic Gains Project National Secretariat, Department of Animal Science, Obafemi Awolowo University, Ile-Ife, Nigeria; ^cDepartment of Animal Science, Usmanu Danfodiyo University, Sokoto, Nigeria; ^dDepartment of Animal Science, Faculty of Agriculture, Nasarawa State University, Shabu-Lafia Campus Lafia, Keffi, Nigeria; ^eDepartment of Animal Science, University of Port-Harcourt, Port-Harcourt, Nigeria; ^fDepartment of Animal Science, Federal University of Technology, Owerri, Nigeria; ^gDepartment of Animal Science, Landmark University, Omu-Aran, Nigeria; ^hDepartment of Animal Breeding and Genetics, Federal University of Agriculture, Abeokuta, Nigeria

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| Model | Equations | Age at inflection point | Weight at inflection point |
|-------------|--|--------------------------------------|----------------------------|
| Gompertz 3P | $Y = a \cdot \exp(-\exp(-b \cdot (\text{age} - c)))$ | $\ln \frac{b}{a}$ | $\frac{c}{e}$ |
| Logistic 3P | $Y = \frac{c}{(1 + \exp(-a \cdot (\text{age} - b)))}$ | $-\ln \left(\frac{1}{b}\right)^{-a}$ | $\frac{c}{2}$ |
| Gompertz 4P | $Y = a + (b - a) \cdot \exp(-\exp(-c \cdot (\text{age} - d)))$ | $\ln \left(\frac{d}{a}\right)$ | $\frac{a}{e} \cdot c$ |
| Logistic 4P | $Y = c + \frac{d - c}{(1 + \exp(-a \cdot (\text{age} - b)))}$ | $\frac{c + d}{2}$ | $\frac{d}{2}$ |

Table 1. Equations of the non-linear regression growth curve models.

| Model | Equations | Age at inflection point | Weight at inflection point |
|-------------|--|--------------------------------------|----------------------------|
| Gompertz 3P | $Y = a \cdot \exp(-\exp(-b \cdot (\text{age} - c)))$ | $\ln \frac{b}{a}$ | $\frac{c}{e}$ |
| Logistic 3P | $Y = \frac{c}{(1 + \exp(-a \cdot (\text{age} - b)))}$ | $-\ln \left(\frac{1}{b}\right)^{-a}$ | $\frac{c}{2}$ |
| Gompertz 4P | $Y = a + (b - a) \cdot \exp(-\exp(-c \cdot (\text{age} - d)))$ | $\ln \left(\frac{d}{a}\right)$ | $\frac{a}{e} \cdot c$ |
| Logistic 4P | $Y = c + \frac{d - c}{(1 + \exp(-a \cdot (\text{age} - b)))}$ | $\frac{c + d}{2}$ | $\frac{d}{2}$ |

Notes: Y is the estimated weight at age x ; a is the maturity index; b is the scale parameter; c is the asymptotic weight; d is the upper asymptote; Gompertz 3P was referenced from Gompertz (1832); Logistic 3P from Darmani et al. (2010); Logistic 4P from Ratwosky and Reddy (1986); Gompertz 4P from Tjørve and Tjørve (2017).

| Model | Equations | Age at inflection point | Weight at inflection point |
|-------------|--|--------------------------------------|----------------------------|
| Gompertz 3P | $Y = a \cdot \exp(-\exp(-b \cdot (\text{age} - c)))$ | $\ln \frac{b}{a}$ | $\frac{c}{e}$ |
| Logistic 3P | $Y = \frac{c}{(1 + \exp(-a \cdot (\text{age} - b)))}$ | $-\ln \left(\frac{1}{b}\right)^{-a}$ | $\frac{c}{2}$ |
| Gompertz 4P | $Y = a + (b - a) \cdot \exp(-\exp(-c \cdot (\text{age} - d)))$ | $\ln \left(\frac{d}{a}\right)$ | $\frac{a}{e} \cdot c$ |
| Logistic 4P | $Y = c + \frac{d - c}{(1 + \exp(-a \cdot (\text{age} - b)))}$ | $\frac{c + d}{2}$ | $\frac{d}{2}$ |

Note: Y is the estimated weight at age x ; a is the maturity index; b is the scale parameter; c is the asymptotic weight; d is the upper asymptote; Gompertz 3P was referenced from Gompertz (1832); Logistic 3P from Darmani et al. (2010); Logistic 4P from Ratwosky and Reddy (1986); Gompertz 4P from Tjørve and Tjørve (2017).

Materials and methods

Experimental site

The on-station test was conducted at Fol-Hope Farms, Ibadan, Oyo State and the Federal University of Agriculture, Abeokuta (FUNAAB), located within the Southern Guinea Savanna, and Dry Lowland Rainforest agro-ecological zones, respectively. The testing of the birds commenced in May 2016. The on-farm test was carried out in five agro-ecological zones as follows: Kebbi State (Sudan and Northern Guinea Savanna), Kwara State (Southern Guinea Savanna), Nasarawa State (Southern Guinea Savanna), Imo State (Wet Lowland Rain Forest and Fresh Water Swamp) and Rivers State (Mangrove Swamp and Fresh Water Swamp).

Management systems

A total of 1939 d-old chicks of both locally sourced breeds (Fulani, FUNAAB Alpha, Noiler and Shika-Brown) and imported breeds (Kuroiler and Sasso) were brooded to 42 days (**Table 2**). The birds were sexed at 42 days, and males and females were grown separately until 140 days under station (intensive production system) conditions. The stocking density was 10 chicks/m², seven birds/m², and five birds/m² during 0–42d, 43–91d and 92–140d, respectively. Commercial feed (Chick mash at 0–42d: 2,993 kcal ME/kg, 22.3% CP and Grower mash at 43–140d: 3013 kcal ME/kg, 17% CP) and water were available *ad libitum*. Birds in both stations were fed the same proprietary feed. Standard biosecurity measures and vaccination schedules were observed at the test centres. Body weight was measured every two weeks. For the on-farm test, a total of 58,639 six-weeks-old pre-vaccinated chickens were distributed to 2100

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Materials and methods
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households across five states representing different agro-ecologies ([Table 3](#)). Standard backyard scavenging management practices were followed by the farmers with the addition of overnight housing, feed supplementation and vaccination programmes. Body weight was taken every four weeks.

All applicable veterinary permits were obtained for the importation and use of the imported breeds for research purposes (Bamidele et al. [2019](#)). Both the on-station and on-farm studies were approved by the International Livestock Research Institute (ILRI) Institutional Research Ethics Committee (IREC) with reference no.: ILRI-IREC2015-08/1, and ILRI Institutional Animal Care and Use Committee (IACUC) with reference number: ILRI-IACUC-RC2016.2. Each farmer gave written informed consent to participate in the study.

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Table 5. Estimated growth curve parameters of female birds raised on-station from 0 to 20 weeks.

| Breed/model | a | b | c | d | Age and weight at inflection point | AIC | BIC | RMSE | AdjR ² | |
|---------------------|------|-------|---------|---------|------------------------------------|------------|---------|---------|-------------------|-------|
| Fulani | | | | | | | | | | |
| Logistic 4P | 0.26 | 11.35 | -38.10 | 1429.00 | 10; 452.42 | 90.350 | 80.340 | 6.779 | 0.995 | |
| Logistic 3P | 0.28 | 11.38 | | 1109.22 | 10; 446.99 | 95.497 | 90.422 | 11.183 | 0.992 | |
| Gompertz 3P | 0.14 | 10.46 | | 1347.86 | 10; 463.54 | 99.351 | 94.276 | 13.324 | 0.989 | |
| Gompertz 4P | 0.16 | 10.41 | | 24.80 | 10; 461.67 | 102.561 | 92.550 | 11.809 | 0.988 | |
| Neural network | | | | | 10; 469.75 | | | 5.933 | 0.992 | |
| FUNAAB Alpha | | | | | | | | | | |
| Logistic 3P | 0.40 | 10.41 | | 2780.35 | 10;1030.22 | 121.831 | 116.756 | 37.019 | 0.994 | |
| Gompertz 3P | 0.20 | 9.07 | | 2735.53 | 10;1070.99 | 125.555 | 120.480 | 43.846 | 0.990 | |
| Logistic 4P | 0.30 | 10.32 | -49.12 | 2247.55 | 10;1037.22 | 126.763 | 116.753 | 35.482 | 0.990 | |
| Gompertz 4P | 0.22 | 9.21 | | 59.16 | 10;1067.04 | 128.188 | 118.177 | 37.856 | 0.989 | |
| Neural network | | | | | 10;1084.91 | | | 38.829 | 0.990 | |
| Sasso | | | | | | | | | | |
| Gompertz 3P | 0.14 | 10.89 | | 3856.63 | 10;1240.85 | 127.704 | 122.629 | 48.346 | 0.997 | |
| Gompertz 4P | 0.14 | 10.88 | | 7.06 | 10;1240.36 | 135.016 | 125.006 | 51.633 | 0.992 | |
| Logistic 3P | 0.28 | 11.76 | | 3155.02 | 10;1192.88 | 138.815 | 133.740 | 80.110 | 0.988 | |
| Logistic 4P | 0.23 | 11.82 | -232.47 | 3412.27 | 10;1221.52 | 139.646 | 129.635 | 63.726 | 0.987 | |
| Neural network | | | | | 10;1261.09 | | | 38.913 | 0.997 | |
| Kuroiler | | | | | | | | | | |
| Gompertz 3P | 0.14 | 10.40 | | 3725.38 | 10;1292.28 | 119.769 | 114.694 | 33.706 | 0.995 | |
| Gompertz 4P | 0.14 | 10.43 | -23.13 | 3777.74 | 10;1293.34 | 126.740 | 116.729 | 35.444 | 0.991 | |
| Logistic 4P | 0.21 | 11.38 | -300.25 | 3385.91 | 10;1276.43 | 132.982 | 122.972 | 47.074 | 0.990 | |
| Logistic 3P | 0.28 | 11.41 | | 3090.09 | 10;1246.74 | 137.304 | 132.228 | 74.792 | 0.985 | |
| Neural network | | | | | 10;1290.89 | | | 17.773 | 0.996 | |
| Shika-Brown | | | | | | | | | | |
| Gompertz 3P | 0.17 | 10.09 | | 2088.27 | 10; 756.01 | 115.702 | 110.627 | 28.016 | 0.998 | |
| Logistic 3P | 0.32 | 11.27 | | 1803.37 | 10; 722.05 | 116.621 | 111.546 | 29.213 | 0.998 | |
| Logistic 4P | 0.29 | 11.19 | -58.62 | 1847.41 | 10; 732.26 | 118.844 | 108.833 | 24.756 | 0.998 | |
| Gompertz 4P | 0.18 | 10.13 | | 32.93 | 2034.90 | 10; 752.68 | 120.354 | 110.343 | 26.514 | 0.995 |
| Neural network | | | | | 10; 712.52 | | | 7.287 | 0.998 | |
| Noiler | | | | | | | | | | |
| Gompertz 3P | 0.18 | 7.97 | | 2917.67 | 13;1955.35 | 105.461 | 92.445 | 58.794 | 0.992 | |
| Logistic 3P | 0.27 | 9.66 | | 30.33 | 2742.17 | 13;1959.03 | 108.221 | 95.206 | 69.867 | 0.990 |
| Logistic 4P | 0.13 | 0.44 | -5131.7 | 3139.89 | 13;1947.59 | 122.384 | 92.781 | 58.948 | 0.989 | |
| Gompertz 4P | 0.12 | 0.22 | -3231.4 | 3204.77 | 13;1946.85 | 122.442 | 92.839 | 59.162 | 0.989 | |
| Neural network | | | | | 13;1916.49 | | | 49.327 | 0.990 | |

Notes: AIC: akaike information criterion; BIC: Bayesian information criterion; RMSE: root mean square error; AdjR²: adjusted coefficient of determination; a is the maturity index; b is the scale parameter; c is the asymptotic weight; d is the upper asymptote. Non-linear model adapted from JMP 13.2 statistical software.

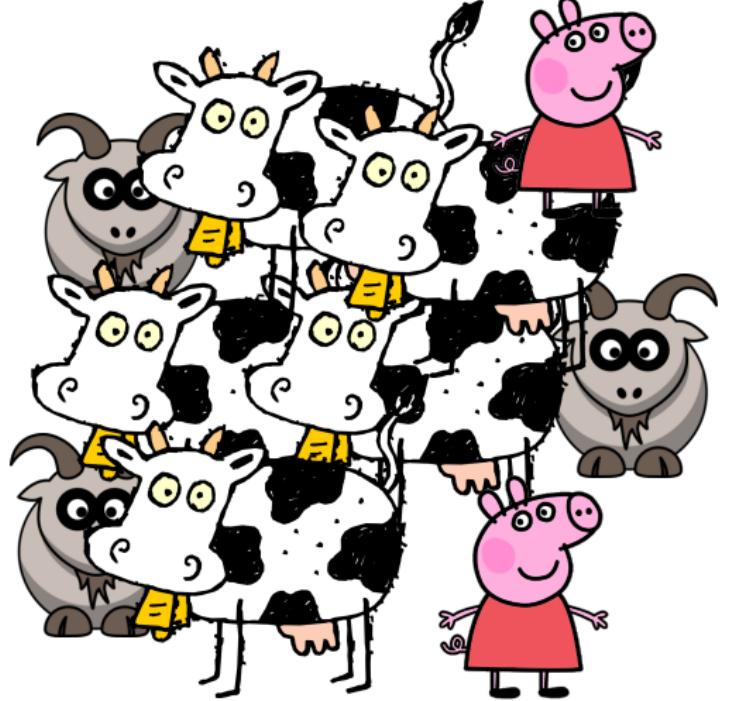
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| | a | b | c | d | Age and weight at inflection point | AIC | BIC | RMSE | AdjR ² | |
|----------------|------|-------|---------|---------|------------------------------------|------------|---------|---------|-------------------|-------|
| Female birds | | | | | | | | | | |
| Logistic 4P | 0.28 | 11.28 | -18.70 | 1409.22 | 10;462.40 | 84.407 | 74.422 | 6.525 | 0.996 | |
| Logistic 3P | 0.28 | 11.28 | -18.70 | 1398.16 | 10;462.40 | 84.407 | 74.422 | 6.525 | 0.996 | |
| Gompertz 3P | 0.16 | 10.41 | | 24.80 | 10;461.67 | 85.481 | 90.530 | 6.080 | 0.996 | |
| Gompertz 4P | 0.16 | 10.41 | | 24.80 | 10;469.75 | 85.481 | 90.530 | 6.080 | 0.996 | |
| Neural network | | | | | 10;469.75 | | | | | |
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| Logistic 3P | 0.28 | 11.41 | | 3090.09 | 10;1246.74 | 137.304 | 132.228 | 74.792 | 0.985 | |
| Neural network | | | | | 10;1290.89 | | | 17.773 | 0.996 | |
| Shika-Brown | | | | | | | | | | |
| Gompertz 3P | 0.17 | 10.09 | | 2088.27 | 10; 756.01 | 115.702 | 110.627 | 28.016 | 0.998 | |
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Note: AIC: akaike information criterion; BIC: Bayesian information criterion; RMSE: root mean square error; AdjR²: adjusted coefficient of determination; a is the maturity index; b is the scale parameter; c is the asymptotic weight; d is the upper asymptote. Non-linear model adapted from JMP 13.2 statistical software.

Heterogeneous herds



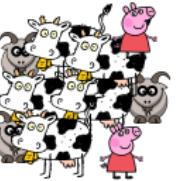
Modelling livestock

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└ Heterogeneous herds

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Heterogeneous herds





Livestock Science 157 (2013) 280–288

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A mathematical model of the dynamics of Mongolian livestock populations



Duncan Shabb^a, Nakul Chitnis^a, Zolzaya Baljinnyam^{a,b}, Sansar Saagii^c,
Jakob Zinsstag^{a,*}

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^b Institute of Veterinary Medicine, Mongolian State University of Agriculture, Ulaanbaatar, Mongolia

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^b Institute of Veterinary Medicine, Mongolian State University of Agriculture, Ulaanbaatar, Mongolia
^c Amin Nutag Orgil NGO, Ulaanbaatar, Mongolia

The state variables for total population at time t of goats sheep, cattle and horses are denoted G , S , C , and H respectively

$$G(t) = \sum_{i=1}^3 X_{ig}(t) + Y_{ig}(t),$$

$$S(t) = \sum_{i=1}^3 X_{is}(t) + Y_{is}(t),$$

$$C(t) = \sum_{i=1}^3 X_{ic}(t) + Y_{ic}(t),$$

$$H(t) = \sum_{i=1}^3 X_{ih}(t) + Y_{ih}(t), \quad (1)$$

with the juvenile population of cattle given by

$$X_{2c} = \sum_{\xi \in \{a,b\}} X_{2c\xi},$$

$$Y_{2c} = \sum_{\xi \in \{a,b\}} Y_{2c\xi},$$

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(1)

with the juvenile population of cattle given by

$$X_{2c} = \sum_{\{\alpha, \beta\}} X_{2c\alpha},$$

$$Y_{2c} = \sum_{\{\alpha, \beta\}} Y_{2c\alpha},$$

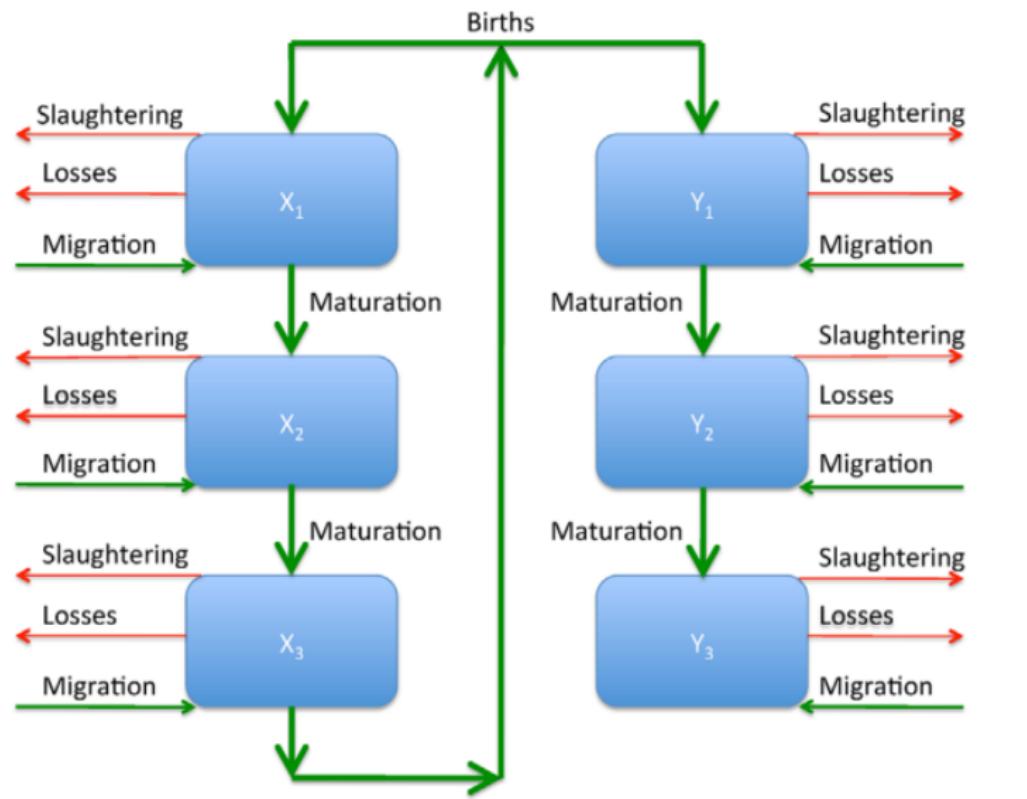


Fig. 1. Goat model dynamics. Arrows represent flows of animals into or out of the population. Boxes represent state variables. The figure is meant to be a simple conceptualization of the model dynamics. Sheep are modeled with an equivalent schematic but cattle and horses have additional juvenile stages.

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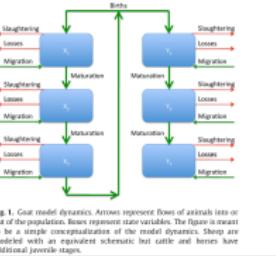


Fig. 1. Goat model dynamics. Arrows represent flows of animals into or out of the population. Boxes represent state variables. The figure is meant to be a simple conceptualization of the model dynamics. Sheep are modeled with an equivalent schematic but cattle and horses have additional juvenile stages.

The model equations for goats are given by

$$X_{1g}(t+1) = b_g X_{3g}(t) + \frac{m_g X_{1g}(t)}{G(t)}, \quad (3a)$$

$$X_{2g}(t+1) = \frac{X_{1g}(t)}{(1 + \alpha_g d_{X_{1g}} A(t)) k_{X_{1g}}} + \frac{m_g X_{2g}(t)}{G(t)}, \quad (3b)$$

$$\begin{aligned} X_{3g}(t+1) = & \frac{X_{2g}(t)}{(1 + \alpha_g d_{X_{2g}} A(t)) k_{X_{2g}}} + \frac{X_{3g}(t)}{(1 + \alpha_g d_{X_{3g}} A(t)) k_{X_{3g}}} \\ & + \frac{m_g X_{3g}(t)}{G(t)}, \end{aligned} \quad (3c)$$

$$Y_{1g}(t+1) = b_g X_{3g}(t) + \frac{m_g Y_{1g}(t)}{G(t)}, \quad (3d)$$

$$Y_{2g}(t+1) = \frac{Y_{1g}(t)}{(1 + \alpha_g d_{Y_{1g}} A(t)) k_{Y_{1g}}} + \frac{m_g Y_{2g}(t)}{G(t)}, \quad (3e)$$

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| Table 1 | | |
|---|---|-------------------|
| Description of types parameters of the model for livestock population dynamics. The model consists of five basic parameters that are further subdivided by species out of which some are further subdivided by age and sex. | | |
| Parameter | Description | Level |
| b | Expected number of female offspring per one adult female over one year. Dimensionless | Species |
| m | Number of new animals entering the area in one year. Dimension: Animals | Species |
| α | Species-dependent resource availability. Dimension: 1/Animals | Species |
| d | Modulation of resource availability by age and sex of animal. Dimensionless | Species, sex, age |
| k | Reciprocal of per capita survival excluding the effects of density-dependent mortality. Dimensionless | Species, sex, age |
| c | Weighting of food consumption by animals. Dimensionless | Species, age |

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Table 1

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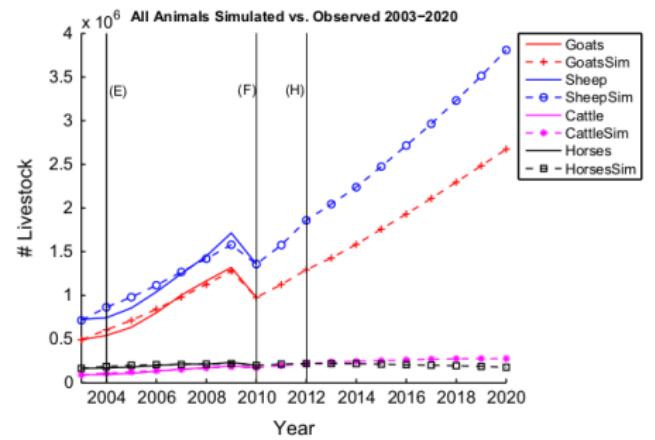
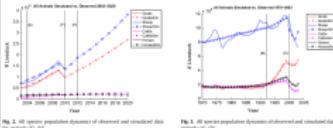


Fig. 2. All species population dynamics of observed and simulated data for periods (E)–(H).

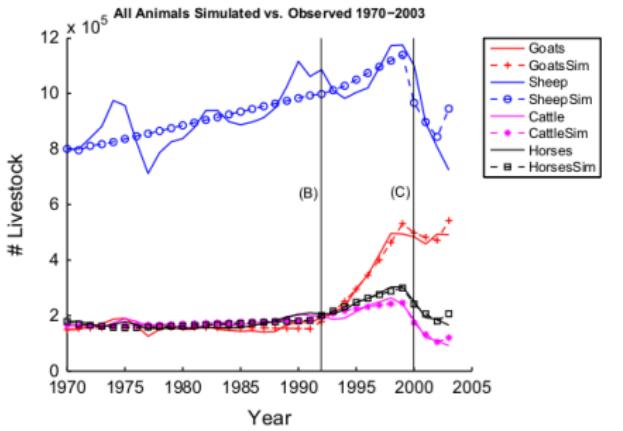


Fig. 3. All species population dynamics of observed and simulated data of periods (A)–(D).

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Models for cattle

Models for other herds

Models for management

Conclusion

REVIEW ARTICLE

Systems approaches to beef cattle production systems using modeling and simulation

Hiroyuki HIROOKA

Graduate School of Agriculture, Kyoto University, Sakyo, Kyoto, Japan

ABSTRACT

Systems approach techniques have been applied to modeling production systems for beef cattle from the relatively micro-level of tissues and organs to the macro-level of farms and geographical regions. This paper reviews the various types of beef cattle production models already in operation in order to analyze beef cattle production systems and their components. It may be theoretically possible to construct system models which describe such complex production systems and can be generally used in various genetic, nutritional, management and economic situations as well as in training, extension and educational programs. Moreover, the systems approach can assist in the organization of information and identification of knowledge gaps and thereby open an avenue to multi-disciplinary research projects.

Key words: beef cattle, model, simulation, systems approach.

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Animal Science Journal (2010) 81, 411–424

REVIEW ARTICLE

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APPLICATION OF MATHEMATICAL MODELLING IN BEEF HERD
MANAGEMENT – A REVIEW*

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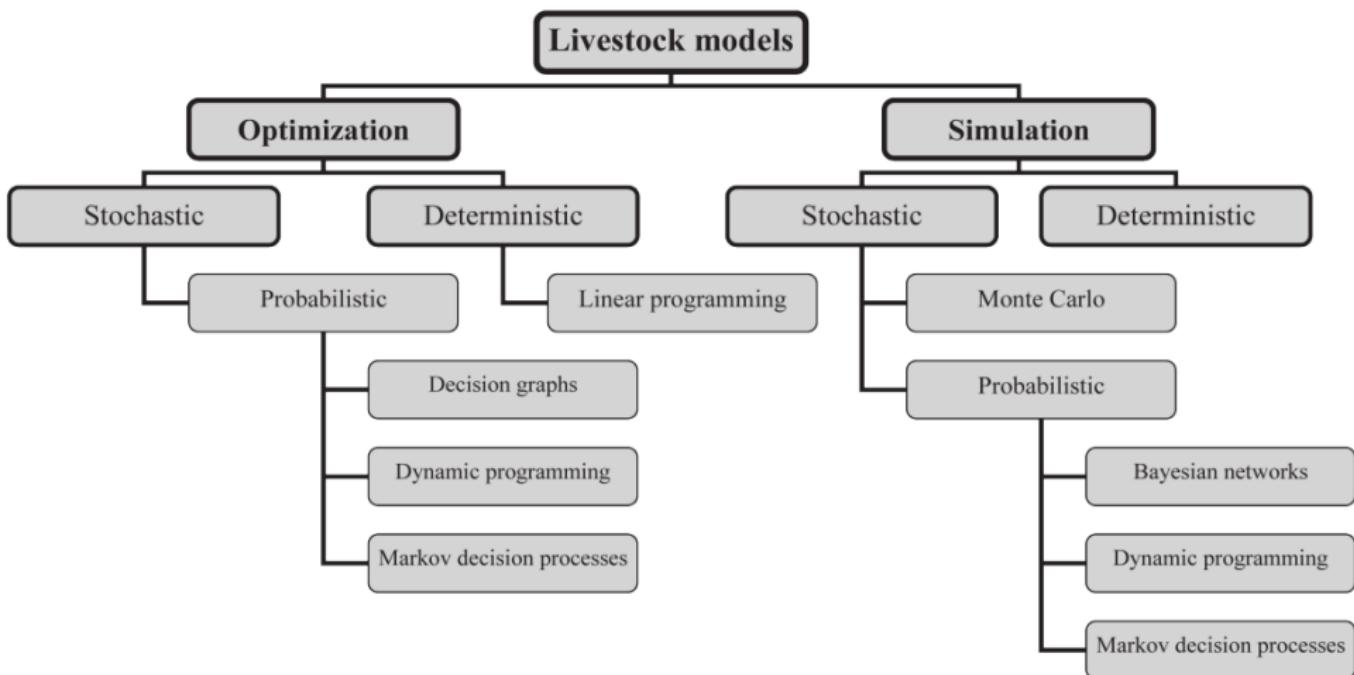


Figure 1. Methodological classification of livestock models

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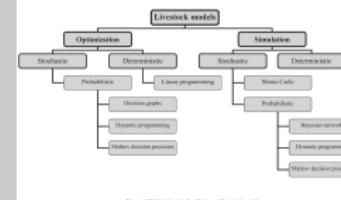


Figure 1. Methodological classification of livestock models



ASAS-NANP Symposium: Mathematical Modeling in Animal Nutrition: Opportunities and challenges of confined and extensive precision livestock production

Hector M. Menendez III,^{†,†,ID} Jameson R. Brennan,[†] Charlotte Gaillard,^{‡,ID} Krista Ehlert,[†] Jaelyn Quintana,[†] Suresh Neethirajan,[§] Aline Remus,^{¶,ID} Marc Jacobs,^{**} Izabelle A. M. A. Teixeira,^{††} Benjamin L. Turner,^{‡‡} and Luis O. Tedeschi^{§,ID}

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^{§,ID}Department of Animal Science, Texas A&M University, College Station, TX 77843-2471, USA

[†]Corresponding author: hector.menendez@sdstate.edu

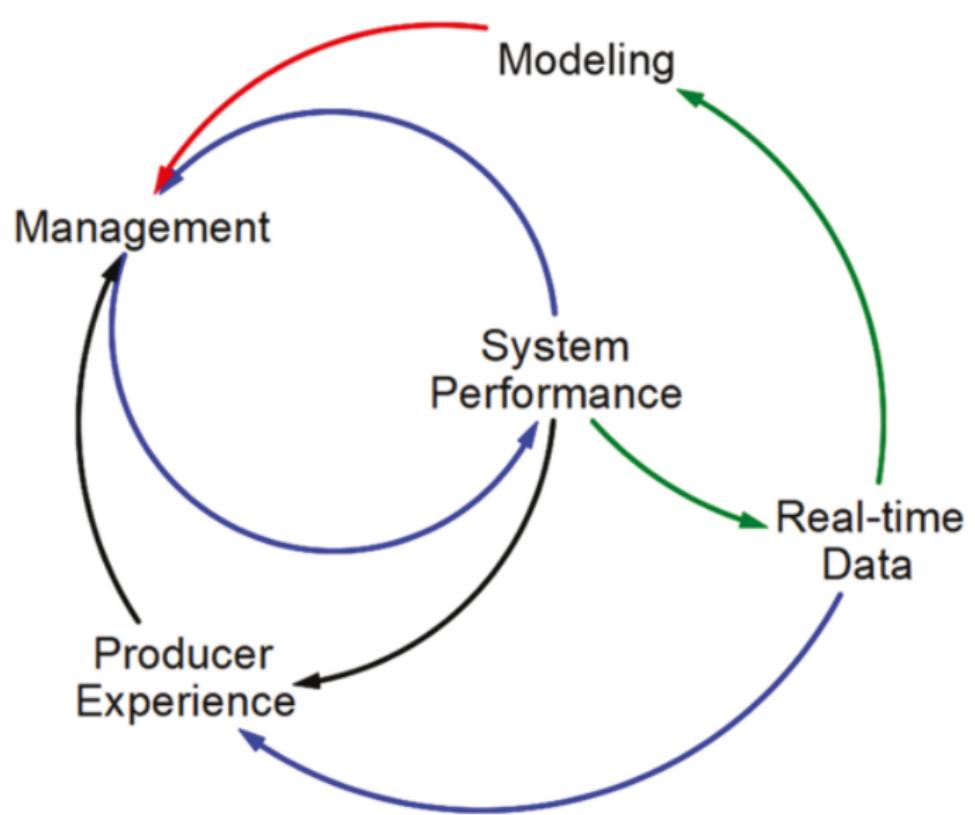


Figure 1. Diagram of the conventional producer decision process, including mental models (producer experience) and the role of modeling in relation to real-time data integration.



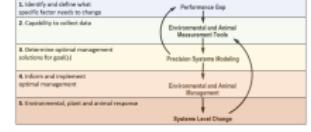


Figure 2. Conceptual diagram of five principles for sustainable precision livestock implementation using precision measurement and management tools integrated with mathematical models.

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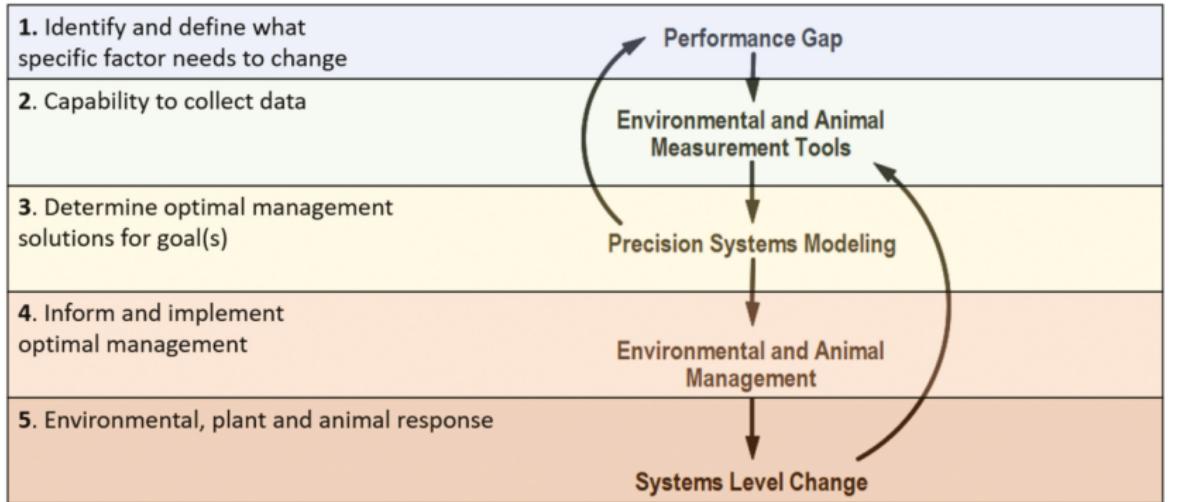


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Models for management

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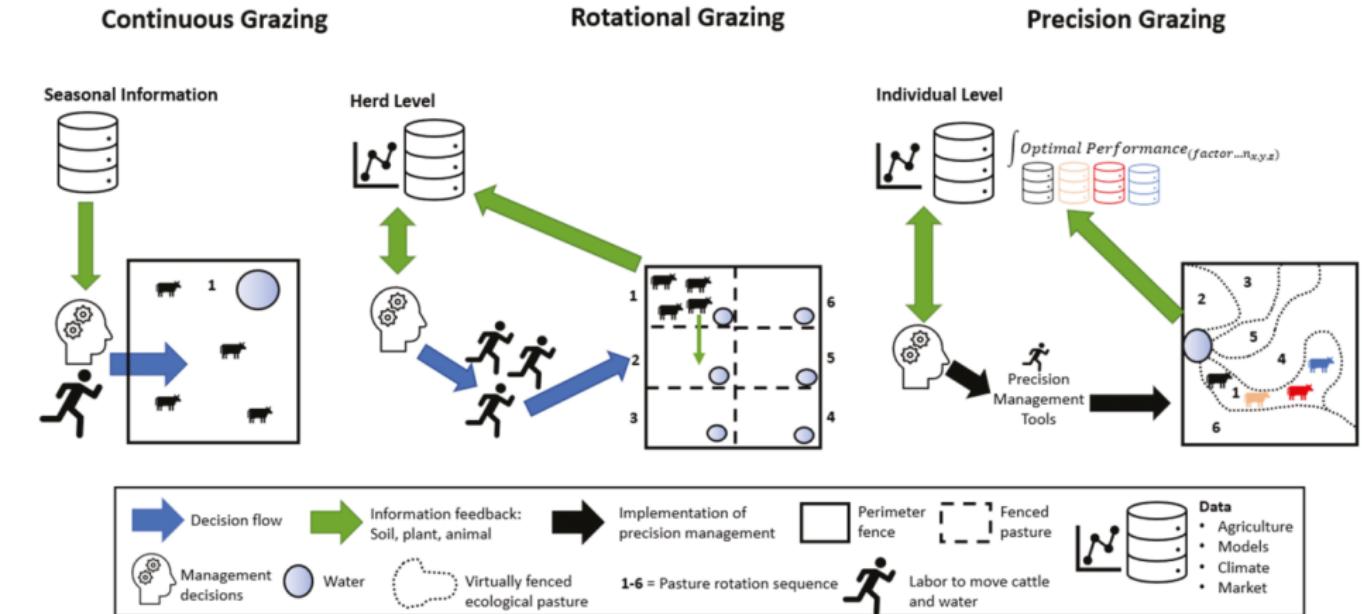
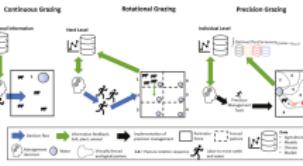


Figure 3. Continuous vs. rotational grazing as each relates to decision flow, information feedback, and management decisions, labor, and data.

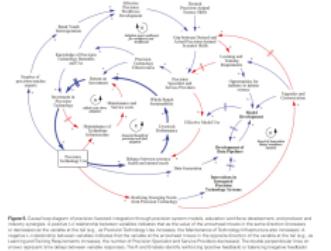


Figure 4. Causal loop diagram of precision livestock integration through precision system models, education workforce development, and producer and industry synergies. A positive (+) relationship between variables indicates that as the value of the arrowhead moves in the same direction (increases or decreases) as the variable at the tail (e.g., as Precision Technology Use increases, the Maintenance of Technology Infrastructure also increases). A negative (-) relationship between variables indicates that the variable at the arrowhead moves in the opposite direction of the variable at the tail (e.g., as Learning and Training Requirements increases, the number of Precision Specialist and Service Providers decreases). The double perpendicular lines on arrows represent time delays between variable responses. The R and B labels identify reinforcing (positive feedback) or balancing (negative feedback) relationships.

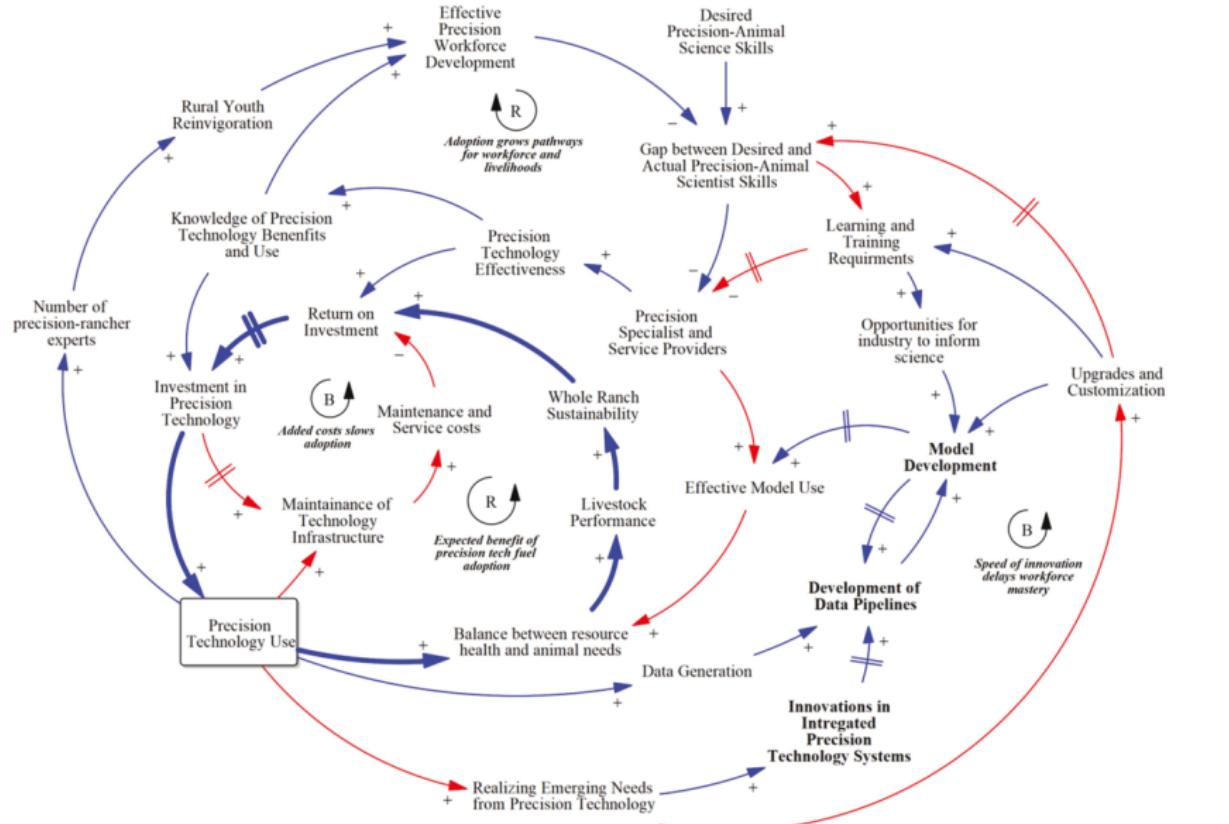


Figure 5. Causal loop diagram of precision livestock integration through precision system models, education workforce development, and producer and industry synergies. A positive (+) relationship between variables indicates that as the value of the arrowhead moves in the same direction (increases or decreases) as the variable at the tail (e.g., as Precision Technology Use increases, the Maintenance of Technology Infrastructure also increases). A negative (-) relationship between variables indicates that the variable at the arrowhead moves in the opposite direction of the variable at the tail (e.g., as Learning and Training Requirements increases, the number of Precision Specialist and Service Providers decreases). The double perpendicular lines on arrows represent time delays between variable responses. The R and B labels identify reinforcing (positive feedback) or balancing (negative feedback) relationships.

Modelling livestock └ Models for management

2023-04-17

Modelling livestock

Models for management

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Table 1. Real-time models found in the literature using the search keywords: real-time, animal science, nutrition, and modeling

| Author | Aim | Target | Type | Response |
|---|---|--------------|--|--|
| Hauschild et al. (2012, 2020); Remus et al. (2020c) | Provide daily tailored diets to individuals | Growing pigs | Gray box (empirical [data-driven] and mechanistic) | Diet composition to sustain observed growth |
| Peña Fernández et al. (2019) | Predict in real-time the indoor particle sizes concentration | Poultry | Data-based mechanistic | Predicted indoor particle sizes concentration |
| Parsons et al. (2007) | Integrated control of pig growth and pollutant emissions | Growing pigs | Data-based mechanistic | Predicted growth response based on diet intake |
| Stacey et al. (2004) | Control of broiler growth and nutrition | Broiler | Semi-mechanistic | Predicted growth response based on diet intake and control nutrient intake |
| Fu et al. (2020) | Predict diet energy digestion | Dairy cows | Kernel extreme learning machine | Predicted digestible energy and energy digestibility |
| Kashiha et al. (2013) | Report malfunctioning in a broiler house to the farmer in real time | Broiler | Empirical (data-driven) | Prediction of the distribution index of broilers |
| Gauthier et al. (2019); Gaillard et al. (2020b) | Provide daily tailored diets to individuals | Sows | Gray box (empirical [data-driven] and mechanistic) | Diet composition to sustain fetus development and milk production |

| Table 1. Real-time models found in the literature using the search keywords: real-time, animal science, nutrition, and modeling | | | | |
|---|---|--------------|--|--|
| Author | Aim | Type | Response | Notes |
| Hauschild et al. (2012, 2020); Remus et al. (2020c) | Provide daily tailored diets to individuals | Growing pigs | Gray box (empirical [data-driven] and mechanistic) | The response is semi-observed |
| Peña Fernández et al. (2019) | Predict in real-time the indoor particle sizes concentration | Poultry | Data-based mechanistic | The indoor particle size concentration |
| Parsons et al. (2007) | Integrated control of pig growth and pollutant emissions | Growing pigs | Data-based mechanistic | Predicted growth response based on diet intake |
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Modelling livestock

└ Conclusion

2023-04-17

Models for cattle

Models for other herds

Models for management

Conclusion

Many interesting models

(Many models have never been “properly” studied mathematically and are “waiting” for an analysis)

I showed only one statistical model, but the majority of published work used to be statistical

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Modelling livestock └ Conclusion

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