Int Microbiol (2001) 4: 187–202 DOI 10.1007/s10123-001-0037-9

## **REVIEW ARTICLE**

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# The branching order and phylogenetic placement of species from completed bacterial genomes, based on conserved indels found in various proteins

Received: 3 August 2001 / Accepted: 5 September 2001 / Published online: 20 December 2001 © Springer-Verlag and SEM 2001

Abstract The presence of shared conserved inserts and deletions (indels or signature sequences) in proteins provides a powerful means for understanding the evolutionary relationships among the Bacteria. Using such indels, all of the main groups within the Bacteria can be defined in clear molecular terms and it has become possible to deduce that they branched from a common ancestor in the following order: Low G+CGram-positive  $\rightarrow$  High G+C Gram-positive  $\rightarrow$  Deinococcus-Thermus  $\rightarrow$  Cyanobacteria  $\rightarrow$  Spirochetes  $\rightarrow$  $Aquifex-Chlamydia-Cytophaga \rightarrow$  Proteobacteria-1  $(\epsilon, \delta) \rightarrow$  Proteobacteria-2  $(\alpha) \rightarrow$  Proteobacteria-3  $(\beta) \rightarrow$ Proteobacteria -4 ( $\gamma$ ). The usefulness of this approach for understanding bacterial phylogeny was examined here using sequence data from various completed bacterial genomes. By using 12 indels in highly conserved and widely represented proteins, the species from all 41 completed bacterial genomes were assigned to different groups; and the observed distribution of these indels in different species was then compared with that predicted by the signature sequence model. The presence or absence of these indels in various proteins in different bacteria followed the pattern exactly as predicted; and, in more than 450 observations, no exceptions or contradictions in the placement of indels were observed. These results provide strong evidence that lateral gene transfer events have not affected the genes containing these indels to any significant extent. The phylogenetic placement of bacteria into different groups based on signature sequences also showed an excellent correlation with the 16 S rRNA with 39 of the 41 species assigned to the same group by both methods. These results strongly vindicate the usefulness of the signature

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Department of Biochemistry, McMaster University, Hamilton L8N 3Z5, Ontario, Canada E-mail: gupta@mcmaster.ca Tel.: +1-905-5259140 Fax: +1-905-5229033 sequence approach to understanding phylogeny within the Bacteria and show that it provides a reliable and internally consistent means for the placement of bacterial species into different groups and for determining the relative branching order of the groups.

**Keywords** Indels · Signature sequences · Bacterial genomes · Lateral gene transfer · Phylogeny

## Introduction

Our current understanding of evolutionary relationships among the Bacteria, which comprise the vast majority of the known prokaryotes, is almost entirely based on the 16 S rRNA sequences [4, 40, 51]. Based on oligonucleotide signatures and the branching pattern of bacteria in the 16 S rRNA trees, 11 main groups (or divisions) among the Bacteria were originally proposed [69, 70, 72]. These included: Thermotogales, green nonsulfur bacteria, Deinococci and relatives, Spirochetes, green sulfur bacteria, Cyanobacteria, Gram-positive bacteria, purple bacteria and relatives (Proteobacteria), Bacteriodes-Flavobacteria-Cytophaga and relatives, Planctomyces and relatives, and chlamydiae. At the time when these divisions were proposed, the rRNA sequence database was quite limited and clear distinctions between these groups was possible on the basis of oligonucleotide signatures or long "naked" branches that separated these groups in the trees [69, 72]. However, in the past 15-20 years, as the sequence database for rRNA has rapidly expanded [42], distinguishing between these divisions on the basis of either of these criteria has become increasingly difficult and imprecise [40, 41]. In recent years, in addition to the above groups, many additional groups or divisions within the Bacteria have been suggested (i.e., Aquificales, Desulfurobacterium, Dictyoglomus, Fibrobacter, Flexistipes, Fusobacteria, Holophaga, Nitrospira, Verrucomicrobium) [40, 41]. In the absence of well defined criteria for the major divisions, it is unclear how many of these newly

described groups actually comprise new divisions within the Bacteria. To place the bacterial phylogeny on a firmer base, it is essential to develop clear molecular criteria by which the different major groups (phyla or divisions) within the Bacteria can be defined and distinguished from each other. Another issue central to bacterial phylogeny is to determine how the different main groups or divisions within the Bacteria are related to each other and how they branched from a common ancestor [21]. Such relationships are not resolved in phylogenetic trees based on rRNA or various proteins [6, 11, 40, 51, 69]. This has led to a growing acceptance of the notion that such relationships are unresolvable and that all the main groups within the Bacteria probably branched off simultaneously from the common ancestor [11, 40, 41, 71].

We recently described a new approach that makes use of conserved inserts and deletions (referred to as indels or signature sequences) found in various proteins, which provides valuable information regarding the issues that are not resolved in the rRNA trees [19, 23]. Based simply on the presence or absence of specific signature sequences, all of the major groups within the Bacteria can be clearly defined and distinguished from each other. Further, this approach also permits a logical deduction of the relative branch order of different main groups from a common ancestor [19, 23, 26], which has been a major impediment in understanding bacterial phylogeny. In the past few years, the entire genomes of many bacterial species have been sequenced, representing all major groups within the Bacteria (http://www.ncbi.nlm.nih.gov:80/PMGifs/ Genomes/micr.html). This provides us with a valuable means to test in an objective manner the usefulness and validity of the signature sequence approach for determining the phylogenetic placement and branching order of the bacterial species. Results of these studies presented here strongly evidence that this approach provides a reliable and internally consistent means for the phylogenetic placement of species into different groups and for determining their relative branching order. Importantly, the assignment of bacterial species into different groups using this new approach shows a very high degree of correlation to that based on the 16 S rRNA trees. Therefore, this new approach is not contradictory to the 16 S rRNA analyses but complements the latter studies in important respects, by providing information regarding issues that are not resolved in such phylogenies.

#### **Results and Discussion**

Bacterial genomes and signature sequence

The information for various bacterial species whose complete genomes have been sequenced to date is given in Table 1. The sequence information is presently available for 41 bacterial genomes, representing all of the main groups within the Bacteria including:  $\alpha$ -,  $\beta$ -,  $\gamma$ -, and *e*-Proteobacteria, Aquificales, Chlamydia, Cyanobacteria, Deinococcus-Thermus group, Spirochetes, Thermotoga, several members of the low G+C Grampositive bacteria including the mycoplasmas, and high G+C Gram-positive species (http://www.ncbi.nlm.nih.gov/PMGifs/Genomes/micr.html). In our earlier work, a large number of sequence signatures in different proteins were identified [19, 23]. Some of these signatures are specific for the particular groups and they provide no information regarding relationships to other groups [19]. However, of the identified signatures, a group of 12 signatures has proven most useful for distinguishing between the major groups within the Bacteria and for determining their branch order (Fig. 1). The sequence information for these signatures for various bacterial species whose genomes have been sequenced was obtained by basic local alignment search tool (BLAST) searches (http://www.ncbi.nlm.nih.gov) on either the non-redundant database or on individual genome sequences.

The rationale for using conserved indels for phylogenetic studies

The rationale for using conserved indels for evolutionary studies has been discussed in detail in earlier work [19, 20]. When a conserved indel of defined length and sequence (referred to as a signature sequence) that is flanked by conserved regions to ensure its reliability is found at the same position within a given protein (or gene) from different species, then the simplest and most parsimonious explanation for this observation is that the indel was introduced only once during the course of evolution and then passed on to all descendants [19, 56]. Thus, based on the presence or absence of a signature, the species containing or lacking the signature can be divided into two unambiguous groups. The well defined indels in different genes/proteins also provide useful milestones for evolutionary events, since all species emerging from the ancestral cell in which a given indel was first introduced are expected to contain the indel, whereas all species that existed prior to this event or which did not evolve from this ancestor will not contain the indel [19, 20]. Thus, by using well defined indels in proteins that were introduced at various stages in evolutionary history, it should be possible to deduce the branching order of different groups of species from a common ancestor.

Testing the signature sequence model using completed bacterial genomes

Figure 1 shows the signature sequences in proteins that have proven most useful for distinguishing the major groups within the Bacteria and to determine their relative

 Table 1
 Details of bacterial

 species whose genomes have
 been sequenced

Bacterial species	Accession number	Bacterial group/division	Reference
Aquifex aeolicus	NC000918	Aquificales	[10]
Bacillus halodurans C-125	NC002570	Low G+C Gram-positive	[65]
B. subtilis	NC000964	Low G+C Gram-positive	[36]
Borrelia burgdorferi	NC001318	Spirochaetales	[14]
Buchnera sp. APS	NC002528	γ-Proteobacteria	[59]
Campylobacter jejuni	NC002163	$\epsilon$ -Proteobacteria	[53]
Caulobacter crescentus	NC002696	α-Proteobacteria	[50]
Chlamydia muridarum	NC002182	Chlamydiales	[55]
C. trachomatis	NC000117	Chlamydiales	[62]
Chlamydophila pneumoniae CWL029	NC000922	Chlamydiales	[31]
C. pneumoniae AR39	NC002179	Chlamydiales	[55]
C. pneumoniae J138	NC002491	Chlamydiales	[60]
Deinococcus radiodurans	NC001263	Deinococcus/Thermus	[68]
Escherichia coli K12	NC000913	γ-Proteobacteria	[5]
E. coli OI57:H7 EDL933	NC002655	y-Proteobacteria	[54]
E. coli OI57:H7	NC002695	y-Proteobacteria	[43]
Haemophilus influenzae	NC000907	y-Proteobacteria	[13]
Helicobacter pylori 26695	NC00915	$\epsilon$ -Proteobacteria	[67]
H. pylori J99	NC000921	$\epsilon$ -Proteobacteria	[1]
Lactococcus lactis	NC002662	Low $G + C$ Gram-positive	(unpublished)
Mesorhizobium loti	NC002678	α-Proteobacteria	[32]
Mycoplasma genitalium	NC000908	Low $G + C$ Gram-positive	[15]
Mycobacterium leprae	NC002677	High $G + C$ Gram-positive	[9]
M. tuberculosis H37Rv	N000962	High G+C Gram-positive	[8]
M. tuberculosis CDC1551	NC002755	High $G + C$ Gram-positive	(unpublished)
M. pneumoniae	NC000912	Low $G + C$ Gram-positive	[28]
M. pulmonis	NC002771	Low $G + C$ Gram-positive	[7]
Neisseria meningitidis MC58	NC002183	$\beta$ -Proteobacteria	[66]
N. meningitidis Z2491	NC002263	$\beta$ -Proteobacteria	[52]
Pasteurella multocida	NC002663	y-Proteobacteria	[44]
Pseudomonas aeruginosa	NC002516	y-Proteobacteria	[64]
Rickettsia prowazekii	NC000963	α-Proteobacteria	[2]
Staphylococcus aureus N315	NC002795	Low $G + C$ Gram-positive	[37]
S. aureus Mu50	NC002758	Low $G + C$ Gram-positive	37
Streptococcus pyogenes	NC002737	Low $G + C$ Gram-positive	[12]
Synechocystis sp. PCC6803	NC000911	Cyanobacteria	[33]
Thermotoga maritima	NC000853	Thermotogales	[49]
Treponema pallidum	NC000919	Spirochaetales	[16]
Ureaplasma urealyticum	NC002162	Low G + C Gram-positive	[17]
Vibrio cholerae	NC002506	γ-Proteobacteria	[27]
Xylella fastidiosa	NC002488	γ-Proteobacteria	[61]

branch orders. Based upon our analyses, these signatures have been introduced in these proteins at the indicated stages of the evolution of the bacterial groups. Hence, by using them, it should be possible to assign any given bacterial species into one of these groups and to determine its branching order, relative to the other groups.

To test in an objective manner the validity of the evolutionary model based on these signatures, we have analyzed the sequence data from various completed bacterial genomes using this approach. For these purposes, an alignment of the corresponding proteins from bacterial species whose complete genomes have been sequenced was carried out; and the presence or absence of the indicated signatures was determined. This information was then used for the phylogenetic placement of the species into different groups and to determine whether the distribution of these signatures in different species followed the pattern, as predicted by the model, or whether the results obtained were more readily explained by other mechanisms, such as either independent occurrence of the indels in different species, or lateral gene transfer (LGT) between species.

According to the model, once an indel has been introduced in an ancestral lineage, various groups of species emerging after that point should all contain the indel, whereas all species from different groups that existed prior to the introduction of the indel should lack the indel. However, if such indels have been introduced either independently in various species or if the genes containing these indels have been frequently horizontally transferred from one species to another, then the presence or absence of these indels in different species will not follow the predicted pattern. In such a case, different groups of species or even individual species from different groups will either contain or lack the indels. Thus, by determining how closely the results of the indel data follow the predictions of the model and how many exceptions to this are observed, it should be possible to objectively determine whether the inferences based on these indels are reliable and to what extent they



Fig. 1 Phylogenetic placement and relative branching order of bacterial species from completed genomes, based on the indel model developed in earlier work [19, 23]. The arrows above the line indicate the specific stages where the indicated signatures in various proteins have been introduced. The model predicts that all bacterial groups to the right of these arrows should contain the indicated signatures whereas all groups to the left should lack them. The sequences from various bacterial genome conform to the expected patterns, with no exceptions observed. The phylogenetic assignment of bacterial species whose genomes have been sequenced into different groups based on these signatures is indicated below the line

have been corrupted by other factors. The results for these signatures for the bacterial species whose genomes have been sequenced are discussed below.

#### Ribosomal S12 protein

Ribosomal S12 protein is an essential protein found in all sequenced microbial genomes. A 13-amino-acid indel

in a highly conserved region of this protein has been shown to distinguish the low G+C Gram-positive bacteria from all other bacteria [19, 20]. Among the completed microbial genomes, this indel was present in all of the low G+C Gram-positive species, i.e. *Bacillus subtilis*, *B. halodurans*, *Lactococcus lactis*, *Mycoplasma genitalium*, *M. pneumoniae*, *Staphylococcus aureus* (N315, MU50 strains), *Streptococcus pyogenes*, and *Ureaplasma urealyticus*, but not in any other bacteria (Fig. 2, see Appendix). Thus, as indicated in Fig. 1, this signature is a distinctive characteristic of the low G+CGram-positive group and, based upon it, the species belonging to this group can be clearly distinguished from all other bacteria.

#### Hsp70 protein

The Hsp70/DnaK family of proteins, which carry out an essential molecular chaperone function in protein-folding



Fig. 2 Alignment of ribosomal S12 protein sequences from completed bacterial genomes showing a 13-amino-acid insert (*boxed*) that is distinctive of the low G+C Gram-positive bacteria. *Dashes* in all sequence alignments show identity with the amino acid on the top line

and other cellular processes, are found in all completed bacterial genomes. A prominent signature, consisting of an indel of 21-23 amino acids, has been identified in the Hsp70 protein that distinguishes Gram-positive bacteria from Gram-negative bacteria [19, 20, 24]. The large indel in the Hsp70 protein is present in homologues from different Gram-negative bacteria but is absent from those of the Gram-positive bacteria (Fig. 3). The Gram-negative bacteria are defined in our work by the presence of both an inner and outer cell membrane, rather than on the basis of the Gram-staining reaction, which is a variable characteristic [19, 20]. Among the completed genomes, this indel, as expected, was found in all Gram-negative bacteria, but was not present in any of the Gram-positive bacteria, nor was it present in Thermotoga maritima or various mycoplasma species, supporting their grouping with the Gram-positive bacteria. In Synechocystis sp., multiple homologues for Hsp70 were found [33] and all of these contain the large insert (Fig. 3) [26]. Two different homologues for Hsp70 were also found in the genome of the spirochete species Borrelia burgdorferi [14]. One of these homologues, which contained the large insert (GenBank no. 2688438), was closely related to the other spirochete species, *Treponema pallidum*. In contrast, a second Hsp70 homologue in *B. burgdorferi* (GenBank no. 2688201) lacked the large insert. BLAST searches on this homologue indicated that all of the top scores in this case consisted of various Gram-positive bacteria and archaeobacteria. Thus, it is likely that this homologue is derived from Gram-positive bacteria by means of LTG. The Hsp70 sequences are available in the databases for more than 150 bacterial homologues. Of these, this insert is not found in any Gram-positive bacteria and, with the single exception of *B. burgdorferi* noted here, it is a distinctive characteristic of all Gram-negative bacteria [19, 26].

Since the indel in Hsp70 divides the Bacteria into two structurally distinct groups, the question arises whether this indel is an insert in the Gram-negative or a deletion in the Gram-positive. Several lines of evidence support the former of these two possibilities. First, based on the accepted rooting of the prokaryotic tree using duplicated elongation factor EF-1/EF-2 sequences [29], the root of the prokaryotic tree has been shown to lay between archaebacteria and Grampositive bacteria [19]. The Hsp70 homologues from both these groups of prokaryotes lack this indel, which strongly suggests that this indel is an insert in the Gram-negative bacteria that evolved at a later stage. A second argument supporting this inference is based on the sequence similarity between Hsp70 and another

		54	·	ו	133
	-E.coli K12	AKRQAVTNPQNTLFAIKRLIG	RRFQDEEVQRDVSIMPFKIIAADN	GDAWVEV	KGQKMAPPQISAEVLKKMKKTAEDYLGE
	E.coli 0157:H7				
	E.coli 0157:H7 EDL933				
V-Proteo	Past. multocida	KK	E-V	G-	EF
7-110100	X. fastidiosa	KFY-V	-K-G-AK-LDLV-YTQH	ATA	DAK-LQEKEF
	Vibrio cholerae	•••••	EIKYVK	A	F
	Hae. influenzae Rd	IK	ESIKE-TR	N-	D-LF
	Buchnera sp. APS	IK	-K-K-DIKYN-VNS	ID-	-K
	L Pse. aeruginosa	Y-V	EENVK-IQMV-YS-VK		
0.0.	🖵 Nei. meningitidis Z2491	AKIY-A	HK-E-KIESEK-N-	KA	Q-KELSREAA
p-Proteo	Nei. meningitidis MC58	AKIY-A	HK-E-KIESEK-N-	KA	Q-KELSREAA
	<sub>r</sub> Mesorhizobium loti	EIV	YD-PVTEK-KKLV-YVKG	A	G-K-QS-SMI-QEA
W Drotoo	Ri. prowazekii	RIY-V	-N-T-PM-RK-QGLV-YN-VK	A	DNH-YS-SFI-QEN
u-110100	L C. crescentus	T	- TAS - PV - EK - KGMV - YE - VKGPT	KA	H-KDYS-QEVFI-QEAAH
	<sub>r</sub> He. pylori 26695	EK-IYSIM-	LM-NEDKAKEAEKRL-YVDR -	-ACAI-I	S-KVYT-QEKI-M-L-EDS
C Drates	He. pylori J99	EK-IYSIM-	LM-NEDKAKEAEKRL-YVDR -	-ACAI-I	S-KIYT-QEKI-M-L-EDS
e-rroteo	L Camp. jejuni	EK-IYSIM-	LMINEDAAKEAKNRL-YH-TER -	- ACAI - I	A-KIYT-QEKM-L-EDAF
	<sub>r</sub> Chl. muridarum	EKASTF	-K- SESEIKTV-Y-VAPNSK	VF	ENKLYT-EE-G-QI-MEA
······	Chl. pneumoniae J138	EKGSTF	-KY SASEIQTV-YTVTSGSK	VF	D-KQYT-EE-G-QI-MEA
Chlamydia	Chlam. pneumoniae AR39	EKGSTF	-KY SASEIQTV-YTVTSGSK	VF	D-KQYT-EE-G-QI-MEA
	Chlam. pneumoniae CWL029	EKGSTF	-KY SASEIQTV-YTVTSGSK	VF	D-KQYT-EE-G-QI-MEA
	Chl. trachomatis	EKASTF	-K- SESEIKTV-Y-VAPNSK	VFD -	EQKLYT-EE-G-QI-MEA
	Aqu. aeolicus	R-ILD-EVYESF	-KKEEAKRVSY-VVPDEK	AFDIPM	A-KLVR-EEVG-HR-L-EAAF
Spirachatas	r Tre. pallidum	N-MEH-IYSF	S N-LTGEAKKV-YV PQG	D-VR	E-KLYSTQEFI-Q
Spriochetes	Bo. burgdorferi	N-MEIYSFM-	ASEIKMV-YEKGL-	R-NISM	IKKQ-SEAT-TEA
	Synechocystis PCC6803-1	MGFYSVF	-K- D-ITNEATEVAYSVVKDG-	-NVKLDCPA	Q - KQF EE Q R - LVDD - SK
	Synechocystis PCC6803-2	SAEVYSF	W DDTVEER-RV-YNCVKGRD	DTVS-SI	RSYT-QEMI-Q-L-ADS-AF
	Deino. radiodurans	-RALAAEVF	W DKEEAARSTVKEGPS	-SVRI	N-KDLE-VR-LVSD-SAKN
	Thermotoga maritima	MILER-IKSKM-		T-YK-RI	DDKEYT-QEFIL-NDAG
	┌ Myc. tuberculosís H37Rv	NVDR-VRSVHM-		S-WSI-I	D-K-YTA-ERI-M-L-RDA
High C+C	Myc. tuberculosis CDC1551	NVDR-VRSVHM-		S-WSI-I	D-K-YTA-ERI-M-L-RDA
ingi di c	Myc. leprae	NVDR-IRSVHM-		S-WSI-I	D-K-YTAQERM-L-RDA
	г Bac. subtilis	SIN -IMSHM-		T-YKI	E-KDYT-QEVII-QHL-SYS
	Bac. halodurans	IN -VISHM-		TNHKENI	E-KEYT-QEII-Q-L-SDA
	Sta. aureus MU50	IN -VQSHM-		T-YK-DI	E-KSYT-QEMI-QNL-NS
	Sta. aureus N315	IN -VQSHM-		T-YK-DI	E-KSYT-QEMI-QNL-NS
Low G+C	L. lactis	E -IISSKM-		TSEK - SA	N-KEYT-QEMI-QNL-AA
	Strep. pyogenes	E -VISSKM-		TSEK-SA	N-KEYT-QEMI-QYL-GY
	U. urealyticum	KQIN -ISSM-		TKEK-T-	LNKDYT-EEKI-TYI-EYKKI-A
	M. pulmonis	LEDT IASM-		ΤΤΚΤΥΚΑ	N-KTYK-EEMI-SHL-EYKKV-K
	M. genitalium	MN -IVSM-		TSNK-K-§	TTKELS-E-VQI-SYL-DFKKI-K
	L M. pneumoniae	MN -IVSM-		TSNK-T-§	STKELT-EEVQI-SYL-DYKKI-K

Fig. 3 Alignment of Hsp70 homologues from completed bacterial genomes, showing the large insert (*boxed*) characteristic of Gramnegative bacteria

protein, MreB, which corresponds to the N-terminal half of Hsp70 [25]. Since the MreB protein, which is believed to have evolved independently from an ancestor of the Hsp70 family of proteins, does not contain this indel, the form of Hsp70 lacking the indel is indicated to be ancestral [24, 25]. Another argument in support of this view can be made on the basis of the cell structure of the prokaryotic organisms. In the formation of the ancestral prokaryotic cell, membrane enclosure must have been a key event [45]. The initial membrane enclosure probably consisted of a single unit membrane, as found in Gram-positive bacteria and archaebacteria, rather than of two different membranes separated by an intervening compartment, as found in Gram-negative bacteria [19, 22]. All of these observations indicate that the Gram-positive group lacking the large indel in Hsp70 is ancestral, in comparison with Gram-negative bacteria. The rooting based on these observations provides a useful reference point for interpreting the signature sequences in various other proteins and for deducing the relative branching orders of different groups. Based on this rooting, it could now be inferred that the 13-aminoacid indel in the S12 protein (Fig. 2), which is present in the low G+C Gram-positive bacteria (also archaebacteria) [19], but absent from both high G+C Grampositive bacteria and different Gram-negative bacteria, is a deletion in the common ancestor of the latter groups of species. This in turn indicates that, in comparison with the high G+C group, the low G+Cgroup is ancestral [19].

# Hsp60/GroEL protein

The Hsp60/GroEL family of proteins found in all sequenced bacterial genomes contain a 1-amino-acid insert in a highly conserved region which is indicated to have been introduced after the branching of various Gram-positive bacteria and the Deinococcus-Thermus groups (Fig. 1) [19]. Among the completed bacterial genomes, this insert was not found in any of the Gram-positive bacterial homologues or in D. radiodurans, but it was present in all other bacteria (Fig. 4). Several Gram-positive bacteria contain multiple Hsp60 homologues and this insert was not present in any of them. Similarly, Mesorhizobium loti and other members of the Rhizobiaceae family contain multiple Hsp60 homologues and this insert is present in all of them. The indicated position of this signature is highly reliable as, of more than 300 bacterial Hsp60 sequences that are available in databases, no exceptions are observed [23].

Fig. 4 Alignment of Hsp60 homologues from bacterial genomes, showing a 1-aminoacid insert (*boxed*) that was introduced after the branching of Gram-positive bacteria and the *Deinococcus–Thermus* groups

		144 r	1/8
	_E.coli	IAQVGTISAN	SDETVGKLIAEAMDKVGKEGVITVE
	E.coli 0157:H7		
	E. coli 0157:H7EDL933		
	Pse. aeruginosa	• • • • • • • • • • • •	SI-QIE
V Proteo	Pas. multocida	-E	SIQIQ
Y-110100	V. cholerae chr. I		SSNIERD
L	V. cholerae chr. II	-TA	HAI-EIQERN
	Xylella fastidiosa	A	SI-NIKI-
	Hae. influenzae Rd	-E	SIQSQE
	Buchnera sp. APS	-T	AKSEND
B-Proteo	-N. meningitidis MC58	S	QAIEE
F	N. meningitidis Z2491	s	QAIEE
	-Meso, loti (1)	V	GSMQN
	Meso, loti (2)	V AG -	GSMQN
W Protoo	Meso, loti (3)	A	G-AAMKND
4-110100	Meso, loti (4)		G-AEI-RFLQN
	Ri. prowazekii	S -	G-KEI-EKKEE
	C. crescentus		G-KEEMKN
	-Camp. jejuni	A	KI-NDED
€-Proteo	Hel. pylori 26695	-TA	HNIDED
	He, pylori J99	-TA	HNIDED
	-Chl. pneumoniae AR39	A -	N-SEI-NEN-S
1	Chl. pneumoniae J138	A	N-SEI-NEN-S
Chlamydia	Chl. pneumoniae CWL029	A -	N-SEI-NEN-S
	Chl. muridarum	A	N-AEI-NEN-S
	Chl. trachomatis	A	N-AEI-NEN-S
	Agu. aeolicus	-EA	N-PEIIDEED
Spirochates	-Bor. burgodferi	AS	N-SYI-EKDD
Spirochetes	Tre. pallidum	V-H-ASV	N-KEI-RIL-S-IENDD-D
	Sv. sp. PCC6803	A-V-S G	TNPEAMDT-D
	D.radiodurans	- KK - AG 🖵	NI-
	T. maritima	H-AA	NSAEI-EED
	-Myc, leprae (1)	ATAA	G-QSI-DN
	Myc. leprae (2)	-TA-V-S	RQI-A-VG-G-NTDVS
High G+C	Mvc. tuberculosis CDC(1)	ATAA	G-QSI-DNN
	Myc, tuberculosis CDC(2)	A-V-S	RQI-D-VGSHDVS
	Myc, tuberculosis H37Rv	A-V-S	RQI-D-VGSHDVS
	Strep, pyogene	AAV-S	RS-KEY-SERNDI-
	L. lactis	A-V-S	RS-KEY-SDERSDI-
	Sta, aureus MU50	<b>-</b> A	AEI-RY-SENDI-
	Sta. aureus N315	A	AEI-RY-SENDI-
Low G+C	Bac. halodurans	AAS	A-DEIERNDI-
	Bac. subtilis	AA	AESERNDI-
	M. genitalium	- E AA S	GSKEIQALNTD
	M. pneumoniae	- E AA S	GSKEIQALNTD

### FtsZ protein

The homologues of the FtsZ protein, which is involved in bacterial cell division, are found in all completed bacterial genomes, except those of the mycoplasma and *Chlamydiae* spp, which are intracellular pathogens [15, 17, 28, 55, 62]. A 1-amino-acid insert in a highly conserved region of this protein is indicated to have been introduced after the branching of Gram-positive bacteria, the *Deinococcus-Thermus* group, and Cyanobacteria (Fig. 1). As expected, this insert was not found in any Gram-positive bacteria, *D. radiodurans* or *Synechocystis* sp., but it was present in all other bacterial species, including *Aquifex*, Spirochetes, and different groups of proteobacteria (Fig. 5).

#### Alanyl-tRNA synthetase

Alanyl-tRNA synthetase contains a 4-amino-acid insert which is commonly shared by all proteobacteria and by the *Aquifex*, *Chlamydia*, and the *Cytophaga– Flavobacteria*–green sulfur bacteria groups, but is absent from all other Bacteria and Archaea (Fig. 6) [26]. This insert is indicated to have been introduced in a common ancestor of the above groups after the branching of Gram-positive bacteria, *Deioncoccus-Thermus*, Cyanobacteria, and Spirochetes (Fig. 1). Alanyl-tRNA synthetase is found in all sequenced bacterial genomes and the presence or absence of this signature in various species followed the expected pattern, with no exceptions observed (Fig. 6).

# Signature sequences for proteobacteria in Hsp70 and CTP synthase

The Hsp70 protein discussed above contains a 2-aminoacid insert, within the large insert found in the Gramnegative bacteria, which is commonly shared by all proteobacteria but not found in any other bacteria [19]. In the completed bacterial genomes, this insert was present in the Hsp70 homologues from all 17 proteobacterial species, but none of the other bacteria (Fig. 7). The sequences from Gram-positive bacteria lacking this region are not shown in this figure. The enzyme CTP synthase, found in all sequenced bacterial genomes except for the mycoplasma species, contains a 10-amino-acid insert which is specific for proteobacteria (Fig. 8). This insert was found in all sequenced proteobacterial genomes but not in any other species. A smaller 4-amino-acid insert in CTP synthase that is specific for the mycobacterial species Fig. 5 Alignment of FtsZ homologues, showing a 1-amino-acid insert (boxed) that was introduced after the branching of Gram-positive bacteria, the Deinococcus-Thermus group and the cyanobacteria

		232	269
	- E.coli K12	GEDRAFEAAFMATSSPLLE	TDI SGARGVI VNTTAGED
	E. coli 0157:H7		
	E. coli 0157:H7 EDL933		
	Vibrio cholerae		AL-
v-Proteo	Buchnera sp. APS	NS-I	
, 110000	Xylella fastidiosa	-DQAA-VQND -	VN-AN-IS-
	Pas. multocida	GTRI-VK-D R	VKS-M-
	H. influenzae	GRL-VRND I	KN-Q-IM-
	L Pse.aeruginosa	- PN R T - A RN	VN-QIP-
B Protoo	- N.meningitidis Z2491	- I RM - TDQ D -	VT-DTAPG
p-rioleo	N. meningitidis MC58	- I RM - TDQ D -	VT-DTAPG
	- Ri. prowazekii	IKSND H	SSMCIG-P-
α-Proteo	Meso. loti	SMKAAND E	VSMKKS-SG-R-
	└C. crescentus	ALMQNAND E	VS-KKAV-G-M-
	— Camp. jejuni	NAILSNED G	M-IKKILHFKTSSN
€-Proteo	Hel. pylori 26695	ES-KL-VQNQD -	ASIEKSII-FFEHHP-
	Hel. pylori J99	ES-KL-VQNQD -	ASIEKSII-FFEHHP-
	Aqu. aeolicus	-DEK-DI-V-K-VT G	NT-ERLT-WTSE-
pirochetes	┌─ Tre. pallidum	NVDTANN E	TRIETRLAVRGSEN
	└ Bor.burgdorferi	NVDRRTSN   E	VRIE-SK-LV-G-D-
	Sy.sp.PCC6803	-KSKTA 🖵	SSIQKVF-V-G-T-
	D. radiodurans	-DKMMSH	RGIERIVTG-Y-
	T. maritima	HRKK-ME-K-I-	HPVEN-SSIVFPSN
	∟Myc. leprae	-DG-SLKI-N	ASMEQMS-AG-S-
ligh G+C	Myc.tuberculosis H37 Rv	G-SLKIN	ASMEQMS-AG-S-
	└ Myc.tuberculosis CDC1551	G-SLKIN	ASMEQMS-AG-S-
	∟ Sta.aureus Mu50	NVKK	TSIVQMG-ES
	Sta. aureus Z2491	NVKK	TSIVQMG-ES
	Bac.subtilis	N A KK	AAIDQMG-TN
Low G+C	Bac. halodurans	NGKK	TS-DQMG-SN
	M. pulmonis	-KVKIHII-	TSIQSHTIIGSAN
	Lac. lactis	E-VITRKY	TTIEENL-V-G-M-
	∟ Strep. pyogenes	E-IVRKY	TTIDQD-IV-G-L-

is found in the same position as the proteobacterial insert (Fig. 8). However, this insert, because of its size and specificity, is of independent origin and it does not confuse or affect the specificity of the proteobacterial signature.

Spiroch

High G

Low G

# Signature sequences indicating the branch order of the proteobacterial groups

Signature sequences in a number of proteins have been shown to make clear distinctions among different groups of proteobacteria [23]. A 1-amino-acid conserved insert in the Lon protease is commonly shared by all  $\alpha$ -,  $\beta$ -, and y-proteobacterial species but not present in any other species. Lon protease homologues are present in all bacterial genomes, except a few Gram-positive bacteria. The insert in Lon protease, as expected, was found in all  $\alpha$ -,  $\beta$ -, and  $\gamma$ -proteobacterial species but not in any other species (Fig. 9). Another signature introduced at a similar stage is found in the SecA protein. The SecA homologues are found in all sequenced bacterial genomes and the 7-amino-acid insert is seen in all of the  $\alpha$ -,  $\beta$ -, and  $\gamma$ -proteobacteria but not in any other bacteria (Fig. 10). A smaller insert in this position is also seen in the two spirochete species but, based on its size and species specificity, this insert was probably introduced independently. The genomes from chlamydial species contain another SecA related protein (not shown), which contains a very large insert in this region, quite different from the insert found in  $\alpha$ -,  $\beta$ -, and  $\gamma$ -proteobacteria.

The Hsp70 family of proteins contains another useful signature that is distinctive of the  $\beta$ - and  $\gamma$ -proteobacteria. This signature, consisting of a 4-amino-acid insert in a highly conserved region, is found in all of the  $\beta$ - and y-proteobacterial species from sequenced genomes but

not in any other species (Fig. 11). The  $\beta$ - and  $\gamma$ -proteobacterial species, in addition to the orthologous Hsp70 protein, also contain a protein, Hsc66, which is distantly related to Hsp70 and carries out unrelated functions [34, 57]. The Hsc66 homologues, do not contain the  $\beta$ - or y-insert, but they are readily distinguished from the Hsp70 homologues because of extensive sequence divergence in different regions, particularly towards the C-terminal end. Another signature, a 1-amino-acid insert, distinctive of the  $\beta$ - and  $\gamma$ -proteobacteria, has been identified in the protein, phosphoribosyl pyrophosphate synthetase. Among the sequenced bacterial genomes, this signature is found in all  $\beta$ - and  $\gamma$ -proteobacteria but in none of the other species (Fig. 12). The  $\gamma$ -proteobacterial group differs from other proteobacteria by a 2-amino-acid deletion in the enzyme, 5'-phosphoribosyl-5-aminoimidazol-4-carboxamide transformylase. This deletion was found in all of the y-proteobacterial genomes (Fig. 13), but in none of the other species where the homologues of this protein are found. In T. maritima, a large deletion of 12-13 amino acids is present in this position which probably originated independently.

The distribution of indels in genomic sequences strongly supports the indel model

The question could now be asked whether the observed results from genomic sequences support the evolutionary model based on indels, or whether these results can be explained by any other reasonable mechanism. In the evolutionary model based on indels, there are two potential problems that could give misleading results. First, it is possible that a given indel, rather than being derived from a common ancestor, was introduced on Fig. 6 Alignment of Ala-tRNA synthetase sequences, showing a 4-amino-acid insert (*boxed*) that is common to only the *Chlam-ydiae–Aquifex* group and proteobacterial species and is not found in any other groups of bacteria



Fig. 7 Alignment of Hsp70 homologues from bacterial genomes, showing a 2-aminoacid insert (*boxed*) that is commonly found in all proteobacterial species. The Hsp70 homologues from Gram-positive bacteria lack this region and hence are not shown

multiple occasions in different species/groups due to similar functional constraints operating on the protein. Second, the shared presence of an indel in different species could also occur if the indel was originally introduced in one species (or group of species) and then transferred to others by LGT. The analyses of genomic sequences in the past few years have led to the view that LGT among prokaryotic species is quite common and that it poses a major problem in deducing evolutionary relationships among prokaryotes [3, 11, 30, 39, 71]. The basic premise on which the indel model is based is that, once an indel has been introduced in an ancestral lineage, various species emerging from that ancestor henceforth should all contain the indel, whereas all species from different groups that either existed prior to the introduction of the indel or which did not evolve from this ancestor should lack the indel. In contrast, if these indels have been introduced into various groups independently or if the genes containing these indels have undergone frequent LGT from one species to Fig. 8 Sequence alignment of CTP synthetase from bacterial genomes, showing a 10-aminoacid insert (boxed) common to all proteobacterial groups

		388				ſ	1 442
	<sub>Г</sub> H. influenzae	EYARN	/AGLTKANSSE	FDK	DCEQPVVALITE	WQDAEGNTEV	RTDESDLGGTMRLG
	E. coli K12	DH-	NMEN T -	- VP	KY	-R-ENV	-SEK
	E. coli 0157:H7	DH-	-NMEN T -	- VP	KY	-R-ENV	-SFK
	E. coli 0157:H7EDL933	DH-		- VP	KY	-B-ENV	-SEK
γ-Proteo	Buchnera sp. APS	- F - Q	V-TKET-	P	0-KY-TTDKN	RENNSSKNVN	KTENRTN
	Yv fastidiosa	D H.	EGT.	N. P		DTTT EV D	DEK
	Rec. multonide DUZO	D 11 -	D T	n - n			-DEK
	Pas. multociua PM/0			H	1-010	1-1	A
	Pse. aeruginosa		L-WSDI-	• • •	SSGH G	I	••EA
	└ Vib. cholerae		MEG-H-T-	- N -	NTKYG	-V-GV-E	- SEK
B-Proteo	<sub>Г</sub> Nei. meningitidis MC58	D -	••••KG•••T•	L	K-AAD-	T-D-SV-T	-DESA
P	L Nei. meningitidis Z2491	D -	· KG T -	L	K-AAD-	T-D-SV-T	-DESA
	<sub>r</sub> C. crescentus	• TL • • •	IKD-S	- G	PTERGIM	- IKGNE-VQ	-RAND
α-Proteo	Ri, prowazekii	- I - Q - L	I-IQD-VTE-	-KI	KGTKIIEKINKN	CETIKI	FBNMTEK
	Meso loti	- A SI	VEH-S-T-	- GP	T-FG-M	-LKGN ML-K	-RETG
	- Hel pylori 26695	- EC		- NO			
C Broton	Hel pyloni 100	-10		- NG		FM-QNHQKQ-	··· [N-F
C-Froico	Her, pytori Jaa		LKGI-	- NG	HYYEU	FM-QNHQKQ-	YN-P
	L Camp. jejuni	• • • • • •	LK-KDV	-NE	K-QNYD-	FM-TN-EKQI	AKTP
	<sub>C</sub> Chl. trachomatis		LDKPLM-	MNP	ETPDCMMEG		QDSVVK
1.	Chl. pneumoniae AR39		LN-DQL-	M - P	NTPH-I-YVMEG		QDPLVAT
Chlamydia	Chl. pneumoniae J138		LN-DQL-	M - P	NTPH-I-YVMEG		QDPLVAT
L	Chl. pneumoniae CWL029		LN-DQL-	M-P	NTPH-I-YVMEG		QDPLVAT
	Chl muridarum	YA	S-PI I -	M-P	NTPDCMMQG		0.TMTK
		- 5	L-ESNT-				OKKNDK
	Aqu. acorreus						
Spirochetes	Bor. burguorieri		U-ILU-DIE-I	NLAND	KPLKSIH-LP-		QKGIK-K-A
	∟ Tre. pallidum		LL-AS-H-H-	-AV	- TPH D - LPG		CV- IPISL
	Sy. sp. PCC6803	- W	-K-PEA-	-ET	ETPNIN-LP-		QQ - VV
	D. radiodurans	H-	IEDA-	E	YAKNK-ID-MP-		QLEVAGM
	T. maritima	- F	F-YKET-	P	NTPYD-ME-		QKRILK
	– Mvc. leprae	- AT - S -	VQA-	- EP	ATPDISTMAD	QK-I	VAG-A-F
High G+C	Myc. tuberculosis H37 Rv	- A S -	NA-	P	- TPD I - TMPD	QF-T	VAG-A
	Myc tuberculosis CDC1551	-AS-	NA-	P	- TPD T - TMPD	OF-T	VAG-4
	- Sta aurous N315	- 59	1 FG-H-A-I	. P	ATPV-TID-LP-	uc 1	
		-13			ATDY ITD LD		
	Sta. aureus M050		LEG-H-A-L	L • P	ATPT-IID-LP-		UK-IEL
	L Bac, subtilis		LKG-H-A-	I-P	STQY-IID-LP-		QK-VEL
Low C+C	Duot Cubcillo						
Low G+C	Bac. halodurans	•F••••	LEG-H-A-	INP	-TPH-IID-LP-		QK-VE -ML
Low G+C	Bac. halodurans Strep. pyogenes	-F	LEG-H-A- LNMEGF-	INP LEP	-TPH-IID-LP- STKY-IIDIMRD		QK-VE -ML QI-IE-ML
Low G+C	Bac. halodurans Strep. pyogenes L. lactis	-F -FH- -F	L EG - H - A - LNMEG F - L EG - H - FA	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD		QK-VE -ML QI-IE-ML QV-VE-M
Low G+C	Bac. halodurans Strep. pyogenes L. lactis	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD		QK-VE -ML QI-IE-ML QV-VE-M
Low G+C	Bac. halodurans Strep. pyogenes L. lactis	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYD	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD		QK-VE -ML QI-IE-ML QV-VE-M 501
Low G+C	E. coli K12	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD	IPAPLLDRMEV	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI
Low G+C	Bac. halodurans Strep. pyogenes L. lactis E. coli K12 E.coli 0157:H7	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYD	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI
Low G+C	E. coli K12 E.coli 0157:H7 E.coli 0157:H7EDL	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI
Low G+C	E. coli K12 E.coli 0157:H7 E.coli 0157:H7 E.coli 0157:H7 Buchnera sp. APS	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI
Low G+C	E. coli K12 E. coli K12 E. coli 0157:H7 E. coli 0157:H7 Buchnera sp. APS Xylella fastidiosa	-F -FH- -F 933 a	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI 
Low G+C γ-Pr	eteo eteo	-F -FH- -F 933 a	LEG-H-A- LIMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN	IPAPLLDRMEV IPAPLLDRMEV G	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP
Low G+C	E. coli K12 E. coli K12 E. coli 0157:H7 E. coli 0157:H7 E.coli 0157:H7EL Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV IPAPLLDRMEV G	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP
Low G+C	E. coli K12 E. coli K12 E. coli 0157:H7 E. coli 0157:H7 E. coli 0157:H7 E. coli 0157:H7EDL Buchnera sp. APS Xylella fastidiosi H. influenzae Vib. cholerae Past. multocida	-F -FH- -F 933 a	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSQYTEDEKLNI IP
Low G+C	oteo Bac. halodurans Strep. pyogenes L. lactis E. coli K12 E.coli 0157:H7 E.coli 0157:H7EDL Buchnera sp. APS Xylella fastidios; H. influenzae Vib. cholerae Past. multocida Pest. arutocida	-F -FH- -F	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP
Low G+C	E. coli K12 E. coli K12 E. coli 0157:H7 E. coli 0157:H7 E. coli 0157:H7EL: Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa	-F -FH- -F 933 a	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV IPAPLLDRMEV G	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI 
Low G+C γ-Pr	oteo Bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL3 Buchnera sp. APS Xylella fastidiosa H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis M	-F -FH- -F 933 a	LEG-H-A- LMMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD	IPAPLLDRMEV 	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP
Low G+C γ-Pr β-Pro	oteo Bac. halodurans Bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL3 Buchnera sp. APS Xylella fastidios; H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis MM N. meningitidis Z2	-F -FH- -F 933 a 258 2491	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD HFVATSNSMN 	IPAPLLDRMEV 	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP
Low G+C γ-Pr	Bac. halodurans Strep. pyogenes L. lactis E. coli K12 E.coli 0157:H7 E.coli 0157:H7EDL Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis Zi Ri. prowazekii	-F -FH- -F 933 a 2491	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYD 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD HFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prc α-Pr	oteo Bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7 E. coli 0157:H7EDL3 Buchnera sp. APS Xylella fastidiosa H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis ZZ Ri. prowazekii Mesorhizobium lotz	-F -FH- -F 933 a 2491 i	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP 
Low G+C γ-Pr β-Prc	bac. halodurans Strep. pyogenes L. lactis E. coli K12 E.coli 0157:H7 E.coli 0157:H7EDL9 Buchnera sp. APS Xylella fastidiosa H. influenzae Past. multocida Pse. aeruginosa N. meningitidis Zi A. meningitidis Zi Ri. prowazekii Geodo	-F -FH- -F 933 a 258 2491 i	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD HFVATSNSMN 	IPAPLLDRMEV 	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Pro α-Pro	Bac. halodurans         Strep. pyogenes         L. lactis         E. coli K12         E.coli 0157:H7         E.coli 0157:H7EDL3         Buchnera sp. APS         Xylella fastidios:         H. influenzae         Vib. cholerae         Past. multocida         Pse. aeruginosa         N. meningitidis Z:         Ri. prowazekii         Mesorhizobium lot:         C. crescentus         Camp. jejuni	-F -FH- -F 933 a 2491 i	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYD 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD HFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prc α-Prc	Bac. halodurans         Bac. halodurans         Strep. pyogenes         L. lactis         E. coli 0157:H7         E.coli 0157:H7ELL         Buchnera sp. APS         Xylella fastidios;         H. influenzae         Vib. cholerae         Past. multocida         Pse. aeruginosa         N. meningitidis Mi         N. meningitidis Z;         Ai. prowazekii         Mesorhizobium lot;         C. crescentus         Camp. jejuni         Heil puloci 199	-F -FH- -F 933 a 2491 i	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P E	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP 
Low G+C γ-Pr β-Prc α-Prc €-Prc	Bac. halodurans         Bac. halodurans         Strep. pyogenes         L. lactis         E. coli K12         E.coli 0157:H7         E.coli 0157:H7EL:         Buchnera sp. APS         Xylella fastidios:         H. influenzae         Vib. cholerae         Past. multocida         Pse. aeruginosa         oteo         N. meningitidis M         N. meningitidis Zi         oteo         Ri. prowazekii         Mesorhizobium loti         C. crescentus         Camp. jejuni         Hel. pylori J99         Webleree	-F -FH- -F 933 a 2058 2491 i	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDD 	INP LEP L-P E	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD HFVATSNSMN 	IPAPLLDRMEV 	0K-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prc ε-Prc	Bac. halodurans         Strep. pyogenes         L. lactis         E. coli 0157:H7         E.coli 0157:H7EDL3         Buchnera sp. APS         Xylella fastidios:         H. influenzae         Vib. cholerae         Past. multocida         Pse. aeruginosa         N. meningitidis Z;         Ai. prowazekii         oteo         Camp. jejuni         Hel. pylori J99         Hel. pylori 2695	-F -FH- -F 933 a 2491 i	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDI 	INP LEP L-P 	-TPH-IID-LP- STKY-IIDIMRD ETKYIDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M 501 IRLSGYTEDEKLNI IP
Low G+C γ-Pr β-Prc ε-Prc	Bac. halodurans         Bac. halodurans         Strep. pyogenes         L. lactis         etc. coli 0157:H7         E.coli 0157:H7EL1         Buchnera sp. APS         Xylella fastidios;         H. influenzae         Vib. cholerae         Past. multocida         Pse. aeruginosa         oteo         N. meningitidis Mi         Mesorhizobium lot;         C. crescentus         Camp. jejuni         Hel. pylori J99         Hel. pylori J99         Hel. trachomatis	-F -FH- -F 933 a 2491 i	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M DV-VE-M IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prα ε-Prα	bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL3 Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis Zi Astronomic Construction N. meningitidis Zi Ri. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori 26695 Chl. trachomatis Chlamydia muridaru	-F -FH- -F 933 a 2491 i um	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDD 	INP LEP L-P	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M SO1 IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prc α-Prc	bac. halodurans Strep. pyogenes L. lactis	-F -FH- -F 933 a 2491 i um R39	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDI 	INP LEP L-P 	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M DV-VE-M DIRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prc ε-Prc Chlam	Bac. halodurans         Bac. halodurans         Strep. pyogenes         L. lactis         etcoli 0157:H7         E.coli 0157:H7         E.coli 0157:H7EL:         Buchnera sp. APS         Xylella fastidios;         H. influenzae         Vib. cholerae         Past. multocida         Pse. aeruginosa         oteo         N. meningitidis Mi         N. meningitidis Zi         oteo         Ai. prowazekii         Mesorhizobium lot;         C. crescentus         Camp. jejuni         Hel. pylori J99         Hel. pylori 26695         Chlamydia muridari         Chlamydia muridari         Chl. pneumoniae Ai	-F -FH- -F 9333 a 2491 i i um R39 138	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDI 	INP LEP L-P E	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD HFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M DV-VE-M IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Pro ε-Pro Chlam	bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL3 Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis Zi Ast. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori J99 Hel. pylori 26695 Chl. trachomatis Chlamydia muridari Chl. pneumoniae Al Chl. pneumoniae Al	-F -FH- -F 933 a 2491 i i	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDD 	INP LEP L-P E	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M QV-VE-M IRLSGYTEDEKLNI  
Low G+C γ-Pr β-Prc ¢-Prc Chlam	bac. halodurans Strep. pyogenes L. lactis etc. coli K12 E. coli 0157:H7 E. coli 0157:H7 E. coli 0157:H7EDL Buchnera sp. APS Xylella fastidios; H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis Mi N. meningitidis Z: A. meningitidis Z: Ri. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori J99 Hel. pylori J99 Hel. pylori J6695 Chl. trachomatis Chl. pneumoniae A Chl. pneumoniae J Chl. pneumoniae C	-F -FH- -F 933 a 2491 i i um R39 138 WL029	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P 	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE - M L QI - IE - M L QV - VE - M QV - VE - M IRLSGYTEDEKLNI IP I I I I - I
Low G+C γ-Pr β-Prc ε-Prc Chlam	bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL9 Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis Zi oteo Fi. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori J99 Hel. pylori J99 Hel. pylori 2695 Chl. trachomatis Chlamydia muridari Chl. pneumoniae J Chl. pneumoniae Ch	-F -FH- -F 933 a 2491 i i um R39 138 WL029	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDI 	INP LEP L-P E	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M QV-VE-M IRLSGYTEDEKLNI 
Low G+C γ-Pr β-Pro ε-Pro Chlam	bac. halodurans Strep. pyogenes L. lactis E. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL9 Buchnera sp. APS Xylella fastidios: H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis Zi Ri. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori J99 Hel. pylori 26695 Chl. trachomatis Chlamydia muridari Chl. pneumoniae AI Chl. pneumoniae Ci Aqu. aeolicus Bor. burgdorferi	-F -FH- -F 933 a 2491 i i um R39 138 WL029	LEG-H-A- LNMEGF- LEG-H-FA DHYLEVDYDI 	INP LEP L-P L-P 	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M QV-VE-M IRLSGYTEDEKLNI  
Low G+C γ-Pr β-Prc ε-Prc Chlam	bac. halodurans Strep. pyogenes L. lactis etc. coli K12 E. coli 0157:H7 E. coli 0157:H7EDL9 Buchnera sp. APS Xylella fastidios; H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis MM N. meningitidis ZZ Al. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori J99 Hel. pylori 26695 Chl. trachomatis Chamydia muridaru Chl. pneumoniae Al Chl. pneumoniae Chl. Aqu. aeolicus Bor. burgdorferi Tre. pallidum	-F -FH- -F 933 a 2491 i i um R39 138 WL029	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P LSDVN E- E- 	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD ETKY-IDIMRD IFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M DV-VE-M DIRLSGYTEDEKLNI 
Low G+C γ-Pr β-Prc α-Prc €-Prc Chlam	bac. halodurans Strep. pyogenes L. lactis E. coli K12 E.coli 0157:H7 E.coli 0157:H7EDL' Buchnera sp. APS Xylella fastidios. H. influenzae Vib. cholerae Past. multocida Pse. aeruginosa N. meningitidis M. N. meningitidis Z. oteo [N. meningitidis Z. oteo [N. meningitidis Z. oteo [N. meningitidis Z. oteo [N. meningitidis Z. fi. prowazekii Mesorhizobium lot: C. crescentus Camp. jejuni Hel. pylori 26695 Chl. trachomatis Chl. pneumoniae J. Chl. pneumoniae G. Aqu. aeolicus Bor. burgdorferi Tre. pallidum D. radiodurans	-F -FH- -F 933 a 258 2491 i i um R39 138 WL029	LEG-H-A- LNMEGF- LEG-H-FA 454 DHYLEVDYDI 	INP LEP L-P 	-TPH-IID-LP- STKY-IIDIMRD ETKY-IDIMRD ETKY-IDIMRD HFVATSNSMN 	IPAPLLDRMEV 	QK-VE -ML QI-IE-ML QV-VE-M QV-VE-M IRLSGYTEDEKLNI 

M. pulmonis

U. urealyticum Bac. subtilis Bac, halodurans

Low G+C

M. genitalium M. pneumoniae

Fig. 9 Alignment of Lon protease sequences from bacterial genomes, showing a 1-aminoacid insert (boxed) that is commonly shared by all  $\alpha$ -,  $\beta$ -, and  $\gamma$ -proteobacteria

another, then the presence or absence of these indels in different species will not follow any predicted pattern. In such a case, different groups of species, or even individual species from different groups, will either contain or lack the indels.

A summary of the results for the various indels studied in this work is presented in Table 2. For each of the proteins containing these indels, the number of species where the protein was found is indicated, together with the number of species in which the indel was expected to be present or absent according to the model. The last column indicates the number of exceptions observed where the presence or absence of an indel was not in accordance with the indel model. As seen from Table 2, the proteins containing these indels are widely represented in different bacteria and many of them were found in all sequenced bacterial genomes. A few of these proteins are absent from species such as

-N-V-E-----K---I-A-YIE D --EA-----I-E-TS--Q--IE--N-V-E-----K-----A-YIE D --EA------E-TS--Q--Q--

		263	· · · · · · · · · · · · · · · · · · ·	328
	∟ E. coli K12	FSVDEKSRQVNLTERGLVLIEELL	VKEGIMD	EGESLYSPANIMLMHHVTAALRAHALFTRDVDYIV
	E.coli 0157:H7			
	E.coli 0157:H7EDL933			
	Buchnera sp. APS	IK-IYIKV-KI-	FDKKL-N	TSNILSKV-NL-
v-Proteo	Pas. multocida	-TL-L-TK-AHQEKV-QW-	TEQ-L-S	AESK-S-LYTE
,	Xylella fastidiosa	GKH-S-V-MERA	HQA LG	-EDAAQ-LSVVLNYQ
	H. influenzae	- TL - L K - AH QEKV - DW -	IAQ-L-P	DSR-V-LMTEK
	Vib. cholerae	-TKHT-QEFV	N-M-Q	DTS-LNVEKN
r	L Pse. aeruginosa	Y-ITE-N-Q-HQFD	SQN-LLG	AH-LS-LTYTH-N-E
B-Proteo	r N. meningitidis MC58	YWAHI-S-A-HEHA-QI-	TQM-LLA	-NDAALMTHK-QH-VI
piroteo	🛛 N. meningitidis Z2491	YWAHI-S-A-HEHA-QI-	TQM-LLA	-NDAALMTHK-QH-VI
	<sub>r</sub> Ri. prowazekii	-EKLKTIA-ITHS	I-DIK	PDTGDFE-LN-V-Y-NQHMIL-
α-Proteo	Mesorhizobium loti	YEIQKTSIFE-TEKL-N	RDADLL	KDVE-VAIVNNKRQK-K
	L C. crescentus	-DHQKID-QEKI-	MSANLAE	DSAGRRVSVVNQNI-IK
	<sub>r</sub> Camp. jejuni	-VN-NILIA-IAKA-K-F		GV-NLD-AI-A-QLDQKNEKH-VL
€-Proteo	Hel. pylori, J99	-TIN-AILIE-IKKA-N-F		GVDNKIE-AA-SLDQK-NYFI-K
	L Hel. pylori 26695	-TIN-AILIE-IKKA-N-F		GVDNKIE-AA-SLDQK-NYFI-K
	<sub>C</sub> Chl.Pneumoniae AR39	RGL-SFLDVDI-PKDKKEGISE		FCR WLVSKG - PLNR - LRRV - E - PDLRAMI - KWD
Chi in Pro-	Chl.Pneumoniae J138	RGL-SFLDVDI-PKDKKEGISE		FCR WLVSKG - PLNR - LRRV - E - PDLRAMI - KWD
Chiamydia	Chl.Pneumoniae CWL029	RGL-SFLDVDI-PKDKKEGISE		FCR WLVSKG - PLNR - LRRV - E - PDLRAMI - KWD
	Chl.trachomatis	RGL-SFLDVDI-PKDKKEGISE		FCRWLVSKG-PLNR-LRRV-E-PDLRAMI-KWD
	L Chl. muridarum	KTL-PFLGMDV-PKDRKIMEGISE		ACRA-WLVSKG-PLNR-LRRV-E-PDLRAMKWD
	Aqu. aeolicus	-TN-TQ-IKKV-KM-		GIDNDLKHVD-L-AILQSIHKKH
Spirochetes	<sub>r</sub> Tre. pallidum	YIN-K-SFSGP-MLH-QDV-	THAGL	IQGFDEE-FKYI-YF-QL-YRAV-
	L Bor. burgdorferi	YTAKRISF-AKNNL-Q	VSKGI	ISG-M-TDS-FNYV-YM-QKLLKNRE
	Sy. sp. PCC6803	YEG-N-LDQ-FINA-Q		GVSD-FDSNDPWA -YIFN-IK-KEIKN
	D. radiodurans	YTIEKA-HQ-ITKR		SLKDE-MDKA-MI-Q-IRE-YH-EK
	T. maritima	-TA-TIIE-VAKA-KII		GV-ND-G-VS-LY-LINK-LHKKV-
	<sub>Г</sub> Myc. tuberculosis H37Rv	YELRK-T-GVH-K-VEFV-DQ-		GIDNETSP-VSYLNNK-KESK
High G+C	Myc. tuberculosis CDC1551	YELRK-T-GVH-K-VEFV-DQ-		GIDNETSP-VSYLNNK-KESK
-	L Myc. leprae	YDT-SDN-HDV-ARKV-KA-		G-IDEEHVGTTLTEVNHV-LQH
	<sub>F</sub> Sta. aureus N315	YKYTKA-HQ-ADKA-RMF		KV-NDVQ-VDVIS-INTVTLQM-
	Sta. aureus MU50	YKYTKA-HQ-ADKA-RMF		KV-NDVQ-VDVIS-INTVTLQM-
	Bac. halodurans	YTLTKS-QE-VNKA-RAF		NIDN-FDQRHVQ-LINQ-MKVVMHAV-
	Bac. subtilis	YTY-I-TKA-QE-MTKA-KAF		GIDN-FDVKHVA-NINQKVAMQKV-
Lawrence	Lactococcus lactis	YKI-LQ-KTISE-IDKA-KFF		QI-NDME-VA-T-F-DNNFIMLH-IM-
LOW G+C	U. urealyticum	-TL-PE-QS-AS-VEKAQKFF		NTKNY-NFE-SDII-KNNFT-FNGRE
·	Strep. pyogenes	YVI-VPTKTIG-SDS-IDKA-SYF		NLSNDIE-VA-T-FIDNNYIMLL-IV-
	M. pneumoniae	YKI-PEAPAL-IKHA-KNF		KTDN-FALE-SD-F-KIINT-VKV-EQGKE
	M. pulmonis	YEIE-KTIK-VDS-IDKANKFF		TLSNDIK-SE-V-RIQNNFIMKKE
	L M. genitalium	-KI-PEAASL-IKKA-QTF		KK-N-FALE-SD-F-KIMNG-T-VKV-EQGKE

**Fig. 10** Alignment of SecA homologues from bacterial genomes, showing a 7-amino-acid insert (*boxed*) that is common to all  $\alpha$ -,  $\beta$ -, and  $\gamma$ -proteobacteria

mycoplasms or chlamydiae, which are intracellular pathogens, where the corresponding genes may have been lost because the cellular functions of these proteins are likely provided for by the host [15, 62]. For all of these proteins, the distribution of indels in various proteins was found to be exactly as predicted by the model, with no exceptions observed. Of a total of 450 indels whose distribution in different species was examined in the present work, all of them showed the expected distribution, as predicted by the model. The only possible exception was the presence of a second Hsp70 homologue in B. burgdorferi, which lacked the large insert in the protein, distinctive of Gram-negative bacteria. The BLAST searches indicate that this gene is likely acquired from Gram-positive bacteria by means of LGT. However, *B. burgdorferi* contains another Hsp70 homologue with the expected characteristics. Hence, the presence of this laterally transferred gene, which is readily identified as such and which is absent from other spirochete species, does not in any way confuse or affect the inference concerning the phylogenetic placement of this species. In a few cases, some species were found to contain a different kind of indel (differing in length, amino acid composition, species specificity) in a similar position as the indicated signature. Such indels, which are probably of independent origin, again do not confuse or affect the inference from specific indels. For all of the studied proteins, in addition to the data from completed bacterial genomes, sequence information is available from a large number of other species and, in almost all cases, the distribution of these indels in various species follows the pattern as predicted by the indel model [19, 23]. These results provide strong evidence that the inferences derived from indel data are reliable [48] and they are not affected to any significant extent by other factors, such as LGT [63] or independent occurrence of these indels in different species.

The evolutionary relationship that emerges based on indels, in addition to its high degree of internal consistency in the placement of species into different groups and in determining their relative branching order, is also quite appealing from other perspectives:

- 1. The model is consistent with and accounts for the major ultrastructural differences seen among the Bacteria. The model indicates that the bacterial groups surrounded by a single membrane (i.e., Grampositive or monoderm bacteria) are phylogenetically distinct from those surrounded by both an inner and outer membrane and containing a periplasmic compartment (i.e., all true Gram-negative bacteria or diderm bacteria) [19, 47]. Of these two structurally and phylogenetically distinct groups of bacteria, the monoderm bacteria are indicated to be ancestral.
- 2. The model places *Deinococcus-Thermus* in an intermediate position between monoderm and diderm bacteria. This placement is consistent with the observation that *Deinococcus* contains a thick

**Fig. 11** Sequence alignment of Hsp70 homologues showing a 4-amino-acid insert (*boxed*) that is distinctive of  $\beta$ - and  $\gamma$ -proteobacteria

	190	233
⊢E. coli K12	IAVYDLGGGTFDISIIEI DEVD	GEKTFEVLATNGDTHLGGEDED
E.coli 0157:H7		
E.coli 0157:H7 EDL933		
Pasteurella multocida		
γ-Proteo Xylella fastidiosa		0FF
Vib. cholerae	F	s
Buchnera, sp. APS		К
Pse. aeruginosa	VI A	HQFF
H. influenzae		QG-N
- Nei, meningitidis MC58	V ANL -	-D-QF
β-Proteo Nei, meningitidis Z2491	V ANI -	-D-QF
-Bi, prowazekij	····	SDGVKSF
C. crescentus	V	-DGVKSF
α-Proteo	VI	-DGVKSF
Camp jejunj	-VVTVI -T	
Hel pylori 26695	-MVTVI - T	
€-Proteo Hel pylori /99	-MVTVL-T	
_ Chl trachomatis		-DGVSD
Chlamydia muridarum	F	-DGVSD
Chlamudia Chlm preumonice CWI 029		-DGVSID
Chim. pheumoniae AR39		
	al as Francis GVS-1 F	
- Tre pallidum		-DGVKSD
Spirochetes Bor burgdorferi	V	-DGVKSDN
	-L-EVL-V	
Sy sp PCC6803 (2)	-I-FV-II0	-NGVS-S-NND
D radiodurans	VI - E VT - I - I	-DGVKS-AA
T maritima	VI V I	GVTTA-NND
- Myc tuberculosis CDC1551	-I-FV-LL	GVVBSND-W-
High C+C Myc. tuberculosis H37 By	-L-FV-LL	GVVRSND-W-
Myc. leprae	-L-FV-LL	GVVRSND-W-
- Bac, halodurans	-LVL-L	-DGFKSNKD
Bac subtilis	-LVL-L	-DGVRS-ANRD
Strep pyogenes	-L-FVL-L	-DGV-DANKD
Sta, aureus MU50	VL-FVL-L	-DGVS-ANKD
Sta, aureus Z2491	VL-FVL-L	-DGVS-ANKD
Low G+C	-L-FVL-L	-DGV-DA-NNKD
U. urealvticum	- L - F V - V L DM	ADGS-SND-W-
M. genitalium	VLV-LLD-	A-GANRD-W-
M. pulmonis	VLV-VL-L	ENGS-SND-W-
L <sub>M</sub> , pneumoniae	VLV-LLD-	A-GANRD-W-

peptidoglycan layer characteristic of Gram-positive bacteria and shows a positive Gram-staining reaction [46]. However, this species contains both inner and outer membranes, which is the main defining characteristic of Gram-negative bacteria. Thus, *Deinococcus* is indicated to be an intermediate in the transition between monoderm and diderm bacteria and it provides suggestive evidence that, in the development of Gram-negative bacteria from Grampositive bacteria, the outer membrane evolved first, before the changes in the cell wall occurred [19].

3. For 39 of the 41 bacterial species whose genomes have been sequenced, their placement into different groups based on indel data is in agreement with that based on the 16 S rRNA. The two species (i.e., Aquifex aeolicus, T. maritima) whose phylogenetic placements differed somewhat from that based on rRNA, show very deep branching in the rRNA trees [40, 51]. Indel data places Aquifex in a similar position as the Chlamydia and Cytophaga-Bacteriodes groups. This inference is based on a number of different signatures, all of which place it in the same position. It is difficult to account for these results by LGT from other species [3]. The branching of Aquifex in a similar position as Chalmydia is also observed in phylogenetic trees based on a number of different proteins including: RNA polymerase  $\beta$ - and  $\beta'$ -subunits [35] and group I sigma factor [18]. The other difference seen between the indel data and

rRNA trees concerns the branching position of *T. maritima*. The rRNA phylogenies place this species in a distinct deep-branching group, whereas the indel data groups this species with other Gram-positive bacteria. Note that, although *T. maritima* (based on the absence of a large insert in Hsp70) has been grouped with the Gram-positive group, the signature sequences in ribosomal S12 protein and DNA gyrase A subunit indicate that it is distinct from both the traditional low G+C and the high G+C Grampositive bacteria [19]. It is thus probable that *T. maritima* forms a separate, deep lineage within the Bacteria, showing a close affinity to the Gram-positive bacteria.

Phylogenetic analysis based on indel data complements the major limitations of the 16 S rRNA trees

An important point that emerges from these studies is that the evolutionary inferences based on indel data are not contradictory to those based on 16 S rRNA trees, but complement such studies in important respects. The two main recognized weaknesses of the rRNA phylogenies are: (1) it has proven difficult to define the main groups within the Bacteria in clear molecular terms and (2) the rRNA trees cannot resolve the relative Fig. 12 Alignment of phosphoribosylpyrophosphate synthetase, showing a 1-amino-acid insert (boxed) distinctive of  $\beta$ - and  $\gamma$ -proteobacteria

		75	- 131
	<sub>r</sub> E. coli K12	DALRRASAGRITAVIPYFGYARQDRRVRS	A RVPITAKVVADFLSSVGVDRVLIVDLH
	E. coli 0157:H7		-   T
	E. coli 0157:H7EDL933		•   <b>- - - -</b>
	Buc. sp. APS	-S	-   T
γ-Proteo	X. fastidiosa	KVSSVSM I	LA-KMI-AI-ATI
	Hae. influenzae		TC
	Pse. aeruginosa	FSTP	A - S M - TV N T
	Vib. cholerae	M	-   TI
	Past. multocida		-   TC
B-Proteo	<sub>r</sub> Nei. meningitidis Z2491	KTAP \	VSLNM-Y-A-IT
p=110tco	Nei.meningitidis MC58	KTAP \	/SLNM-Y-A-IT
	<sub>r</sub> C. crescentus	KGKKTGG	-┘-TSLNLITRS-ATM
α-Proteo	L Meso. loti	FM-SKASG	-TSLNMITRATL
	<sub>r</sub> Camp. jejuni	KANP	LNLIQAA-IATI
E-Proteo	Hel. pylori 26695	KAAP	MNLMQEIE-IITM
C I I OLLO	└ Hel. pylori J99	KAAP	MNLMQEIE-IITM
	Aqu. aeolicus	VK-S-PKEVYA-GQDKP	-TSLLIQKA-ANIV
a	<sub>F</sub> Trep. pallidum	V-H-GV-L-L-TYP-SHKK CG	-EGLGLLGSVYEYLSHIVTL
Spirochetes	L Bor. burgdorferi	CMQ-K-NSVSVISYP-SKKHS	-ECLSLIGREEL-IRHI-TL-I-
	Sy. sp. PCC6803	CRQLYAKTAG	-ES-SLNLITGA-AQAM
	D. radiodurans	AKSVYSS-KKDSP	-IS-AGRLL-QEA-ATMT
	T. maritima	FKNT-AVYKAKG	-DSLNLITVA-ATT
	<sub>r</sub> Myc. tuberculosis CDC1551	K-GKM-FYPKKH-G	-ES-RLIL-KTA-AIVT
High G+C	Myc. leprae	K-GKFYPKKH-G	-ES-RLL-KTA-AIVT
Ť	└ Myc. tuberculosis H37Rv	K-GKM-FYPKKH-G	-ES-RLIL-KTA-AIVT
	<sub>r</sub> Strep. pyogenes	KEK-SV-MYKA	-ES-LNM-EVAL-T
	Bac. subtilis	KKT-NIYKA	-ELF-NL-ETA-ATIAL
	Bac. halodurans	VKKT-NVYKA-A	-ELNL-ETA-ATTL
	Lac. lactis	KA-A	-ES-LNM-QIA-ALITF
Law CIC	Sta. aureus	CKAT-NI-VYKA	-ELNLIETA-AT-MIAL
LOW G+C	Sta. aureus NCTC 8325	CKAT-NI-VYKA	-ELNLIETA-AT-MIAL
	M. pulmonis	-SKT-NVILS-YKAEG	-QALLL-QVA-IS-IVV
	U. urealyticum	-SIKKA-SVYKAKP	-ERLKMIE-A-ATSTW-I-
	N nnoumoniae	KTMG	-ES-LL-TTASALT-T-

M. genitalium

Fig. 13 Sequence alignment of 5'-phosphoribosyl aminoimidazole-4-carboxamide transformylase, showing a 2-amino-acid deletion that is distinctive of the y-proteobacteria

		59	108
	<sub>Γ</sub> Ε. coli K12	GFPEMMDGRVKTLHPKVHGGILGRRGQDDAI	MEEHQIQPIDMVVVNLYPF
	E. coli 0157		
	E. coli 0157:H7EDL933		
	Pse. aeruginosa	G-	-AQ-GI
v-Proteo	Vib. cholerae	· · · · · · · · · · · · · · · · · · ·	-NT-G
1 110000	Pas. multicoda	TEV	- SQQG - EG
	Xylella fastidiosa	A-IV	-AK-G-ALLIL
	Buchnera sp. APS	IIMQKQK-QE-	-KLYN-CI-IF
	H. influenzae	·····T····	
G Ductoo	r Nei. meningitidis MC58	DLPEHV	AKG-GNL-C
p-Proteo	Nei. meningitidis Z2491	DLPEHV	AKG-GNL-C
	г C. crescentus	LV-DAA-HA	KA - AD-G-GGILY
a c-Proteo	Mesorhizobium loti	ISALV-DDPEHA	AA -RKYG-ELL-S
w,e moteo	L Camp. jejuni	HK-SDENH-	KQ AK-NEXLGL-C
	Agu, aeolicus	F-DWVEKDK	EE I-K-G-KV
	Sv. sp. PCC6803	-AILGRIADLPSDQ	AD L-AND-R-L-L
	D. radiodurans	AEAG HL	GQ LAAQD-GTL-C
	T. maritima	ENLLG-LEIFAPEPR	W-V-F-DP
	г Myc. leprae	VLRA-L-ADLRKPEHA	AA L-QLG-EAFEL
High G+C	Myc. tuberculosis H37 Rv	VLRA-L-ADLRKSEHA	AA L-QLG-EAFEL
ing die	Myc. tuberculosis CDC1551	VLRA-L-ADLRKSEHA	AA L-QLG-EAFEL
	г Bac. subtilis	ILNIL-AVNEEHM	AQ INGL
	Bac. halodurans	ILNIL-AM-ER-EHL	AQ LNH-RF
	Lactococcus lactis	DLESHM	KS -TH-SL
Low G+C	Sta. aureus N315	IAAD-NKPQHL	NE LS-QH-DL
	Sta. aureus MU50	IAAD-NKPQHL	NE LS-QH-DL
	L Strep. pyogenes	NIL-ADA-SHL	QAAKDNN-ELL

---K-G--KS---IL--Y-----KTKG

branching order of the main groups. These are in fact strong points for the signature sequence approach. The main reason for the success of the signature sequence approach in these regards is that the derived inferences in this case are based on minimal assumptions [19, 56, 58, 63]. The sole assumption involved in these analyses is that when a shared conserved indel is present in different groups of species, it is assumed to have been introduced only once in a common ancestor of these groups, rather than on multiple occasions in different species. This is the most parsimonious way to explain these results. In contrast, the branching patterns of species in phylogenetic trees are dependent upon and affected by a large number of variables and assumptions (e.g., sequence regions that are retained or excluded, the number and range of species examined, differences in the evolutionary rates between species, base compositional differences between species, phylogenetic methods employed, order in which different species are added to the alignment, etc.) and hence are not clearly resolved [19, 38, 70].

Based on the various indels described here, it is now possible to define in clear molecular terms most of the major groups within the Bacteria that were previously

-E---S-LI--M-TKA-AN--VLT-I-

200

Protein	Signature description	No. of genomes with protein	Genomes lacking the protein	No. of genomes with insert (expected/found)	No. of genomes lacking the insert (expected/found)	Exceptions observed
Hsp70/DnaK	21–23-a.a. $G + /G$ – insert	41	None	27/27	14/14	0
Ribosomal S12 protein	13-a.a. low $G + C$ signature	41	None	37/73	31/31	0
Hsp60/GroEL	1-a.a. insert after <i>Deinococcus</i>	39	mp, uu	26/26	37/68	0
FtsZ protein	1-a.a. insert after cyanobacteria	33	ct, cp, cm, mn, mg, uu	20/20	37/68	0
Ata-tRNA synthetase	4-a.a. common to <i>Chlamydia</i> /proteobacteria	41	None	23/23	18/18	0
Hsp70/DnaK	2-a.a. proteobacterial insert	41	None	17/17	24/24	0
CTP Synthetase	10-a.a. proteobacterial insert	37	mp, mg, uu, mn	17/17	20/20	0
Lon protease	1-a.a. $\alpha\beta\gamma$ -proteobacterial deletion	33	ll, mt, ml, sa, sp	19/19	14/14	0
SecA protein	7-a.a. $\alpha\beta\gamma$ -proteobacterial insert	41	None	14/14	27/27	0
HSP70/DnaK	4-a.a. $\beta\gamma$ -proteobacterial insert	41	None	37/05	30/30	0
PRPP synthetase	1-a.a. $\beta\gamma$ -proteobacterial insert	35	cp, ct, cm, rp	37/05	24/24	0
PAC-transfor mylase	2-a.a. γ-proteobacterial deletion	27	bb, cp, cm, ct, hp, mp, mg, tp, uu, rp	18/18	37/42	0

The abbreviations used are: a.a., amino acid; bb, *Borrelia burg-dorferi*; cm, *Chlamydia muridarum*; cp, *Chlamydia pneumonia*; ct, *Chlamydia trachomatis*; G+, Gram-positive; G-, Gram-negative; hp, *Heliobacter pylori*; ll, *Lactococcus lactis*; mg, *Mycoplasma* 

genitalium; mn, M. pneumonia; mp, M. pulmonis; ml, Mycobacterium leprae; mt, Myc. tuberculosis; rp, R. prowazekii; sa, Staphylococcus aureus; sp, S. pyogenes; tp, Treponema pallidum; uu, Ureaplasma urealyticum

identified solely on the basis of their branching pattern in the 16 S rRNA trees. For example, the low G+CGram-positive group can be defined by the presence of the large insert in the S12 protein. The high G+CGram-positive group can be defined by the lack of the large inserts in both the Hsp70 protein and the S12 protein. A flow chart detailing how these indels could be used to taxonomically define the different main groups within the Bacteria and for assigning any given species to one of these groups has been described in earlier work [23]. The branch orders of different groups as deduced, based on these signatures, is internally highly consistent and it is difficult to explain these results by any other reasonable mechanism [63]. It should be recognized, however, that the number of main groups within the Bacteria that can presently be identified by signature sequence represents the minimal number. As additional signature sequences are identified in future work, the relative branching orders of species within some of the presently defined groups should become clearer; and this may lead to further divisions of these groups. We expect this to be the case for the low and high G+C Grampositive bacteria and for the Aquifex, Chlamydiae, and Cytophaga groups, which have not been studied in detail for the presence of signature sequences. It is expected, however, that any newly identified group should be placed in an adjoining position to the presently assigned position and it should not affect the overall branching order of the other groups.

#### Conclusions

Results presented here show that the conserved indels that have been identified in various proteins provide a powerful new approach for understanding bacterial phylogeny. Based on these signatures, most of the main groups within the Bacteria can be identified in clear molecular terms and any given bacterial species could be assigned to one of these groups in an unambiguous manner. The phylogenetic assignment of different bacteria whose genomes have been sequenced using this approach showed an excellent correlation to that based on the 16 S rRNA, with 39 of the 41 species similarly assigned. Thus, the inferences deduced based on this new approach are not contradictory to the 16 S rRNA trees, but complement it in important respects. One distinct advantage of this new approach is that it permits a logical deduction of the relative branching order of different groups of bacteria from a common ancestor (Fig. 1), which could not be resolved from phylogenetic trees based on the 16 S rRNA or various proteins and constituted a major unresolved problem in bacterial phylogeny. The deduced branching order of different groups shows a very high degree of internal consistency and it is strongly supported by the analyses of completed bacterial genomes. As sequence information from other bacterial genomes becomes available, it should be possible to further determine: (1) whether the results

obtained are in accordance with this model and (2) the ability of this model to help explain and integrate different observations.

#### Appendix

The following figures illustrate the alignment of various protein sequences from completed bacterial genomes, as discussed in the text.

The first five groups of proteins in this Appendix cover the ribosomal S12 protein (2), Hsp70 protein (3), Hsp60/GroEL protein (4), FtsZ protein (5), and alanyl-tRNA synthetase (6).

The remaining seven groups cover signature sequences for proteobacteria in Hsp70 and CTP synthase (7, 8), signature sequences indicating the branch order of the proteobacterial groups (9, 10), and useful signatures for the  $\beta$ - and  $\gamma$ -proteobacteria in the Hsp70 family of proteins (11, 12, 13).

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