Seasonal variation in expression pattern of genes under HSP70

Seasonal variation in expression pattern of genes under HSP70 family in heat- and cold-adapted goats (*Capra hircus*)

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Abstract Heat shock protein 70 (HSP70) is one of the most abundant and best characterized heat shock protein family that consists of highly conserved stress proteins, expressed in response to stress, and plays crucial roles in environmental stress tolerance and adaptation. The present study was conducted to identify major types of genes under the HSP70 family and to quantify their expression pattern in heat- and cold-adapted Indian goats (*Capra hircus*) with respect to different seasons. Five HSP70 gene homologues to HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2 were identified by gene-specific primers. The cDNA sequences showed high

similarity to other mammals, and proteins have an estimated molecular weight of around 70 kDa. The expression of HSP70 genes was observed during summer and winter. During summer, the higher expression of HSPA8, HSPA6, and HSPA1A was observed, whereas the expression levels of HSPA1L and HSPA2 were found to be lower. It was also observed that the expression of HSPA1A and HSPA8 was higher during winter in both heat- and cold-adapted goats but downregulates in case of other HSPs. Therefore, both heat and cold stress induced the overexpression of HSP70 genes. An interesting finding that emerged from the study is the higher expression of HSP70 genes in cold-adapted goats during summer and in heat-adapted goats during winter. Altogether, the results indicate that the expression pattern of HSP70 genes is species- and breed-specific, most likely due to variations in thermal tolerance and adaptation to different climatic conditions.

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Introduction

Ambient temperature plays a crucial role in determining the distribution and performance of animals. Despite having well-developed mechanisms of thermoregulation, ruminants do not maintain strict homeothermy under heat stress (Lu 1989). Heat stress is the major constraint on animal productivity in tropical climatic conditions. Growth, production, and reproduction are impaired as a result of the drastic changes in biological functions caused by heat stress.

The ability to survive and adapt to thermal stress is a fundamental requirement of cellular life. Thermal stresses trigger a complex program of gene expression and biochemical

adaptive responses (Fujita 1999; Lindquist 1986). Heat shock proteins (HSPs), a large protein family, allow cells to adapt to gradual environment changes and are considered to play crucial roles in environmental stress tolerance and thermal adaptation (Feder and Hofmann 1999; Frydenberg et al. 2003; Hoffmann et al. 2003; Sørensen et al. 2003). These proteins are highly conserved across evolutionary lines (Lindquist 1986; Parsell and Lindquist 1993) and represent between 2–15 % of total cellular proteins expressed by all living organisms (Morimoto et al. 1994). Their expression follows both constitutively expressed and inducible patterns. Classified according to their molecular weight (Kregel 2002), one of the most abundant and the best characterized HSP is the 70-kDa family (HSP70), which has both inducible (HSP72) and cognate (HSP73) forms (Kiang and Tsokos 1998). Changes in gene expression in response to heat shock that are associated with acclimation include elevation of the constitutive HSP system and reduced threshold for systemic heat shock responses (Horowitz 2001). The mechanism by which the HSPs confer stress tolerance is not completely understood but may relate to the important role of HSPs in the processing of stress-denatured proteins (Mizzen and Welch 1988).

Goat, a small ruminant, is popularly known as the “poor man's cow” because of their immense economic importance (Mac Hugh and Bradley 2001). These animals are found in almost every corner of the world as they are well adapted under different geographical and environmental conditions including extreme and harsh climates. In India and other developing countries, goat is mainly reared by the small and marginal farmers, including landless agricultural laborers (Rekib 1998). Goats adapted to a harsh environment perform better than other domesticated ruminants (Devendra 1990; King 1983; Shkolnik and Silanikove 1981). These animals have developed adaptive mechanisms that allow their survival at very high temperatures (45 to 50 °C) as well as cold temperatures (−20 to −40 °C). However, despite their extreme tolerance (Al-Tamimi 2007), the productivity of these animals often declines due to thermal stress. However, a large number of information is available on physiological changes in heat and cold stress, but information in cellular and genetic levels are still very few. Elucidating the structure and expression profiling of major HSP70 genes might provide some insight into the mechanisms underlying the heat and cold adaptation in goats. The information generated will be of immense value for understanding the adaptive response of goats to heat and cold shock and also for the development of therapeutics, since adaptation to one stress often leads to cross protection. Therefore, the present study was undertaken to identify the major HSP70 genes and their expression pattern in heat- and cold-adapted goats (*Capra hircus*).

Materials and methods

Location, animals, and experimental protocol

All the experiments were carried out at the National Dairy Research Institute, Karnal, India. Two heat-adapted (Sirohi and Barbari) and two cold-adapted breeds (Gaddi and Chegu) were selected for the study. Sirohi and Barbari goat breeds are well adapted to the dry hot climatic conditions of Indian states of Rajasthan, Uttar Pradesh, and Gujarat, whereas Gaddi and Chegu breeds are found at high altitude and cold climatic conditions of Himachal Pradesh state. Sampling from the Sirohi and Barbari goat breeds was conducted at the Central Institute for Research on Goats (Mathura, India; 27°10'N, 70°02'E) and from Gaddi and Chegu breeds at Himachal Pradesh Krishi Viswavidyalaya (Palampur, India; 32.12°N, 76.53°E).

Twenty nonlactating and nongestating females (doe) from each breed were randomly selected for this study. The goats were 1.5 to 2 years old and had an average body wt of 25–30 kg. All the animals were clinically healthy and free from any physical or anatomical abnormalities. The experiments were carried out during three distinct phases coinciding with three seasons of the year, viz. winter (mid-December–mid-February), spring (mid-February–mid-April), and summer (mid-April–mid-June). All the animals were closely monitored and were provided similar managerial inputs during experimental period. The data on meteorological variables (ambient temperature, relative humidity) were recorded by weather station installed at CIRG, Mathura and HPKV, Palampur. Temperature humidity index (THI) was calculated from dry and wet bulb temperature by using the following formula (Johnson et al. 1963).

$$\text{THI} = 0.72 (\text{Dbt} + \text{Wbt}) + 40.6$$

where Dbt is the dry bulb temperature in degrees Celsius, and Wbt is the wet bulb temperature in degrees Celsius.

Blood collection and RNA extraction

About 10 ml of blood samples was collected from each animal in lithium heparin-coated vacutainer tubes (BD Biosciences, USA) under aseptic conditions and was immediately transported to laboratory under refrigeration. A total 240 blood samples (20 animals from four breeds in three seasons) were collected. Within a season (i.e., summer, winter, or spring), blood sample collection from all the animals in a breed was not possible in a single day, so samples were collected in three consecutive days in each breed in a season, and the environmental conditions in each date of collection were recorded and considered during data analysis. Total RNA from blood was extracted using Trizol method. The

concentrations were measured in NanoDrop (Thermo Scientific, USA), and purity was analyzed by agarose (1.5 %) gel electrophoresis. To exclude any genomic DNA contamination, total RNA was treated with RNase-free DNase I (Kappa Biosystem, USA).

cDNA synthesis, cloning, and sequencing of HSP70 genes

The primers for amplification of full-length cDNA of HSP70 genes (HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2) were designed on the basis of prior sequence information available at National Center for Biotechnology Information (NCBI) on other mammalian species (Table S1, Supplementary data). First-strand cDNA was prepared using SuperScript III Kit (Life Technology, USA). Gene-specific PCR amplifications were carried out as per standard cycling parameters [95 °C for 2 min, (95 °C for 30 s, 56–58 °C for 30 s, and 72 °C for 1 min) × 35 cycles, and 72 °C for 10 min], and the products were analyzed by agarose (1.5 %) gel electrophoresis (Fig. 1).

Purified PCR products were inserted into the pGEM-T Easy plasmids (Promega) and amplified by culturing in XL1-Blue super competent cells. Plasmid DNA was then isolated and sequenced using forward, reverse, and internal primers in automated ABI 377 Sequencer (PerkinElmer Applied Biosystem, Foster City, CA, USA) by Xcelris Privet Limited (Ahmadabad, India). The predicted protein was done by ExPASy Translate Tool and BioEdit. The predicted molecular weight, isoelectric points (pI), and charges of the translated HSP70 proteins were estimated using Protein Calculator v3.3.

Phylogenetic analysis

For phylogenetic analysis, available sequences of the five HSPs (HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2) of different mammalian species were obtained from the NCBI protein database. Phylogenetic analyses were conducted in MEGA5.1 (Tamura et al. 2011). The evolutionary history

was inferred using the maximum parsimony method (Swofford 1998). The bootstrap consensus tree inferred from 1,000 replicates is taken to represent the evolutionary history of the taxa analyzed (Felsenstein 1985). A test of the homogeneity of substitution patterns between sequences was performed by disparity index test (Kumar and Gadagkar 2001). A Monte Carlo test (1,000 replicates) was used to estimate the *P* values (North et al. 2002), and values smaller than 0.05 were considered significant.

Real-time PCR (qPCR)

Relative expression level of mRNA transcripts of HSP70 genes was measured by quantitative real-time PCR (qPCR) in Roche LC480 using the SYBR Green I (Roche Diagnostics, Switzerland) chemistry. The primers for qPCR were designed on the basis of prior sequence information from other animals (Table S2, Supplementary data). Along with the target genes (HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2), two housekeeping (glyceraldehyde 3-phosphate dehydrogenase (GAPDH) and β -actin) genes were amplified for relative expression measurements (Fig. 2). Each sample had triple replicates, and in all cases, samples of total RNA were used as negative control. Two housekeeping genes, GAPDH and β -actin, were used in this study. Proper validation was done to determine that its expression is unaffected by the experimental treatments (heat and cold). The results indicated that, for our study, β -actin was better than GAPDH. So expression data of β -actin were used for analysis of relative expression data. Relative quantification of a target gene was done by comparing the expression levels of reference gene (β -actin), as per the method of Livak and Schmittgen (2001).

Statistical analysis

All the experiments were replicated three times. Results were expressed as the means $\pm$ SEM. A difference with value *P* <

Fig. 1 cDNA-amplified products of HSP70 genes on agarose gel (1.5 %). *I* (a) cDNA-amplified product of HSPA8. *I* (b) cDNA-amplified product of HSPA6. *I* (c) cDNA-amplified product of HSPA1A. *I* (d) cDNA-amplified product of HSPA1L. *I* (e) cDNA-amplified product of HSPA2

Fig. 2 Real-time PCR-amplified products of HSP70 genes (*HSPA8*, *HSPA6*, *HSPA1A*, *HSPA2*, and *HSPA1L*) and housekeeping genes (*GAPDH* and β -*actin*) on agarose gel (3 %)

0.05 was considered statistically significant. Data were analyzed by analysis of variance, and statistical differences between the various treatment group means were determined by Duncan's multiple range test (DMRT) using the statistical Product and Service Solutions, Version 17.0.1 software (SPSS Inc., Chicago, IL, USA).

Results

Climatic conditions

The average climatic parameters recorded during the experimental periods are presented in Table 1. In both the places during spring, the THI was below 72, reflecting that the animals were in thermoneutral condition. During summer, THI was above 72, representing heat stress, whereas during winter, THI was nearly 50 in both places, indicating cold stress.

Identification of HSP70 gene homologues in goat

Five HSP70 cDNAs (*HSPA8*, *HSPA6*, *HSPA1A*, *HSPA1L*, and *HSPA2*) were cloned and sequenced. *HSPA8* cDNA was 1,953 bp long and encoding 650 amino acids. The cDNA sequence exhibited the highest identity and similarity to heat shock 70-kDa protein 8 (*HSPA8*) from cattle (97 %), followed by pig (92 %), horse (91 %), and human (90 %) as well as substantial homology with 70-kDa heat shock proteins from

many other species (Table S3, Supplementary data). *HSPA6* cDNA was 1,912 bp long, encoded for 637 amino acids and showed substantial identity and homology to 70-kDa heat shock protein (*HSPA6*) from many different organisms, e.g., cattle (97 %), camel (92 %), horse (91), pig (90 %), human (90 %), etc. (Table S3, Supplementary data). Similarly, other three genes, i.e., *HSPA1A*, *HSPA1L*, and *HSPA2* cDNA, were 1,926, 1,926, and 1,911 bp long, encoded for 641, 641, and 636 amino acids, respectively. The amino acid translation of these fragments exhibited the highest identity and similarity to heat shock 70-kDa protein 1A (*HSPA1A*), heat shock 70-kDa protein 1-like (*HSPA1L*), and heat shock 70-kDa protein 2 (*HSPA2*), respectively, from the cattle, horse, pig, and human as well as substantial homology with 70-kDa heat shock proteins from many other animals (Table S3, Supplementary data). The sequences of HSP70 genes (*HSPA8*, *HSPA6*, *HSPA1A*, *HSPA1L*, and *HSPA2*) in goat were submitted in NCBI GenBank with accession nos. JN616380, JN616381, JN656104, JN656105, and JN656106, respectively. The predicted molecular weights of HSP70 proteins estimated from amino acid sequences by using Protein Calculator v3.3 showed that the proteins have an estimated molecular weight around 70 kDa and with pI around 6.0 (Table S4, Supplementary data).

Phylogenetic analysis of mammalian HSPs (Fig. 3) showed three different clusters with *HSPA2* and *HSPA8* in one group, and *HSPA1A* and *HSPA6* in another. *HSPA1L* formed a separate cluster. All goat HSPs were found to be grouped with other ruminants and porcine species, except *HSPA6*, where

Table 1 Environmental condition during the experimental period

	Place	Season	Average temperature (°C)	Relative humidity (%)	THI
Winter (mid-December to mid-February), spring (mid-February to mid-April), summer (mid-April to mid-June)	Mathura	Winter	11.98	63.5	51.38
		Spring	22.8	59.4	69.78
		Summer	42.10	45.0	93.81
	Palampur	Winter	9.84	59.0	48.87
		Spring	19.6	52.2	64.91
		Summer	32.10	60.5	83.13

ruminants and porcine species were grouped differently. Monte Carlo analysis for the test in homogeneity of substitution patterns revealed that the HSPA6, after its divergence from other HSPs, has evolved independently in mammals (Table S5, Supplementary data). A similar trend was also found for sheep and goat HSP1L that was found to evolve differently than cattle (*Bos taurus*). The substitution patterns of amino acids of other HSPs among different mammals were found to be nonsignificant.

Relative expression profile of HSP70 genes

The expression of HSP70 genes (HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2) showed temperature sensitivity and seasonal variation. Relative expression of HSP70 genes varied markedly among the heat- and cold-adapted goat breeds with a moderate variation between breeds and showed a good response to increased or decreased ambient temperature (Fig. 4). Statistical analysis revealed a significant variation between different seasons ($P < 0.01$) for all HSP70 gene expression. The expression of HSPA8, HSPA6, and HSPA1A was higher, but the relative mRNA expression of HSPA1L and HSPA2 was very low. Results indicated that the expression level of HSPA8 and HSPA1A was higher during both winter and summer. The expression level of HSPA6 and HSPA1L was higher only during summer. HSPA2 was observed to be downregulated during the summer and winter seasons. During summer, the relative expressions of all the HSP70 genes were higher in cold-adapted breeds (Gaddi and Chegu) than the heat-adapted breeds (Sirohi and Barbari) (Fig. 4). In Chegu, i.e., a cold-adapted breed, the expression of all the genes was higher as compared to the other cold-adapted breed of goat, Gaddi, during summer. The HSPA8 and HSPA1A expression during winter in heat-adapted breeds was observed to be higher than cold-adapted breeds.

Discussion

The cloning and sequencing of five full-length cDNAs encoding the HSP70 family genes (HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2) in goat were carried out for the first time. The BLAST results indicated that the goat HSP70 gene sequences were highly conserved and shared a high similarity (up to 91 %) with heat shock proteins of other mammals. The highest nucleotide sequence identities of goat HSP70 genes were observed with cow and pig followed by horse, camel, and human, indicating a close evolutionary relationship among HSP70 family genes. Similarly, high homology with other HSP70 genes has also been studied with the HSP70 gene from goat (Gade et al. 2010), where maximum identity with cow HSP70 (97.8 %) was observed.

Fig. 3 Phylogenetic relationship of five HSP70 (HSPA8, HSPA6, HSPA1A, HSPA1L, and HSPA2) amino acid sequences from different mammalian species obtained from the NCBI protein database using MEGA5.1

The phylogenetic analysis of mammalian HSPs indicated that goat HSPs have close relationship with other ruminants including cattle (*Bos indicus* and *B. taurus*), buffalo, and sheep. They share homology with pig, camel, and horse, but are fairly distant from HSPs of human, chimpanzee,

Fig. 4 Relative expression profile of HSP70 genes by real-time PCR during winter and summer in comparison to spring in heat-adapted (Sirohi and Barbari) and cold-adapted (Gaddi and Chegu) goats. Relative quantities of mRNA were normalized to beta actin. Values are the means $\pm$ SEM of three experiments performed. Columns with different superscript letters (a, b, c) indicate significant difference ($P < 0.05$). Statistical differences between various treatment group means were determined by Duncan's multiple range test

orangutan, dog, and mouse. The amino acid substitution pattern of different HSPs indicated that HSPA6 and HSPA1L have evolved independently in mammals compared to other HSPs. It will be interesting to find the biological significance of this divergence of HSP homologs in different mammalian species.

During summer and winter, animals were exposed to ambient temperature beyond the comfort zone, so they were under thermal stress in both the seasons. The findings of the present study indicated that HSP70 genes were expressed in both heat- and cold-adapted goats during summer as well as winter. The expression level of HSP70 genes, except HSPA2,

was observed to be higher during summer in both heat- and cold-adapted breeds than that of winter. Our findings are in accordance with those of previous studies where heat stress-induced HSP70 expression was observed in bovine lymphocytes (Lacetera et al. 2006; Patir and Upadhyay 2007, 2010). Among all HSP70 genes studied, a significant increase was observed with respect to HSPA1A, HSPA6, and HSPA8. Similarly, heat stress-induced upregulation of HSPA8, HSPA6, and HSPA1A gene expression was observed in human blood (Sonna et al. 2002, 2004). In Brown Swiss cattle, increased HSP72 or HSPA1A mRNA levels in peripheral blood mononuclear cells due to heat stress have also been reported by Lacetera et al. (2006). Our results were also supported by earlier studies in camel showing that temperature elevation increases the level of constitutively expressed HSP70 in camel (Garbuz et al. 2011; Ulmasov et al. 1993).

A nonsignificant upregulation of HSPA1A and HSPA8 was also observed in response to cold exposure during winter. It is not surprising to see the HSP70 upregulation in response to cold as the cold-induced denaturation of proteins is a general phenomenon (Kostal and Tollarová-Borovanská 2009; Privalov 1990; Tsai et al. 2002), and in turn, partially denatured or misfolded proteins are potent triggers of rapid HSP70 accumulation (Feder and Hofmann 1999; Welch 1993). An earlier study on *Drosophila* suggested that the induction of HSPs following cold exposure (0 °C) was due to the shift from 0 °C back to the control temperature of 25 °C, i.e., due to heat shock, rather than to the cold treatment itself (Burton et al. 1988). Other studies on mammalian species observed that the induction of HSP70 gene expression after cold treatment (4 °C) occurred upon recovery at control temperature (37 °C). The magnitude and the kinetics of the response were, however, related to the duration of cold stress (Liu et al. 1994). However, HSPs have been implicated in cold survival, but this link has not been established conclusively (Michaud and Denlinger 2004). Therefore, further studies are highly required to differentiate between the cold exposure and the subsequent recovery phase in order to establish the role of HSPs during cold stress in goats. Low temperature affects many different structures and processes simultaneously, and therefore, cold tolerance is a highly complex adaptation and must be explained as a combination of different mechanisms (Danks 1996; Lee 1991; Murray et al. 2007). From the findings of this study, we suggest that the HSP70 is closely related to the cold tolerance in animals.

Increase of HSP70 gene expression during summer was observed to be higher in cold-adapted goat breeds (Gaddi and Chegu) as they experience higher stress during summer than heat-adapted goats (Sirohi and Barbari). In contrast, during winter, the expression of HSP70 genes was higher in heat-adapted goats than the cold-adapted goats. In different species of *Drosophila*, decreased HSP70 expression after acute heat stress seems to be the evolutionary consequence if the lines are

exposed to chronically stressful high-temperature conditions or originate from natural populations that inhabit warm environments (Sørensen et al. 2003). The heat-adapted goats are adapted to dry hot environment in Rajasthan and parts of the U.P. region in India, while the cold-tolerant breeds are native to the relatively cold and hilly regions (Jammu-Kashmir and Himachal Pradesh) of India. Stress-induced HSP accumulation has been observed to be associated with thermotolerance and the ability to survive otherwise lethal heat stress. The difference observed in relative expression of HSP70 genes in cold-tolerant and heat-tolerant goats could be attributed to breed difference and their adaptation to different environmental conditions. The ability of the HSPs to confer thermotolerance in both cultured cells and in animals is well documented (Li 1985; Weshler et al. 1984). Stress tolerance is a complex characteristic, and the mechanism of stress tolerance by HSPs is not yet completely understood, but according to Mizzen and Welch (1988), this may be related to the important role of HSPs in the processing of stress-denatured proteins.

Among the two cold-adapted breeds, Chegu is more susceptible to heat stress than Gaddi breed because Chegu is adapted to higher altitude and low temperature. Both heat-adapted breeds, Sirohi and Barbari, showed almost similar pattern of expression, but the expression level in Sirohi goat was observed to be relatively lower. There is a growing body of literature supporting the role for HSPs in the whole organism's adaptation to heat other than through thermotolerance. A survey of lizard species inhabiting a variety of environments, including highlands, forests, and deserts, demonstrated a remarkable diversity of constitutive HSP70 levels that were correlated with the lizard's environmental temperatures (Ulmasov et al. 1992). The present study demonstrates that though HSP70 genes are highly conserved across evolutionary lines, the HSP70 gene expression is species- and breed-specific. The species-specific differences in HSP70 isoforms are most likely due to variations in thermal tolerance (Yamashita et al. 2004), and isoform expression may vary with regard to thermotolerance (Hightower et al. 1999). The role of HSP70 in thermal tolerance and adaptation to heat and cold should be uncovered in the future by exploring various cellular and animal models, enabling to directly investigate the interactions between these vitally important genomic loci.

Conclusions

The sequences of HSP70 genes in goat are well conserved and have similarities with other mammalian species. The major findings that emerged from this study include the higher expression of HSPA1A and HSPA8 and lower expression of HSPA1L and HSPA2 during both winter and summer seasons, whereas the expression of HSPA6 was higher only during summer. The expression level of HSP70 genes was higher in

cold-adapted goats, and expression during winter was higher in heat-adapted goats. Regardless of reasonably high evolutionary conservation, the expression of HSP70 isoforms is species-specific, most likely due to variations in thermal tolerance. HSP70 gene expression can be used as a marker for thermal adaptation in different species, and information generated will have significant implications in the development of strategy to cope up with the challenges of climate change.

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