

Identification of a Highly Conserved Pro-Gly Doublet in Non-animal Small Heat Shock Proteins and Characterization of Its Structural and Functional Roles in *Mycobacterium tuberculosis* Hsp16.3

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Abstract—Small heat shock proteins (sHSPs) are highly divergent in primary sequences, with short conserved motifs found in various subfamilies. Here a Pro-Gly doublet was found to be conserved in most non-animal sHSPs by sequence analysis of a total of 344 unique sHSPs (covering the subfamilies: bacterial class A, bacterial class B, archae, fungi, plant, and animal) placed in data banks. In contrast, the residues corresponding to this Pro-Gly doublet in most of animal sHSPs are often charged. Site-directed mutagenesis studies of *Mycobacterium tuberculosis* Hsp16.3 replacing the Gly (at position 59) residue by Cys or Trp demonstrate that this Gly is likely involved in subunit interactions, which is consistent with that in *Methanococcus jannaschii* Hsp16.5 and wheat Hsp16.9. Our data suggest that this Pro-Gly doublet in Hsp16.3 is not directly involved in binding of denatured substrate proteins, whereas the corresponding charged residues in bovine α -crystallin were instead proposed before to be involved in substrate binding. These observations indicate that the highly conserved Pro-Gly doublet is critical to discriminate between non-animal and animal sHSPs.

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Small heat shock proteins (sHSPs), as one subclass of molecular chaperones, have been found in almost all organisms [1]. Studies performed *in vitro* have demonstrated that sHSPs are able to exhibit chaperone-like activities and even to enhance the refolding of denatured proteins with help from other chaperone systems, such as Hsp70/Hsp40 [2-4]. In primary sequence comparison, sHSPs are characterized by having a relatively conserved α -crystallin domain [5-7]. Nevertheless, sHSPs have diverged much more during evolution than the large HSPs like Hsp90, Hsp70, and Hsp60 [1, 5, 7].

Conserved sequence motifs have been revealed in the whole family or various subfamilies of sHSPs. Such

motifs conserved in the whole family include the IXI/V motif in the C-terminal extension [8] and consensus I region represented by the sequence of Pro-(X₁₄)-Gly-Val-Leu in the C-termini of the α -crystallin domain [7, 9-11], those conserved in plant sHSPs include the consensus II region represented by the sequence of Pro-(X₁₄)-X-Val/Leu/Ile-Val/Leu in the N-termini of the α -crystallin domain [9, 10], and those conserved in animal sHSPs include the sequence of SRLFDQFFG in the N-terminal region [5, 11]. Biochemical studies have implicated important structural and functional roles of these motifs for sHSPs [8, 12-14].

The Pro-Gly doublet, located at the N-terminal end of the α -crystallin domain, was previously revealed to be conserved in plant and bacterial sHSPs [11, 15]. Here the sequence analysis of a total of 344 unique sHSP sequences demonstrates that this Pro-Gly doublet is actually highly conserved in almost all non-animal sHSPs. Site-directed mutagenesis studies through *Mycobacterium tuberculosis* Hsp16.3 demonstrate that the Pro-Gly doublet is very likely involved in subunit

Abbreviations: DTT) dithiothreitol; DTNB) 5,5'-dithiobis(2-nitrobenzoic acid); sHSPs) small heat shock proteins; L57C, G59C, and D61C) the Hsp16.3 mutant proteins with Leu57, Gly59, and Asp61, respectively, substituted by cysteine; G59A and G59W) the Hsp16.3 mutant proteins with Gly59 substituted by alanine and tryptophan, respectively.

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interaction but is not directly involved in binding of denatured substrate proteins. These observations will enrich our insights into the evolution of sHSPs from the perspective of protein structure, function, and action mechanism.

MATERIALS AND METHODS

Materials. Dithiothreitol (DTT), 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB), and insulin were all obtained from Sigma (USA). MutantBest Kit was purchased from TaKaRa Biotechnology (Dalian, China). DEAE-Sepharose FastFlow and Q-Sepharose High Performance for chromatography were obtained from Amersham Pharmacia Biotech (Sweden). All other chemical reagents were of analytical purity.

Sequence analysis. All the available amino acid sequences of sHSPs were downloaded from the mirror website of SWISSPROT and TREMBL databases in China (available at <http://cn.expasy.org>). The fragments and the duplicated sequences were discarded manually and the remaining 344 unique proteins, named by either SWISS_PROT or TREMBL accession number, were grouped as bacteria class A and B [16], archaea, fungi, plant, animal, and other non-animal eukaryotes. The 344 amino acid sequences were aligned using Clustalx (version 1.7).

Plasmid construction and protein purification. The single-site mutants of Hsp16.3 (L57C, G59C, D61C, G59A, G59W) were constructed according to methods described before [17]. The wild type, L57C and D61C mutant Hsp16.3 proteins were purified according to procedures previously described [17, 18]. For the purification of the G59C mutant Hsp16.3 protein, the supernatant of cell lysates was loaded onto a DEAE-Sepharose FastFlow column pre-equilibrated with 50 mM imidazole-HCl (pH 6.5) and was then eluted with a salt gradient of 0.15–0.35 M NaCl. The fractions containing G59C were pooled, dialyzed against 20 mM Tris-HCl (pH 8.5), and loaded onto a Q-Sepharose high performance column before elution with a salt gradient of 0.15–0.4 M NaCl. The G59W and G59A mutant proteins were purified similarly, with both being eluted with a salt gradient of 0.12–0.25 M NaCl from the DEAE column, and the G59W was eluted with a 0.15–0.3 M and G59A with a 0.15–0.4 M salt gradient from the Q-Sepharose column. The purified proteins were dialyzed in deionized water, lyophilized, and stored at -20°C before further analysis. Protein concentrations were determined using the Bio-Rad Protein Assay.

DTNB modification. The exposure status of Cys59 in the G59C mutant Hsp16.3 protein was detected by using DTNB modification as described previously [19]. The optical density of the mixture of protein (1 mg/ml), urea at various concentrations, and DTNB (0.2 mM), dis-

solved in 50 mM sodium phosphate buffer (pH 7.0), was recorded on a UV-8500 spectrophotometer (Shanghai Techcomp, China) at 412 nm.

Tryptophan intrinsic fluorescence assay. The intrinsic fluorescence of tryptophan in the G59W mutant Hsp16.3 protein was scanned between 300–400 nm while excited at 279 nm on a Hitachi F-4500 fluorescence spectrophotometer, which was connected to a water bath (Boyikang Inc., China). To monitor the urea-induced exposure of tryptophan, the G59W mutant protein (0.2 mg/ml) unfolded by various concentrations of urea from 0.2 to 7 M for 15 h was subjected to fluorescence assay. To monitor the heat-induced exposure of tryptophan, the G59W protein (0.1 mg/ml) was incubated (ranging from 25 to 73°C) at each temperature for 5 min before fluorescence assay was performed.

Chemical cross-linking by glutaraldehyde. Glutaraldehyde (0.5%) and Hsp16.3 proteins (0.2 mg/ml) were reacted at 25°C in 50 mM sodium phosphate buffer (pH 7.0) for 1 or 10 min. The reactions were stopped by quenching with 1 M Tris-HCl for 10 min. The cross-linked samples were analyzed by SDS-PAGE (12%).

Chaperone-like activity assay. Chaperone-like activities of the Hsp16.3 wild type and mutant proteins (0.3 mg/ml, wild type, G59A, or G59W) were assayed by measuring their capacity to suppress the DTT-induced (20 mM) aggregation of insulin (0.3 mg/ml) at different temperatures (from 25 to 65°C). The measurements were performed with the UV-8500 spectrophotometer as described previously [20, 21].

RESULTS

Sequence alignment reveals a Pro-Gly doublet highly conserved in non-animal sHSPs. The sequence alignment of 344 sHSPs unanimously identified a highly conserved doublet Pro-Gly in the non-animal sHSPs (data presented in Table 1, Fig. 1a, and supplementary data in Fig. 2). The 344 unique sHSP sequences were obtained from the SWISSPROT and TREMBL databases, covering the seven subfamilies of sHSPs, i.e., bacterial class A (32 sequences), bacterial class B (64), archaea (35), fungi (10), plant (115), animal (79), and other eukaryotes (9) (see supplementary data in Table 2). The corresponding positions of the Pro-Gly doublet are, instead, often occupied by charged residues in animal sHSPs (Table 1 and Fig. 2). One exception among the non-animal sHSPs is that most of the members of the bacteria class A sHSPs have an Ala at the position of the highly conserved Pro (Fig. 2), somehow suggesting a distinct evolutionary history of bacterial class A sHSPs from the rest of the non-animal sHSPs. Additionally, the sequence alignments (Fig. 2) also revealed that the two residues sandwiching the Pro-Gly doublet in almost all the sHSPs are hydrophobic.

Table 1. The conserved Pro-Gly doublet in non-animal sHSPs^a

Subfamily	Bacterial class A	Bacterial class B	Archaea	Fungi	Plant	Animal	Other eukaryotes ^b
Total number	32	64	35	10	115	79	9
Pro	4 ^c	64	32	10	114	1 ^d	9
Gly	32	64	33	10	113	5 ^d	9

^a The data in this table were obtained from the sequence alignment results displayed in Fig. 2.

^b These eukaryotes contain three kingdoms: Alveolata (seven sequences), Entamoebidae (one sequence), Rhodophyta (one sequence).

^c The corresponding residue is Ala, instead of Pro, in 26 out of 32 members of bacterial class A sHSPs.

^d The residues corresponding to Pro and Gly are charged residues, respectively, in 38 and 51 out of 79 members of animal sHSPs.

Inter-subunit disulfide bond is formed spontaneously in the G59C mutant Hsp16.3 protein having a buried Cys59 residue. What are the structural and functional roles of the highly conserved Pro-Gly doublet in non-animal

sHSPs? Hints were first sought by examining the spatial positions of this Pro-Gly doublet in the crystal structures of archaea *M. jannaschii* Hsp16.5 and plant wheat Hsp16.9 [22, 23]. This reveals that they are located in a

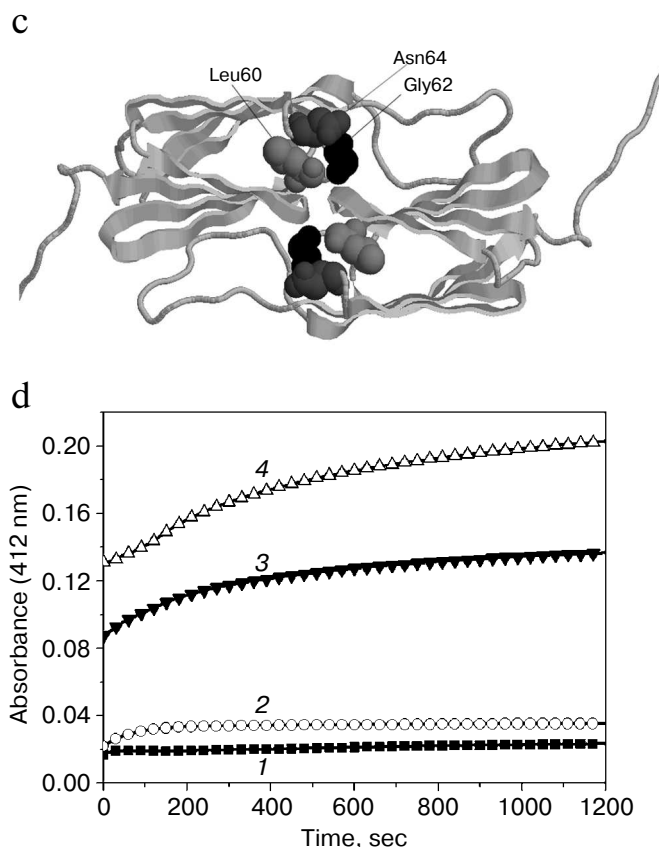
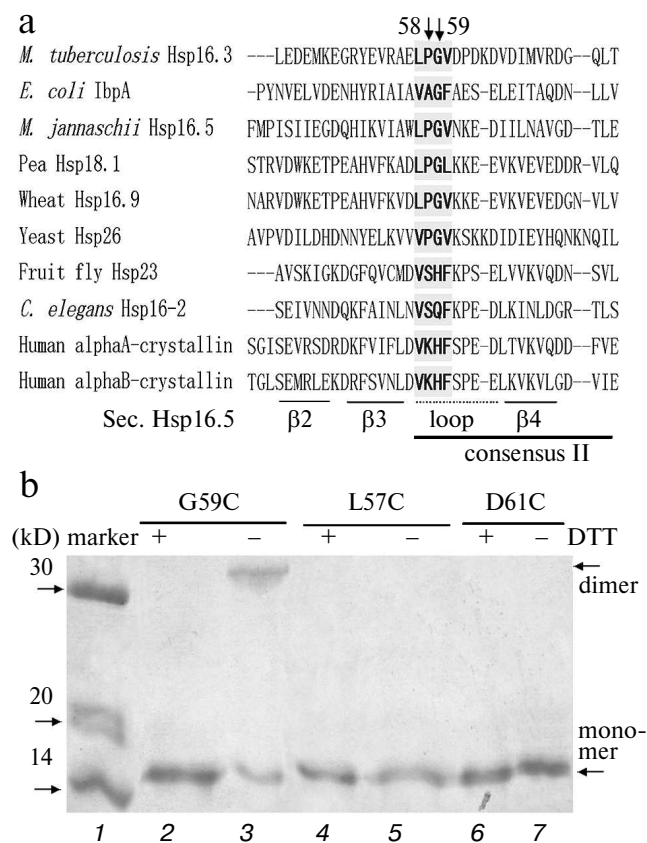


Fig. 1. Characterization of the G59C mutant Hsp16.3 protein. a) Sequence alignment of representative sHSPs, covering bacterial class B (*M. tuberculosis* Hsp16.3), bacterial class A (*E. coli* IbpA), archaea (*M. jannaschii* Hsp16.5), plant (pea Hsp18.1 and wheat Hsp16.9), fungi (yeast Hsp26), and animal (the other four sequences). For simplicity, only a part of the residues are shown here. The secondary structure elements were arbitrarily assigned according to the crystal structure of *M. jannaschii* Hsp16.5 [22]. The consensus II region identified in plant sHSPs [10] is shown here. b) SDS-PAGE analysis results for the G59C, L57C, and D61C mutant Hsp16.3 proteins with DTT added in half of the loaded samples (lanes 2, 4, and 6). Positions of the dimers (lane 3) generated by the spontaneous formation of disulfide bonds in the G59C mutant protein and monomers are indicated by the arrows on the right. c) The crystal structure of Hsp16.5 dimer (cartoon model). The black, dark gray, and light gray balls are Gly62, Asn64, and Leu60 in Hsp16.5, respectively, which correspond to Gly59, Asp61, and Leu57 in Hsp16.3, respectively. d) The time-dependent change in the absorption (at 412 nm) of DTNB (added to a final concentration of 0.2 mM) during its reaction with sulfhydryl groups of the G59C mutant Hsp16.3 protein (1 mg/ml) monitored at room temperature in the presence of different urea concentrations (0, 1, 4, and 8 M urea for the curves 1, 2, 3, and 4, respectively).

bacterial class A			bacterial class B			archaea		fungi	
HSPA_BRAJA	ALAGFS	HS19_RICCN	EVPGFD	Q8PBS2	DLPGID	HSPS_METJA	WLPGVN	HS16_SCHPO	ELPQVK
HSPB_BRAJA	AVAGFS	HS19_RICPR	EVPGFD	Q87C16	DLPGID	Q9C4M2	ELPQVR	HS26_YEAST	VVPQVK
HSPD_BRAJA	AVAGFS	ASP1_STRTR	ELPGIP	Q82T55	DLPGVK	059514	ELPGVR	HS30_EMENI	ELPGIP
HSPH_BRAJA	ALAGFS	052190	ELPGIP	Q83CE9	DIPGVD	Q9YFZ9	DLPGHM	HS30_NEUCR	ELPGID
HSPF_BRAJA	ALVGFY	Q9X9C3	ELPGIP	Q9ZFD1	EIPGID	Q9YAJ0	DIPGAN	HS42_YEAST	ALPGAN
IBPA_ECOCI	AVAGFA	ASP2_STRTR	ELPGIP	Q9WYK7	EIPGID	Q9HSS6	DLPQVE	Q8NJ78	ELPGSK
Q8XG77	AVAGFA	Q9X9N3	ELPGIP	Q98ITS	EIPGME	Q9HSD8	DTEGAD	Q8TG38	ELPGID
Q7VQU7	VVAGFM	Q81QE9	DLPGFO	Q92V80	ELPGLD	Q9HP75	DVAGVT	Q74984	EVPQAD
Q87TQ1	AVAGFS	Q81DY2	DLPGFO	Q98IT6	EIPGLD	Q9HP50	EVPQFS	Q7S0S7	EVPQAD
Q7M9J3	AVAGFA	HS16_ONYPE	ELPGFK	P70919	ELPGLE	Q9HKX2	ELPGVS	Q7RW96	ELPGVD
Q9KXV0	AVAGFG	Q8G6R2	DMPGFK	HSPC_BRAJA	ELPGLE	Q9HJY9	EMPGFD		
Q8EET9	AVAGFS	HS18_CLOAE	DLPGVK	069243	ELPGIE	Q9HHW1	EMPGFD		
Q99H48	AVAGFS	Q890A7	DVPGID	HSPF_BRAJA	ELPGIE	Q97W19	EVPQVN		
Q8FXL4	AVAGFS	Q88SP7	DIPGID	Q92Z87	ELPGME	Q97VL9	DLAGFN		
Q8U943	AVAGFD	14KD_MYCTU	ELPGVD	Q8R745	ELPGVD	Q97AM1	EMAGFD		
Q89CQ6	AVSQA	Q8KY67	EIPGVD	Q9KXF9	DIPGLE	Q979Z6	ELPGVS		
Q92VA9	SVAGFA	18K1_MYCTT	DLPGID	067316	DLPGVK	Q974V6	EVPQVN		
Q8UKP8	AVAGFE	18K2_MYCLE	DLPGIK	Q7NYU5	ELPGVG	Q9ZV10	DMPGIR		
Q98TI5	AVAGFG	18K2_MYCTT	DLPGID	Q8AAA0	AAPGMT	Q8ZTS4	DMPGLE		
Q98P38	ALAGFS	18K1_MYCAV	DLPGID	Q8A8R3	AAPGMT	Q8ZTN4	DMPGVE		
Q8ZL03	ALAGFR	18K2_MYCAV	DLPGVN	Q7NH34	LVPGLD	Q9P9K5	ELPQVR		
Q8Z2L8	ALAGFR	Q82QP6	DLPGVS	Q88U58	ELPGFD	028308	DLPGFS		
IBPB_ECOCI	ALAGFR	Q82Q29	DLPGID	HS15_LEPIN	DLPGVE	028973	EMPGVN		
IBPB_AZOVI	AAAGFQ	HS18_STRAL	DLPGVD	Q98DGO	DLPGVG	026947	DLPGVK		
Q9AMN8	AAAGFE	Q7WSY3	DLPGVD	053673	ELPGID	Q9V1L0	ELPGVR		
Q99GS9	AVAGFA	Q9RVB5	DIPGVK	Q7U3R4	ELPGVQ	Q8TUD5	DLPGIN		
Q8UIC2	AVAGFS	Q8Z017	ELPGID	Q81DE3	ELPGIQ	Q8TK35	DLPGIN		
Q89WG1	AVAGFT	Q8DKI6	ELPGMD	Q8EMZ6	ELPGYT	Q8THE4	EPPEIE		
IBP_BUCAL	SIPPGYQ	Q8F5V3	LIPGID	YOCM_BACSU	YLPQYR	Q8THE3	EMPGIE		
IBP_BUCAP	SIPPGYE	Q7UF81	ELPGFN	COTP_BACSU	EMPGVY	Q8TZC8	EVPGAR		
IBP_BUCTS	SVPGYL	SP21_STIAU	DLPGVD	Q7WZ1	ALPGVA	Q8PEK9	DLPGID		
Q8ZPY6	SVPGVK	Q8PNC2	DLPGID	Q9RTR5	DVPGVD	Q8PXG4	EPPEIE		
						Q8PXG3	EMPGID		
						Q8PX03	DLPGIN		

plant			animal		
HS11_ARATH	DVPGLK	HS2M_SOYEN	DMPGLA	Q39930	DLPGLK
HS11_CHERU	DLPGVK	HS41_PEA	DVPGLK	Q23640	DMPGLS
HS11_DAUCA	DLPGLK	HS41_SOYEN	DMPGLK	Q8L470	DMPGHT
HS11_HELAN	DLPGMK	Q8W541	DLPGLK	Q40852	DMPGLK
HS11_LYCES	DLPGLK	Q43543	DMPQVR	Q94KN0	DMPGLG
HS11_MEDSA	DLPGLK	Q8L7T2	DMPGHT	Q9LS37	EVPGLN
HS11_ORYSA	DLPGVK	Q40698	DVPGLK	Q96458	DLPGVK
HS11_PEA	DLPGLK	Q9ZSX9	DLPGLK	Q43544	DMPGVG
HS11_SOYBN	DIPGLK	Q82545	DMPGLK	Q9ZSY1	DLPGLT
HS11_WHEAT	DLPGVK	P93440	DVPGLK	Q943E7	DLPGVK
HS12_ARATH	DLPGLR	P93439	DVPGLK	Q80432	DMPGLD
HS12_DAUCA	DLPGLK	Q40057	DLPGVK	Q9ZSZ0	DLPGLK
HS12_MEDSA	DLPGMK	Q96489	DMPGLK	Q48398	DMPGLS
HS12_ORYSA	DVPGLK	Q9SRD6	RIPDIV	Q9SE11	DMPGLG
HS13_ARATH	DLPGLK	Q64564	YLPQVD	Q943Q3	DLPGVR
HS13_SOYBN	DIPGLK	Q9LGS1	DMPGLK	Q8W007	DVPGLK
HS14_SOYBN	DIPGLK	Q9M566	DMPGLD	Q8LCB4	DMPGLS
HS15_SOYBN	DIPGLK	Q82149	DVPGLK	Q943E6	DLPGVK
HS16_SOYBN	DIPGLK	Q9XGS6	DMPGLK	Q8LMQ7	QVPDDC
HS21_ARATH	DMPGIK	Q40056	DLPGVK	Q81961	DMPGLS
HS21_HELAN	DMPGLK	Q40826	DMPGLK	Q8H288	DLPGVK
HS21_MAIZE	DMPGLG	Q39820	DVPGLK	Q8H1A6	DLPGLK
HS21_PEA	DMPGVK	Q9ZP25	DMPGLG	Q8GWH1	DMPGIK
HS21_PHANI	DMPGLK	Q9ZW31	DLPGLK	Q8GV39	DMPGLS
HS21_SOYBN	DMPGLK	Q64960	DMPGLG	Q8GV37	DMPGLS
HS22_MAIZE	DMPGLG	Q49710	DLPGLR	Q8GV36	DMPGLS
HS22_PHANI	DMPGVK	Q24082	DMPGLK	Q8GV35	DMPGLS
HS23_MAIZE	DMPGLS	Q9ZSX7	DVPGLO	Q81822	DMPGLK
HS2C_ARATH	DMPGLS	Q39819	DLPGMK	Q9FHQ3	NVPGYN
HS2C_CHERU	DMPGLA	Q40847	DMPGLG	Q9LW00	DLPGMK
HS2C_CHLRE	DAPGMG	Q9M6K2	DVPGLK	Q38806	DIPGLK
HS2C_LYCES	DMPGLS	Q43545	DMPGLE	Q84J50	DVPGLK
HS2C_PEA	DMPGVK	Q9ZSX8	DMPGLK	Q24247	DMPGLK
HS2C_PETHY	DMPGLS	Q9ZP24	PMPGLT	Q9SYU9	DLPGVK
HS2C_SOYBN	DMPGLA	Q9ZSY0	DLPGLK	Q9SYV0	DLPGVK
HS2C_WHEAT	DMPGLS	Q9SE12	DMPGLS	Q9SYU8	DLPGVK
HS2M_ARATH	DMPGLS	Q41028	DLPGVK	Q9FGM9	DMPGLS
HS2M_PEA	DMPGLG	Q49962	DLPGLK	Q7XY72	DLPGLK
				Q7XS46	DLPGFA

other non-animal eukaryote	
Q9BHCO	ELPGFS
Q9EMFO	ELPGFS
Q8IES0	DVPGLN
Q8TB02	DMPGFK
Q7YWK6	DMPGFK
Q7RS01	DMPGFK
Q7RLW1	NVPGFS
Q8MZU6	EVPQID
Q7XZ71	ELPGVG

Fig. 2. Sequence alignment of seven sHSPs subfamilies. The full sequences of 344 sHSPs (with names shown in Table 2 including bacterial class A and B, archaea, fungi, plant, animal, and other eukaryote sHSPs) were aligned using Clustalx. For simplicity, only the Pro-Gly doublet and its neighboring residues are shown here.

Table 2. Total sHSPs (344 unique sequences)**(Bacterial class A, 32^a)**

HSPA_BRAJA^b HSPB_BRAJA HSPD_BRAJA HSPH_BRAJA HSPE_BRAJA IBPA_ECOLI Q8XGW7 Q7VQU7 Q87TQ1 Q7MQJ3 Q9K VX0 Q8EET9 Q98H48 Q8FXL4 Q8U943 Q89CQ6 Q92VA9 Q8UKP8 Q98II5 Q98P38 Q8ZL03 Q8Z2L8 IBPB_ECOLI IBPB_AZOVI Q9AMN8 Q98GS9 Q8UIC2 Q89WG1 IBP_BUCAI IBP_BUCAP IBP_BUCTS Q8ZPY6

(Bacterial class B, 64)

HS19_RICCN HS19_RICPR ASP1_STRTR O52190 Q9X9C3 ASP2_STRTR Q9X9N3 Q81QZ9 Q81DY2 HS16_ONYPE Q8G6R2 HS18_CLOAB Q890A7 Q88SP7 14KD_MYCTU Q8KY67 18K1_MYCIT 18KD_MYCLE 18K2_MYCIT 18K1_MYCAV 18K2_MYCAV Q82QP6 Q82Q29 HS18_STRAL Q7WSY3 Q9RVB5 Q8Z017 Q8DKI6 Q8F5V3 Q7UF81 SP21_STIAU Q8PNC2 Q8PBS2 Q87C16 Q82T55 Q83CE9 Q9ZFD1 Q9WYK7 Q98IT5 Q92VB0 Q98II6 P70919 HSPC_BRAJA O69243 HSPF_BRAJA Q92Z87 Q8R745 Q9KFJ9 O67316 Q7NYU5 Q8AAA0 Q8A8R3 Q7NH34 Q88U58 HS15_LEPIN Q98DG0 O53673 Q7U3R4 Q81DZ3 Q8EMZ6 YOCC_BACSU COTP_BACSU Q7WMZ1 Q9RTR5

(Archaea, 35)

HSPS_METJA Q9C4M2 O59514 Q9YFZ9 Q9YAJ0 Q9HSS6 Q9HSD8 Q9HP75 Q9HP50 Q9HKX2 Q9HJU9 Q9HHW1 Q97W19 Q97VL9 Q97AM1 Q979Z6 Q974V6 Q970D9 Q8ZV10 Q8ZTS4 Q8ZTN4 Q9P9K5 O28308 O28973 O26947 Q9V1L0 Q8TUD5 Q8TK35 Q8THE4 Q8THE3 Q8TZC8 Q8PZK9 Q8PXG4 Q8PXG3 Q8PX03

(Fungi, 10)

HS16_SCHP HS26_YEAST HS30_EMENI HS30_NEUCR HS42_YEAST Q8NJ78 Q8TG38 O74984 Q7S0S7 Q7RW96

(Plant, 115)

HS11_ARATH HS11_CHERU HS11_DAUCA HS11_HELAN HS11_LYCES HS11_MEDSA HS11_ORYSA HS11_PEA HS11_SOYBN HS11_WHEAT HS12_ARATH HS12_DAUCA HS12_MEDSA HS12_ORYSA HS13_ARATH HS13_SOYBN HS14_SOYBN HS15_SOYBN HS16_SOYBN HS21_ARATH HS21_HELAN HS21_MAIZE HS21_PEA HS21_PHANI HS21_SOYBN HS22_MAIZE HS22_PHANI HS23_MAIZE HS2C_ARATH HS2C_CHERU HS2C_CHLRE HS2C_LYCES HS2C_PEA HS2C_PETHY HS2C_SOYBN HS2C_WHEAT HS2M_ARATH HS2M_PEA HS2M_SOYBN HS41_PEA HS41_SOYBN Q8W541 Q43543 Q8L7T2 Q40698 Q9ZSX9 O82545 P93440 P93439 Q40057 Q96489 Q9SRD6 O64564 Q9LGS1 Q9M566 O82149 Q9XGS6 Q40056 Q40826 Q39820 Q9ZP25 Q9ZW31 O64960 O49710 O24082 Q9ZSX7 Q39819 Q40847 Q9M6R2 Q43545 Q9ZSX8 Q9ZP24 Q9ZSY0 Q9SE12 Q41028 O49962 Q39930 O23640 Q8L470 Q40852 Q94KM0 Q9LS37 Q96458 Q43544 Q9ZSY1 Q943E7 O80432 Q9ZSZ0 O48898 Q9SE11 Q943Q3 Q8W007 Q8LCB4 Q943E6 Q8LMQ7 O81961 Q8H288 Q8H1A6 Q8GWH1 Q8GV39 Q8GV37 Q8GV36 Q8GV35 O81822 Q9FHQ3 Q9LNW0 Q38806 Q84J50 O24247 Q9SYU9 Q9SYV0 Q9SYU8 Q9FGM9 Q7XY72 Q7XE46

(Animal, 79)

CRA2_MESAU CRAA_HUMAN CRAA_ALLMI CRAA_ANAPL CRAA_CHICK CRAA_ARTJA CRAA_BALAC CRAA_RANCA CRAA_RANES CRAA_ASTFA CRAA_SQUAC CRAA_ZALCA CRAB_MESAU CRAB_HUMAN CRAB_ANAPL CRAB_BOVIN CRAB_CHICK CRAB_RANCA CRAB_SPAJD CRAB_SQUAC EFL2_DROME HS11_CAEEL HS12_CAEEL HS16_CAEEL HS17_CAEEL HS20_NIPBR HS22_DROME HS23_DROME HS26_DROME HS27_DROME HS30_ONCTS HS3C_XENLA HS3D_XENLA HS6A_DROME HS6C_DROME HSB1_CANFA HSB1_CHICK HSB1_CRIL0 HSB1_HUMAN HSB1_MOUSE HSB1_POELU HSB1_RAT HSB2_HUMAN HSB2_MOUSE HSB2_RAT HSB3_HUMAN HSB3_MOUSE HSB3_RAT HSB6_HUMAN HSB6_RAT HSB7_HUMAN HSB7_MOUSE HSB8_HUMAN HSB8_MOUSE HSB8_RAT HSB9_HUMAN HSB9_MOUSE OV21_ONCVO OV22_ONCVO P40_SCHMA YKZ1_CAEEL Q96E17 O44112 Q93141 Q27922 O01718 Q17268 Q9GT43 Q20165 Q93889 Q86NK3 Q86GU1 Q86G69 Q7YZT0 Q7YWK0 Q7YT58 Q7YT57 Q864Y6 O13225

(Other eukaryote, 9)

Q9BHC0 Q9BMF0 Q8IES0 Q8IB02 Q7YWK6 Q7RS01 Q7RLW1 Q8MZU6 Q7XZ71

^a The total number of sHSPs in each subfamily is indicated.

^b The name of each sHSP is either TREMBL or SWISS_PROT number.

loop that connects two anti-parallel β -strands (as indicated in Fig. 1a) and is localized at the interface of subunit–subunit interactions in the oligomers [22] (or implicated in Fig. 1c).

The Gly59 in *M. tuberculosis* Hsp16.3 (belonging to bacterial class B subfamily [16]) existing as nonamers and exhibiting chaperone-like activities *in vitro* [18, 24] was then replaced via site-directed mutagenesis to understand the roles of this Pro-Gly doublet. For this purpose, we took advantage of the fact that the wild type Hsp16.3 protein contains neither cysteine nor tryptophan residues,

two of the residues whose side chains can be effectively probed in the protein due to the special chemical reactivity of the former and high fluorescence intensity of the latter [25].

First, Gly59 in the Pro-Gly doublet of Hsp16.3 was replaced by cysteine. Data presented in Fig. 1b clearly demonstrate that an intersubunit disulfide bond was spontaneously formed in the G59C mutant Hsp16.3 protein (lane 3). Such an intersubunit disulfide bond, however, was not detected in the L57C and D61C mutant Hsp16.3 proteins (lanes 4–7 in Fig. 1b). Given that disul-

disulfide bonds are only formed between two cysteine residues being within about 6 Å from each other [26], these observations suggest that Gly59 residues on a certain pair of subunits are close to each other in the wild type Hsp16.3 protein. Supporting this speculation is that the distance between the side chains of a pair of Gly62 residues in Hsp16.5 dimer (residue corresponding to Gly59 in Hsp16.3) is slight closer than that between Leu60 residues, and is much closer than that between Asn64 residues (Fig. 1c).

To probe the topological location of Gly59 in the wild type Hsp16.3 oligomers, we examined the exposure status of the cysteine residues in the G59C mutant protein using DTNB, which is able to access and react with those exposed sulfhydryl groups [19]. Data presented in Fig. 1d demonstrate that the Cys59 residue in the G59C mutant protein is buried and not accessible to DTNB under normal conditions, but becomes exposed and thus accessible to DTNB in the presence of urea at a relatively high concentration. Examination of the exposure status of Cys59 upon heating was not successful due to the formation of aggregates of the G59C mutant protein at high temperatures. The cysteine modification curves of 3 and 4 (Fig. 1d) obviously contain two phases (fast and slow). A reasonable explanation for this observation is that the fast phase might represent a fast unfolding and/or dissociation of the mutant protein in the presence of high concentrations of urea, and the slow phase a limited subunit exchange process between the dissociated mutant protein oligomers since the time scale of this phase is close to that of subunit exchange as previously observed [27].

The tryptophan residue in the G59W mutant Hsp16.3 protein is also buried and is located within or near the subunit interfaces. Second, Gly59 in the Hsp16.3 protein was replaced by tryptophan with its topological location in the protein examined by recording its intrinsic fluorescence. The introduction of a tryptophan residue indeed dramatically increases the apparent intrinsic fluorescence of the Hsp16.3 protein (insert in Fig. 3a). The fact that the maximum emission wavelength of fluorescence is around 321 nm (insert in Fig. 3a) suggests that Trp59 is buried in a hydrophobic environment. The exposure of Trp59 in the presence of urea or upon heating is indicated by the red shift of the maximum emission wavelengths (Figs. 3a and 3b).

A very striking observation is that the urea-induced exposure of Trp59 in the G59W mutant protein was surprisingly accompanied with a dramatic increase in the fluorescence intensity (Fig. 3a). The most likely explanation for this is that the urea induces the oligomers to dissociate and this will in turn reduce the self-quenching of the tryptophan residues located close to each other at the subunit interfaces [28]. Nevertheless, the priority of Trp fluorescence red shift to its intensity upon urea increase (Fig. 3a) may imply an unfolding process leading to the exposure of Trp occurring ahead of the subunit dissociation.

The G59W mutant Hsp16.3 protein exists predominantly as dimers rather than nonamers and exhibits efficient chaperone-like activity even at low temperature. It was subsequently examined whether the replacement of Gly59 in the Hsp16.3 protein would alter its oligomeric structure and chaperone-like activity. Results of chemical cross-linking analyses (Fig. 3c) demonstrate that the G59W mutant Hsp16.3 protein predominantly exists as dimer, whereas the G59A mutant protein (being a control mutant protein) still appears as a nonamer similar to the wild type protein. This observation indicates that Gly59 in Hsp16.3 is indeed located in a position that is important for the assembly of the protein.

Given that the Trp59 residue is buried under native conditions and exposed upon heating (Fig. 3b), it is then asked whether such exposure correlates with any increase in chaperone-like activity, which would implicate the involvement of the Pro-Gly doublet in substrate binding for the wild type Hsp16.3. Data presented in Fig. 3d show that the G59W mutant protein exhibits chaperone-like activity at 25°C as efficiently as it does at 60°C, in contrast to the wild type protein, which exhibited little at 25°C and high activity only at a higher temperature (see also ref. [18]).

Parallel examination on the G59C mutant protein revealed that it exists as polydisperse oligomers rather than as nonamers, is different in secondary structure content, and exhibits lower chaperone-like activity when compared with the wild type protein (data not shown). However, it is not very easy to explain these observations since disulfide bonds are known to be critical for the folding and assembly of proteins in general [29] and cysteine itself may interfere with the chaperone-like activity of chaperones [17].

DISCUSSION

The Pro-Gly doublet is often involved in the formation of β -turn structure in proteins due to the unique properties of the Pro and Gly residues [30, 31]. Our site-directed mutagenesis studies of Gly59 for Hsp16.3 of *M. tuberculosis* (a residue in a Pro-Gly doublet highly conserved in non-animal sHSPs) strongly demonstrate its important direct role in the subunit-subunit interactions. This conclusion is also well supported by the analysis on the crystal structures of Hsp16.5 and Hsp16.9 [22, 23] (Fig. 1c).

Our observations, summarized below, however, suggest the Gly59 residue of Hsp16.3 is not directly involved in binding denatured substrate proteins. First, the G59W mutant Hsp16.3 protein does not expose Trp59 at room temperature but exhibits efficient chaperone-like activity (Figs. 3b and 3d), and the exposure of Trp upon heating is not correlated with a change in chaperone-like activity. Second, the wild type Hsp16.3 protein exhibits efficient

chaperone-like activity and maintains its nonameric structure in the presence of low concentrations of urea (0.8 M) [27, 32], at which Trp59 in the G59W mutant protein is not exposed (Fig. 3a). Additionally, supporting this claim is the previous demonstration that the N-terminal region (2-35 residues) of the wild type Hsp16.3 protein is critical for substrate binding [33]. This claim, however, should not be taken as exclusive, given that it is based on observations of the behavior of the G59W mutant Hsp16.3 protein, where the degree of the exposure of the Trp residue was examined in the absence of denatured substrate proteins, which may induce such an exposure.

The substitution of this Pro-Gly doublet by charged residues in the animal sHSPs may suggest a different role

of this region. For instance, one of the substrate-binding sites for bovine α A- and α B-crystallin was identified as a region containing these charged residues [34, 35]. These charged residues, on the other hand, may also be important for protein oligomerization of animal sHSPs since the region containing these charged residues in α B-crystallin was found to be involved in an interaction with α A-crystallin [36].

In our preliminary study [37], far-UV CD spectroscopy analysis revealed that the replacement of Gly59 by Trp leads to a significant disturbance of the secondary structure of the Hsp16.3 protein, consistent with its effect on oligomeric structure. More importantly, the behavior of G59W mutant Hsp16.3 protein is very similar to that of

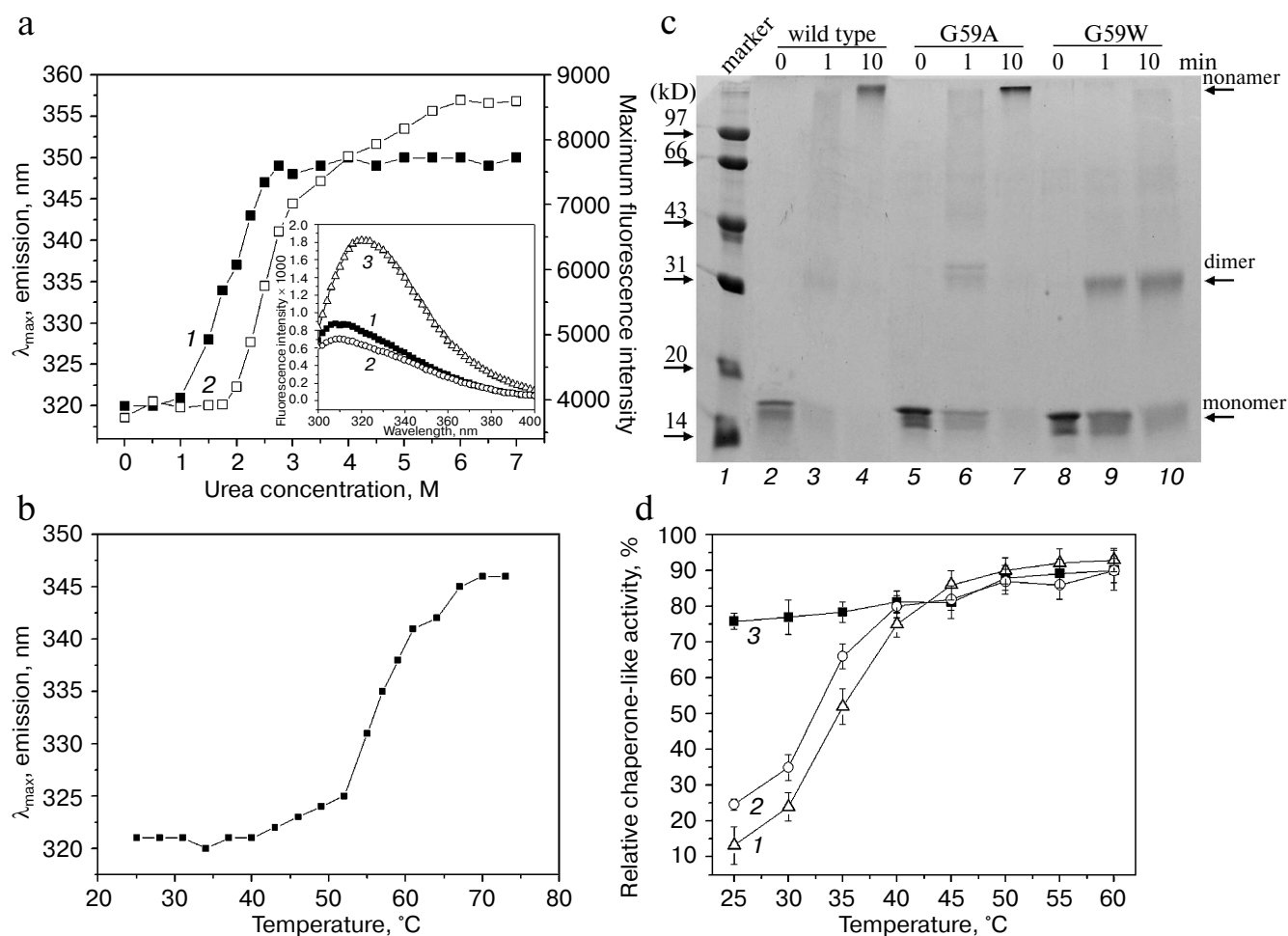


Fig. 3. Characterization of the G59W mutant Hsp16.3 protein. a) The tryptophan maximum emission wavelength (curve 1) and fluorescence intensity at the spectral maximum (curve 2) of the G59W mutant Hsp16.3 protein (0.2 mg/ml) are plotted against the urea concentration. The inset represents the intrinsic fluorescence spectra of the wild type (curve 1), G59A (curve 2), and G59W (curve 3) mutant Hsp16.3 proteins (all being at 0.1 mg/ml). b) The tryptophan emission maximum wavelength of the G59W mutant Hsp16.3 protein (0.1 mg/ml) is plotted against temperature increase applied to unfold the protein. c) SDS-PAGE analysis results of the wild type (lanes 2-4), G59A (lanes 5-7), and G59W (lanes 8-10) mutants of Hsp16.3 protein after cross-linking by glutaraldehyde for the indicated time intervals. d) The relative chaperone-like activities of the wild type, G59A, and G59W mutants of Hsp16.3 protein (curves 1-3, respectively) in suppressing the aggregation of insulin B molecules at the indicated temperatures. Each value represents the average of two independent experiments.

another mutant Hsp16.3 protein lacking nine residues from the C-terminus [33] in that both predominantly exist as dimers and exhibit efficient chaperone-like activities even at room temperature. These observations strengthened the view that sHSPs, like Hsp16.3, exhibit chaperone-like activities via oligomeric dissociation [18, 21, 27, 38]. Moreover, the existence of both as dimers is in line with the recent report that Hsp16.3 protein was a dodecamer using dimers as the assembling units [39].

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