



Relationships between subclinical ketosis, BCS, fat-protein-ratio and other diseases in Fleckvieh

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Background

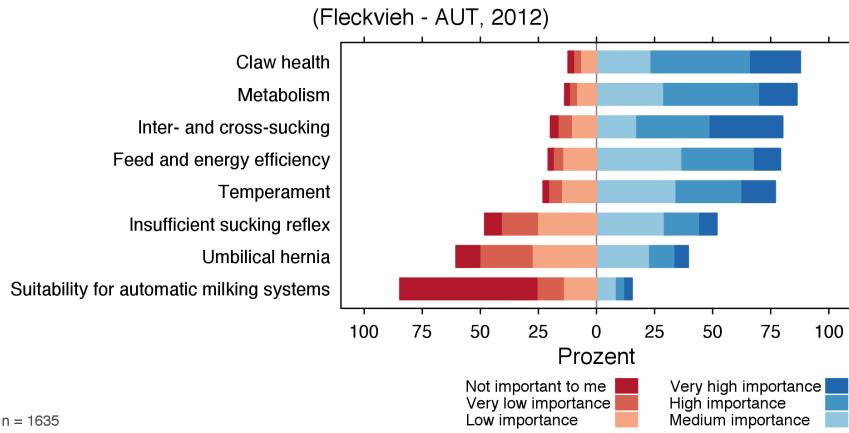
- In 2006, Austrian health monitoring project was launched (Egger-Danner et al., 2012)
- Since 2010, breeding values for mastitis, early fertility disorders, cystic ovaries and milk fever have been published (Fuerst et al., 2011)
- Metabolism is currently covered only by the trait milk fever



"Wish list" for new traits from farmers

– Fleckvieh (Steininger, 2013)

New traits





Approach – field data for novel traits

- Study based on data of Austrian project "Efficient Cow"
- Extended data recording on-farm on 161 farms in Austria with app. 6,500 cows for one year (2014)

Data recorded:

- general information about the farm
- various data related to health (e.g. veterinarian diagnoses, claw trimming, farmer observations, **subclinical ketosis**)
- feeding information
- body weight and body measures, linear scoring
- body condition score, lameness
- infrared-spectra

- ...



Aim

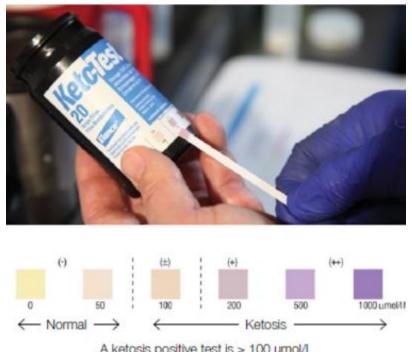
- Phenotypic and genetic analysis of subclinical ketosis
- Associations with BCS, ketosis, fat-protein-ratio and other diseases





Subclinical ketosis

Detected by using the milk Keto-test from ELANCO (measures milk ß-hydroxybutyrate) at days 7 and 14 after calving



A ketosis positive test is ≥ 100 µmol/l



Phenotypic analysis

- Keto-Test results 7 days after calving considered (N=1,920)
- Keto-Test results were grouped into 3 classes according to their milk ß-hydroxybutyrate:
 - healthy = $<100 \mu mol/l$
 - suspicious = 100-< $200 \mu mol/l$
 - subclinical ketosis = ≥ 200 μmol/l

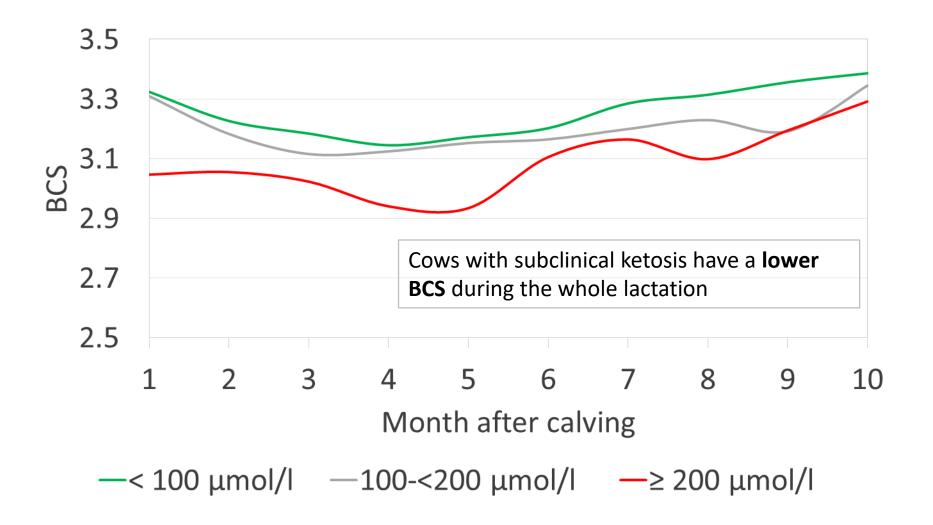


Frequency of subclinical ketosis (%)

Keto-test	Parity				
(N=1,920)	All	1	2	3+	
< 100 μmol/l	64.2	71.9	68.5	58.0	
100-<200 μmol/l	28.5	23.4	24.0	33.2	
≥ 200 µmol/l	7.3	4.7	7.5	8.8	

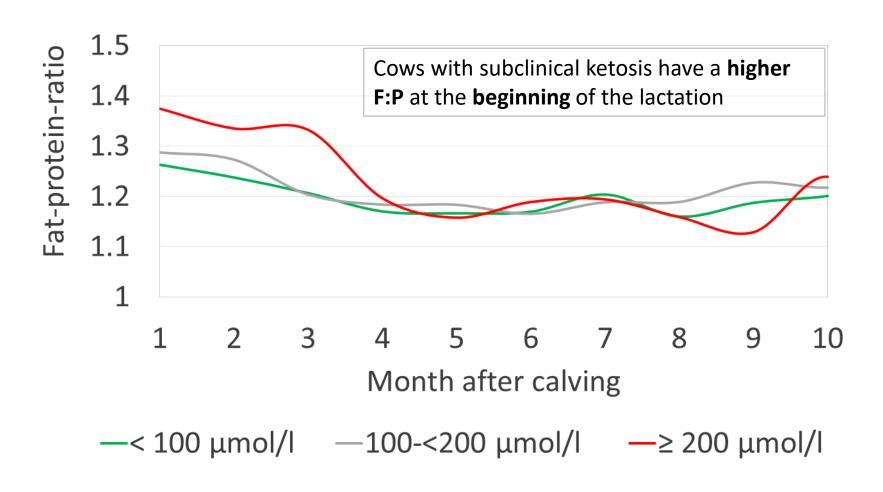


Subclinical ketosis and BCS



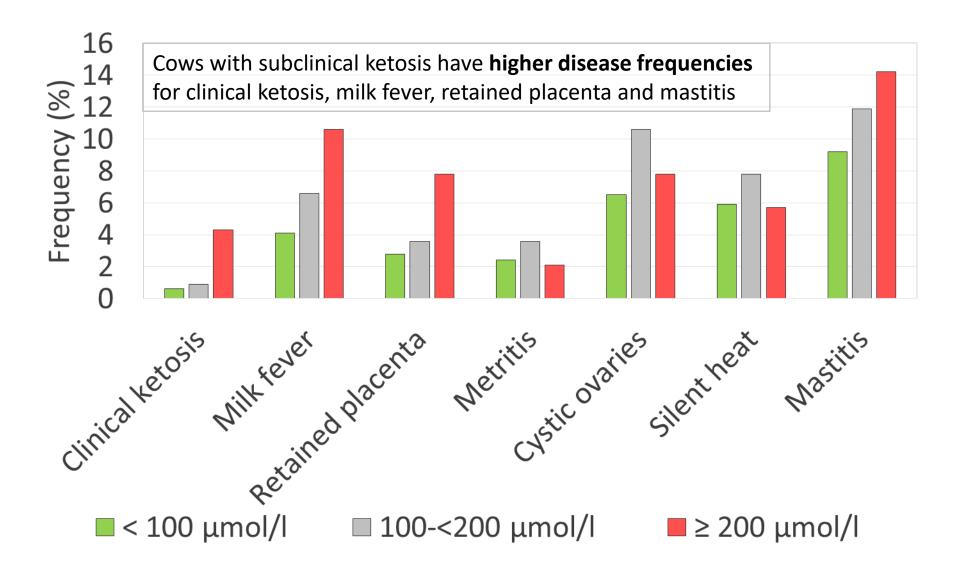


Subclinical ketosis and fat-protein-ratio





Subclinical ketosis and other diseases





Genetic analysis - Data

- Keto-test results 7 and 14 days after calving were combined (N=1,805); grouping as for phenotypic analysis (0/1/2)
- Ketosis (1-100 DIM) and Milk fever (-10 to 10 DIM) in years 2012-2014 (0/1)
- BCS and Fat:Protein ratio on 1st test-day



Summary statistics of data set

	Number of records	Mean
Subclinical Ketosis	1,805	0.56
Ketosis, %	5,670	0.70
Milk Fever, %	5,670	4.10
BCS	2,492	3.31
F:P	7,187	1.28



Genetic parameters (VCE6, Groeneveld et al., 2008)

	SUB KET	KET	MF	BCS	F:P
SUB KET	0.05 (0.029)				
KET		0.01 (0.006)			
MF			0.02 (0.027)		
BCS				0.17 (0.039)	
F:P	Heritabilities studies	in accordance t	to previous		0.14 (0.026)



Genetic parameters (VCE6, Groeneveld et al., 2008)

	SUB KET	KET	MF	BCS	F:P
SUB KET	0.05 (0.029)	0.89 (0.62)	0.51 (0.31)		
KET		0.01 (0.006)	0.79 (0.32)		
MF			0.02 (0.027)		
BCS				0.17 (0.039)	
F:P		between metal uding sublinical			0.14 (0.026)



Genetic parameters (VCE6, Groeneveld et al., 2008)

	SUB KET	KET	MF	BCS	F:P
SUB KET	0.05 (0.029)	0.89 (0.62)	0.51 (0.31)	-0.45 (0.22)	0.16 (0.26)
KET		0.01 (0.006)	0.79 (0.32)	-0.99 (0.17)	-0.33 (0.41)
MF			0.02 (0.027)	-0.61 (0.36)	0.43 (0.26)
BCS	Animals with lower BCS are more susceptible to metabolic disorders			0.17 (0.039)	-0.26 _(0.16)
F:P		P difficult to into	•		0.14 (0.026)



Conclusions

- BCS is a valuable tool for the prevention and early detection of metabolic diseases
- Metabolism with subclinical and clinical symptoms is complex
- Different information sources and traits could be used to improve metabolic disease resistance
- Development of Metabolism Index?



Acknowledgements

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Genetic analysis – Model (VCE6, Groeneveld et al., 2008)

Fixed effects/covariates:

- Year*season of calving
- Parity-age at calving
- Type of recording (for clinical ketosis and milk fever)
- Days in milk (for BCS, F:P)
- Day of first ketotest (for subclinical ketosis)
- Number of ketotests (for subclinical ketosis)

• Random effects:

- Herd*year
- Animal (genetic effect)
- Permanent environmental effect