

Index for Mastitis Resistance and Use of BHBA for Evaluation of Health Traits in Canadian Holsteins

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Genetic evaluation for mastitis resistance

- In August 2014, first official run of genetic evaluations for mastitis resistance
- Multiple-trait linear animal model (Jamrozik et al., 2013)
 - First vs. later parities: clinical mastitis, mean SCS, standard deviation of SCS, excessive test-day SCC
 - First parity cows: udder depth, fore udder attachment, body condition score
- Genetic evaluations expressed as relative breeding values (RBV) with a mean of 100 and a SD of 5 (higher values are desirable)



Objective

Development of a Mastitis Resistance Index

Clinical mastitis in first lactation (CM-F)

Clinical mastitis in later lactations (CM-L)

SCS from Canadian Test Day Model



Why an index?

- Why not using just mastitis EBV?
- Mastitis EBV are indicators of clinical mastitis
- SCS EBV are indicators of subclinical mastitis





Boettcher et al., 1998

Udder health index

- Subclinical mastitis (measured by SCS) in lactations 1 and ≥2
- ➢ Clinical mastitis in lactations 1 and ≥2
- Milking time
- Estimated economic weights were -\$12, -\$31, -\$15, -\$59 and -\$11, respectively, per genetic standard deviation.
- At that time clinical mastitis was not recorded, thus traits in the selection index were milking speed, udder conformation and SCS in first and later lactations



Mastitis Resistance Index

Based on the work by Boettcher et al. (1998)

Mastitis Resistance (MR) = 1/3 CM-F + 1/3 CM-L - 1/3 SCS

where;

CM-F = Clinical Mastitis in First lactation CM-L = Clinical Mastitis in Later lactations SCS = overall SCS evaluation as officially published whereby low values are desired



Selection response

Assumptions

- Heritability for CM-F, CM-L and SCS = 0.03, 0.05, 0.20, respectively
- Genetic correlations among the three traits: CM-F with CM-L = 0.60 and 0.55 for the other 2 combinations
- Reliability of RBV for MR traits = 0.30, and for SCS = 0.50 (conservative estimates)
- Selection only on Mastitis Resistance (with various combinations/emphasis among 3 traits)



Selection response

| Weights | | | Genetic gain per year (RBV points) | | |
|---------|------|-----|---------------------------------------|------|------|
| CM-F | CM-L | SCS | CM-F | CM-L | SCS |
| 1/3 | 1/3 | 1/3 | 0.14 | 0.19 | 0.44 |
| 0.5 | 0.5 | 0 | 0.13 | 0.18 | 0.24 |
| 0 | 0 | 1 | 0.13 | 0.17 | 0.63 |
| 1 | 0 | 0 | 0.15 | 0.12 | 0.21 |
| 0 | 1 | 0 | 0.11 | 0.23 | 0.25 |
| 1/6 | 3/6 | 2/6 | 0.13 | 0.20 | 0.44 |
| 0.5 | 0 | 0.5 | 0.15 | 0.16 | 0.50 |



Genetic trends





Conclusions

- New index for Mastitis Resistance
- Equal weights for CM-F, CM-L and SCS
 - >2/3 on clinical mastitis and 1/3 on SCS
 - > Equal weight between clinical mastitis in first vs. later
- Expressed on RBV scale, mean of 100, SD of 5
 - > Higher value is desirable
 - At least 45 REL with 10 daughters in 10 herds
- Higher accuracy of selection for both clinical and subclinical mastitis





Use of BHBA for Evaluation of Health Traits in Canadian Holsteins





Milk ß-hydroxybutyrate (BHBA)

- Hyperketonemia or ketosis is one of the most frequent diseases in dairy cattle
- Level of milk
 ß-hydroxybutyrate (BHBA) is an indicator of subclinical ketosis
- Since October 2011 screening for hyperketonemia based on a BHBA analysis by MIR of test-day milk samples is offered in Canada by Valacta



Objective

 Estimate genetic parameters for milk BHBA in first lactation Holstein cows

 Determine genetic correlations between milk BHBA and metabolic diseases (clinical ketosis and displaced abomasum)



Mean milk BHBA





Proportion (%) of cows with a positive (milk BHBA \geq 0.20 mmol/L) test result





Analysis of milk BHBA

| Trait | DIM | Records, no. | Mean |
|----------------------------|--------|--------------|-------|
| BHBA ₁ , mmol/L | 5-20 | 20,845 | 0.115 |
| BHBA ₂ , mmol/L | 21-40 | 26,871 | 0.094 |
| BHBA ₃ , mmol/L | 41-60 | 27,404 | 0.075 |
| BHBA ₄ , mmol/L | 61-80 | 27,233 | 0.068 |
| BHBA ₅ , mmol/L | 81-100 | 26,811 | 0.067 |



Heritabilities and genetic correlations

| Trait | $BHBA_1$ | BHBA ₂ | BHBA ₃ | BHBA ₄ | BHBA ₅ |
|-------------------|----------|-------------------|-------------------|-------------------|-------------------|
| BHBA ₁ | 0.13 | 0.96 | 0.84 | 0.75 | 0.67 |
| BHBA ₂ | | 0.13 | 0.99 | 0.85 | 0.77 |
| BHBA ₃ | | | 0.16 | 0.98 | 0.96 |
| BHBA ₄ | | | | 0.22 | 0.99 |
| BHBA ₅ | | | | | 0.29 |



Associations between milk BHBA and metabolic diseases

- Milk BHBA at the first test-day (5-40 DIM)
- Ketosis
- Displaced abomasum

| Trait | Records, no. | Mean |
|------------------|--------------|------|
| BHBA, mmol/L | 7,635 | 0.10 |
| KET frequency, % | 3,437 | 3.61 |
| DA frequency, % | 6,894 | 2.74 |



Frequency of clinical ketosis and displaced of negative, suspect and positive tested cows





Heritabilities and genetic correlations

| Trait | BHBA | KET | DA |
|-------|-------------|-------------|-------------|
| BHBA | 0.13 (0.01) | 0.50 (0.26) | 0.21 (0.16) |
| KET | | 0.03 (0.03) | 0.63 (0.43) |
| DA | | | 0.05 (0.02) |



Conclusions

- Heritabilities for milk BHBA ranging from 0.13 to
 0.29
- Higher milk BHBA in early lactation was genetically associated with a higher frequency of clinical ketosis and displaced abomasum
- Milk BHBA can be routinely analyzed in milk samples on test-days, and, therefore, provide a practical tool for breeding cows with a lower susceptibility to hyperketonemia



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