Serial Analysis of Gene Expression (SAGE) in the Skeletal Muscle of Pig

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ABSTRACT

Skeletal muscle growth represents one of the main economic traits in pig production. To gain a better understanding of expression profile in pig muscle, serial analysis of gene expression (SAGE) was performed in Longissimus dorsi of two pigs at 3 and 9 months of age. A total of 53,120 long tags were obtained and sequenced from the four muscle SAGE libraries, representing 17,902 different tags, or putative transcripts, 0.64% (+0.09) of which had a relative expression level higher than 1‰. Overall, a total of 218 tags were highly expressed and 31 had a frequency higher than 3‰.

Functional characterisation of the expression profiles was performed using Kyoto Encyclopedia of Genes and Genomes metabolic maps and 139 pathways were identified for swine skeletal muscle. Focal adhesion, Mitogen-Activated Protein Kinase signalling, oxidative phosphorylation, ribosomal proteins, regulation of actin cytoskeleton and insulin signalling pathways showed an abundance of genes greater than 1.5% of all the expressed transcripts. A comparison with human SAGE library indicated no statistical differences for the frequency of genes involved in these metabolic pathways.

Key words: Growth, SAGE, Skeletal muscle, Sus Scrofa.

RIASSUNTO

ANALISI SERIALE DELL’ESPRESSIONE GENICA (SAGE) NEL MUSCOLO SCHELETRICO DI SUINO

L’accrescimento muscolare costituisce uno dei principali aspetti economici della produzione del suino. Al fine di aumentare la conoscenza del profilo di espressione genica nel suino, è stata applicata la tecnica SAGE (Serial Analysis of Gene Expression) a campioni biotipi di Longissimus dorsi ottenuti da 2 suini a 3 e a 9 mesi di età. Dalle 4 librerie SAGE, sono stati ottenuti e sequenziati 53120 tags complessivamente, corrispondenti a 17902 tags differenti, o trascritti putativi, di cui 0,64% (+0.09) con un livello relativo di espressione superiore a 1‰. I tags altamente espressi sono risultati pari a 218, di cui 31 presentano una frequenza superiore al 3‰.

La caratterizzazione funzionale dei profilo di espressione è stata condotta utilizzando le mappe metaboliche della Kyoto Encyclopedia of Genes and Genomes (KEGG) e i trascritti hanno interessato 139 cammini di muscolo scheletrico del suino. L’adesione focale, il segnale delle protein-chinasi attivate da mitogeni, la
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fosforilazione ossidativa, le proteine ribosomiali, la regolazione dell’actina citoscheletrica e il segnale dell’insulina hanno evidenziato un numero di geni superiore a 1,5% dei tutti i trascritti espressi. Il confronto con il profilo di espressione ottenuto con la tecnica SAGE nell’uomo è risultato simile, con una frequenza di geni coinvolti nelle stesse mappe metaboliche non statisticamente significativo.

Parole chiave: Accrescimento, SAGE, Muscolo scheletrico, Sus Scrofa.

Introduction

Breeding goals for pigs are directed towards retail carcass yield and meat quality because of the high economic value of these traits and the identification of genes expressed in skeletal muscle could enhance the selection and evaluation of candidate genes associated with QTL (Fahrenkrug et al., 2002). These gene markers can be integrated with performance information in the marker-assisted selection and used by commercial pig industry to improve traits of economic importance in pig production.

The growth and the development of skeletal muscle is a multistage process involving functional interaction between the intrinsic genetic program and the extrinsic regulation mediated by hormones and growth factors (Wilson et al., 2004). Techniques that allow to investigate tissue-specific expression profile of mRNAs or pattern of mRNAs expression may contribute to identify genes which play a major role for the condition under investigation (Dal Monego et al., 2007; Yoshioka et al., 2007), providing tools for the identification of candidate genes for association studies.

Serial analysis of gene expression (SAGE, Velculescu et al., 1995) allows qualitative and quantitative transcriptome-wide analysis of gene expressions within a tissue during discrete physiological stages (Tutej and Tuteja, 2004). Furthermore, SAGE does not require extensive knowledge of genome information or access to cDNA libraries and clones, allowing the discovery of novel genes.

The ability of SAGE to characterize the molecular mechanisms responsible for the modification of physiological or pathological conditions in muscle, has been already demonstrated (Welle et al., 2000; Yoshioka et al., 2003; Schaff et al., 2005; Wang, 2007).

In this study, SAGE was employed to describe and quantify gene expression in pigs at 3 and 9 months of age. The aim was to obtain a gene expression profile of the most representative skeletal muscle transcript of Longissimus dorsi. These genes provide a base inventory of the main metabolic pathways of pig muscle.

Material and methods

Animals and sample preparations

Twenty castrate pigs, 10 purebred Large White (LW) and 10 Landrace X Duroc (DU), were reared from 3 to 12 months of age in the same environmental and feeding conditions. The experimental procedures were carried out in accordance with the Italian legislation on experimental animals (DL n.116, 27/01/1992).

At the age of 70 days, piglets were introduced in the experimental farm for the adaptation to the new environmental and feeding conditions; the experiment started at 90 days of age and live weights were recorded every month until slaughter, using conventional procedures in a local abattoir. During the trial, animals were given a restricted amount of feed and were kept in two separate boxes, with free access to water and to an external paddock. Feed consumptions
were recorded per group on a weekly basis.

At the age of 3 and 9 months, Longissimus dorsi (LD) biopsy was collected from all the animals, using a Magnum Biopsy System (Bard, Covington, GA USA), under subcutaneous anaesthesia. Biopsies were sampled on the left side, between the third and fourth last ribs, at about 20-40 mm (3 months) and 60-80 mm (9 months) from the median line.

Samples were immediately homogenized in monophasic solutions of phenol and guanidine isothiocyanate (RNAtidy G, Applichem, Darmstadt, Germany). The homogenates were frozen at -80°C within 1 day and processed to extract total RNA within 1 week after sampling, following the manufacturer’s instructions.

The amount of total RNA extracted from the tissues was determined by absorbance of ultraviolet light at 260 nm, with background compensation for the absorbance at 320 nm, by using a spectrophotometer. Total RNA integrity was evaluated by the observation of 28S to 18S fluorescence of ribosomal bands after electrophoresis on 1% denaturing formaldehyde agarose gel in the presence of ethidium bromide. The mean total RNA extracted from the biopsies used for the 4 SAGE library was 25.3±12.8 µg, with a 260/280 ratio of 2.02±0.10.

**Construction and sequencing of SAGE libraries**

For SAGE libraries, two pigs were selected on the basis of the live weight and daily gains, to represent the Large White and LandraceXDuroc phenotypes and the corresponding biopsies sampled at 3 and 9 months were processed. Pooling samples can potentially hide biological variance and give false confidence concerning data significance (Jolly et al., 2005).

The 4 SAGE libraries were constructed starting from about 10 µg total RNA using a commercial kit (I-SAGE™ Long Kit, Invitrogen, Carlsbad, CA USA), according to the manufacturer’s recommendations. Protocol was based on a modified (Claverie, 1999; Lash et al., 2000) method of Velculescu et al. (1995), with further modifications.

Polyadenylate RNA (mRNA) was isolated by means of oligo(dT) magnetic beads, retrotranscribed to cDNA and digested with restriction enzyme NlaIII. To test the efficiency of cDNA synthesis and NlaIII digestion, *Sus scrofa* β-actin primers surrounding NlaIII 3’ restriction site were designed (AF1or GGACTTCG AGCAGGA GATGG, ARev1 GCACCGTGTTGGCGTCGTAGAGG, ARev2 GTGAGAACGTTAGGCCCCAG). The 3’ terminal fragments were isolated with the anchored streptavidin-coated magnetic beads, ligated to LS adapters, modified with an amino group at the 3’ end to prevent self-ligation. Adjacent tags were released by cleavage with the type II restriction endonuclease MmeI e linked together to obtain ditags of ~130bp. These ditags were first amplified by PCR (HotMaster TAQ Polymerase, Qiagen, Hilden, Germany) and products were analyzed by agarose gel electrophoresis. Large-scale PCR amplification and polyacrylamide gel (6%, 19:1) purification were performed followed by secondary NlaIII digestion. The ditags were isolated with polyacrylamide gel (10%, 19:1), excised and self-ligated to produce long concatamers. The concatamers between 500 and 1500 bp were isolated by agarose gel (1.5%) and extracted by DNA and gel purification kit (GFX purification kit, GE Healthcare, Uppsala, Sweden). Three different pools of concatamers were prepared: 500-750 bp, 750-1000 bp and 1000-1500 bp. These products were cloned into SphI site of pZErO®-1 vector using the Quick Ligation kit (NEB, Ipswich, MA). Clones were PCR screened to select inserts for sequencing, PCR products were sequenced by BMR sequencing service (BMR genomics, 2008).
The entire SAGE dataset was submitted to the GEO repository (Gene Expression Omnibus, 2008), and have been assigned GEO series accession number GSE6062.

**Bionformatic analysis of SAGE tag sequences**

The sequence and occurrence of each of the transcript tags were determined using the software SAGEparser, a Perl program, developed by Dinel et al. (2005). This software is freely available on the net for use in research. Taq annotations and Sus scrofa - Homo sapiens correlations were investigated by means of several in-house Perl scripts (BioPerl, 2007).

**Annotation of SAGE tag via NCBI tables**

The flow chart of annotation process is reported in Figure 1. In the first attempt, the annotation of the transcripts was carried out by matching the 17 bp SAGE tags with a public database (Gene Expression Omnibus, 2008). Sus scrofa files from SAGEmap (SAGEmap_tag_ug-full.zip, release of 2007-03-01) and from UniGene (Ssc.data.gz, Ssc.seq.all.gz and Ssc.sequniq.gz; build 28, 2007-03-01) were downloaded. These databases were merged in order to construct a local database (MySQL, 2007) containing all information necessary for the annotations. Using this tool, all 17 bp tags were linked to a UniGene cluster. This process resulted in tags either associated with unique, multiple or not associ-
Gene expression profiles. All tags were merged by organism and KEGG pathways comparative analysis was carried out using the Fatigo+ web tool of Babelomics suite (Al-Shahrour et al., 2006).

Significant differentially expressed genes between 3 and 9 months were identified by analysing SAGE libraries using the IDEG6 software, a web tool for detection of differentially expressed genes, with Bonferroni correction and a significance Chi square level of 0.05 (Romualdi et al., 2003).

Results

SAGE results

A total of 53120 long tags were obtained and sequenced from the 4 muscle SAGE libraries, 13590 tags for DU_03 (GSM140696), 12980 for DU_09 (GSM140697), 13130 for LW_03 (GSM140698) and 13420 for LW_09 (GSM140699) (Table 1). Combining together, the 4 libraries represented 17902 different tags, or putative transcripts, and 5158, 4437, 5241 and 7665 putative transcript unique for GSM140696, GSM140697, GSM140698 and GSM140699, respectively. Considering the skeletal muscle growth, an high percentage of tags were specific for 3 and 9 months (74.35 and 78.71%, respectively) and 11.64% of the non redundant tags were common.

The relative tags frequencies obtained via SAGE reflected the expression level of the transcripts represented by those tags. The rate of occurrence of unique tags revealed that 0.64% (+0.09) of the transcripts have a relative expression level of more than 1‰.

The SAGEmap NCBI database repository contains an outdated table and several mismatches were found in the annotation fields. For this reason and for better mining of data for ontology and physiology, the porcine SAGE tags were linked directly to human UniGene clusters.
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The annotation of SAGE tag via NCBI tables resulted in 4947 Sus scrofa UniGene links to our not redundant tag database, 3,370 of which associated to a human cluster. The second approach, through Blastn similarity search, increased the number of tag linked to Sus scrofa UniGene to 7,377. Of these Sus scrofa UniGene 72% matched to human UniGene clusters via TBlastx association. This corresponds to a 29.8% of total number of porcine tags with human annotation, a quite low level, but considering only the transcripts expressed more than 1‰, this value increased to 90.2%.

To reduce the problem of multiple tags linked to a single transcript and to make the ontology mining more reliable, a final table of swine clusters with human linked annotation and the corresponding tags count was created. This newly created dataset, an important tool in itself, was used for the following functional analysis. A list of genes with a frequency higher than 3‰ (in the skeletal muscle biopsies of pigs is given in Table 2.)

Comparing human and pig skeletal muscle SAGE tag profiles

Of 146 and 139 KEGG pathways identified in human and swine skeletal muscle respectively, 12 in humans showed a relative abundance greater than 1% and an overlapping rank was found for the pig gene data (Figure 2). The most represented classes of pathway were focal adhesion, involving major proteins responsible for signal transduction and sarcomere assembling, followed by MAPK, which is also strictly integrated to the former in the regulation of gene transcription and metabolism in response to oxidative, energetic, and mechanical stress in skeletal muscle. Other pathways with a large number of gene transcripts, apart from oxidative phosphorylation, were insulin signalling and regulation of actin cytoskeleton pathways, that are downstream regulated by the MAPK cascade.

Skeletal muscle SAGE tags differentially expressed between 3 and 9 months ages.

To analyse swine muscle growth the sum of GSM140696/GSM140698 tags (3 months

Table 1. Number of tags counted, redundant and specific for the SAGE projects.

| SAGE libraries | Mean | 3 months | 9 months | Overall |
|----------------|------|----------|----------|---------|
| GSM 140696     | 13,590 | 12,980   | 13,420   | 27,620  | 26,400   | 13,280   | 53,120   |
| GSM 140697     | 13,130 | 13,340   | 13,420   | 26,860  | 26,740   | 13,100   | 49,850   |
| GSM 140698     | 13,280 | 13,130   | 13,620   | 26,860  | 26,740   | 13,100   | 49,850   |
| GSM 140699     | 53,120 | 49,850   | 49,850   | 99,700  | 99,700   | 49,850   | 199,400  |

1SAGE libraries: GEO series accession number GSE6062.
2GSM140696: DU_03; GSM140697: DU_09; GSM140698: LW_03; GSM140699: LW_09.
33 months: GSM140696: +GSM140698.
49 months: GSM140697+GSM140699.
5all: GSM140696+GSM140697+GSM140698+GSM140699
old) and of GSM140697/GSM140699 tags (9 months old) were considered. To have a deeper insight into the genes differentially expressed during the two ages, statistical evaluation and functional analysis was performed.

The comparison of transcripts abundance between 3 and 9 months was performed with IDEG6 using the chi square test (Romualdi et al., 2003). A set of 190 differentially expressed genes (47 without annotation) with a P value lower than 0.05 was identified, 114 of which featured the 3 months. The functional analysis described for the human-swine comparison was applied to classify the age-specific genes to the KEGG pathway (Table 3).

Discussion

The number of unique putative transcripts accounted for 5158, 4437, 5241 and 7665 for GSM140696, GSM140697, GSM140698 and GSM140699, respectively (Table 1). The putative transcripts common to the four SAGE projects were 548. A limited number of common tags between SAGE projects and a large number of tags with very low abundance in muscle samples were also reported by Welle et al. (2000), comparing young and old muscle in man in a database cataloguing more than 100,000 tags. Moreover, the limited number of transcripts sequenced can have also contributed to the reduced number of common tags between the 4 libraries.

The almost complete overlapping of KEGG pathways between human and swine skeletal muscle transcriptomes (Figure 2) indicates that the number of tags sequenced in the present study can represent, at least, the more abundant transcripts in this tissue.

Bortoluzzi and Danieli (1999) described
Table 2. The most abundant tags (>3% of total tags) in *Longissimus dorsi* muscle of pig and the corresponding annotation to human UniGene clusters (SAGE libraries in the GEO series accession number GSE6062).

| Transcript | Homo sapiens | Tag frequency |
|------------|--------------|---------------|
|            | Cluster      | Name          | All tags | GSM 140696 | GSM 140697 | GSM 140698 | GSM 140699 |
| CTTTTCCTCACTCAAAAG | 334347 | CKM | 35.8 | 41.7 | 44.3 | 29.9 |
| GAGGCGGGTGGCAGCCAAG | 632642 | PGAM2 | 29.3 | 41.1 | 31.7 | 29.5 | 14.7 |
| TACCATCAATAAGTAC | 534770 | PKM2 | 23.6 | 13.9 | 26.0 | 37.9 | 17.2 |
| CCTACTAGCGGCGGAGG | 513490 | ALDOA | 22.9 | 24.6 | 31.6 | 20.6 | 15.1 |
| TGGGCAGCTTCCCCTCC | 50889 | MYLPF | 22.9 | 27.9 | 30.4 | 21.1 | 12.1 |
| TGGGCAGCTTCCCCCCC | 50889 | MYLPF | 22.2 | 27.0 | 28.4 | 22.2 | 11.4 |
| AGGATGAGGAGGCTCTT | 224171 | PKM2 | 23.6 | 13.9 | 26.0 | 37.9 | 17.2 |
| GCCGGCTACATCGAT | 182421 | TNN12 | 8.5 | 6.8 | 12.2 | 9.9 | 5.2 |
| CCCATGTACTCGAAGCTT | 73454 | TNNT3 | 8.3 | 6.4 | 15.1 | 6.3 | 5.7 |
| ACCACTTACCGGAGGCC | Mitochond.| COX2 | 9.7 | 5.7 | 12.1 | 11.3 | 9.6 |
| AAGATCAAGATCATCGC | 1288 | ACTA1 | 9.7 | 5.7 | 9.2 | 16.5 | 7.4 |
| GGGCTTGAGGAGAAG | 2795 | LDHA | 8.8 | 7.8 | 14.2 | 7.8 | 5.4 |
| GAGGCGCCGCAAAGAGAT | 523403 | TNN2 | 8.5 | 6.8 | 12.2 | 9.9 | 5.2 |
| AACGTTCGGCGCAGAT | 73454 | TNNT3 | 8.3 | 6.4 | 15.1 | 6.3 | 5.7 |
| ACCACTTACCGGAGGCC | Mitochond.| COX3 | 6.8 | 2.4 | 11.4 | 8.4 | 5.1 |
| ACTTCAACCGGGATGTG | 523443 | HBB | 6.2 | 17.3 | 2.6 | 1.8 | 2.8 |
| TGGATGCAAGGAAATAA | 499745 | MYH2 | 5.9 | 1.2 | 7.8 | 9.5 | 5.3 |
| GCGCCCTACAGGGGGG | 445037 | ACTN3 | 5.6 | 6.7 | 8.0 | 4.6 | 3.2 |
| GACAGCGGAACACTAA | 175473 | AK1 | 5.6 | 6.8 | 6.5 | 5.8 | 3.2 |
| TGATTAAAGTGTCAAC | 334629 | SLN | 5.5 | 4.9 | 5.8 | 6.5 | 5.0 |
| ACTAATGTCACCATCT | 187338 | MYL1 | 5.1 | 3.5 | 6.5 | 6.7 | 3.7 |
| AGGCCCTGGCCGGCCG | 534404 | RPL10 | 5.0 | 8.3 | 4.6 | 4.0 | 3.1 |
| TCCAAATAACACAGGAG | 247831 | MYLC2PL | 4.1 | 3.8 | 1.7 | 8.0 | 3.1 |
| CAGGGTTAACTTCGAT | 134602 | TTN | 4.1 | 1.5 | 3.9 | 7.7 | 3.4 |
| AATTGGAGGACAACCGC | Mitochond.| CYTB | 3.9 | 1.9 | 6.2 | 4.9 | 2.8 |
| GGCGCCAACCTACAGCAG | 73454 | TNNT3 | 3.9 | 3.2 | 4.1 | 6.4 | 1.9 |
| GACACTGAGTTTGTCTT | 505662 | PPP1R1A | 3.6 | 3.9 | 4.0 | 4.0 | 2.5 |
| GTCCCTACTTCCCCAG | 594952 | DES | 3.6 | 3.4 | 5.3 | 3.2 | 2.5 |
| ATGGCGGAGGAGCTGAA | 499745 | MYH2 | 3.5 | 1.0 | 5.5 | 5.1 | 2.5 |
| AAAAATCTCATTTGCTAC | Mitochond.| tRNA-Glu | 3.3 | 1.0 | 4.2 | 4.6 | 3.4 |
| AAAATAAACATCCTCGG | 533040 | PDLIM7 | 3.3 | 4.0 | 2.4 | 3.1 | 3.6 |
| TCGCGTTGCTAGGGGCG | 320890 | TNN11 | 3.2 | 4.6 | 3.5 | 3.4 | 1.2 |

All tags: GSM140696+GSM140697+GSM140698+GSM140699.

GSM140696: DU_03; GSM140697: DU_09; GSM140698: LW_03; GSM140699: LW_09.
the transcriptional profile of human muscle using an in silico and a SAGE database of over 53,875 tag sequences, of which 295 tags were considered highly expressed, since they were detected more than 20 times. Applying the same criteria to the present database, a total of 218 tags were found to be highly expressed and those with a frequency higher than 3‰ are reported in Table 2.

The functional analysis of the 190 differentially expressed genes (P<0.05) between 3 and 9 months of age was performed allocating the genes to the same KEGG pathways using for the human-swine comparison, in order to find those which can elicit the physiological status of muscle at the two analysed ages (Table 3).

Analysing both Figure 2 and Table 3 one can observe three KEGG pathways, insulin signalling, focal adhesion and regulation of actin cytoskeleton closely interacting between them and that are involved in the regulation of protein synthesis in mammals (Dardevet et al., 1996; Goel and Dey, 2002). The initiation phase of translation for the formation of 80S initiation complex is a limiting step of protein synthesis (Pain, 1996) and is controlled by some major proteins, such as eIF2, eIF2B, 4E-BPI and ribosomal protein S6 kinase (S6K1). S6K1 has been reported to be among the major proteins involved in the control of initiation phase of mRNA translation (Tesseraud et al., 2003) and is phosphorylated by PIK3, a gene of the phosphatidyl inositol signalling system, involved in the insulin signalling, focal adhesion and regulation of actin cytoskeleton pathways. This gene is down-stream to IRS in the insulin signalling pathway and the activation of the PI3K-mTOR signal transduction results in a more intense protein synthesis, through modulation of multiple steps involved in the initiation of mRNA translation and ribosome biogenesis (Bolster et al., 2004). This process and the following elongation and protein synthesis is very energy demanding.

The observed up regulation of PIKR1 at 3 months of age (Table 3) agrees with the increased abundance of all the significant ribosomal and oxidative phosphorylation proteins related transcripts in younger pigs, as a sign of the intense metabolic activity. The age-related decline of genes involved in electron transport and ATP synthesis is consistent with the study of Welle et al. (2000), in which SAGE method was used to compare transcriptional profile of young and old Vastus lateralis muscles. Similar results were also obtained with microarray method applied to young and old mice (Lee et al., 1999).

Within the focal adhesion pathway, COL3A1 and ITGA7, other than PIK3R1, were up regulated at the 3 months of age. PIK3R1 promotes ACTN2, a F-actin cross-linking protein which is thought to anchor actin to a variety of intracellular structures. Collagen type 1 alpha (COL3A1), is involved in the ECM receptor interaction and takes part of the regulation of integrins, as integrin alpha-7 (ITGA7), that is a primary laminin receptor on skeletal myoblasts and adult myofibers (Song et al., 1993).

ITGA7, together with PIK3R1 and ACTN2 genes, are also part of the regulation of actin cytoskeleton KEGG pathway. During myogenic differentiation, ITGA7 may induce changes in the shape and mobility of myoblasts, and facilitate their localization at laminin-rich sites of secondary fiber formation, indicating that myofibers cytoarchitecture is still active at a younger age (Ziober et al., 1993).

Older animals exhibited increased abundance of CSDE1, an NRAS upstream gene protein, which in turn regulated RAF1, a proto-oncogene serine/threonine-protein kinase. RAS and RAF1 are also part of the transduction of mitogenic signals (MAPK...
According to Coolican et al. (1997), cell proliferation is mediated primarily by the RAS/RAF1 MAP kinase pathway, whereas stimulation of differentiation utilizes the PI3-kinase/p70S6k pathway. Extracellular signals can lead to a simultaneous activation of both the RAF-MEK-ERK and PI3K-AKT pathways. The effects of these two signalling pathways can be opposite on muscle cells, since the PI3K-AKT pathway was shown to inhibit the RAF-MEK-ERK pathway and cross-regulation depends on the differentiation state of the cell (Rommel et al., 1999). In particular, AKT activation inhibits RAF-MEK-ERK pathway in differentiated myotubes, but not in their myoblast precursors (Wilson et al., 2004). This observation is consistent with opposite behaviour of the two components of microtubule protofilaments. At 3 months a predominant expression of the beta subunit (TUBB) can lead to a more instable cytoskeleton filament, which agrees with the

| KEGG pathways                        | 3 months                          | 9 months                          |
|--------------------------------------|-----------------------------------|-----------------------------------|
|                                      | GSM140696 + GSM140698             | GSM140697 + GSM140699             |
|                                        |                                   |                                   |
| Focal adhesion                        | ITGA7 PIK3R1 FLNA ACTN2 COL3A1 MYL6 | RAF1 PPP1CC                       |
| MAPK signaling pathway                | FLNA HSPA5                        | RAF1 CSDE1                        |
| Oxidative phosphorylation             | COX8A COX4I1 ATP6V1G1 AT- P5G3 NDUV3 COX7C NDUFA12 NDUFB9 | No genes                        |
| Regulation of actin cytoskeleton      | MYH9 ITGA7 PIK3R1 ACTN2 RPL10 RPL36 RPL27A RPL29 | RAF1 CSDE1 PPP1CC |
| Ribosome                              | RPS23 RPL35 RPL37 RPS15 RPL18 RPL10A RPL13A RPL14 | No genes                        |
| Insulin signaling pathway             | PIK3R1 PKM2                       | RAF1 CSDE1 FASN PPP1CC            |
| Calcium signaling pathway             | No genes                          | No genes                          |
| Wnt signaling pathway                 | No genes                          | NFAT5                             |
| Tight junction                        | MYH9 ACTN2 MYL6                   | CSDE1                             |
| Axon guidance                         | No genes                          | CSDE1 NFAT5                       |
| Proteasome                            | PSMA6                             | No genes                          |
| Purine metabolism                     | AK1 PKM2                          | No genes                          |
| Glycolysis / Gluconeogenesis          | PGAM2 ENO3 PKM2                   | PGM1 ALDH2                        |
| Gap junction                          | TUBB                              | TUBA1 RAF1 CSDE1                  |

GSM140696: DU_03; GSM140697: DU_09; GSM140698: LW_03; GSM140699: LW_09.
function of microtubules during the skeletal muscle growth and differentiation (Pizon et al., 2005). In the later phase of growth (9 months) a stabilization of skeletal muscle structure is expected, justifying the prevalence of alpha subunits (TUBA) of tubulin isoforms.

A similar IDEG6 analysis crossing data between the two breeds do not produce consistent results. The lack of transcripts featuring this two breeds mean that the differences between them are to ascribe to genes transcribed at lower level than this work can discriminate. This observation moreover is confirmed by microarray analysis on the same population (manuscript in preparation).

Conclusions

This first SAGE database of pig muscle of more than 50,000 tags may be a useful resource for researchers interested in functional genomics and in the relative mRNA expression in muscle tissues. For the most abundant mRNA species, the data provided can be considered accurate. But for the less abundant transcripts, an increase of tag database may be required. However, these results demonstrate the power of transcriptional profiling for gene discovery and provide opportunities for investigating new proteins potentially involved in different aspects of growth factor action in muscle tissues.

In the paper, data mining was mainly based on human KEGG maps, assuming the correspondence with swine cellular regulation and metabolism, but many of the differences identified by SAGE involve protein products or full mRNA sequences that have not been characterized in swine. The possibility to compare the activation or inhibition of specific pathways during growth would also offer the opportunity to identify candidate genes with regulatory activities in skeletal muscle. As soon as the pig genome project will be completed, the species-specific pathways for pigs will be available and a critical reconsideration of these results will probably required, to provide information about the expression profile of muscle in pigs.

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REFERENCES

Al-Shahrour, F., Minguez, P., Tárraga, J., Montaner, D., Alloza, E., Vaquerizas, J.M.M., Conde, L., Blaschke, C., Vera, J., Dopazo, J., 2006. Babelomics: a systems biology perspective in the functional annotation of genome-scale experiments. Nucleic Acids Res. 34:W472-W476.

BioPerl, 2007. Home page address: http://www.bioperl.org/

BMR genomics, 2008. Home page address: http://www.bmr-genomics.it/

Bolster, D.R., Jefferson, L.S., Kimball, S.R., 2004. Regulation of protein synthesis associated with skeletal muscle hypertrophy by insulin, aminoacid and exercise induced signalling. P. Nutr. Soc. 63:351-356.

Bortoluzzi, S., Danieli, G.A., 1999. Towards an in silico analysis of transcription patterns. Trends Genet. 15:118-119.

Claverie, J.M., 1999. Computational methods for the identification of differential and coordinated gene expression. Hum. Mol. Genet. 8:1821-1832.

Coolican, S.A., Samuel, D.S., Ewton, D.Z., McWade,
F.J., Florini, J.R., 1997. The mitogenic and myogenic actions of insulin-like growth factors utilize distinct signaling pathways. J. Biol. Chem. 272:6653-6662.

Dal Monego, S., Colitti, M., Pallavicini, A., D’Andrea, M., Pilla, F., Graziosi, G., Stefanon, B., 2007. Evaluation of gene expression profiles of pig skeletal muscle in response to energy content of the diets using human microarrays. Ital. J. Anim. Sci. 6:45-59.

Dardevet, D., Sornet, C., Vary, T., Grizard, J., 1996. Phosphatidylinositol 3-kinase and p70 S6 kinase participate in the regulation of protein turnover in skeletal muscle by insulin and insulin-like growth factor 1. Endocrinology 137:4087-4094.

Dinel, S., Bolduc, C., Belleau, P., Boivin, A., Yoshioka, M., Calvo, E., Piedboeuf, B., Snyder, E.E., Labrie, F., St-Amand, J., 2005. Reproducibility, bioinformatic analysis and power of the SAGE method to evaluate changes in transcriptome. Nucleic Acids Res. 33:1-8.

Fahrenkrug, S.C., Smith, T.P., Freking, B.A., Cho, J., White, J., Vallet, J., Wise, T., Rohrer, G., Pelt, G., Sultana, R., Quackenbush, J., Keele, J.W., 2002. Porcine gene discovery by normalized cDNA-library sequencing and EST cluster assembly. Mamm. Genome 13:475-478.

Goel, H.L., Dey, C.S., 2002. Insulin stimulates spreading of skeletal muscle cells involving the activation of focal adhesion kinase, phosphatidylinositol 3-kinase and extracellular signal regulated kinases. J. Cell. Physiol. 193:187-198.

Goel, H.L., Dey, C.S., 2002. Insulin stimulates spreading of skeletal muscle cells involving the activation of focal adhesion kinase, phosphatidylinositol 3-kinase and extracellular signal regulated kinases. J. Cell. Physiol. 193:187-198.

Jolly, R.A., Goldstein, K.M., Hong Gao, T.W., Chen, P., Huang, S., Colet, J.M., Ryan, T.P., Thomas, C.E., Estrem S.T., 2005. Pooling samples within microarray studies: a comparative analysis of rat liver transcription response to prototypical toxicants. Physiol. Genomics 22:346-355.

Lash, A.E., Tolstoshev, C.M., Wagner, L., Schuler, G.D., Strausberg, R.L., Riggins, G.J., Altschul, S.F, 2000. SAGEmap: a public gene expression resource. Genome Res. 10:1051-1060.

Lee, C.K., Klopp, R.G., Weindruch, R., Prolla, T.A., 1999. Gene expression profile of aging and its retardation by caloric restriction. Science 285:1390-1393.

MySQL, 2007. Home page address: http://www.mysql.com/

Pain, V.M., 1996 Initiation of protein synthesis in eukaryotic cells. Eur. J. Biochem. 236:747-771.

Pizon V., Gerbal, F., Diaz C.C., Karsenti, E., 2005. Microtubule-dependent transport and organization of sarcomeric myosin during skeletal muscle differentiation. EMBO J. 24:3781-3792.

Rommel, C., Clarke, B.A., Zimmermann, S., Nunez, L., Rossman, R., Reid, K., Moelling, K., Yancopoulos, G.D., Glass, D.J., 1999. Differentiation stage specific inhibition of the Raf-MEK-ERK pathway by Akt. Science 286:1738-1741.

Romualdi, C., Bortoluzzi, S., D’Alessi, F., Danieli, G.A., 2003. IDEG6: a web tool for detection of differentially expressed genes in multiple tag sampling experiments. Physiol. Genomics. 12:159-162.

Schaaf, G.J., Ruijter, J.M., van Ruisen, F., Zwijsen, D.A., Waaier, R., Valenti, L.J., Benit-Deekman, J., van Kampen, A.H.C, Baas, F., Kool, M., 2005. Full transcriptome analysis of rhabdomyosarcoma, normal and fetal skeletal muscle: statistical comparison of multiple SAGE libraries. FASEB J. 19:404-406.

Tesseraud, S., Bigot, K., Taouis, M., 2003. Amino acid availability regulates S6K1 and protein synthesis in avian insulin-insensitive QM7 myoblasts. FEBS Lett. 540:176-180.

Tutej, R., Tuteja, N., 2004. Serial Analysis of Gene Expression: Applications in Human Studies. J. Biomed. Biotech. 2:113-120.

Veledescu, V.E., Zhang, L., Vogelstein, B., Kinzler, K.W., 1995. Serial analysis of gene expression. Science 270:484-487.

Wang, S.M., 2007. Understanding SAGE data.
Gene expression of pig skeletal muscle

Trends Genet. 23:42-50.
Welle, S., Bhatt, K., Thornton, C.A., 2000. High-abundance mRNAs in human muscle: comparison between young and old. J. Appl. Physiol. 89:297-304.
Wilson, E.M., Tureckova, J., Rotwein, P., 2004. Permissive role of phosphatidylinositol 3-kinase and Akt in skeletal myocyte maturation. Mol. Biol. Cell 15:497-505.
Yoshioka, M., Tanaka, H., Shono, N., Shindo, M., St-Amand, J., 2003. Serial analysis of gene expression in the skeletal muscle of endurance athletes compared to sedentary men. FASEB J. 17:1812-1819.
Yoshioka, M., Tanaka, H., Shono, N., Shindo, M., St-Amand, J., 2007. Gene expression profile of sprinter’s muscle. Int. J. Sports Med. 28:1053-1058.
Ziober, B.L., Wu, M.P., Waleh, N., Crawford, J., Lin, C.S., Kramer, R.H., 1993. Alternative extracellular and cytoplasmic domains of the integrin α7 subunit are differentially expressed during development. J. Biol. Chem. 268:26773-26783.