

Figure S1. Amino acid alignment of MSTN of medaka (*Oryzias latipes*; <u>AB520935.1</u>), tilapia (*Oreochromis mossambicus*; <u>AF197193.3</u>), gilthead seabream (*Sparus aurata*; <u>AF258448.1</u>), carp (*Cyprinus carpio*; <u>GQ214770.1</u>), catfish (*Ictalurus punctatus*; <u>AF396747.1</u>), turbot (*Scophthalmus maximus*; <u>EF683115.1</u>), zebrafish (*Danio rerio*; <u>AY258034.1</u>), fugu (*Takifugu rubripes*; <u>AY445322.1</u>), salmon (*Salmo salar*; <u>NM 001123549.1</u>) and pejerrey (*Odontesthes bonariensis*; <u>HM061693.1</u>). Identical amino acids are in black. The proteolytic processing site RXXR is boxed. Asterisks show the 9 conserved cysteines in the bioactive TGF-β2 domain, which is indicated by a black line over the sequence.

His / Cys-rich



Figure S2. Amino acid alignment of MyoG of tilapia (*Oreochromis niloticus*; <u>GU246717.1</u>), gilthead seabream (*Sparus aurata*; <u>EF462192.1</u>), carp (*Cyprinus carpio*; <u>AB012881.1</u>), catfish (*Ictalurus punctatus*; <u>AY534329.1</u>), sole (*Solea senegalensis*; <u>EU934044.1</u>), zebrafish (*Danio rerio*; <u>NM 131006.1</u>), fugu (*Takifugu rubripes*; <u>AY566282.1</u>), pufferfish (*Tetraodon nigroviridis*; <u>AY822074.1</u>), salmon (*Salmo salar*; <u>NM_001123600.1</u>) and pejerrey (*Odontesthes bonariensis*; <u>HM061694.1</u>). Identical amino acids are in black. The His / Cys-rich domain is indicated by a black line over the sequence and the bHLH domain is boxed.



Figure S3. Molecular Phylogenetic analysis by Maximum Likelihood method carried out with the partial nucleotide sequence of MSTN open reading frame (500 bootstrap replicates, nucleotide pdistance and 864 informative sites) with Mega5 program (Tamura et al., 2011). The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (500 replicates) is shown next to the branches. The human α -inhibin (<u>NM 002191.3</u>) was used as out-group. The accession numbers of the MSTN sequences are as follows: Acanthopagrus schlegelii (DQ303480.1), Dicentrarchus labrax (AY839106.1), Equus caballus (NM 001081817.1), Gallus gallus (NM 001001461.1), Morone americana (AF290911.1), Morone saxatilis (AF290910.1), Mus musculus (NM 010834.2), Odontesthes bonariensis (HM061693.1), Oncorhynchus mykiss (<u>NM 001124282.1</u>), Oncorhynchus mykiss MSTN-2 (<u>NM 001124283.2</u>), Oreochromis mossambicus (AF197193.3), Oryzias latipes (NM 001201499.1), Paralichthys adspersus (EU443627.1), Paralichthys olivaceus (**DQ412048.1**), Salmo salar MSTN-1a Salmo salar MSTN-1b (NM 001123634.1), Salvelinus fontinalis (NM 001123549.1), (AF247650.2), Sparus aurata (AF258448.1), Sus scrofa (NM 214435.2), Takifugu rubripes MSTN-2 (NM 001032672.1), Verasper variegatus (JN226745.1).



Figure S4. Molecular Phylogenetic anaylsis by Maximum Likelihood method carried out with the partial nucleotide sequence of MyoG open reading frame (500 bootstrap replicates, nucleotide pdistance and 394 informative sites) with Mega5 program (Tamura et al. 2011). The percentage of replicate trees in which the associated taxa clustered together in the bootstrap test (500 replicates) is shown next to the branches. The mouse follistatin (NM_008046.2) was used as out-group. The accession numbers of the MSTN sequences are as follows: *Ameiurus catus* (<u>AY562557.1</u>), *Cyprinus carpio* (<u>AB012881.1</u>), *Danio aequipinnatus* (<u>DQ219849.1</u>), *Danio rerio* (<u>NM 131006.1</u>), *Epinephelus coioides* (<u>HM190251.1</u>), *Gallus gallus* (<u>NM 204184.1</u>), *Ictalurus furcatus* (<u>AY540993.1</u>), *Mus musculus* (<u>NM_031189.2</u>), *Odontesthes bonariensis* (<u>HM061694.1</u>), *Oncorhynchus mykiss* (<u>Z46912.1</u>), *Oreochromis aureus* (<u>GU246726.1</u>), *Oreochromis niloticus* (<u>GU246725.1</u>), *Pelodiscus sinensis* (<u>AB480162.1</u>), *Pelteobagrus fulvidraco* (<u>HQ246723.1</u>), *Rattus norvegicus* (<u>NM 017115.2</u>), *Salmo salar* (<u>DQ294029.2</u>), *Sparus aurata* (<u>EF462191.1</u>), *Sternopygus macrurus* (<u>AY396565.1</u>).









Figure S6: Scatter plots showing distribution of muscle data points used for correlation analysis.



Figure S7: Scatter plots showing distribution of adipose tissue data points used for correlation analysis.