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Functional insights from the distribution and role of homopeptide repeat-containing proteins

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Expansion of “low complex” repeats of amino acids such as glutamine (Poly-Q) is associated with protein misfolding and the development of degenerative diseases such as Huntington’s disease. The mechanism by which such regions promote misfolding remains controversial, the function of many repeat-containing proteins (RCPs) remains obscure, and the role (if any) of repeat regions remains to be determined. Here, a Web-accessible database of RCPs is presented. The distribution and evolution of RCPs that contain homopeptide repeats tracts are considered, and the existence of functional patterns investigated. Generally, it is found that while polyamino acid repeats are extremely rare in prokaryotes, several eukaryote putative homologs of prokaryote RCP—involved in important housekeeping processes—retain the repetitive region, suggesting an ancient origin for certain repeats. Within eukarya, the most common uninterrupted amino acid repeats are glutamine, asparagines, and alanine. Interestingly, while poly-Q repeats are found in vertebrates and nonvertebrates, poly-N repeats are only common in more primitive nonvertebrate organisms, such as insects and nematodes. We have assigned function to eukaryote RCPs using Online Mendelian Inheritance in Man (OMIM), the Human Reference Protein Database (HRPD), FlyBase, and Wormpep. Prokaryote RCPs were annotated using BLASTp searches and Gene Ontology. These data reveal that the majority of RCPs are involved in processes that require the assembly of large, multiprotein complexes, such as transcription and signaling.

[Supplemental material is available online at www.genome.org.]

Single amino acid repeats are regions within proteins that comprise a single homopolymeric tract of a particular amino acid. Uncontrolled genetic expansions of such regions have been shown to lead to the development of serious debilitating human diseases. For example, expanded poly-Q and poly-A tracts are associated with the development of neurological disorders such as Huntington disease and Oculopharyngeal Muscular Dystrophy (OPMD), respectively. Several studies have also demonstrated that many nondisease-linked polyamino acid tracts are toxic to cells and/or lead to protein aggregation or misfolding (Dorsman et al. 2002; Fandrich and Dobson 2002).

Of the polyamino acid repeats characterized to date, poly-Q repeats are the most extensively studied. Nine poly-Q-linked diseases have been identified, and the proteins believed to be responsible for the disease contain expanded poly-Q tracts that have been shown to possess an enhanced tendency to aggregate and form fibrils both in vitro and in vivo (Scherzinger et al. 1997; Skinner et al. 1997; Becher et al. 1998; Holmberg et al. 1998; Li et al. 1998; Warrick et al. 1998; Chow et al. 2004c). The pathogenic length of the poly-Q tract appears to be specific to each protein family (Cummings and Zoghbi 2000). For example, Huntington’s

disease only develops when the poly-Q repeat within the Huntingtin protein is 38 amino acids (generally encoded by 36 CAG repeats + CAA + CAG). In contrast, Machado-Joseph disease develops when the poly-Q repeat in ataxin-3 is 45 amino acids in length (Cummings and Zoghbi 2000; Chow et al. 2004a). The accumulation of the aggregated state in vivo appears to correlate with cell death and the onset of degenerative disease; however, the mechanism of poly-Q toxicity remains controversial. While the aggregated or fibrillized conformation of poly-Q tracts is postulated to be β -sheet rich, the precise structure and mechanism of fibril formation remains to be characterized (Wetzel 2002; Chow et al. 2004b). Furthermore, the function (if any) of a typical poly-Q tract remains to be determined. In contrast, studies focusing on the function of poly-Q RCPs have revealed interesting functional patterns; an investigation of *Drosophila melanogaster* poly-Q RCPs revealed that many of these proteins are transcription factors involved in development (Karlin and Burge 1996; Alba and Guigo 2004).

More recently, proteins containing expanded alanine tracts have been linked to several human diseases (Brown and Brown 2004). Like poly-Q RCPs, many of the disease-linked alanine RCPs are transcription factors (Brown and Brown 2004), and proteins containing lengthened alanine tracts (>10) also demonstrate an enhanced tendency to aggregate and form fibrils (Fan et al. 2001). Structural studies have revealed that while short poly-alanine peptides form α -helices (Giri et al. 2003), the secondary structure of longer polyalanine tracts is predicted to be predominantly β -strand like (Giri et al. 2003). It is thus suggested that

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longer alanine-tracts possess an enhanced tendency to form β -sheet-rich fibrillar structures.

Several other repeat types have also been investigated. A polyglycine tract in the plant protein Toc-75 (a component of the protein import machinery in the chloroplast) has been shown to be important for targeting this protein to the outer envelope of the chloroplast (Inoue and Keegstra 2003) and several viral polyarginine-rich proteins have been shown to be involved in RNA binding (Calnan et al. 1991; Nam et al. 2001). Finally, a recent biochemical study (Oma et al. 2004) revealed that long amino acid tracts of almost all types possess a general tendency to aggregate.

To date, the role of many amino acid repeats and RCPs remains somewhat obscure, and it is likely that numerous disease-linked RCPs remain to be identified. To begin to address this problem, a global genome survey has been performed to identify all homopeptide RCPs; these data have been stored in an online database. Resources such as Online Mendelian Inheritance in Man (OMIM) and FlyBase were used to map function onto eukaryote RCPs. BLASTp and Gene Ontology were used to functionally annotate where possible prokaryote RCPs. When considered as a whole, striking functional patterns, independent of amino acid type, can be observed across all RCPs; these data reveal that the majority of RCPs perform roles in processes that require the assembly of large multiprotein or protein/nucleic acid complexes.

Results

A Web-accessible database of RCPs

We identified all homopeptide repeats in GENPEPT greater than six amino acids in length; these data are available at <http://repeats.med.monash.edu.au>. The homepage includes a table listing the number of RCPs for each amino acid type. A variety of search options are available (accession numbers, de-

scription of the protein, etc.) and searches based upon a repeat pattern (for example, poly-A followed by poly-Q) within a single RCP can be performed for multirepeat-containing proteins. A graphical display of the results is presented and all proteins are linked to their National Centre for Biotechnology Information (NCBI) record and an OMIM record, if applicable.

Within GENPEPT (2,677,049 proteins) 1.4% of proteins are RCPs; a total of 54,566 homopeptide repeats could be identified in 37,355 RCPs (Table 1; Fig. 1A). RCPs from environmental sequences (Venter et al. 2004) and viral sequences are listed in Table 1; however, these have not been further considered in this present study.

Several general trends are apparent across all the data. The vast majority (87%) of all RCPs are from eukaryotes; prokaryote RCPs are rare (4%) (Table 1). This is in agreement with previous studies (Karlin and Burge 1996; Marcotte et al. 1999; Huntley and Golding 2000). Within GENPEPT, poly-Q repeats are the most common (16%), whereas poly-W repeats are extremely rare (only three poly-W RCPs could be identified) (Fig. 1A). In eukaryotes, poly-Q, poly-N, poly-A, poly-S, and poly-G are the most common repeat types. In prokaryotes, poly-S, poly-G, poly-A, and poly-P are most common; however, poly-Q and poly-N repeats are relatively rare. In both eukaryotes and prokaryotes, Poly-I, Poly-M, Poly-W, Poly-C, and Poly-Y repeats are either absent or very rare.

When classified according to their physicochemical properties and normalized for the overall frequency of single amino acids within GENPEPT, there is an overrepresentation of polar repeats in comparison to hydrophobic repeats and of acidic repeats in comparison to basic repeats (Fig. 1B). These data are in agreement with a previous study that suggested that long stretches of hydrophobic residues possess greatly enhanced toxicity in comparison to similar stretches of hydrophilic residues (Dorsman et al. 2002; Oma et al. 2004) and are thus selected against.

Table 1. Number of homopeptide repeats and RCPs in GENPEPT, Eukaryotes, and Prokaryotes

	GENPEPT		Eukaryote		Prokaryote		Other (viruses/environmental sequences)	
	Repeats	Proteins	Repeats	Proteins	Repeats	Proteins	Repeats	Proteins
Alanine	6132	5045	5465	4425	251	250	416	370
Valine	149	117	94	83	9	9	46	25
Leucine	1638	1602	1446	1426	70	70	122	106
Isoleucine	57	56	34	33	3	3	20	20
Proline	4837	3931	4157	3333	217	184	463	414
Methionine	27	22	19	18	0	0	8	4
Phenylalanine	196	186	175	172	1	1	20	13
Tryptophan	3	3	3	3	0	0	0	0
Glycine	5981	5020	5002	4168	310	281	669	571
Serine	6383	5463	5424	4742	378	258	581	463
Threonine	2997	2415	2492	1984	63	59	442	372
Cystine	64	52	38	38	0	0	26	14
Asparagine	7126	3731	6962	3597	31	29	133	105
Glutamine	8334	5699	8022	5464	52	51	260	184
Tyrosine	56	51	39	38	4	4	13	9
Aspartic Acid	1835	1707	1554	1451	34	34	247	222
Glutamic Acid	4779	4302	4334	3912	67	61	378	329
Lysine	2081	1926	1920	1774	25	25	136	127
Arginine	751	714	462	443	60	57	229	214
Histidine	1140	1061	1049	971	32	32	59	58
Total	54,566	37,355	48,691	32,628	1607	1388	4268	3339

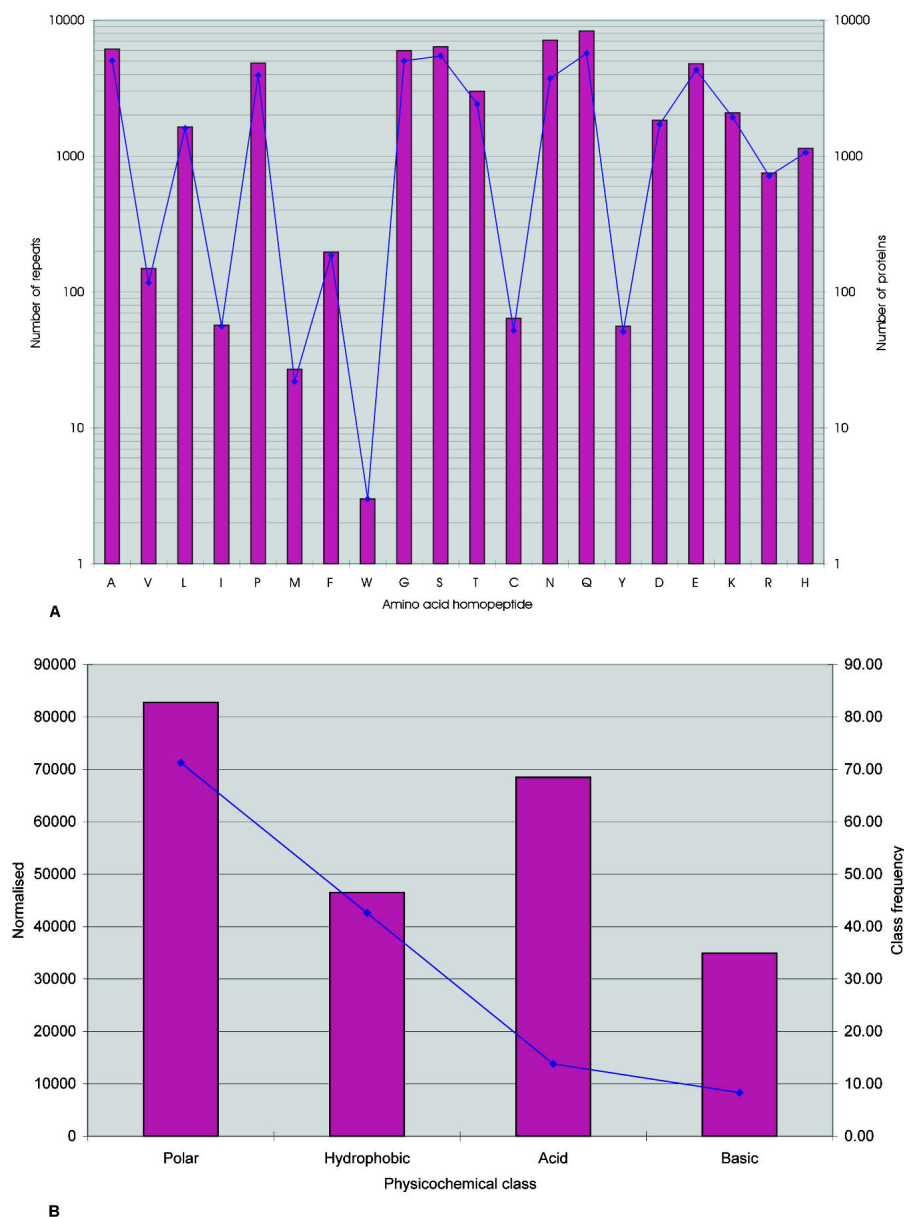


Figure 1. Distribution of RCPs in GENPEPT. (A) The distribution of the RCPs in GENPEPT and the total number of repeats in GENPEPT. The bars represent the total number of repeats and the solid diamonds the number of RCPs. (B) Distribution of the repeats based on physicochemical class (polar, hydrophobic, acidic, and basic). Red bars represent the number of repeats for the amino acid class normalized for the amino acid frequencies in GENPEPT for that amino acid class (i.e., number of repeats in class X/[amino acid frequency for class X, not including the RCPs]). The solid blue diamonds represent frequency of the amino acid class in the RCPs.

Homopeptide length

Figure 2 shows a graph of homopeptide repeat length against frequency for each amino acid type. Several general trends are apparent. Most hydrophobic or rare repeats (poly-I, poly-F, poly-Y, poly-V, poly-L, poly-M, poly-R, and poly-W) tend to be short (<15 amino acids in length). In contrast, more extensive repeats of amino acids such as poly-Q, poly-N, poly-T, poly-S, and poly-E are common. Certain very long (>50 amino acids) repeats can be identified; however, these are very rare (148 repeats).

Proteins with more than one homopeptide repeat

Within GENPEPT, 23% of all RCPs contain more than one repeat tract (Table 1). In eukaryotes, 24% of RCPs contain multiple repeat tracts and only 9% of proteins in prokaryotes (two archaeal and 113 bacterial proteins) are multirepeat-containing proteins.

The most common pattern in GENPEPT after a single amino acid repeat is the doublet GG (752) followed by QQ (736), PP (596), SS (505), and NN (485). The propensity of one repeat type to occur with another in the same protein was investigated. Repeat pairs were tallied according to the number of related sequence families in which they were found; Table 2 shows the frequency with which a repeat of one type occurs with another. Strikingly, for all repeats except poly-L, poly-R, and poly-V, the strongest association was with either poly-N or poly-Q tracts (excluding self-self pairs).

Distribution of repeats within eukaryote organisms

Figure 3, A and B, shows the distributions of RCPs in eukaryotes whose genomes are either complete or near completion. These data highlight several interesting anomalies. *Drosophila melanogaster* possesses an overabundance of poly-Q RCPs, >3.5-fold more than that of *Homo sapiens* and sixfold more than another insect, the mosquito *Anopheles gambiae* (Fig. 3B). In contrast, poly-Q repeats are extremely rare in *Plasmodium falciparum*; this organism instead possesses an overabundance of poly-N RCPs. Analysis of other complete eukaryote genomes revealed that poly-Q repeats are absent in *Encephalitozoon cuniculi* (an intracellular parasite). Another striking difference is in the distribution of poly-N RCPs. Nonvertebrate organisms all contain asparagine RCPs, whereas poly-N tracts are either absent or extremely rare in vertebrates (Fig. 3A).

The human genome contains 233 poly-Q RCPs, but only eight poly-N RCPs, all of which are 8-residue repeats in the N terminus of the insulin receptor substrate 2. The genome of *Mus musculus* contains 170 poly-Q RCPs and only 13 poly-N RCPs (seven from thioredoxin interacting factor, one in the insulin receptor substrate 2, one in a transcription factor, and four in unknown proteins). The genomes of *Gallus gallus* and *Xenopus laevis* do not contain asparagine RCPs.

In order to include an avian representative in our analysis, we also examined the distribution of RCPs in the chicken. These data reveal an apparent paucity of RCPs in *G. gallus* as compared

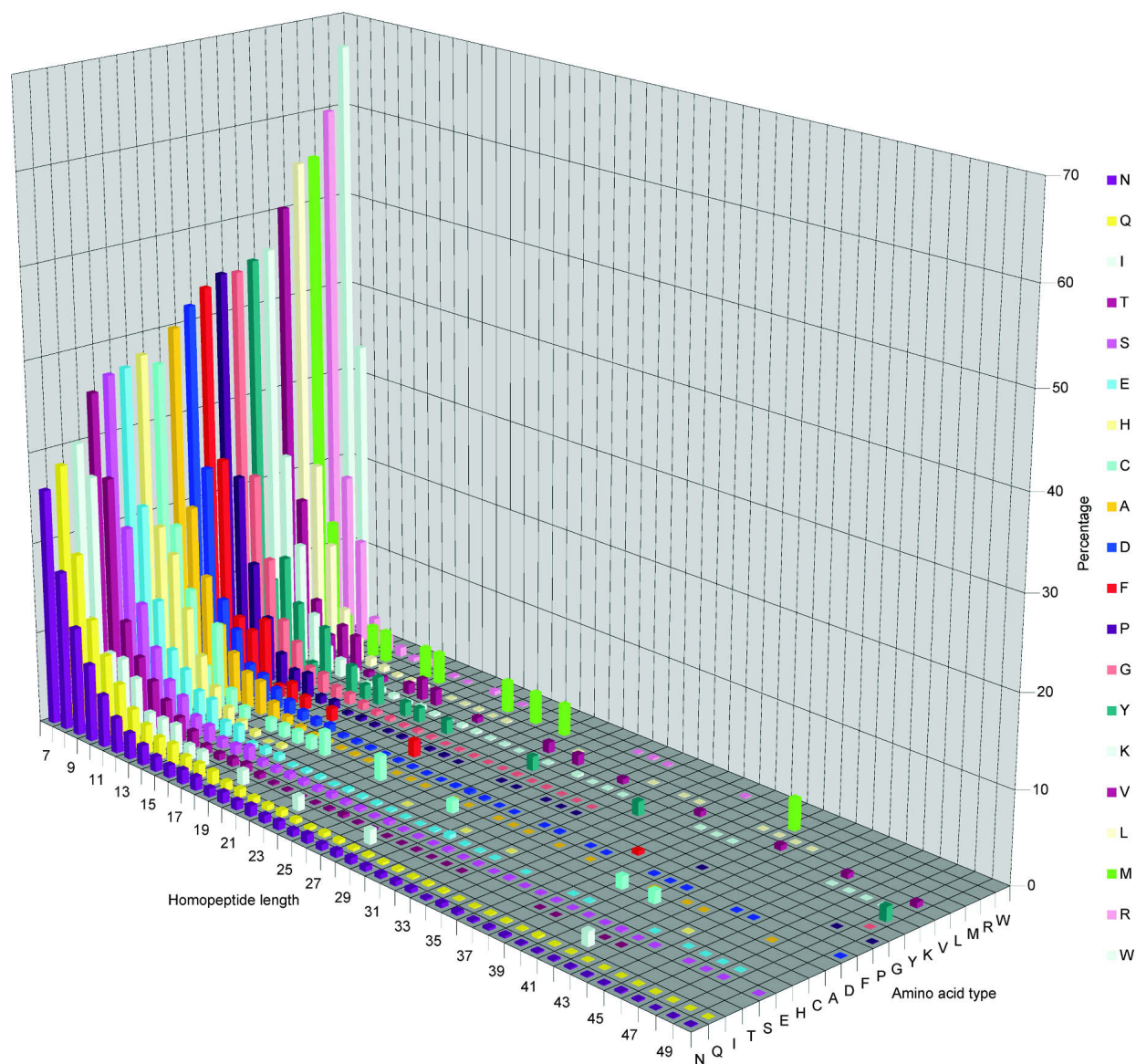


Figure 2. Length of homopeptide repeats. Three-dimensional plot showing repeat length (x-axis) versus amino acid type (y-axis; also highlighted in key) versus percentage of each repeat class of a particular length limited to those repeats <51 amino acids in length. A blank square indicates that no repeat of that length and type exists in GENPEPT. There are repeats >50 amino acids in length; however, these are infrequent, with lengths up to 410 amino acids and a sporadic distribution.

with *H. sapiens* and *M. musculus* (Fig. 3A); however, we cannot exclude the possibility that this observation is a result of the preliminary nature of the available genomic data. Finally, we note that repeats are completely absent in the nucleomorph of *Guillardia theta* (Chromophyte algae).

Evolution of RCPs

In order to investigate the evolutionary context of RCPs, prokaryote RCPs were clustered into 1435 families (www.genome.org). Of these families, the majority (1056) are “orphan” RCPs (i.e., a single member). In order to search for eukaryote homologs of RCPs, PSI-BLAST searches of GENPEPT using probes from all clusters with more than five members (47) and eight randomly chosen clusters with more than five members (a total of 55 clusters)

were performed (Table 3). The largest cluster considered contains 117 members from the xylanase family. Eukaryote putative homologs of this family were identified, and these putative homologs did not contain the repeat tract. Three other large families, the Ribonuclease E (25 members), membrane carboxypeptidase (25 members), and the β -propeller domain of methanol dehydrogenase (23 members), were also identified. Eukaryote homologs of the Ribonuclease family did not contain repeat tracts, and no convincing eukaryote putative homolog of methanol dehydrogenase could be identified. These results are summarized in Table 3. We were able to detect 20 prokaryote RCP families that extended into eukaryotes. Of these, only three families contained analogous repeat regions or an “interrupted” single amino acid-rich repeat tract in both prokaryote and eukaryote members, the heat-shock protein DnaJ (a molecular chaperone;

Table 2. Frequency of repeat pairing

Amino acid (no. of proteins)	%	Self	Decreasing frequency (% , first column) of pairs (<5% not shown)															
			→															
A (1117)	18.7	A	22.4	Q	15.4	G	12.6	S	7.8	P	5.0	N						
D (511)	15.3	D	20.9	N	12.0	E	10.2	Q	8.0	T	7.6	S	7.6	G	6.5	A	5.5	T
E (681)	28.9	E	9.8	Q	9.6	N	9.0	D	8.1	S	6.3	P	6.0	A	5.6	K		
F (22)	13.6	F	22.7	N	13.6	K	9.1	D	9.1	L	9.1	P						
G (1126)	28.0	G	16.1	Q	15.2	A	11.3	S	7.6	P	5.8	N						
H (309)	8.9	H	21.5	Q	14.2	A	12.4	S	12.3	N	10.4	G	7.2	T	5.2	P		
K (382)	23.8	K	36.0	N	10.0	E	8.5	S	6.0	D								
L (96)	11.5	L	16.7	A	14.6	E	13.5	Q	9.4	P	7.3	G	6.2	S				
N (2098)	37.2	N	15.8	Q	12.6	T	9.9	S	6.6	K	5.1	D						
P (718)	28.3	P	13.0	Q	12.1	A	11.9	G	10.0	S	6.0	E						
Q (2172)	30.3	Q	15.2	N	11.5	A	11.0	S	8.8	T	8.8	G						
R (126)	15.1	R	14.3	G	11.1	P	11.1	E	10.3	S	8.3	A	7.1	T	5.2	D		
S (1460)	22.1	S	16.3	Q	14.3	N	10.8	T	9.7	A	8.7	G						
T (1064)	20.3	T	24.8	N	17.9	Q	14.8	S										
V (21)	28.6	V	19.0	S	9.5	A	9.5	P	9.5	R								

HSP40) (Fig. 4), and the two ribosomal proteins L10 and L12 (Figs. 5 and 6).

Functional groups in *H. sapiens*, *D. melanogaster*, *C. elegans*, and prokaryote RCPs

We used the OMIM database and related resources to functionally group human RCPs. (Fig 7A). Sixty percent of the human RCPs have an OMIM record, and 120 diseases are associated with these records (Supplemental Table 1). In addition, all *D. melanogaster* RCPs were mapped onto FlyBase (FlyBase Consortium 2003). These were functionally grouped using Gene Ontology (GO) (Fig. 7B; Supplemental Table 2). These data reveal that the majority (85%) of *D. melanogaster* RCPs are intracellular proteins. Of the remainder, 13% are predicted to be membrane associated, and only 2% are predicted to be extracellular.

Interestingly, clear functional trends are apparent throughout the data set, the majority of both human and fruit-fly RCPs performing roles in transcription/translation and signaling processes. Enzymes, transport proteins, adhesion proteins, and structural proteins also commonly contain homopeptide repeats. We performed a similar analysis of *C. elegans* RCPs using Wormpep, and observed similar trends (Fig. 7C). Finally, we functionally annotated, where possible, prokaryote RCPs (Fig. 7D).

Discrete domains within RCPs

In order to investigate the functional context of homopeptide repeats within RCPs, we performed a survey of protein domains associated with repeat tracts, using the OMIM data set. Ninety-three percent of RCPs listed in OMIM contained a domain listed in pfam (Bateman et al. 2004) and/or SMART (Letunic et al. 2004). Of these, 43% contained an N-terminal repeat in relation to a characterized domain, 10% contained a repeat between domains, and 30% contained a C-terminal repeat in relation to a characterized domain. Also, 16% of RCPs contained a repeat within a characterized domain. Many RCPs are transcription factors or involved in transcriptional processes; hence, many repeat tracts are associated with a variety of nucleic acid-binding domains. However, no obvious pattern of association between repeat type and precise domain type could be detected.

Discussion

Our data reveal that RCPs are far more abundant in eukaryotes than in prokaryotes. In addition, based upon analysis of the *D. melanogaster* data set, the majority of eukaryote RCPs are predicted to be intracellular proteins. Furthermore, in agreement with the studies of Marcotte et al. (1999), there is an overrepresentation of polar repeats in comparison to hydrophobic repeats, and most hydrophobic repeats are relatively short in comparison to their polar counterparts. These data are consistent with previous studies that suggest hydrophobic RCPs aggregate and form toxic fibrils (Dorsman et al. 2002; Oma et al. 2004).

Glycine, serine, and proline repeats are common in both prokaryotes and eukaryotes; however, common eukaryote repeats such as glutamine, asparagines, and glutamic acid are relatively rare in prokaryote organisms; of 29 asparagine RCPs in prokaryotes, 11 are orphans, 12 do not have eukaryotic homologs, and six have putative eukaryote homologs. However, these homologs do not contain the repeat or an equivalent amino acid-rich region. Of the 51 glutamine RCPs in prokaryotes, 23 are orphans, 21 do not have eukaryotic homologs, and seven have putative eukaryote homologs, but again, these homologs do not contain a repeat or an equivalent amino acid-rich region.

Certain discrepancies are clearly apparent when considering repeat distribution within eukaryotes. For example, glutamine RCPs (the most common eukaryote repeat) are rare or absent in *P. falciparum*, *E. cuniculi*, and *G. theta*. Furthermore, our data reveal that while asparagine repeats are common in nonvertebrates such as insects and nematodes, such repeats are extremely rare in vertebrates. Kreil and Kreil (2000) previously observed the rarity of asparagine repeats in mammals. This is surprising, since glutamine and asparagine are chemically and structurally similar. We could not identify any function specific to poly-N repeats; indeed, an analysis of all *D. melanogaster* poly-N RCPs reveals that, like other RCPs (see below), a large proportion (>60%) are transcription factors or proteins involved in transcriptional processes (Table 2; Supplemental data). Interestingly, in most multirepeat-containing RCPs, glutamine or asparagine is the most likely associated partner polyamino acid (Table 2). The functional significance of these data remains to be understood.

Very rarely, repeats are conserved across entire protein families, and only three families (DnaJ and the ribosomal proteins L10 and L12) could be identified with repeat regions in eukaryote

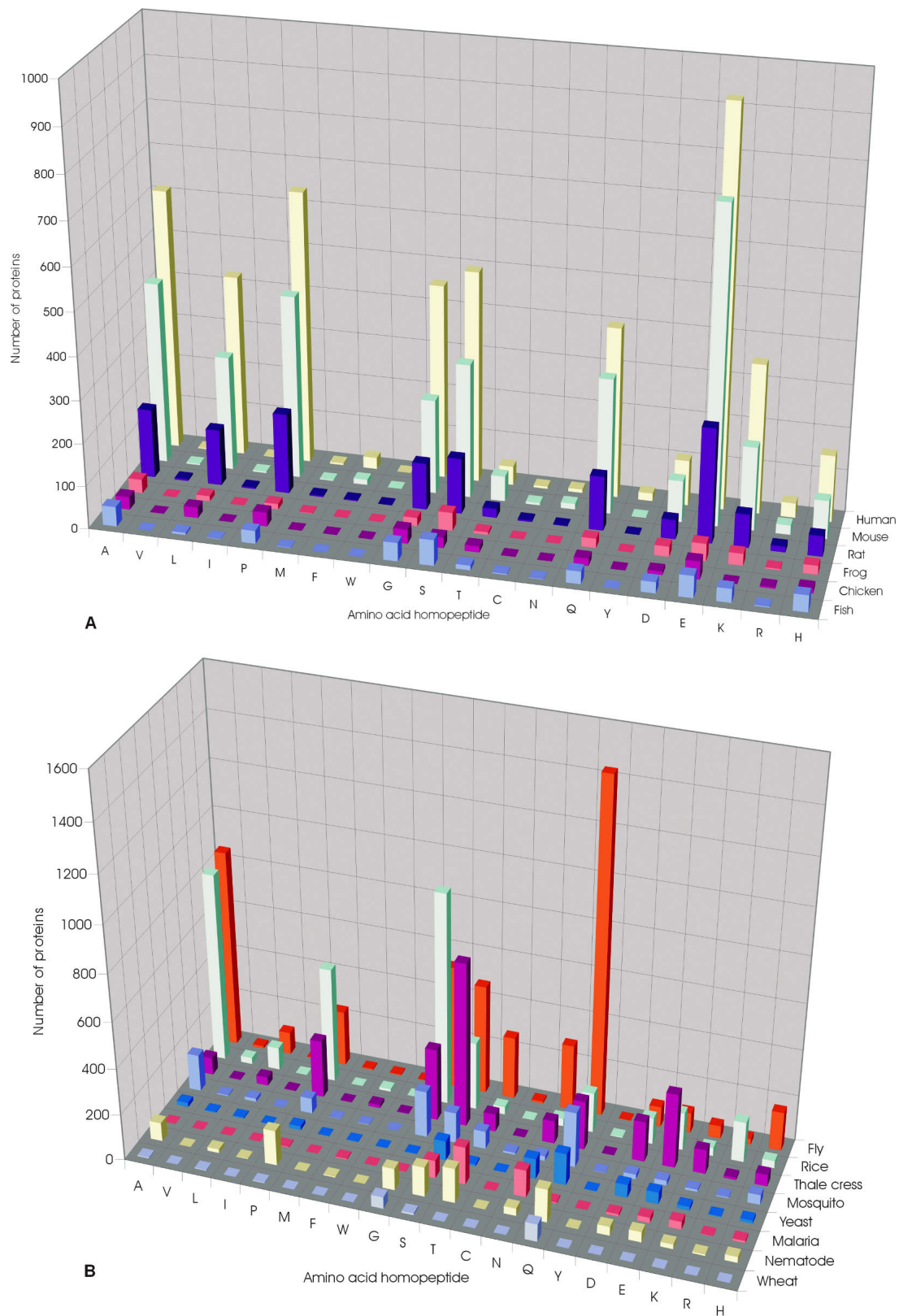


Figure 3. Distribution of RCPs in eukaryotes. (A) Distribution of the RCPs in vertebrate species, *Homo sapiens* (human), *Mus musculus* (mouse), *Rattus norvegicus* (rat), *Xenopus laevis* (frog), *Danio rerio* (fish), and *Gallus gallus* (chicken). (B) Distribution of RCPs in nonvertebrate species, *Drosophila melanogaster* (fly), *Oryza sativa* (rice), *Arabidopsis thaliana* (thale cress), *Saccharomyces cerevisiae* (yeast), *Plasmodium falciparum* (malaria), *Anopheles gambiae* str. PEST (mosquito), *Caenorhabditis elegans* (nematode), and *Triticum aestivum* (wheat). The genomes of the chosen species have been completed or are near completion.

Table 3. RCP's representatives from the prokaryota clusters with more than four members and the seven randomly chosen clusters with less than five members

gi accession	Species	Repeat type (length)		Number of members in the cluster	Protein/function
26987138	<i>Pseudomonas putida</i>	G(7)		5	Hypothetical protein Unknown
2983200	<i>Aquifex aeolicus</i>	A(7) & R(9)	*	25	Ribonuclease E
33602211	<i>Bordetella bronchiseptica</i>	P(11)		8	Proline-rich inner membrane protein Unknown
33331898	<i>Mycoplasma hyopneumoniae</i>	Q(13)		5	P216 surface protein Unknown
32473910	<i>Rhodopirellula baltica</i>	G(7)	*	6	Polyribonucleotide nucleotidyltransferase RNA/DNA binding
32474384	<i>R. baltica</i>	R(7)		10	Probable serine/threonine-protein kinase Enzyme
27375632	<i>Bradyrhizobium japonicum</i>	P(7) & P(8) & P(7)		11	Hypothetical protein blr0521 Unknown
33864553	<i>Synechococcus</i> sp. WH 8102	G(7)		9	RNA-binding region RNP-1 RNA binding
27378819	<i>B. japonicum</i>	S(7)		7	Efflux protein Transport
29830831	<i>Streptomyces avermitilis</i>	G(7)		6	Putative single-stranded DNA-binding protein DNA binding
34499217	<i>Chromobacterium violaceum</i>	E(7)		8	RNA polymerase σ factor RpoD Transcription
32041487	<i>Pseudomonas aeruginosa</i>	S(17)		23	β -propeller domains of methanol dehydrogenase type Enzyme
27367908	<i>Vibrio vulnificus</i>	Q(89)		5	TPR repeat containing protein Unknown
23015533	<i>Magnetospirillum magnetotacticum</i>	G(7)	*+	9	DnaJ-class molecular chaperone with C-terminal Zn finger domain
23129192	<i>Nostoc punctiforme</i>	G(8)		13	Periplasmic protein TonB, links inner and outer membranes Structural
20094262	<i>Methanopyrus kandleri</i>	E(7) & E(10)	*+	9	Ribosomal protein L10 Translation
16124685	<i>Caulobacter crescentus</i>	A(7)		6	Methyl-accepting chemotaxis protein McpA Signaling
23027882	<i>Microbulbifer degradans</i>	G(7)		5	Cation/multidrug efflux pump Transport
23472931	<i>Pseudomonas syringae</i>	A(7) & A(7)	*	5	Tfp pilus assembly protein FimV Structural
21226168	<i>Methanosarcina mazei</i>	G(7)		13	Hypothetical protein MM0066 Unknown
4887174	<i>Neisseria meningitidis</i>	T(7)		5	Porin
22991761	<i>Enterococcus faecium</i>	S(9)	*	19	Muramidase (flagellum-specific) Enzyme
15839665	<i>Mycobacterium tuberculosis</i>	A(8)		10	PPE family protein Unknown
399792	<i>Erwinia chrysanthemi</i>	S(7)		9	PECTIC ENZYMES SECRETION PROTEIN OUTD Transport
22968935	<i>Rhodospirillum rubrum</i>	P(7)	*	10	Mg-chelatase subunit ChII Enzyme
22125459	<i>Yersinia pestis</i>	G(7) & S(8)		13	Hypothetical Unknown
580740	<i>Azotobacter vinelandii</i>	A(8)		6	Dihydrolipoyltransacetylase Enzyme
8163676	<i>Streptococcus pneumoniae</i>	S(7)		7	Hypothetical protein Rv3088 U
7481905	<i>Streptomyces</i> sp.	A(7)		7	Polyketide synthase Enzyme
14602144	<i>Aeropyrum pernix</i>	G(7)		10	Hypothetical protein APE2556
23050783	<i>Methanosarcina barkeri</i>	E(14)		10	CO dehydrogenase/acetyl-CoA synthase β subunit Enzyme
23051653	<i>M. barkeri</i>	E(7)	*+	5	Ribosomal protein L12E Translation
22968174	<i>R. rubrum</i>	G(9)	*	8	Membrane protease subunits, stomatin/prohibitin homologs Enzyme
15604160	<i>Rickettsia prowazekii</i> str. Madrid E	P(8)		7	VIRB10 protein (virB10) U
17549401	<i>Ralstonia solanacearum</i>	G(7)	*	117	Xylanase Enzyme
23465971	<i>Bifidobacterium longum</i>	A(7)		7	Cell division protein FtsK-DNA translocase DNA binding
15806123	<i>Deinococcus radiodurans</i>	P(8)		7	ABC transporter, ATP-binding protein, EF-3 family Transport
20149845	<i>Archaeoglobus fulgidus</i>	A(32)		7	Chain B, Reverse Gyrase DNA binding
29840119	<i>Chlamydomonas reinhardtii</i>	S(7)	*	16	ATP-dependent Clp protease, ATP-binding subunit ClpC Enzyme
11095653	<i>Treponema pallidum</i>	A(8)		15	TprK U Unknown
7442918	<i>Mycoplasma pneumoniae</i>	S(7)		8	Adhesin P1, group 2 variant precursor Adhesion
15639317	<i>Treponema pallidum</i> subsp. <i>pallidum</i> str. Nichols	S(11)	*	7	Outer membrane protein Unknown
7443057	<i>Salmonella typhimurium</i>	P(9)		8	Virulence-associated protein mkaA Unknown
22979423	<i>Ralstonia metallidurans</i>	H(7)	*	7	Putative GTPases Enzyme
23037072	<i>Oenococcus oeni</i>	S(8)		25	Membrane carboxypeptidase Enzyme
21231078	<i>Xanthomonas campestris</i> pv. <i>campestris</i> str. ATCC 33913	G(13)	*	6	Unknown acidic aa rich protein Unknown
19745410	<i>Streptococcus pyogenes</i>	G(9)	*	4	Hypothetical protein spyM18_0265 Unknown
23108546	<i>Novosphingobium aromaticivorans</i>	P(12)	*	4	Translation initiation factor 2 Translation
15596494	<i>Pseudomonas aeruginosa</i>	H(14)	*	2	Probable metal transporter Transport
548937	<i>Clostridium thermocellum</i>	G(19)	*	2	Cell surface glycoprotein 1 precursor Adhesion
22978658	<i>Ralstonia metallidurans</i>	G(7)	*	2	Translation initiation factor 2 Translation
15606084	<i>Aquifex aeolicus</i>	H(7)	*	2	Hydrogenase expression/formation protein B Enzyme
15922873	<i>Sulfolobus tokodaii</i>	T(17)	*	2	Hypothetical protein ST2539 Transport

*Putative eukaryote homologs, (+) the putative eukaryote homologs contain the repeat or an equivalent amino acid rich region.

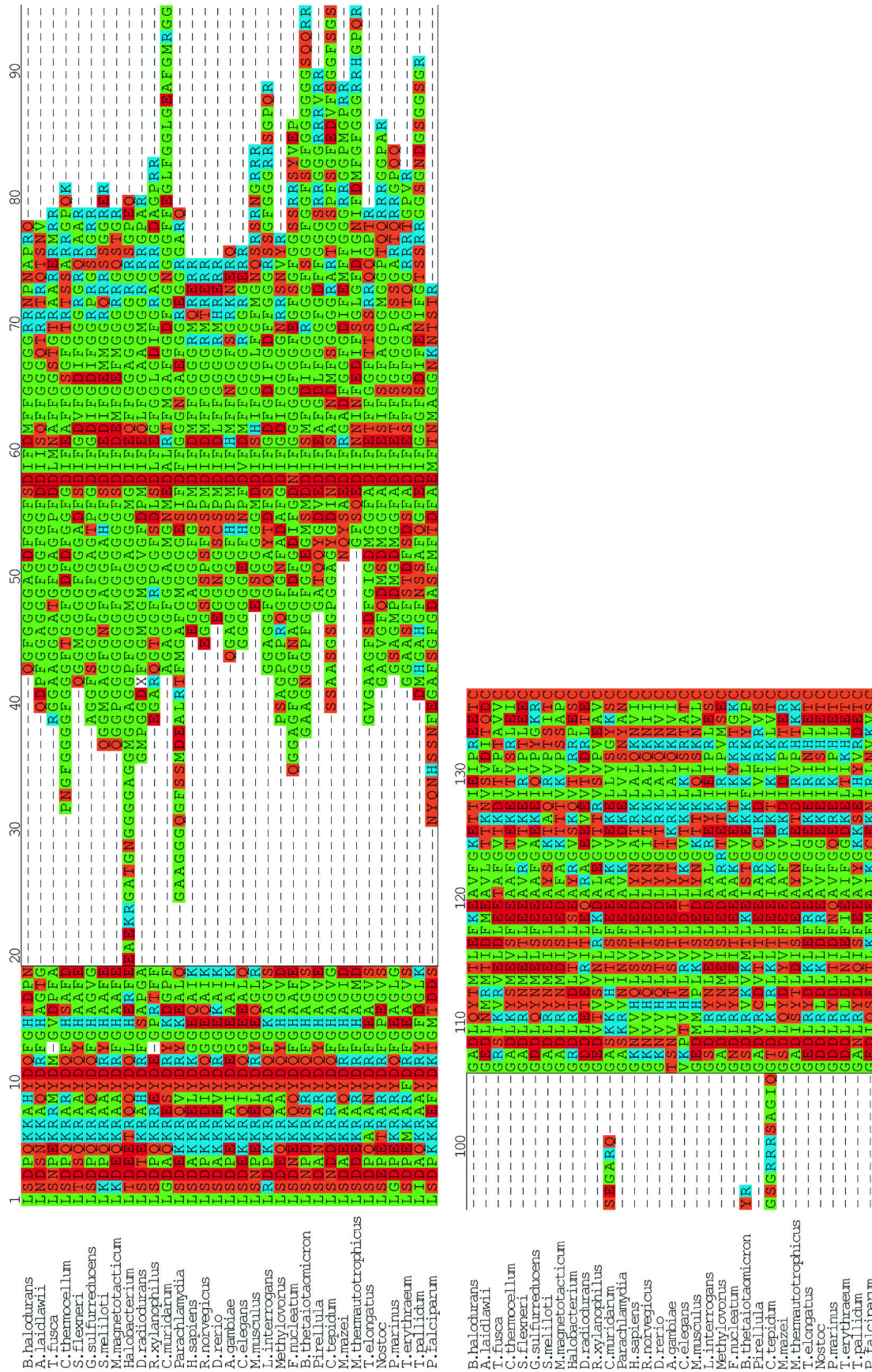


Figure 4. Multiple sequence alignment of DnaJ. A multiple alignment of the glycine repeat in DnaJ (Hsp40). From prokarya: *Bacillus halodurans*, *Clostridium thermocellum*, *Sinorhizobium mellioli*, *Shigella flexneri*, *Magnetospirillum magnetotacticum*, *Geobacter sulfurreducens* PCA, *Leptospira interrogans* serovar lai str. 56601, *Thermosynechococcus elongatus* BP-1, *Nostoc* sp. PCC 7120, *Prochlorococcus marinus*, *Trichodesmium erythraeum* IMS101, *Methanosarcina mazei* Goel, *Treponema pallidum*, *Rubrobacter xylanophilus* DSM 9941, *Fusobacterium nucleatum* subsp. *polymorphum*, *Methanobacterium thermoautotrophicum* str. Delta H, *Pirellula* sp., *Chlamydia muridarum*, *Parachlamydia* sp. UWEZ5, *Bacteroides thetaiotaomicron* VPI-5482, *Acholeplasma laidlawii*, *Chlorobium tepidum* TLS, *Thermobifida fusca*, *Halobacterium* sp. NRC-1, and *Deinococcus radiodurans*. From eukarya: *H. sapiens*, *R. norvegicus*, *M. musculus*, *D. rerio*, *A. gambiae* str. PEST, *C. elegans*, and *P. falciparum*. The boxed region from positions 20–106 highlights the glycine/phenylalanine-rich region. The sequences were manually positioned with respect to the first instance of the highly conserved motif DxF (boxed, position 58–60). The regions to the left and right of the boxed region were aligned with T-COFFEE (Notredame et al. 2000) and the final figure was generated with ALSCRIPT (Barton 1993).

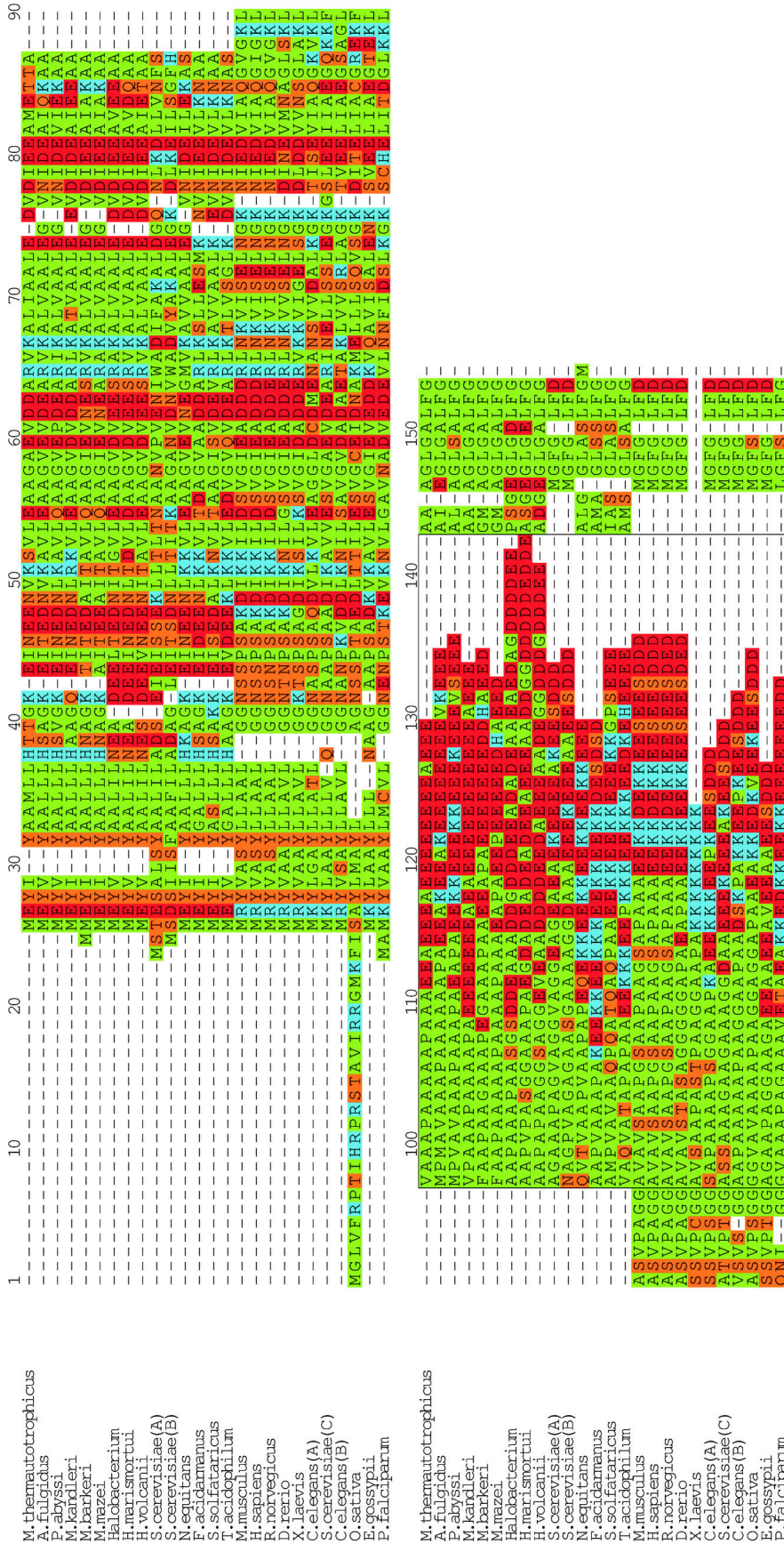


Figure 5. Multiple sequence alignment of the ribosomal protein L12. A multiple sequence alignment of the glutamic acid repeat in L12 from the prokaryotic species *M. thermautotrophicus* str. Delta H, *Archaeoglobus fulgidus* DSM 4304, *Pyrococcus abyssii*, *M. kandleri* AV19, *Methanosarcina barkeri*, *Methanosarcina mazeri* Goel, *Halo bacterium* sp. NRC-1, *Haloarcula marismortui*, *Haloferax volcanii*, *Nanoarchaeum equitans* Kin4-M, *Ferroplasma acidarmanus*, *Sulfolobus solfataricus*, *Thermoplasma acidophilum*, and the eukaryotic species *M. musculus*, *H. sapiens*, *R. norvegicus*, *D. rerio*, *C. elegans* (A) gi 25141400, (B) gi 17543850, *S. cerevisiae* (A) gi 171813, (B) gi 171815, (C) gi 236358, *O. sativa*, *Eremothecium gossypii*, *P. falciparum*. The boxed region from positions 98–144 highlights the two amino acid-rich regions, the N-terminal alanine-rich region and the C-terminal glutamic acid-rich region. Since no obvious alignment could be built of this region, the sequences were flushed left. The regions to the left and right of the boxed region was aligned with T-COFFEE (Notredame et al. 2000) and then manually adjusted. The final figure was generated with ALSCRIPT (Barton 1993).

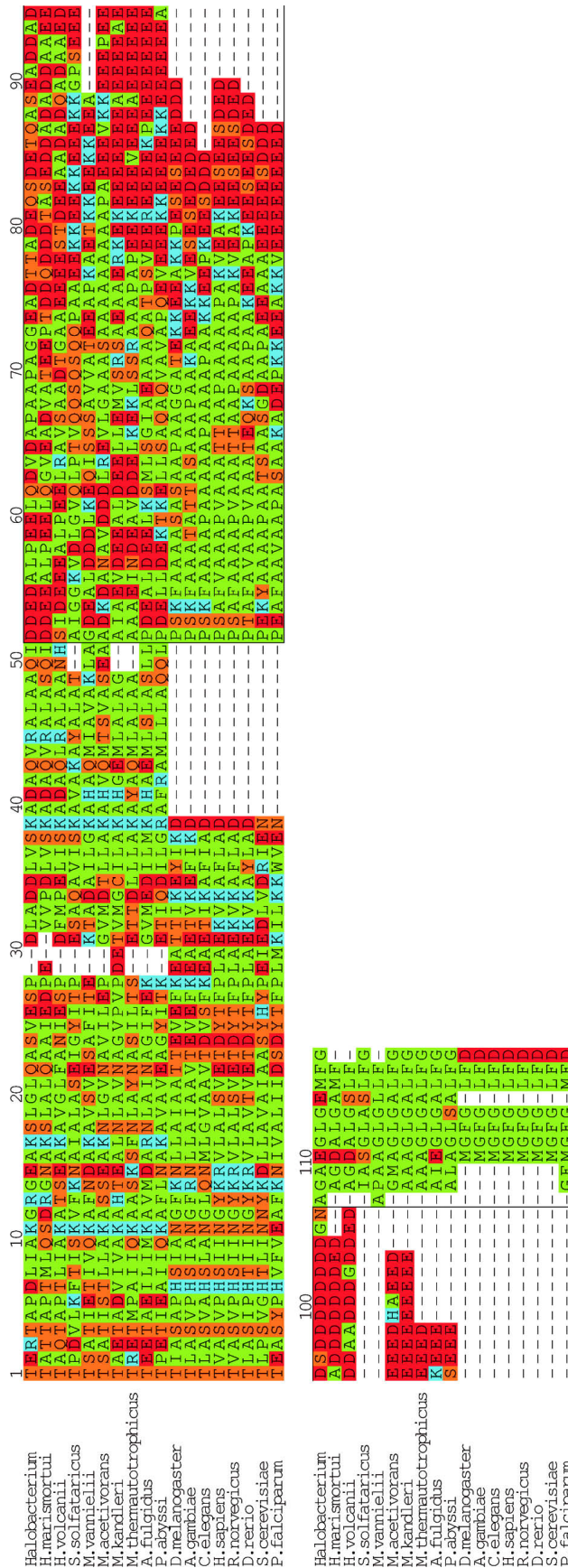


Figure 6. Multiple sequence alignment of the ribosomal protein L10. A multiple alignment of the glutamic acid-rich region in L10 from the following prokaryotic species: *Halobacterium* sp. NRC-1, *Halorcula marismortui*, *Haliferax volcanii*, *Sulfolobus solfataricus*, *M. vannielii*, *Methanosarcina acetivorans* str. C2A, *M. kandleri* AV19, *M. thermautotrophicus* str. Delta H, *Archaeoglobus fulgidus* DSM 4304, *P. abyssii*, and the eukaryotic species *H. sapiens*, *R. norvegicus*, *D. riero*, *A. gambiae* str. PEST, *C. elegans*, *P. falciparum*. The boxed region from positions 52–117 highlights the acidic tail. Since no obvious alignment could be built of this region, the sequences were flushed left. The regions to the left and right of the boxed region was aligned with T-COFFEE (Notredame et al. 2000) and then manually adjusted. The final figure was generated with ALSCRIPT (Barton 1993).

Homopeptide repeats in whole genomes

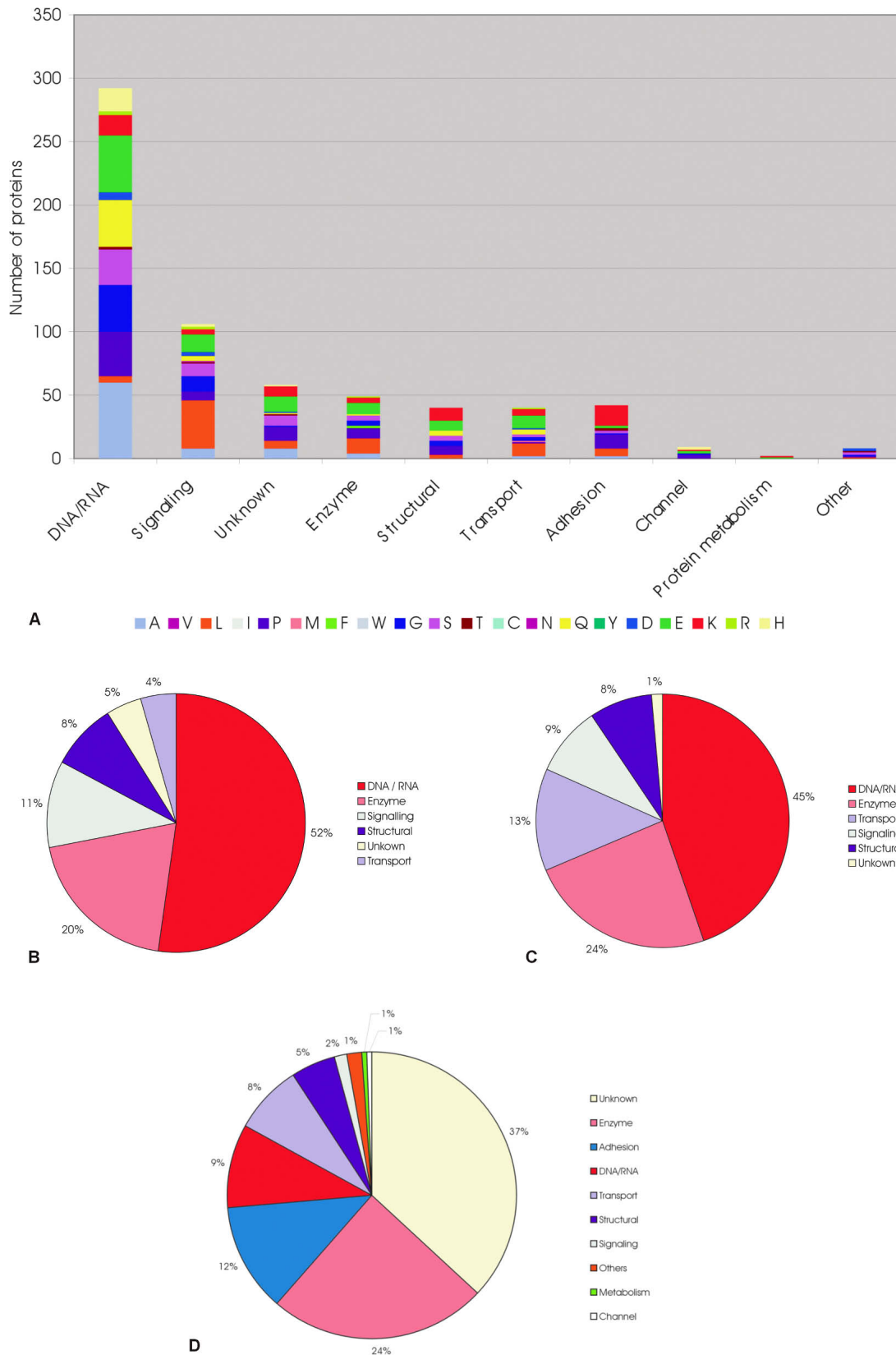


Figure 7. Function of RCPs. (A) Bar graph showing the function of human RCPs, based upon OMIM. Amino acid types are represented by different colors (see key). On the x-axis, the functional classes DNA/RNA (i.e., Transcription/chromatin binding/DNA binding/RNA binding/translation), signaling, unknown, Enzyme, structural, transport protein, adhesion, channel, metabolism, and other are shown. (B) Pie chart showing the function of *D. melanogaster* RCPs, (C) pie chart showing the function of *C. elegans* RCPs, (D) pie chart showing the function of prokaryote RCPs.

and prokaryote putative homologs. A sequence alignment of the DnaJ family (Fig. 4) reveals that an extensive glycine repeat is present in most putative homologs. However, in the majority of these, the repeat is interrupted (typically with an alanine or phenylalanine residue), and this region is contracted in many eukaryotic counterparts. DnaJ functions in complex with at least two other proteins (DnaK and GrpE) to control processes such as protein folding, apoptosis, and the degradation of misfolded proteins (Gragerov et al. 1992; Hendrick et al. 1993; Craig et al. 1994; Gotoh et al. 2004). Deletion of the glycine/phenylalanine-rich motif decreases the binding affinity of the substrate σ^{32} to DnaK (Wall et al. 1995), and it has been proposed that this region may act as a flexible linker to allow DnaJ/K complex formation and subsequent activation of DnaK (Wall et al. 1995).

Both archaeal and eukaryotic L10 and L12 proteins contain a C-terminal region that comprises an alanine-rich region, termed the hinge, followed by a glutamic acid-rich repeat (Figs. 5, 6). L10 and L12 form part of a complex in the large ribosomal subunit termed the stalk protuberance. The hinge, as well as the acidic region in both proteins, are postulated to function as flexible regions that mediate a variety of protein–protein interactions and are important for processes such as elongation (Remacha et al. 1995; Uchiumi et al. 2002a,b; Gonzalo and Reboud 2003). The acidic region in L10 and L12 is severely truncated in certain eukaryote orthologs; for example, this region is absent in L12e from *X. laevis*. Bacterial L12 and L10 proteins do not contain the acidic tail. The absence of this region has been observed previously (Ramirez et al. 1989), and it is suggested that the evolution of this region arose after the archaea/bacteria split, but prior to the emergence of eukaryotes.

A major question in repeat-related research fields is the role of RCPs and, in particular, the role of the repeat region itself. Evolutionary pressure on repeat regions is likely to include functional requirement, mutability of the underlying nucleotide sequence, and potential toxicity. Our analysis allows us to begin to address these questions from a functional perspective. Assigning function to protein sequences is nontrivial, since many proteins perform overlapping functions (for review, see Whisstock and Lesk 2003). However, using the OMIM database and other annotation resources, we have attempted to classify RCPs under as broad a class as possible. Most eukaryote RCPs are involved in transcription/translation or interact directly with DNA, RNA, or chromatin, irrespective of repeat type (Fig. 7A). Other common classes of RCPs include signaling molecules, structural proteins, transport molecules, and enzymes (Fig. 7). Previous research on smaller eukaryote data sets also note that repeat tracts are over-represented in transcription factors and DNA-binding proteins (Karlin and Burge 1996; Mar Alba et al. 1999; Alba et al. 2002; Alba and Guigo 2004). The results of this study are consistent with these findings and identify other functional classes rich in RCPs.

We performed a functional analysis of RCPs from prokaryotes (Table 3; Fig. 7D). While 38% of these families perform as yet uncharacterized roles, several of the functional themes apparent in the eukaryote data set are also noticeable in prokaryotes. The most common functional class (accounting for 24% of classified molecules) are RCPs that perform roles as enzymes. Transport proteins, structural proteins, and transcription/translation-related RCPs are also common; however, in contrast to eukaryotes, the dramatic bias toward transcription/translation-related processes is not observed. One possible explanation for these data is that bacterial genomes are smaller in relation to eukary-

otes and are not packaged and controlled in such a complex fashion. RCPs involved in signaling-related processes are also relatively rare in prokaryotes in comparison to eukaryotes. Again, we argue that this may be a result of the increased complexity of eukaryote processes; while bacteria utilize intracellular signaling processes to communicate intracellularly and with their environment, these processes are relatively modest in comparison to eukaryote-signaling cascades.

The repeats database provides a basis for understanding the function of RCPs and their associated repeats in all organisms. Strikingly, the majority of RCPs considered are involved in processes that require the assembly or association of large multiprotein and/or nucleic acid complexes (Fig. 7). For example, the ribosome (itself a large protein/RNA complex) requires a large number of additional factors (e.g., elongation factors) to properly function. Processes such as transcription involve the assembly of multiprotein complexes (e.g., RNA polymerase) and the binding of discrete sequences of DNA that may be kilobases apart (Tolhuis et al. 2002); chromatin condensation requires bundling and folding of nucleosome arrays by protein factors such as Nucleoplasmin (for review, see Akey and Luger 2003; Grigoryev 2004); signaling requires the assembly of a large number of proteins into a signalosome or transducosome (e.g., the MAP Kinase complex) (for review, see Burack and Shaw 2000) and ion channels such as the cGMP-gated channels in rod photoreceptors are involved in large protein complexes that link the plasma membrane with the outer segment disk (Korschen et al. 1999). In many of these processes, the involvement of repeats in mediating protein–protein interactions have been noted (for example, the glutamic acid-rich proteins of cGMP-gated channels and the acidic tails of ribosomal proteins) (Remacha et al. 1995; Korschen et al. 1999; Poetsch et al. 2001).

The data presented in this study reveal that the vast majority of the human repeat tracts present in the OMIM data set (83%) are located N-terminal to, as well as C-terminal to, or in-between discrete domains. Alba and Guigo (2004) have also noted that certain repeats (L, A, P, and Q) appear to be preferentially located at the N terminus. Structural studies reveal that most repeat regions do not adopt discrete well-ordered structures, and instead, are often disordered (Huntley and Golding 2002).

The function of RCPs, as well as the interdomain or terminal location of the repeat tract within these molecules and the disordered structure of these repeat sequences, supports the idea that repeats play roles as flexible spacer elements/“tethers” between individual folded domains in molecules that mediate protein–protein or protein–nucleic acid interactions (Karlin and Burge 1996).

Based upon this work, it is suggested that a general function of the majority of repeat sequences is to mediate the assembly of protein complexes, and that RCPs may act as molecular “fishing lines”, mediating interactions either through tethered distant domains, or indeed, through interactions with the repeat itself (e.g., Gerber et al. 1994; Korschen et al. 1999). Such a flexible structure would allow a single protein within a complex to recruit additional factors from the cytoplasmic or nuclear milieu (Fig. 8). A repeat tract would also be able to bridge large distances, such as is required in chromatin packaging or transcription. While many specialized proteins are able to achieve both distance and conformational mobility via an ordered three-dimensional fold (e.g., myosin) (Pollard 2000) or the serpin superfamily of proteinase inhibitors (Whisstock et al. 1998), a flexible unstructured region is a simple way of achieving a similar, albeit conformationally

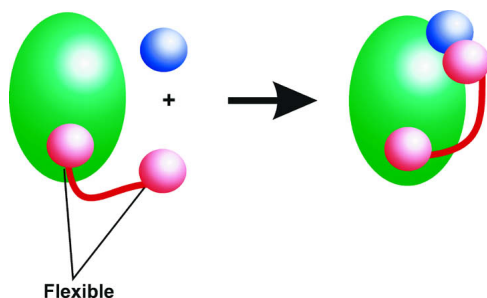


Figure 8. General role of repeats regions. It is suggested that RCPs (red) function from within large multiprotein and/or nucleic acid complexes (green circle). An example is shown where a two-domain protein (pink circles) functions via a flexible repeat to recruit an additional binding partner (blue circle).

uncontrolled, outcome. A repeat does not require complicated molecular machinery to achieve flexibility and, because it is unstructured, appropriately sized repeats would be predicted to interfere minimally with other molecules in a complex.

Several studies have revealed that long stretches of hydrophobic amino acids are more toxic than hydrophilic counterparts (Dorsman et al. 2002; Oma et al. 2004). Thus, in order to bridge large distances, long tracts of polar repeats such as poly-Q, poly-N, and poly-E can be used as suggested by the distribution of repeat length and amino acid type (Fig. 2). The consequence of utilizing single amino acid repeats is that repeat expansion can result in the formation of ordered aggregates or fibrils. Such an event could result in cell death through a variety of mechanisms; the toxic nature of the fibril, loss or gain of function, the destruction of a large essential multiprotein complex, and/or the sequestering of nonspecific factors.

The majority of proteins sequenced to date do not contain repeats. While certain repeats are common throughout entire protein superfamilies (such as the DnaJ family), the data gleaned in this study reveals that repeat proteins are often “orphans.” We suggest that the putative role of repeats is ancient; however, the relatively sporadic distribution of these regions suggests that repeats often evolve to perform in specialized processes unique to a particular organism or set of organisms.

Methods

The March (2004) version of GENPEPT (from <ftp://ftp.ncbi.nih.gov/blast/db>) was used in this study.

Homeopeptide searches

All homeopeptides longer than 4 amino acids were discovered using the regular expression “[AaVvLlIiPpMmFfWwGgSsTtCcNnQqYyDdEeKkRrHh]\1+” across the GENPEPT database. This expression will find all occurrences of an amino acid repeat such as “aa,” “aaa,” and “gggg” (Friedl 2002). A minimum length threshold of 7 amino acids was set. The results from the study can be queried and visualized via the Web site <http://repeats.med.monash.edu.au>.

Evolution of RCPs

All RCPs from prokaryotes were used in an all-against-all BLASTp search with the following parameters: -f T (on), e = 0.001. Single-linkage clustering of the BLASTp results was then performed using

a similar approach to the package GeneRAGE (Enright and Ouzounis 2000). That is, an NxN matrix was created from the BLASTp results and forced into symmetry. Clustering was performed across the matrix.

When considering eukaryote RCPs, only completed or near-complete genomes were used, so as to avoid potential bias due to overrepresentation of commonly studied protein families.

Thus, the following species were considered: *Homo sapiens*, *Mus musculus*, *Rattus norvegicus*, *Gallus gallus*, *Danio rerio*, *Xenopus laevis*, *Drosophila melanogaster*, *Anopheles gambiae*, *Caenorhabditis elegans*, *Saccharomyces cerevisiae*, *Plasmodium falciparum*, *Oryza sativa*, *Triticum aestivum*, and *Arabidopsis thaliana*.

The longest sequence from each of the prokaryote clusters was used as probes to search GENPEPT using PSI-BLAST. The following parameters were used: j = 5, b = 100,000, e = 0.001, and -f T. All sequences with significant expect scores (<0.001) (Park et al. 1998) from the sequenced eukaryote organisms (listed above) were selected. Sequences that were >98% identical were then removed using the nrdb90 algorithm (Holm and Sander 1998). For each eukaryote species, the top 10 putative homologs (with the lowest expect scores) were used for the multiple-sequence alignment. The alignments were initially generated using CLUSTALW (Thompson et al. 1994) with default parameters. Inspection of the alignments enabled identification of eukaryote putative homologs that contained a repeat or an equivalent amino acid-rich tract; these were subject to further analysis. The alignment shown in Figures 4–6 was initially generated with T-Coffee (Notredame et al. 2000), then manually curated.

Analysis of repeat pairs

In order to explore whether certain repeat types have a tendency to appear together within a given protein, the colocalization of repeat pairs was investigated. PSI-BLAST (three iterations, inclusion threshold $E < 0.02$, reporting threshold $E < 1 \times 10^{-6}$) was used to obtain clusters of sequences from a database of multiple repeat-containing proteins, where each pairwise alignment spanned >50% of the length of the smaller sequence. For each unique repeat pair identified in a protein, a score was given as the reciprocal of the number of related proteins; i.e., a protein with the repeats TNNNNNK and related to four others would increment the scores for repeat pairs TN, TK, NN, and NK by 0.25. This was performed for all multiple repeat-containing proteins.

Functional annotation

Existing information within OMIM (<http://www.ncbi.nlm.nih.gov/omim/>), HPRD (<http://www.hprd.org/>), FlyBase (FlyBase Consortium 2003), and Wormpep (http://www.sanger.ac.uk/Projects/C_elegans/WORMBASE/current/wormpep.shtml) was used to annotate the human, *Drosophila*, and nematode data set, respectively. In order to annotate prokaryote sequences, the longest member from each of the 1435 clusters were used as probes in BLASTp searches of GENPEPT (using the default parameters). The results were manually analyzed, and putative general function assigned if a cluster shared significant similarity with a characterized family (only hits with an expect score <0.001 were considered). Literature searches were used to establish general function where required.

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