An archaeal origin for the actin cytoskeleton
Implications for eukaryogenesis

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A hallmark of the eukaryotic cell is the actin cytoskeleton, involved in a wide array of processes ranging from shape determination and phagocytosis to intracellular transport and cytokinesis. Recently, we reported the discovery of an actin-based cytoskeleton also in Archaea. The archaeal actin ortholog, Crenactin, was shown to belong to a conserved operon, Arcade (actin-related cytoskeleton in Archaea involved in shape determination), encoding an additional set of cytoskeleton-associated proteins. Here, we elaborate on the implications of these findings for the evolutionary relation between archaea and eukaryotes, with particular focus on the possibility that eukaryotic actin and actin-related proteins have evolved from an ancestral archaeal actin gene. Archaeal actin could thus have played an important role in cellular processes essential for the origin and early evolution of the eukaryotic lineage. Further exploration of uncharacterized archaeal lineages is necessary to find additional missing pieces in the evolutionary trajectory that ultimately gave rise to present-day organisms.

Discovery of an Actin-Based Cytoskeleton in Archaea

In eukaryotes, actin filaments constitute key components of the cytoskeleton, involved in pivotal processes such as determination and maintenance of cell shape and cellular junctions, cytokinesis, and vesicle-mediated transport such as endocytosis and phagocytosis. Recently, bona fide archaeal actin orthologs were identified in several crenarchaeal genomes, as well as in Candidatus Korarchaeum cryptofilum (Fig. 1). The archaeal actin ortholog, denoted Crenactin, was shown to polymerize into a cytoskeletal structure in the hyperthermophilic crenarchaeon Pyrobaculum calidifontis, and therefore inferred to be involved in cell shape determination. Immunofluorescence microscopy imaging revealed that Crenactin formed helical filaments that traversed the length of the rod-shaped cells. In a cell subpopulation, these filaments had been remodelled into band-like structures, presumably in preparation for cell division, in this respect resembling the bacterial cell-shape-determining protein MreB, which belongs to the same ATPase superfamily as actin and Crenactin. In addition, the phylogenetic distribution of the Crenactin-encoding gene (cren-1) across archaeal genomes correlated with rod- or filamentous cell morphologies, supporting an involvement in cell shape formation for the archaeal actin orthologs.

The cren-1 gene was found to belong to a conserved gene cluster, which, in agreement with the involvement in cytoskeletal processes, was denoted Arcade (actin-related cytoskeleton in Archaea involved in shape determination; grey panel in Fig. 1). In addition to cren-1, the gene cluster comprised several genes (rkd genes) whose products, Arcadins, were shown also to assemble into helical structures, presumably in conjunction with Crenactin filaments.

The Arcadin-2 gene product, in contrast, displayed a punctuated distribution across P. calidifontis cells, and was found to localize between segregated nucleoids in a cell subpopulation and might, hence,
Crenactin and Eukaryogenesis

Recent phylogenomic studies have provided support for an evolutionary scenario in which the eukaryotic lineage emerged from within the archaeal domain.\textsuperscript{9,11} In particular, evidence has been presented for a model in which eukaryotes are suggested to have originated from a lineage that also gave rise to the Crenarchaeota phylum.\textsuperscript{12} In light of this, the discovery of an actin-based cytoskeleton in archaea, as well as in other deep archaeal lineages, adds momentum to discussions concerning the origin of the eukaryotic cell and early stages of eukaryogenesis. The role of actin as a key-player in invagination-based processes such as endocytosis and phagocytosis is well documented, and a requirement for a phagocytic machinery to engulf a putative alphaproteobacterial ancestor of the mitochondrion has been put forward as a prerequisite in several models to explain the origin of the eukaryotic cell (reviewed in ref. 13).

In certain evolutionary models, it is argued that the proto-eukaryote was relatively complex at the cellular level in order to execute engulfment.\textsuperscript{13,14} The phagocytic machinery present in extant eukaryotes is, indeed, highly complex and comprises a wide variety of associated proteins,\textsuperscript{2} in addition to actin. We envision that Crenactin could have provided primitive phagocytic capabilities to an ancestral archaeal lineage that contributed to the emergence of the eukaryotic cell, by facilitating the fusion with the proto-mitochondrion\textsuperscript{3} in a far less complex process than that carried out by the fully developed phagocytic machineries present in extant eukaryotes. Interestingly, several members of the Thermoproteales, all of which contain the Crenactin-encoding gene, display bent or even branched cells, occasionally with extended cellular protrusions.\textsuperscript{15,16} Assuming these structures are not methodological artifacts, this suggests a certain degree of cytoskeletal flexibility in these species, and, by inference, in ancestral archaeal lineages. It is, thus, conceivable that, in an archaeal lineage from which the eukaryotic lineage would have emerged, such actin-mediated cytoskeletal flexibility may have allowed for primitive

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**Figure 1.** An archaeal actin ortholog. Schematic overview of actin protein family phylogeny, demonstrating strong support (bootstrap value 97) for common ancestry of archaeal Crenactin and eukaryotic actin proteins. The tree was generated as described previously in reference 3, except that distant members (MreB and Hsp70) of the actin protein family were omitted and novel archaeal sequences were added (Crenactin orthologs from Ca. Caldiarchaeum subterraneum, *Vulcanisaeta distrausta*, *Vulcanisaeta moutnovskia* and *Thermoproteus uzonensis*). For clarity, the actin, ParM/ALPs and MamK clades are displayed as triangles. The grey shading highlights the Crenactin clade and, for each species, the gene organization of the archae gene cluster is indicated, with the genes encoding Arcadin-1 (*rkd-1*), Crenactin (*cren-1*), Arcadin-2 (*rkd-2*), Arcadin-3 (*rkd-3*) and Arcadin-4 (*rkd-4*) depicted in orange, blue, green, magenta and yellow, respectively. The tree was rooted according to the topology obtained previously in reference 3, with bootstrap values shown for branches with a support above 70 (out of 100 replicates). Sequences are denoted by species name and gene identifier, with crenarchaeal, korarchaeal and aigarchaeal Crenactin orthologs indicated in green, red and blue fonts, respectively. Note that the Crenactin ortholog of *Ca. Caldiarchaeum subterraneum* represents the deepest branch in the Crenactin tree, although with low support.

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play a cytokinesis-associated role in this organism. The presence of an Arcadin-2 ortholog in the composite genome of *Candidatus Caldiarchaeum subterraneum*, belonging to the recently proposed Aigarchaeota phylum\textsuperscript{3} is, however, puzzling since this genome also encodes components of both the CdvBC and FtsZ cell division machineries, although the latter protein seemingly lacks the typical tubulin motif.\textsuperscript{5} In this light, it is interesting to note that recent studies have reported that CdvA protein has been shown to bind DNA,\textsuperscript{7} and that the FtsZ homolog TubZ has been reported to effectuate plasmid segregation in *Bacillus* species.\textsuperscript{8} These observations open up the possibility that the CdvA function in *Ca. Caldiarchaeum subterraneum* has been taken over by Arcadin-2, in accordance with an involvement in division, together with CdvB and CdvC, while the FtsZ homolog might play a role in the genome segregation process and/or in coordination of mitosis and cell division. However, other scenarios are also possible, and experimental data is required to conclusively corroborate the functions of the respective proteins in these evolutionary distinct archaeal lineages.

Irrespective, a picture emerges from the *P. caldophila* study in which a core cytoskeleton is formed by Crenactin and Arcadin-1, with Arcadin-3 and -4 performing auxiliary roles, while the precise function of Arcadin-2 in the Thermoproteales\textsuperscript{1} needs to be further investigated. Detailed biochemical and molecular characterization of Crenactin and Arcadins is also of potential value for biotechnological exploitation, due to the intrinsic heat-stability of all gene products from the hyperthermophilic *P. caldophila* species.

Importantly, the identification of Crenactin-based shape-determining structures in Archaea indicates that an actin-based cytoskeleton was established prior to the divergence of the archaeal and eukaryotic lineages. In the following, we discuss the implications of this observation with respect to the process of eukaryogenesis and the emergence of the eukaryotic cell.
phagocytosis that occasionally sustained cellular fusions.

In addition, a recent study of the deeply branching microbial eukaryote *Giardia intestinalis* revealed that the actin cytoskeleton was capable of facilitating receptor-mediated endocytosis in absence of canonical actin-binding proteins. Whether the relatively simple cytoskeleton in *Giardia* represents a relic of an ancient past, or whether it is the result of reductive genome evolution remains to be elucidated. Yet, it does indicate that actin is capable of sustaining membrane invagination in absence of the array of proteins usually involved in this process.

### Actin Proliferation and Sub-Functionalization during Early Stages of Eukaryogenesis

Apart from a potential role in cellular fusion events that may have given rise to the eukaryotic cell, actin has functionally diversified during evolution and is, in addition to the well-characterized cytoskeletal roles, implicated in other key processes that may have promoted eukaryogenesis. For example, a number of recent studies have provided evidence for nuclear localization of an actin sub-fraction where, in conjunction with other proteins, it is involved in the assembly of the nuclear envelope and in processes influencing transcriptional activity.

Further support for the importance of actin in the evolution of the eukaryotic cell may be inferred from the excessive proliferation of actin homologs across all branches of the eukaryotic tree. Most eukaryotes contain at least eight actin orthologs in the nuclear envelope and in processes influencing transcriptional activity.

For example, several ARPs have been identified as members of chromatin remodeling complexes that may have promoted eukaryogenesis. For example, several ARPs have been implicated in the assembly of the nuclear envelope and in processes influencing transcriptional activity.

### Additional Eukaryotic Signatures in Archaeal Phyla

The identification of a Crenactin-based cytoskeleton in Archaea necessitates a re-evaluation of the hypothesis of the actin cytoskeleton as specific to the eukaryotic cell, and the presence of actin orthologs in members of the *Crenarchaeota* and other proposed archaeal phyla lends support to scenarios that associate a progenitor of this phylum with the origin of the eukaryotic cell. Interestingly, recent studies have revealed a series of additional presumed eukaryotic signature proteins (ESPs) in different established or inferred, archaeal phyla (Fig. 2). For example, orthologs of the eukaryotic ESCRT-III membrane remodeling system have been found to be a part of the Cdv cell division machinery in *Crenarchaeota* and, in addition, several eukaryotic transcription machinery components (Rcps4, RpoG and Elf1) have been identified in *Crenarchaeota*. More recently, an analysis of the *C. Caldivirga* genome revealed the presence of a presumably fully functional ubiquitin-like protein modifier system. Further exploration of uncharacterized deep archaeal lineages may well reveal additional eukaryotic-like features, and thus provide new insights into events that may have been fundamental for the origin and early evolution of the eukaryotic cell.

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