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Published on: 03 Mar 2018 - bioRxiv (Cold Spring Harbor Laboratory)

Topics: Methanogenesis and Rumen

The Rumen Metatranscriptome Landscape Reflects Dietary Adaptation and Methanogenesis in Lactating Dairy Cows

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Running title: Methanogenesis in the rumen metatranscriptome

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Abstract

Methane eructed by ruminant animals is a main contributor to greenhouse gas emissions and is solely produced by members of the phylum *Euryarchaeota* within the domain *Archaea*. Methanogenesis depends on the availability of hydrogen, carbon dioxide, methanol and acetate produced, which are metabolic products of anaerobic microbial degradation of feed-derived fibers. Changing the feed composition of the ruminants has been proposed as a strategy to mitigate methanogenesis of the rumen microbiota.

We investigated the impact of corn silage enhanced diets on the rumen microbiota of rumen-fistulated dairy cows, with a special focus on carbohydrate breakdown and methanogenesis. Metatranscriptome analysis of rumen samples taken from animals fed corn silage enhanced diets revealed that genes involved in starch metabolism were significantly more expressed while archaeal genes involved in methanogenesis showed lower expression values. The nutritional intervention also influenced the cross-feeding between *Archaea* and *Bacteria*.

The results indicate that the ruminant diet is important in methanogenesis. The diet-induced changes resulted in a reduced methane emission. The metatranscriptomic analysis provided insights into key underlying mechanisms and opens the way for new rational methods to further reduce methane output of ruminant animals.

Keywords: rumen, cow, microbiota, methane, greenhouse effect, metatranscriptome, RNAseq, microbial ecology

41 **Introduction**

42 Reduction of global greenhouse gas (GHG) output is necessary to prevent a further increase in
43 global warming, which is predicted to result in multiple detrimental effects for the
44 environment and human affairs (Schleussner et al., 2016). The necessary measures are
45 focused on the industrial and agricultural sectors in developed countries, with the aim to
46 reduce carbon dioxide, methane and other GHG emissions. One of the predominant sources of
47 methane emission, estimated to be as high as ~35% of the total anthropogenic methane
48 emissions worldwide (IPCC, 2007;McMichael et al., 2007), is the agricultural sector, and
49 especially the eructation by ruminant animals (Murray et al., 2007).

50 Ruminal microbes play a pivotal role in the breakdown of animal feed and contribute between
51 35 to 50% of the animal's energy intake (Bergman, 1990). The ruminal microbial
52 composition is complex, with diverse populations including bacteria, archaea, fungi, and
53 protozoa. Their functional capacity is vast and has not yet been fully elucidated (Hess et al.,
54 2011;Li et al., 2012).

55 Notwithstanding the ruminal microbial complexity, methane is solely produced by a few
56 members of the phylum *Euryarcheota* belonging to the *Archaea* (Hook et al., 2010). It has
57 been shown that a change in diet can have a significant effect on the methane emissions of
58 ruminants (Liu et al., 2011;van Gastelen et al., 2015), but the mechanisms that drive this
59 change are not fully understood. The methanogenic archaea are not directly involved in the
60 breakdown of the feed, but rely on their relationships with other community members that
61 provide the necessary substrates for methanogenesis like hydrogen, formate and methanol.

62 Microbial ecology in cows and other ruminants has been investigated using 16S ribosomal
63 RNA (rRNA) genes as molecular markers (Fernando et al., 2010;Pitta et al., 2014), the sheep
64 rumen microbial metatranscriptome has been investigated (Shi et al., 2014), and in cows
65 specialized and general microbial functions have been examined (Brulc et al., 2009;Hess et

66 al., 2011;Poulsen et al., 2013;Dai et al., 2014;Dassa et al., 2014;Roehe et al., 2016;Jose et al.,
67 2017). Understanding the mechanisms that influence cow rumen methanogenesis requires
68 community-level analysis of active metabolic functions, however, a comprehensive analysis
69 of diet-dependent effects on the functional landscape of the rumen microbiota is lacking. Here
70 we investigated the effect of feed composition on bovine rumen activity patterns with a
71 special focus on methane metabolism. By analysis of the rumen metatranscriptome landscapes
72 in animals fed mixed grass silage (GS) and corn silage (CS) diets, we were able to elucidate
73 the impact of the diet on the expression of methanogenic pathways and on the relationships of
74 methanogens with other community members.
75
76 **Materials and Methods**
77
78 **Study design and sampling**
79 The study design has been described in detail by Van Gastelen et al. (van Gastelen et al.,
80 2015). Briefly, the experiment was performed in a complete randomized block design with
81 four dietary treatments and 32 multiparous lactating Holstein-Friesian cows. Cows were
82 blocked according to lactation stage, parity, milk production, and presence of a rumen fistula
83 (12 cows). Within each block cows were randomly assigned to 1 of 4 dietary treatments. All
84 dietary treatments had a roughage-to-concentrate ratio of 80:20 based on dry matter. In the
85 four diets, the roughage consisted of either 100% GS (**GS100**), 67% GS and 33% CS (**GS67**),
86 33% GS and 67% CS (**GS33**), or 100% CS (**GS0**; all dry matter basis). This study, including
87 the rumen fluid sampling, was conducted in accordance with Dutch law and approved by the
88 Animal Care and Use Committee of Wageningen University.
89
90 **Sample collection and processing**

91 In total, samples from 12 rumen fistulated cows, three per dietary treatment, were used for
92 metatranscriptome analysis. Rumen fluid was collected 3 hours after morning feeding on day
93 17 of the experimental period (for further details regarding the whole experimental period, see
94 (van Gastelen et al., 2015)). The samples were obtained as described previously (van
95 Zijderveld et al., 2011), and collected from the middle of the ventral sac. The rumen fluid
96 samples were immediately frozen on dry ice and subsequently transported to the laboratory
97 where the samples were stored at -80°C until further analysis.
98 For RNA extraction, 1 ml rumen fluid was centrifuged for 5 min at 9000 g, after which the
99 pellet was re-suspended in 500 µl TE buffer (Tris-HCl pH 7.6, EDTA, pH 8.0). Total RNA
100 was extracted from the resuspended pellet according to the Macaloid-based RNA isolation
101 protocol (Zoetendal et al., 2006) with the use of Phase Lock Gel heavy (5 Prime GmbH,
102 Hamburg) (Murphy and Hellwig, 1996) during phase separation. The aqueous phase was
103 purified using the RNAeasy mini kit (Qiagen, USA), including an on-column DNAseI
104 (Roche, Germany) treatment as described previously (Zoetendal et al., 2006). Total RNA was
105 eluted in 30 µl TE buffer. RNA quantity and quality were assessed using NanoDrop ND-1000
106 spectrophotometer (Nanodrop Technologies, Wilmington, USA) and Experion RNA Stdsens
107 (Biorad Laboratories Inc., USA).
108 rRNA was removed from the total RNA samples using the Ribo-Zero™ rRNA removal Kit
109 (Meta-Bacteria; Epicentre, Madison, WI, USA) using 5 µg total RNA as input. Subsequently,
110 barcoded cDNA libraries were constructed for each of the rRNA depleted samples using the
111 ScriptSeq™ Complete Kit (Bacteria; Epicentre) according to manufacturer's instructions in
112 combination with Epicentre's ScriptSeq Index PCR Primers.
113 The barcoded cDNA libraries were pooled and sent to GATC Biotech (Konstanz, Germany)
114 for 150 bp single end sequencing on one single lane using the Illumina HiSeq2500 platform in

115 combination with the TruSeq Rapid SBS (200 cycles) and TruSeq Rapid SR Cluster Kits
116 (Illumina Inc., San Diego, CA, USA).
117

118 **Bioinformatics**

119 The general workflow for data quality assessment and filtering was adapted from (Davids et
120 al., 2016). rRNA reads were removed with SortMeRNA v1.9 (Kopylova et al., 2012) and all
121 included databases. Adapters were trimmed with cutadapt v1.2.1 (Martin, 2011) using default
122 settings except for an increased error value of 20 % for the adapters. The latter was chosen
123 considering that with the default setting of 10% adapter sequences could still be found after
124 trimming. Quality trimming was performed with PRINSEQ Lite v0.20.0 (Schmieder and
125 Edwards, 2011) with a minimum sequence length of 40 bp and a minimum quality of 30 at
126 both ends of the read and as mean quality. All reads with non-IUPAC characters were
127 discarded as were all reads containing more than three Ns. Details on the RNAseq raw data
128 analysis can be found in Supplementary Table 1. The log files with the used commands can
129 be found in supplementary file 1 and the used python script in supplementary file 2. The raw
130 data was deposited at EBI ENA, and can be accessed under accession numbers ERS685245 -
131 ERS685256.
132

133 **Assembly and annotation**

134 All reads which passed the quality assessment were pooled and cross-assembled with
135 IDBA_UD version 1.1.1 with standard parameters (Peng et al., 2012). A second dataset was
136 added to the assembly to increase coverage (see supplementary materials & methods for
137 details on this dataset). Prodigal v2.5 was used for prediction of protein coding DNA
138 sequences (CDS) with the option for meta samples (Hyatt et al., 2010). Proteins were
139 annotated with InterProScan 5.4-47.0 (Hunter et al., 2012) on the Dutch Science Grid. The

140 annotation was further enhanced by adding EC numbers via PRIAM version March 06, 2013
141 (Claudel-Renard et al., 2003). Carbohydrate active modules were predicted with dbCAN
142 release 3.0 (Yin et al., 2012). Further EC numbers were derived by text mining and matching
143 all InterproScan derived domain names against the BRENDA database (download 13.06.13)
144 (Chang et al., 2015). Further details on the text mining can be found in the supplementary
145 materials & methods.
146 Reads were mapped back to the assembled metatranscriptome with Bowtie2 v2.0.6
147 (Langmead and Salzberg, 2012) using default settings. The resulting BAM files were
148 converted with SAMtools v0.1.18 (Li et al., 2009), and gene coverage was calculated with
149 subread version 1.4.6 (Liao et al., 2013). Read mappings to the contigs were inspected with
150 Tablet (Milne et al., 2013). The log files with the used commands for mapping and counting
151 can be found in supplementary file 1 and the used python script in supplementary file 2. The
152 whole read table including all annotations can be found in supplementary file 3.
153
154 **Taxonomic assignments**
155 All assembled contigs were analysed by blastn (Altschul et al., 1990) against the NCBI NT
156 database (download 22.01.2014) with standard parameters, except for an e-value of 0.0001,
157 and against the human microbiome (download 08.05.2014), the NCBI bacterial draft genomes
158 (download 23.01.2014), the NCBI protozoa genomes (download 08.05.2014), the human
159 genome (download 30.12.2013, release 08.08.2013, NCBI *Homo sapiens* annotation release
160 105) and the genomes of *Bos mutus*, *Bos taurus* and *Bubalus bubalis* (download 21.05.2014).
161 Taxonomy was estimated with the LCA algorithm as implemented in MEGAN (Huson et al.,
162 2011), but with changed default parameters. Only hits exceeding a bitscore of 50 were
163 considered, and of these only hits with a length of more than 100 nucleotides and that did not
164 deviate more than 10% in length from the longest hit.

165 For contigs, which did not retain any hits after the filtering described above, another run with
166 blastp of the associated proteins was performed against a custom download of the KEGG
167 Orthology (KO) database (download 25.04.2014). Taxonomic assignment was again
168 performed with the LCA algorithm, but only hits were considered, which did not deviate by
169 more than 10% from the hit with the maximal identity.
170 All taxa, which were attributed to the phylum Chordata, kingdom Viridiplantae or to artificial
171 constructs were considered to be contaminations and were automatically removed, as well as
172 any proteins in which the annotation contained the word "microvirus". Furthermore, contigs
173 that had a length of less than 300 nucleotides and which did not contain any proteins with a
174 functional domain (disregarding the coils database) were discarded. Contigs belonging to the
175 Illumina spike in PhiX phage were manually removed.
176 A compact schematic representation of the workflow is provided in Figure 1.
177
178 **Statistical analysis**
179 Differential expression was calculated in R version 3.1.1 (R Core Team, 2012) with the
180 edgeR package release 3.0 (Robinson et al., 2010). Only genes, which had at least 50 reads
181 mapped in all ten samples together were considered, and only genes with a p-value and q-
182 value <0.05 in any of the comparisons were considered to be significantly differentially
183 expressed. Furthermore, samples from cow #14 and #511 were excluded from the statistical
184 analysis, due to dermal antibiotic treatment and due to feeding aberrations. To examine
185 missing links within pathways, a q-value <0.1 was also considered (referred to as "lenient
186 approach"). The used input file, the R script with the commands, output tables and MA plots
187 can be found in supplementary file 4. To determine whether transcription levels corresponded
188 to the diet components, the differentially expressed genes were sorted for each gene by diet
189 group with increasing GS content, and an increasing or decreasing isotonic regression was fit

190 on the data. An R^2 value of ≥0.8 was considered to be indicative of an increasing or
191 decreasing profile, respectively, and all other values were considered to indicate that gene
192 expression followed another, irregular, profile. Regression values and assignment of profile
193 can be found in supplementary file 3. Isotonic regressions were computed in Python with
194 scikit-learn version 0.15.2 (Pedregoa et al., 2011). Spearman rank correlation between the
195 samples and Mann-Whitney U-test were calculated in Python with Scipy version 1.6.1 and
196 NumPy version 0.9.0 (van der Walt et al., 2011).
197

198 **Metabolic mapping**

199 All derived EC numbers were mapped with custom scripts onto the KEGG database
200 (Kanehisa et al., 2012) and visualized using Python Scipy version 1.6.1 and NumPy version
201 0.9.0 (van der Walt et al., 2011) together with matplotlib version 1.4.3 (Hunter, 2007).
202 Differentially expressed genes were investigated separately for microbial groups, which
203 showed changes over multiple genes per pathway, and changed functions were determined by
204 manual inspection of the KEGG maps.

205 **Availability of data and material**

206 All data has been deposited at the European Nucleotide Archive (ENA) under accession
207 numbers ERS685245 - ERS685256 and ERS710560 - ERS710568

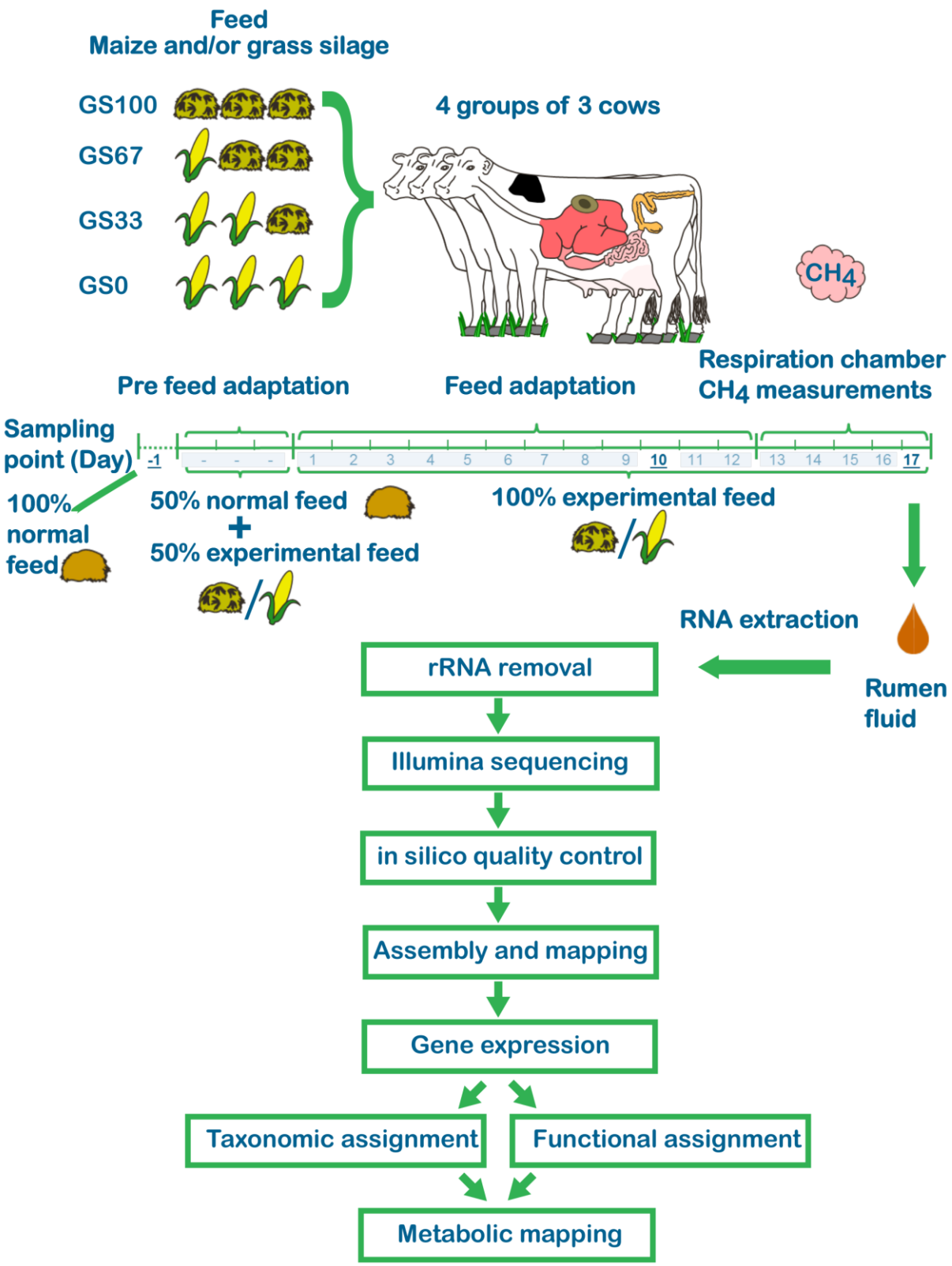
208

209 **Results**

210 Four experimental groups of three cows each were fed a control diet that contained only GS
211 as roughage, and three different CS-enhanced diets for twelve days (Figure 1). From day 13 –
212 17, methane emission was measured using a respiration chamber, showing a significant
213 reduction of methane emission with increasing CS proportion in the diet (van Gastelen et al.,

214 2015). This decrease accounted for approximately 10% of the cows' methane emission. The
215 analysis by van Gastelen et *al*. (van Gastelen et al., 2015) showed that the dry matter feed
216 intake of the different treatment groups did not differ significantly. Therefore the reduction in
217 methane emission was not based on the available energy, but rather on the composition of the
218 different diets.
219 Rumen fluid was collected at day 17, and used for microbial RNA extraction, mRNA
220 enrichment and RNAseq. The complete set of RNAseq reads was cross-assembled into a
221 single metatranscriptome. To determine activity per phylogenetic group the *de novo*
222 assembled transcripts/genes were assigned to a taxonomic rank, and relative expression levels
223 were obtained for four groups of animals fed different diets. Gene functional assignments
224 were subsequently used to assess potential metabolic changes as predicted from the gene
225 expression profiles observed in animals fed the four different diets, with a focus on
226 carbohydrate breakdown, short chain fatty acid (SCFA) production and methane metabolism.

227



228

229 **Figure 1: Study design.** Four groups of three cows were allowed to adapt to one of four

230 different experimental diets for twelve days. From day 13 – 17 methane emission was
231 measured using a respiration chamber. Rumen fluid was collected at day 17 and used for
232 microbial RNA extraction. See Methods section for details.

233

234

235 **Sequence, assembly and annotation metrics**

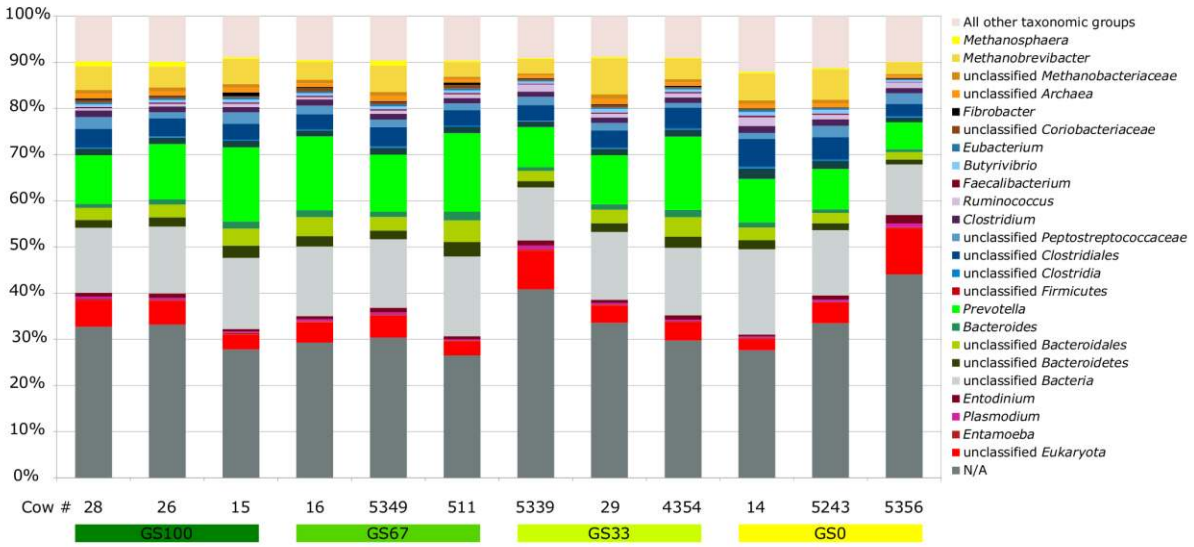
236 In total more than 160 million reads were obtained from twelve rumen fluid samples.
237 On average, 22.5% (Standard Deviation (SD) 6.15%) of all reads obtained per sample passed
238 all filtering steps, retaining 18.5% of the total raw reads. Of these filtering steps, the filtering
239 for rRNA sequences had the most impact, and removed the majority of the reads with an
240 average of more than 63% (SD 8.75%) (all details are given in Supplementary Table 1**).** The
241 majority of these rRNA reads (min. 96%) were matched to sequences from eukaryotes.
242 The assembly yielded 712,246 contigs with in total 866,052 protein coding sequences, a
243 length of 414,768,486 bp and an N50 of 596. While the longest contig had a size of 54,845
244 bp, most contigs (645,026, 90.1%) were smaller than 1000 bp. A total amount of 30 million
245 reads, on average 58% (SD 8.75%) of the reads per sample which passed quality filtering,
246 could be mapped back to the assembly (see Supplementary Table 1; in the following,
247 expression values will be given relative to the amount of mapped reads, referred to as "overall
248 expression").
249 For 556,705 of the predicted protein encoding sequences a domain (excluding "Coils"
250 domains) could be predicted. To 85,404 protein encoding sequences an EC number could be
251 assigned.
252 A taxonomic classification could be obtained for 635,892 protein encoding sequences (73%),
253 of which 282,074 could be classified at genus level. In total 1152 genera were detected, and
254 additional 190 taxonomic assignments above the genus level were retrieved. 24 groups (at

255 different taxonomic ranks) accounted for more than 58% of the total expression data (Figure
256 2). These groups included 13 genera (*Bacteroides, Butyrivibrio, Clostridium*, *Entamoeba,*
257 *Entodinium*, *Eubacterium*, *Faecalibacterium*, *Fibrobacter*, *Methanobrevibacter*,
258 *Methanosphaera*, *Plasmodium*, *Prevotella*, *Ruminococcus*) and 11 sequence clusters (not
259 including the data assigned to the 13 genera) that could only be assigned at higher taxonomic
260 levels (Archaea, Bacteria, Bacteroidales, Bacteroidetes, Clostridia, Clostridiales,
261 Coriobacteriaceae, Eukaryota, Firmicutes, Methanobacteriaceae, Peptostreptococcaceae).
262 Fungal genes could be detected, but accounted for less than 0.1% of the overall expression.
263 184,991 genes without a taxonomic assignment accounted for 32% of the total expression. To
264 only 34,731 of these genes (18.7%) any type of domain (excluding "Coils" domain) could be
265 assigned, and only 2685 of these had an EC number assigned. Most present domains within
266 the proteins encoded by taxonomically not assigned genes were generic domains (e.g.
267 membrane lipoprotein attachment site, MORN repeat, P-loop containing nucleoside
268 triphosphate hydrolase, WD40 repeat, etc.) without more specific functions.
269 Methanogens were represented by sequence assemblies that could be assigned to
270 *Methanobrevibacter smithii, Methanobrevibacter ruminantium* and *Methanosphaera*
271 *stadtmanae.* Reads mapping to protein coding genes assigned to methanogens captured on
272 average 6.2% of the overall expression. In general, the overall taxonomic expression profile
273 of the methanogens did not seem to change considerably between the different diets (Figure
274 2). When expression was summarized at genus level (or otherwise deepest taxonomically
275 assigned group, as given in Figure 2, with minor groups treated together as "all other
276 taxonomic groups"), the lowest correlation between all samples was 0.85. All microbial
277 groups included in Figure 2 were furthermore tested (after exclusion of cows 511 and 14, due
278 to mentioned aberrations) for statistically significant differences between animals fed the
279 different diets (Mann–Whitney *U* test, $p<0.05$, not multi-test corrected), which was rejected

280 for 150 out of 156 tests. None of the differences were statistically significant after multi-test

281 correction (Bonferroni).

282



283

284 **Figure 2: Taxonomic composition of metatranscriptome landscapes observed in animals**

285 **fed one of four different diets.** Diets and cows are indicated on the X-axis, taxonomic

286 groups (at genus level, or otherwise deepest classification) are colour coded, see legend for

287 details. N/A: No taxonomic rank could be assigned.

288

289 **Differential expression analysis of the rumen microbiomes**

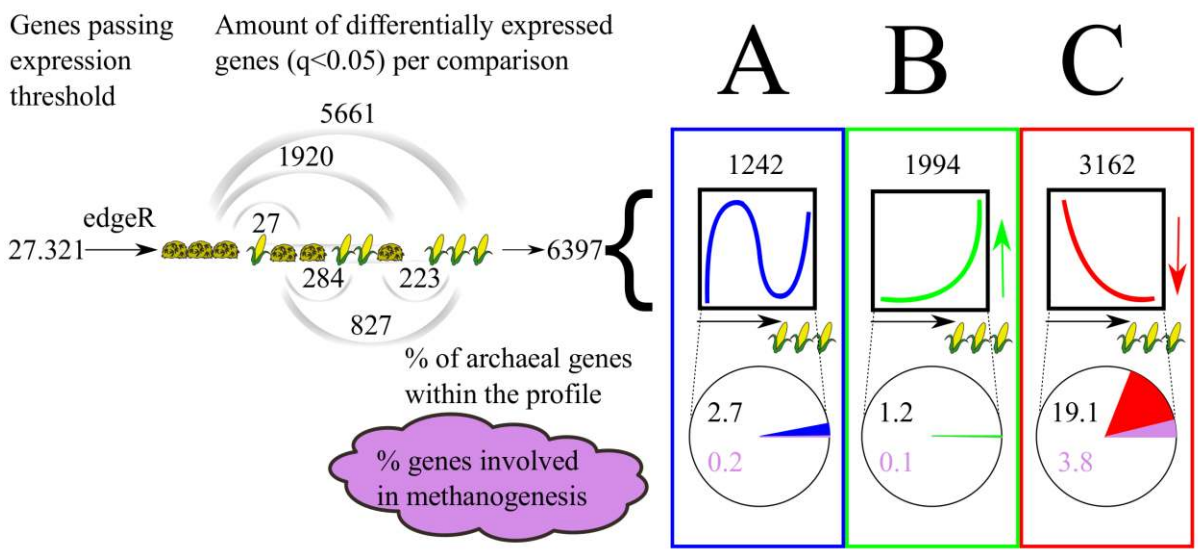


Figure 3: Differential expression analysis of the rumen microbiomes. Overview of the number of genes that were found to be differently expressed in pairwise comparisons of metatranscriptomes derived from animals fed different diets. Three profiles are distinguished: Profile A, genes with an expression, which does not follow a dietary pattern. Profile B, genes which are upregulated with increasing amounts of corn silage (CS). Profile C, genes, which are downregulated with increasing amounts of CS. Furthermore the results show that with the increase of CS, archaeal genes were mainly downregulated considerably affecting methane metabolism.

In total, 27,731 genes, which passed a set threshold for having captured at least 50 reads over all conditions combined, were subjected to the differential expression analysis, and 6397 were differentially expressed in at least one comparison (q<0.05). Three corn silage (CS) enhanced diet-induced expression profiles were distinguished (via regression analysis with isotonic regression), i.e. genes with an unlinked expression profile (profile A, 1241 genes), an induced expression corresponding to the amount of CS in the diet (profile B, 1994 genes), and a

308 reduced expression corresponding to the amount of CS in the diet (profile C, 3162 genes)
309 (Figure 3). Three heatmaps of all genes (per profile) can be found in supplementary file 5,
310 displaying the overall trends within the data.
311

312 **Taxonomic and functional analysis of the three diet induced expression profiles**

313 Genes grouped into the three different expression profiles were investigated for their
314 taxonomic and functional classification.
315 For profile A, i.e. genes that did not follow a diet specific expression profile, most genes were
316 related to general energy metabolism/carbohydrate breakdown and ribosomal protein
317 production, as well as transport reactions. No other major functions seemed to be affected in
318 the diet-unspecific way characteristic of profile A, and most of the genes within this group
319 could be linked to the Clostridiales, but also to Bacteroidales, Actinobacteria and Archaea.
320 Most predominantly represented taxa among genes following transcription profile B were
321 bacteria belonging to the order Clostridiales, and to a lesser extent the genera Prevotella,
322 Proteobacteria and Actinobacteria, but more than half of the differentially expressed genes
323 could not be classified below kingdom level. The most affected functions were ribosomal
324 protein production (mainly Eukaryota), and nucleotide metabolism in different groups,
325 including the Eukaryota. The almost complete lack of genes associated with *Archaea* and/or
326 methanogenesis among the genes with expression profile B indicated that there was hardly
327 any increase of methanogenic activity with the increase of CS.
328 Among the genes exhibiting a lower expression upon increasing the amount of CS in the diet
329 (profile C), the main represented microbial groups included three different methanogens
330 (*Methanobrevibacter smithii*, *Methanobrevibacter ruminantium*, *Methanosphaera*
331 *stadtmanae*), members of the genus *Prevotella*, and many genes, which could not be classified
332 beyond the order Clostridiales. Functional profiling showed that the most downregulated

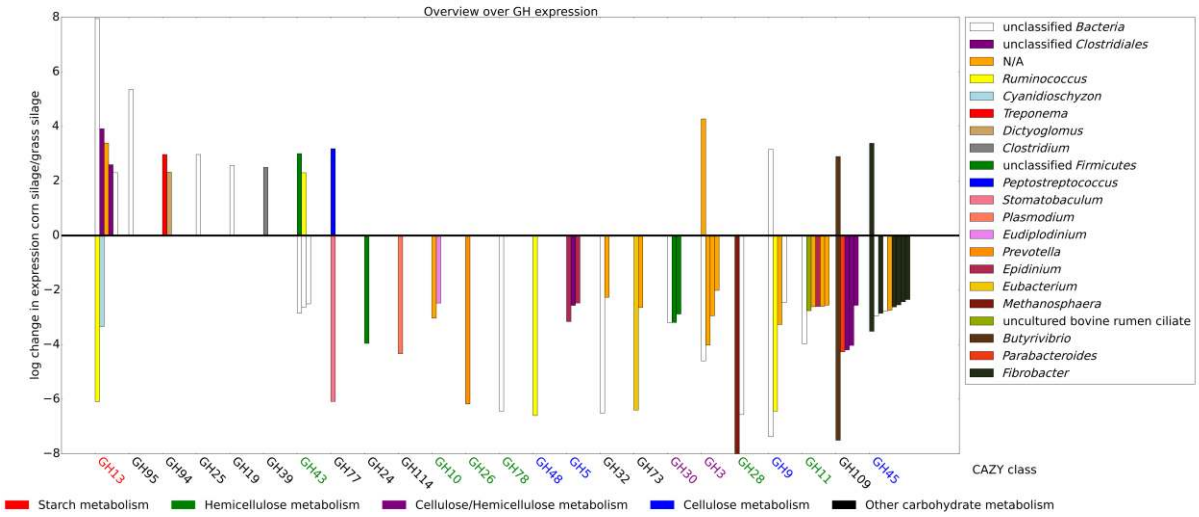
333 processes were related to methanogenesis, electron transport and regulatory processes in the
334 *Archaea*, as well as general metabolic functions like glycolysis, ATP generation or ribosomal
335 protein production in all affected groups. Increased expression could also be observed for nine
336 genes encoding putative non-ribosomal peptide synthase (NRPS) modules, among which
337 three were taxonomically linked to *M. ruminantium* whereas the other six NRPS modules
338 could not be classified beyond the kingdom bacteria.
339 With an increase of CS in the diet, Eukaryota appeared to show a decrease in their expression
340 of genes encoding glycosylhydrolases (GH) and glycosyltransferases (GT). Furthermore, they
341 also showed differential regulation of genes associated with movement abilities and
342 cilia/cytoskeleton assembly, chaperons and ribosomal proteins in response to the diet changes.
343 Most of the sequences (71.9%) assigned to the Eukaryota could not be classified below the
344 kingdom level. For example, of the 85 differentially expressed genes encoding proteins
345 involved in cilia/cytoskeleton assembly, only 12 could be assigned to a rank more specific
346 than the kingdom level. Within all the classified eukaryotic sequences that showed consistent
347 downregulation with increasing CS in the diet, the phylum Apicomplexa was the most
348 represented, whereas the family of Ophyoscolecidae (*Entodinium, Epidinium*) showed a
349 specific downregulation of GH encoding genes.
350

351 **Microbial starch and cellulose metabolism in cows fed with different diets**

352 The expression of genes related to the breakdown of different complex carbohydrates differed
353 considerably between animals fed different diets. Profile A did not include major changes in
354 genes coding for carbohydrate degradation associated enzymes.
355 For genes following expression profile B, an increase of CS in the diet mainly lead to the
356 increased expression of genes encoding different extracellular binding proteins in the genera
357 *Ruminococcus, Bifidobacterium* and *Entodinium*, as well as an increase in the expression of

358 genes coding for starch binding modules (CAZy classes CBM25 and CBM26) and alpha-
359 amylases (GH13).
360 Most carbohydrate-metabolism associated genes affected by an increase in CS in the diet,
361 however, followed expression profile C. With an increase of the CS in the feed, a
362 downregulation of multiple genes involved in the breakdown of plant cell walls and their
363 constituents could be observed, such as all the steps involved in cellulose degradation (Lynd
364 et al., 2002). Expression of genes encoding endocellulases (CAZy classes GH5, GH9, GH45;
365 mainly assigned to *Fibrobacter*), catalysing the first step of cellulose breakdown, was most
366 affected, followed by genes that code for exocellulases (GH48, *Ruminococcus*) and beta-
367 glucosidases (GH3), catalysing the second and the last step of cellulose breakdown,
368 respectively, as well as genes encoding cellulose binding modules (e.g. CBM4, CBM13).
369 Downregulation of the expression of genes encoding proteins involved in the breakdown of
370 hemicellulose constituents (xylan, mannan, galactan/pectate, rhamnose) could also be
371 observed, including genes encoding endo-1,4-beta-xylanases (GH10, GH11), beta-mannanase
372 (GH26), pectate lyase (PL3), alpha-L-rhamnosidase (GH78), beta-1,4-galactan binding
373 (CBM61), and xylan binding modules (e.g. CBM35). Expression of genes related to transport
374 of glucose into the cells was also downregulated (monosaccharide transporters, EC 3.6.3.17).
375 An overview of differentially expressed genes encoding glycosylhydrolases and
376 carbohydrate-binding modules, including their taxonomic distribution, is presented in Figure
377 4 and Figure 5, respectively.

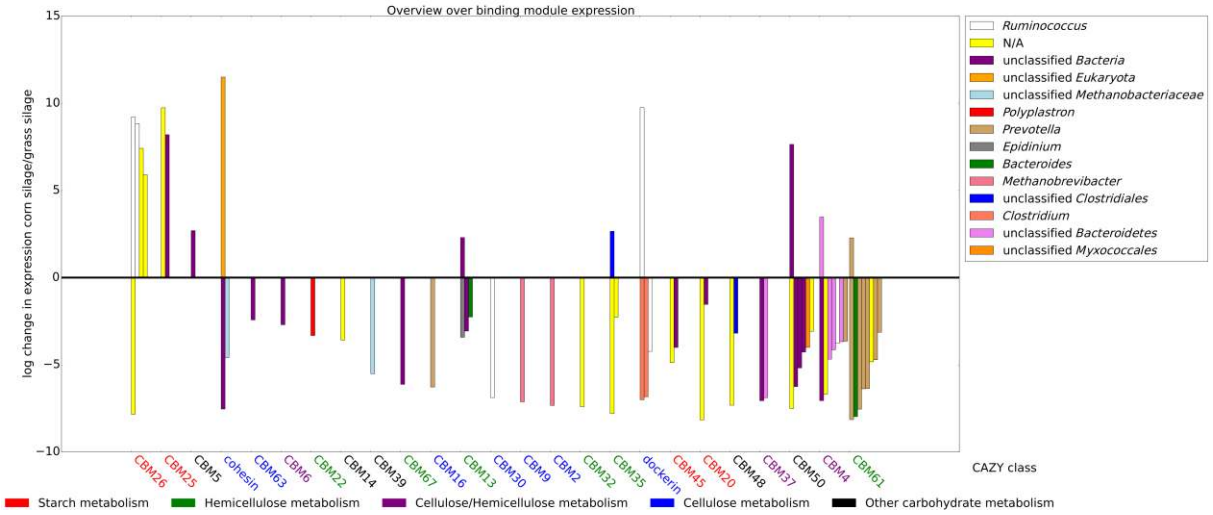
378



379

380 **Figure 4: Log10 fold changes in expression of differentially expressed glycosylhydrolase**
381 **encoding genes in a comparison of the 100% corn silage diet (GS0) versus the 100%**
382 **grass silage diet (GS100).** Positive values indicate an upregulation of gene expression in the
383 corn silage diet. N/A: No taxonomic rank could be assigned. Colour-coding of bars indicate
384 different taxonomic groups, whereas colour-coding of protein families indicate their
385 involvement in the metabolism of different carbohydrates.

386



387

388 **Figure 5: Log10 fold changes in expression of differentially expressed carbohydrate**
389 **binding module encoding genes in a comparison of the 100% corn silage diet (GS0)**
390 **versus the 100% grass silage diet (GS100).** Positive values indicate upregulation of gene
391 expression in the corn silage diet. N/A: No taxonomic rank could be assigned. Colour-coding
392 of bars indicate different taxonomic groups, whereas colour-coding of protein families
393 indicate their involvement in the metabolism of different carbohydrates.

394

395

396 With the increase of CS, a downregulation (profile C) could be observed for *susC* and *susD*
397 genes coding for starch binding proteins, and which could be assigned to the phylum
398 Bacteroidetes, mainly in the genus *Prevotella*. A downregulation of expression of genes
399 encoding proteins involved in cellulose binding was also found, including e.g. sortases,
400 cohesins, dockerins, extracellular binding and calcium binding domains, which potentially
401 could belong to a cellulosome (Bayer et al., 2008;Flint et al., 2008). This was mainly
402 observed for genes assigned to the families Cellulomonadaceae, Clostridiaceae,
403 Lachnospiraceae and Ruminococcaceae. Many functionally similar downregulated protein-
404 coding genes could not be assigned to a taxonomic rank below the superkingdom level,
405 mainly in the bacteria. The downregulation of a gene encoding a cohesin module was also
406 detected in the *Archaea*, as well as the upregulation in the expression of a cohesin and
407 dockerin module with an increase of CS.

408

409 **Microbial short chain fatty acid metabolism in cows fed different diets**

410 The production of SCFAs is an important function of the rumen microbiome. These
411 metabolites are taken up by the host and serve as an energy source (Bergman, 1990), have a
412 considerable effect on methane production (van Kessel and Russel, 1996), and affect the pH,

413 which in turn has an influence on the animal's wellbeing (Kleen et al., 2003). In the study by
414 Gastelen et *al*., only a small significant reduction in the SCFA butyrate was reported, with
415 other SCFAs not changing significantly.
416 Increased expression upon an increase of CS in the diet (profile B) was found for genes
417 coding for proteins which are involved in the conversion of acetyl-CoA to crotonyl-CoA,
418 which is part of butyrate synthesis. This increase was found within the family
419 Lachnospiraceae. The total expression of this family was on average 1.9% in all samples.
420 Reduced expression upon an increase of CS in the diet (profile C) was observed for genes
421 encoding proteins involved in butyrate metabolism, and again mainly for genes assigned to
422 the Lachnospiraceae. Several of the downregulated genes encode proteins catalysing the
423 reactions from pyruvate to crotonyl-CoA, via acetyl-CoA, acetoacetyl-CoA and (S)-3-
424 hydroxybutanoyl-CoA. Genes that code for enzymes catalysing the last steps to butyrate via
425 crotonyl-CoA and butanoyl-CoA were also present in the assembly, but were not found to be
426 differentially expressed in any of the conditions. Thus, the here presented data provide an
427 inconclusive picture regarding the regulation of genes encoding proteins involved in ruminal
428 butyrate production. Furthermore, consistent differential expression patterns could also not be
429 observed for genes involved in the formation of the other SCFAs acetate or propionate. Genes
430 encoding SCFA transporters were present in the assembly, but were not differentially
431 expressed. Overall, these observations are in line with the fact that total SCFA concentration
432 was found to be not affected by increasing CS in the diet, with only a minor, albeit significant
433 increase in the molar proportion of butyrate (van Gastelen et al., 2015).
434

435 **Expression of archaeal genes involved in methane metabolism**

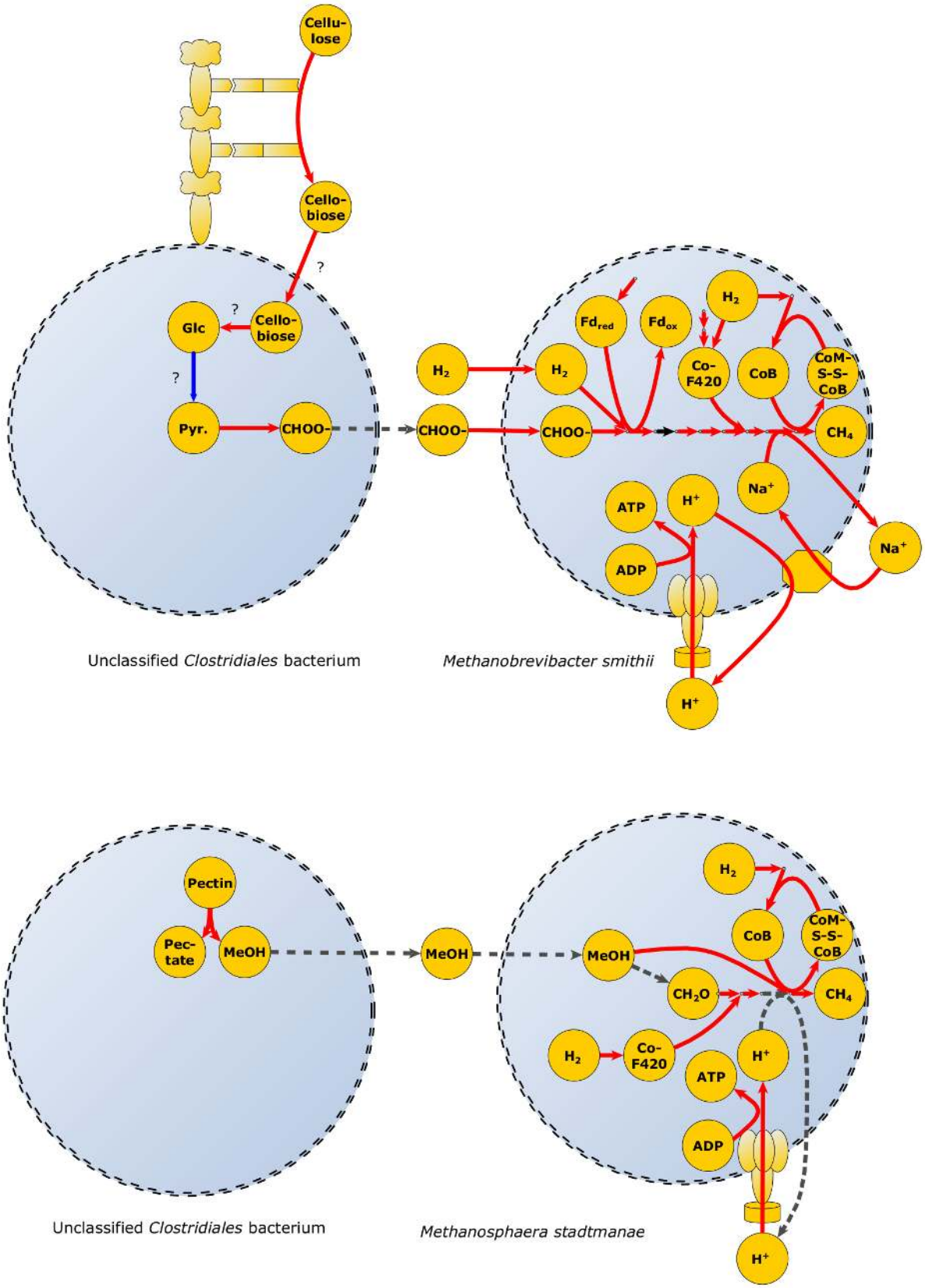
436 A considerable amount of differentially expressed genes in the *Archaea* was found to encode
437 proteins involved in methane metabolism. Based on the RNAseq data almost the complete

438 pathways leading to methanogenesis could be reconstructed (Fig. 6). Closer inspection
439 revealed that with an increase of CS in the diet, nearly all genes of the methanogenesis
440 pathways were downregulated in a subset of the *Archaea* (expression profile C). Of the four
441 possible methanogenic pathways, those for the production of methane from
442 methanol/hydrogen, as well as from formate/carbon dioxide and hydrogen were affected.
443 Proteins for the utilization of trimethylamines into methane could be detected in the dataset,
444 but were not differentially expressed between animals fed the different diets. The pathway for
445 methanogenesis from acetate was absent in the dataset.
446 Among genes assigned to *Methanosphaera stadtmanae,* genes coding for proteins involved in
447 the conversion of methanol to methyl-CoM (methanol-corrinoid protein Co-
448 methyltransferase, EC 2.1.1.90) and of methyl-CoM to methane (methyl-CoM reductase, EC
449 2.8.4.1) showed consistent downregulation with increasing CS in the diet (Figure 6).
450 Compared to changes observed for *Methanosphaera stadtmanae*, the change in the
451 transcription pattern of genes encoding proteins involved in methanogenesis from hydrogen
452 and formate in *Methanobrevibacter smithii* was more extensive. More specifically, the
453 expression of genes associated with the methanogenesis pathway with formate/hydrogen was
454 downregulated in nearly all steps (besides formylmethanofuran-tetrahydromethanopterin N-
455 formyltransferase, EC 2.3.1.101), following expression profile C. In addition, expression of
456 several genes encoding proteins involved in the biosynthesis of coenzyme F420 was
457 downregulated with an increasing amount of CS in the diet (profile C). Some of these
458 reactions could only be assigned to taxonomic levels above species, but the placements of
459 these functions in the metabolic network indicate that they most probably can also be assigned
460 to *M. smithii.*

461 Expression of genes encoding transporters for formate uptake were also downregulated
462 (profile C), as well as genes involved in other processes related to methanogenesis, e.g. the
463 general production of ATP, electron transport via the membrane, and sodium transport.
464 Nearly none of the genes that could be assigned to the third detected major methanogen in the
465 dataset, *Methanobrevibacter ruminantium*, showed considerable downregulation, however, it
466 should be noted that several archaeal genes, including several genes encoding proteins
467 involved in methanogenesis, could not be classified at the species level and therefore it cannot
468 be excluded that some of these in fact also belong to this species. Differential regulation of
469 genes assigned to a potential syntrophic partner of *M. ruminantium, Butyrivibrio*
470 *proteoclasticus* (Leahy et al., 2010), could only be detected in a few genes. Genes assigned to
471 other formate producing organisms were also present in the data, pointing towards their
472 potential involvement as syntrophic partners, however, no differential expression was
473 observed for these genes, making deduction of possible syntrophic connections difficult.
474 Further analysis of the data at the functional level showed downregulation of the expression
475 of genes encoding proteins linked to the production of necessary substrates for
476 methanogenesis. Expression of one of the genes encoding a subunit of pyruvate formate lyase
477 (EC 2.3.1.54) that catalyses the production of formate from pyruvate was downregulated in a
478 bacterium in the order Clostridiales, which could not further be classified, as well as in
479 *Eubacterium hallii*. At the same time, several genes encoding proteins involved in the
480 degradation of cellulose were found downregulated in animals fed CS-containing diets
481 (profile C), and could be assigned to a not further classifiable bacterium in the order
482 Clostridiales as well as *Ruminococcus flavefaciens, Fibrobacter succinogenes*, and several
483 other bacteria/eukaryotes. A downregulation of genes that code for proteins involved in the
484 production of the other substrates needed for methanogenesis, hydrogen and carbon dioxide,
485 could not be detected.

Interestingly, using a more lenient approach (see Methods) a downregulation of expression of a gene for the production of the second major substrate for methanogenesis, methanol, was observed. More specifically, an unspecified Clostridiales bacterium showed decreased expression of a gene encoding pectinesterase (EC 3.1.1.11), catalysing the degradation of pectin to pectate and methanol.

An overview of the metabolic consequences of the observed changes in gene expression profiles is provided in Figure 6. A version of this figure with more details and a table with all reactions and assigned genes can be found in supplementary figure 1 and supplementary file 6.



496

497 **Figure 6: Graphical summary of metabolic consequences of the different diets in the two**

498 **major methanogens and possible syntrophic partners.** Red arrows: genes downregulated

499 with the increase of corn silage in the diet; black arrows: Gene is detected but not
500 differentially expressed; The blue arrow represents glycolysis of which the majority could be
501 detected; punctuated arrow: orphan reactions; ? = Phylogenetic association unclear. Pyr. =
502 Pyruvate, $CHOO^-$ = Formate, FD_{ox} = oxidized Ferredoxin, FD_{red} = reduced Ferredoxin, CH_2O
503 = Formaldehyde.

504

505

506 **Discussion**

507 **How feed affects methanogenesis**

508 The rumen microbiome is a complex ecosystem, and its dynamics are determined by many
509 variables. Most investigations to date have been focussed on the community composition and
510 changes therein in response to different perturbations. In a recent metagenomic study by
511 Roehe *et al*. (Roehe et al., 2016) on animals fed similar diets as the ones tested here the
512 authors found no considerable effect on the composition of the microbiome. Here we show
513 that in response to a diet change, gene expression within a microbiome and consequently the
514 metabolic profile may change. Differential expression analysis revealed that although there
515 were no extensive changes visible within the overall community expression, in line with what
516 has previously been noticed for the sheep rumen (Shi et al., 2014), major effects could be seen
517 regarding the expression of genes related to methane metabolism, which are also in agreement
518 with genes which were prior identified within the metagenomics dataset by Roehe et *al*. and
519 related publications (Wallace et al., 2015;Roehe et al., 2016;Wang et al., 2017). In two of the
520 three methanogens identified in the dataset a coordinated downregulation of genes involved in
521 methanogenesis as response to increased CS in the diet could be observed. Thus not only
522 isolated single nodes involved in methanogenesis, but whole pathways were downregulated.
523 We further found evidence for a possible syntrophy between these methanogens and several

524 yet unidentified members of the rumen community belonging to the order of Clostridiales,
525 which might contribute to the production of the necessary substrates (formate, methanol) for
526 the methanogens, which was also discussed (albeit with potentially different syntrophy
527 partners) in a related setup by Parmar et *al*. (Parmar et al., 2017). Additionally we observed a
528 downregulation of cellulose degradation functions with increased CS in the diet. For *M.*
529 *ruminatium*, we did not see a significant response to the diet changes nor did we see a
530 significant response in possible syntrophic partners. Thus it may be that in addition to diet
531 changes other types of biological effectors are necessary to further influence the process. Our
532 findings are also in contrast to those reported by Shi *et al*.(Shi et al., 2014), who concluded
533 that in the sheep rumen the supply of hydrogen is the determining factor for methane output,
534 whereas in the present study the supply of other substrates seem to have a bigger influence.
535 We further observed community wide responses to the change in the main energy/carbon
536 source, with a shift in the involved glycosylhydrolases over multiple organisms and
537 phylogenetic branches. Nevertheless, we did not observe a response in all members of the
538 microbial community. While there was a definite downregulation of certain processes like
539 methanogenesis, these processes were not affected in all organisms. To this end, it should be
540 noted that the total gene count assigned to members of the Archaea greatly exceeded the size
541 of currently known individual archaeal genomes, suggesting the presence of multiple strains
542 of the same species in this environment (Hudman and Gregg, 1989;Sasson et al., 2017). Not
543 all of these strains seemed to be affected by the different diets, as there were also instances of
544 pathways, which did not show a differential regulation at all. As already observed here for the
545 different species of methanogens, which were potentially affected because their syntrophic
546 partners were affected, this could also be the case for the different strains of the same species,
547 which might inhabit different niches in the rumen. It cannot be expected that e.g.
548 methanogens living intracellularly within protozoa (Finlay et al., 1994) are in the same way

549 affected as free living methanogens are, and that populations living closer to the substrates,
550 i.e. those associated with the fibre fraction, will show the same behaviour as populations in
551 the liquid fraction of the rumen (Mullins et al., 2013). Finally, as overall a reduction of
552 methane production by ~10% was observed in this study when comparing animals fed either
553 the GS or CS diets, it is perceivable that not all pathways and microorganisms are affected to
554 an extend that would be detectable in significant differences in gene expression levels, also
555 considering the relatively small sample size of three animals per experimental group.

556 **Unexpected findings and limitations**

557 Several findings in this study were surprising, at least at first glance.
558 As shown in Figure 2, and also shown by the statistical testing, the overall expression profile
559 did not change significantly. A major change in the supplied feed was expected to result in
560 significant changes though. Also the study of Roehe *et al*. (Roehe et al., 2016) showed no
561 considerable changes in the relative abundance of organisms in a similar setting. We showed
562 that the main changes are not within a taxonomic group, but rather the expression patterns per
563 taxonomic group, which also explains the findings by (Roehe et al., 2016).
564 There are also concerns that differential expression analysis in communities could not reflect
565 actual differential expression, but rather a change in organism abundance, leading to wrongly
566 perceived changes in expression. Since in this dataset the overall expression profile per group
567 did not statistically significantly change (although the small sampling size gives only limited
568 power to detect this change), this is likely not an issue, and genes detected as differentially
569 expressed are probably truly differentially expressed.
570 The overall taxonomic composition itself as shown in Figure 2 in general agrees with
571 previous findings, as most of the major taxonomic groups were reported previously (Jami et
572 al., 2013). This is also the case for the methanogens, which are similar to the ones commonly
573 found the rumen of cows (Carberry et al., 2014) and other ruminants (Li et al., 2014;Seedorf

574 et al., 2015). Despite this, it should be noted that the genes assigned to *Methanobrevibacter*
575 *smithii* most likely belong to a related species/group of *Methanobrevibacter*, since *M. smithii*
576 itself is not a dominant member of the rumen microbiota, but the closest sequenced relative of
577 the species appearing in the rumen (Janssen and Kirs, 2008).
578 As shown in Figure 3, we also recovered changing expression profiles, which did not
579 correspond with the diets. We were not able to find any specific functional background for
580 these profiles, and suspect that some organisms are influenced more by the surrounding
581 community members and not primarily by the diet, or maybe inhabit very specific niches.
582 This would be in agreement with the findings in Figure 4 and 5, which show that a minor
583 amount of carbohydrate active enzymes and binding modules show expression profiles
584 against the expected trend, e.g. increase in expression of some cellulose degrading enzymes
585 while less cellulose is fed (Özcan and Johnston, 1999;Sloothaak et al., 2015). It could also be
586 possible that this change in expression reflects a change in metabolic strategy. As response to
587 e.g. the lower abundance of cellulose in the environment, the affected organisms could
588 attempt to downregulate the expression of genes coding for cellulose binding modules with
589 low affinity, and upregulate the expression for genes coding for modules with high affinity.
590 This mechanism is similar to the regulation of carbohydrate transporters in different
591 organisms (Özcan and Johnston, 1999;Sloothaak et al., 2015). Additionally it needs to be
592 considered that initial annotations might not always be correct. We found an increase in
593 cohesin and dockerin coding modules with an increase of starch in the diet. These
594 components are primary known as cellulosome components, but non-cellulosomal origin of
595 these modules has been reported before (Peer et al., 2009;Ze et al., 2015). Furthermore one of
596 these modules was found in the Archaea, which are not known to harbour either cellulosomal
597 complexes or their starch counterparts. The same issue holds for the downregulation in
598 expression of the genes coding for different starch binding proteins, *susC* and *susD*, which

599 have been found to not only be starch binding, but also cellulose binding (Mackenzie et al.,
600 2012).
601 Another finding, which was obvious in the investigated data, is the substantial decrease in
602 expression of genes coding for proteins involved in cytoskeleton assembly in different
603 Eukaryota. As several Archaea are endosymbionts of Protozoa, it can be speculated that an
604 experimental change, which has an impact on the symbionts, will also affect their host (Finlay
605 et al., 1994) (although this relationship is also not entirely clear (Morgavi et al., 2012)).
606 General cellular processes, like replication, in which the cytoskeleton is involved, will then
607 probably be directly affected, and this has been observed before in a different setting with
608 intracellular Archaea (Holmes et al., 2014). Recently the high abundance of these proteins in
609 the rumen proteome also have been demonstrated (Snelling and Wallace, 2017).
610 At last, the biggest limitation on this study are the lack of sequencing depth and little
611 replication. The former was mainly caused by the inefficiency of the ribosomal rRNA
612 depletion. The method used could not remove all rRNA, due to the diversity of unknown
613 eukaryotic sequences, which resulted in a lower sequencing depth than expected. Also due to
614 the low number of replicates, an arbitrary cutoff for the tested genes had to be applied, which
615 is common practice and can help in some settings to increase power (Bourgon et al., 2010),
616 and therefore it was not possible to find more subtle changes in the expression levels (e.g. a
617 change in transcription levels of the butyrogenic pathway). Therefore this work mainly
618 focused on changes within more highly expressed genes, and most changes were also not
619 dependent on single p-values, but supported by expression changes in multiple genes. This
620 has still lead to the ability to track the impact of diet on methane production, which was the
621 aim of this study, and other effects, which were not initially expected, could still be observed.
622 It still needs to be pointed out that the amount of replication was very small and probably too

small for this type of experiment, and that many changes, including not only subtle ones, were potentially missed due to this setup.

In summary, in this study we found a significant effect of a dietary change on the gene expression in the cow rumen. A substantial fraction of the affected genes was related to methane emission, showing that a decrease in cellulose in the diet decreased the gene expression of methane related pathways. The here presented metatranscriptomic analysis is in agreement with the experimental measurements, which showed a decrease in methane emissions with the diet change (van Gastelen et al., 2015), suggesting that a change in the feed regime can have a positive effect on microbial GHG emissions.

Acknowledgements

The authors thank Jasper Koehorst for his help with the transcriptome annotation, and Jesse van Dam, Maria Suarez-Diez and Edoardo Saccenti for helpful discussions.

This computational work was carried out on the Dutch national e-infrastructure with the support of SURF Foundation.

Author Contributions

BVDB, CMP and HS conceived designed the study. BVDB performed the experiments. BH and MD performed the bioinformatics processing and analysed the data. BH, BVDB, MD, VAPMDS, CMP, PJS and HS interpreted the results. BH, BVDB and PJS wrote the

645 manuscript with input from the other authors. All authors read and approved the final
646 manuscript.

647 **Conflict of interest statement**The authors declare not conflict of interest.

648 **Funding**

649 B. Hornung was supported by Wageningen University and the Wageningen Institute for
650 Environment and Climate Research (WIMEK) through the IP/OP program Systems Biology
651 (project KB-17-003.02-023).
652 This work was furthermore supported by funding from the Top Institute Food and Nutrition,
653 Wageningen, The Netherlands, a public-private partnership on pre-competitive research in
654 food and nutrition. The funders had no role in study design, data collection and analysis,
655 decision to publish, or preparation of the manuscript.

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875
876 **Supplementary information:**

877 **File #1:**

878 - File name: Supplementary_material_and_methods_1.docx
879 - File format including the correct file extension: Word document, .docx
880 - Title of data: Supplementary Materials and Methods
881 - Description of data: Extended description of the used text mining approach;
882 processing description of the second dataset.

883 **File #2:**

884 - File name: Supplementary_table_1.docx
885 - File format including the correct file extension: Word document, .docx

886 • Title of data: Supplementary Table 1: Overview of the processed RNAseq data.

887 • Description of data: Metrics of the quality of the RNAseq data, e.g. trimming metrics

888 **File #3:**

889 • File name: supplementary_file_1_logs.7z

890 • File format including the correct file extension: 7zip archive, .7z

891 • Title of data: Supplementary file 1: Logfiles.

892 • Description of data: Logfiles of RNAseq quality control, read mapping and read

893 counting

894

895 **File #4:**

896 • File name: supplementary_file_2_python_scripts.7z

897 • File format including the correct file extension: 7zip file, .7z

898 • Title of data: Supplementary file 2: Python scripts

899 • Description of data: Python scripts used for quality control, read mapping, read

900 counting and deriving of custom EC numbers

901

902

903 **File #5:**

904 • File name: supplementary_file_3_all_read_counts.with_annotation.7z

905 • File format including the correct file extension: 7zip file, .7z

- Title of data: Supplementary file 3: Read count table with annotation
- Description of data: Read counts for the ten relevant samples, including taxonomy, functional annotation (interpro domains, EC numbers, dbcan identifiers), note if differentially expressed and regression values

File #6:

- File name: supplementary_file_4_edgeR.7z
- File format including the correct file extension: 7zip file, .7z
- Title of data: Supplementary file 4: edgeR analysis
- Description of data: input file for the edgeR analysis, R script with used commands, output tables and MA plots.

File #7:

- File name: supplementary_file_5_heatmaps.7z
- File format including the correct file extension: 7zip file, .7z
- Title of data: Supplementary file 5: Heatmaps
- Description of data: Heatmaps of all genes in the three distinguished expression patterns

File #8:

- File name: supplementary_file_6.methane_related_processes_fig_6_curated.7z
- File format including the correct file extension: 7z file, .7z

926 • Title of data: Supplementary file 6: Overview over methane related reactions.

927 • Description of data: Overview over methane related reactions with assigned genes,
928 corresponding to figure 6/supplementary figure 1.

929 **File #9:**

930 • File name: supplementary_figure_1_rumen_metabolism_with_legend.svg

931 • File format including the correct file extension: scalable vector graphic, .svg

932 • Title of data: Supplementary figure 1: Annotated version of figure 6.

933 • Description of data: Annotated version of figure 6, with details given in supplementary
934 file 6.

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939 **Figure Legends:**

940 **Figure 1: Study design.** Four groups of three cows were allowed to adapt to one of four
941 different experimental diets for twelve days. From day 13 – 17 methane emission was
942 measured using a respiration chamber. Rumen fluid was collected at day 17 and used for
943 microbial RNA extraction. See Methods section for details.

944

945 **Figure 2: Taxonomic composition of metatranscriptome landscapes observed in animals**
946 **fed one of four different diets.** Diets and cows are indicated on the X-axis, taxonomic
947 groups (at genus level, or otherwise deepest classification) are colour coded, see legend for
948 details. N/A: No taxonomic rank could be assigned.

949

950 **Figure 3: Differential expression analysis of the rumen microbiomes.** Overview of the
951 number of genes that were found to be differently expressed in pairwise comparisons of
952 metatranscriptomes derived from animals fed different diets. Three profiles are distinguished:
953 **Profile A**, genes with an expression, which does not follow a dietary pattern. **Profile B**, genes
954 which are upregulated with increasing amounts of corn silage (CS). **Profile C**, genes, which
955 are downregulated with increasing amounts of CS. Furthermore the results show that with the
956 increase of CS, archaeal genes were mainly downregulated considerably affecting methane
957 metabolism.

958

959

960 **Figure 4: Log10 fold changes in expression of differentially expressed glycosylhydrolase**
961 **encoding genes in a comparison of the 100% corn silage diet (GS0) versus the 100%**
962 **grass silage diet (GS100).** Positive values indicate an upregulation of gene expression in the
963 corn silage diet. N/A: No taxonomic rank could be assigned. Colour-coding of bars indicate
964 different taxonomic groups, whereas colour-coding of protein families indicate their
965 involvement in the metabolism of different carbohydrates.

966

967 **Figure 5: Log10 fold changes in expression of differentially expressed carbohydrate**
968 **binding module encoding genes in a comparison of the 100% corn silage diet (GS0)**
969 **versus the 100% grass silage diet (GS100).** Positive values indicate upregulation of gene
970 expression in the corn silage diet. N/A: No taxonomic rank could be assigned. Colour-coding
971 of bars indicate different taxonomic groups, whereas colour-coding of protein families
972 indicate their involvement in the metabolism of different carbohydrates.

973

974 **Figure 6: Graphical summary of metabolic consequences of the different diets in the two**
975 **major methanogens and possible syntrophic partners.** Red arrows: genes downregulated
976 with the increase of corn silage in the diet; black arrows: Gene is detected but not
977 differentially expressed; The blue arrow represents glycolysis of which the majority could be
978 detected; punctuated arrow: orphan reactions; ? = Phylogenetic association unclear. Pyr. =
979 Pyruvate, $CHOO^-$ = Formate, FD_{ox} = oxidized Ferredoxin, FD_{red} = reduced Ferredoxin, CH_2O
980 = Formaldehyde.

981