Bug22p, a Conserved Centrosomal/Ciliary Protein Also Present in Higher Plants, Is Required for an Effective Ciliary Stroke in Paramecium

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Centrioles, cilia, and flagella are ancestral conserved organelles of eukaryotic cells. Among the proteins identified in the proteomics of ciliary proteins in Paramecium, we focus here on a protein, Bug22p, previously detected by cilia and basal-body high-throughput studies but never analyzed per se. Remarkably, this protein is also present in plants, which lack centrioles and cilia. Bug22p sequence alignments revealed consensus positions that distinguish species with centrioles/cilia from plants. In Paramecium, antibody and green fluorescent protein (GFP) fusion labeling localized Bug22p in basal bodies and cilia, and electron microscopy immunolabeling refined the localization to the terminal plate of the basal bodies, the transition zone, and spots along the axoneme, preferentially between the membrane and the microtubules. RNA interference (RNAi) depletion of Bug22p provoked a strong decrease in swimming speed, followed by cell death after a few days. High-speed video microscopy and morphological analysis of Bug22p-depleted cells showed that the protein plays an important role in the efficiency of ciliary movement by participating in the stroke shape and rigidity of cilia. The defects in cell swimming and growth provoked by RNAi can be complemented by expression of human Bug22p. This is the first reported case of complementation by a human gene in a ciliate.

The centriole, a microtubule-based cylindrical structure, is an ancient organelle traced to primitive eukaryotes and present in numerous phyla but lost in some others, such as fungi and higher plants (24). Centriolar function is dual: organization of the centrosome and, as a basal body, nucleation of an axoneme, the backbone of cilia and flagella. The ultrastructure of centrioles and axonemes appears to be highly conserved. In the majority of cases, the centriole is composed of 9 triplets of microtubules, and when the centriole acts as a basal body, the triplets are elongated into 9 doublets of microtubules forming the axoneme, with occurrence of a central pair of microtubules (9-plus-2 pattern) (reviewed by Satir and Christensen [30]). Renewal of interest in cilia in recent years has been prompted by their implication in numerous functions in development and cell physiology through their motile and sensory properties, so that many human genetic diseases, now called ciliopathies, appear to arise from primary defects in cilia proteins (14). The ultrastructural conservation of centrioles and cilia in evolution is correlated with good conservation of the protein constituents.

High-throughput studies (proteomics, transcriptome analysis, and comparative genomics) have been conducted in several species to identify ciliary and centriolar proteins or proteins involved in their biogenesis. The set of proteins identified comprises both proteins specific to cilia (such as outer- and inner-arm dynein subunits or radial-spoke proteins) or to their biogenesis (specific transcription factors and intraflagellar-transport [IFT] proteins) and ubiquitous proteins (such as tubulins).

In order to identify novel conserved proteins and to specify their roles in ciliary function, we built a new knowledge base, Cildb (1) (http://cildb.cgm.cnrs-gif.fr), which integrates orthology links between the whole proteomes of 18 species and high-throughput studies identifying ciliary proteins, as well as the OMIM database. Cildb includes species for which no ciliary studies are available and species that lack centrioles or cilia in order to distinguish proteins specific to these organelles from ubiquitous proteins present in them.

In the present work, starting from the ciliary proteome analyzed in Paramecium (1), the Bug22p protein, highly conserved through evolution, was selected. Apart from its identification in high-throughput studies of various species (6, 8, 17, 21, 23, 25, 32, 36), no analysis dedicated to Bug22p exists to date for any organism. This protein has amazing sequence
conservation, indicating a key role in cellular processes and/or the existence of numerous interacting partners. Intriguingly, the protein is also present in higher plants (e.g., Arabidopsis), a phylum in which centriolar and ciliary structures are absent, which induced us to analyze the relevance of its ciliary function.

We report here the localization of Bug22p in HeLa and Paramecium cells and its functional study in Paramecium. The choice of this organism, beyond the power of functional analysis by RNA interference (RNAi) (13), stems from the fact that we can specifically analyze the motility aspect of ciliary function.

MATERIALS AND METHODS

Strains and culture conditions. Stock d4-2 of Paramecium tetraurelia, the wild-type reference strain, was used in all RNAi experiments. The mutant ndt-1 (31), which carries a recessive monogenic mutation preventing trichocyst discharge, a dispensable function under laboratory conditions, was used for transformations. Cells were grown at 27°C in wheat grass infusion, BHB (L’arbre de vie, Luçay Le Mâle, France), inoculated with Klebsiella pneumoniae and supplemented with 0.8 µg/ml β-sitosterol according to standard procedures (33).

Expression vectors. In order to visualize the cellular localization of Bug22p, the Bug22a paramecium gene, as well as the human (Hs) Bug22 cDNA, were cloned into the Paramecium expression vector pPXV-GFP. The human Bug22 cDNA was also cloned into the plasmid pD2EGFP for expression in HeLa cells. In all three constructs, the green fluorescent protein (GFP) sequence was fused at the N terminus of the Bug22 gene.

(i) pPXV-HsBUG22 expression vector for Paramecium. The human Bug22 gene was amplified by PCR from a cDNA library kindly provided by François Lacroute using the primers 5’-GGTGCTCGAGATGTTCAAAAACACGT TCCAG-3’ and 5’-GGTGGGTACCTCATCTGTTGGCCTGTTGCG-3’ and cloned into the XhoI/KpnI sites of the RNAi expression vector pPXV. The sequences of genes to be knocked down were amplified by PCR and cloned into a modified Litus 26 plasmid (New England Biolabs) in which the EcoRI site had been replaced by an SphI site, using the method developed for PCScript cloning by Stratagene (La Jolla, CA), and transferred to the feeding vector L4440 (37) into the Xhol/HindIII sites of the polylinker between two T promoters.

(ii) Constructs for RNAi of the Paramecium BUG22 genes. Two constructs were used to knock down the two subfamilies of BUG22 genes. For BUG22a and -b knockdowns, Bug22a was amplified from positions 75 to 391 with the primers 5’-GGATAAAGAATGTTGGTGGG-3’ and 5’-CTTATTACACTTTAAGAATGTTGGG-3’ and cloned into the NcoI/KpnI sites of the pPXV-GFP vector.

(iii) Constructs for RNAi of the Paramecium BUG22 genes. The human Bug22 gene was amplified by PCR with primers 5’-GGTGCTCGAGATGTTCAAAAACACGT TCCAG-3’ and 5’-GGTGGGTACCTCATCTGTTGGCCTGTTGCG-3’ and cloned into the KpnI site of the pPXV-GFP vector.

Protein extraction, SDS gels, and Western blots. Paramecium protein extracts were prepared from a pellet of 2 × 10^5 cells homogenized in a potter with trichocysts crude. 10 mM Tris, 1 mM EDTA, 1% SDS, 50 mM NaCl, 2 mg/ml protease inhibitor cocktail (Set1; Calbiochem). Low-speed pellets and superna
tants were separated by centrifugation at 10,000 × g. Total protein extracts of bacteria were obtained by sonication of a bacterial pellet in lysis buffer (50 mM HEPES, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, 0.1% SDS, 0.1% Na deoxycholate). Gel electrophoresis was performed using Criterion XT gels at neutral pH and 12% Bis-Tris (Bio-Rad) according to the manufacturer’s recommendations. The primary antibodies used were anti-GTL3 (1:500) and anti-GFP (1:750). The secondary antibody was anti-rabbit–AP (1:500). Western blots were revealed using the 5-bromo-4-chloro-3-indolylphosphate (BCIP)-ni troblue tetrazolium (NBT) liquid substrate system from Sigma.

Electron microscopy. For morphological observations, cell pellets were processed as previously described (28) using the anti-ciliary tubulin (1:500), the monocular ID5 (1:500), or the anti-ciliary rootlet (1:400) antibody. For both HeLa cells and paramecia, appropriate fluorescent secondary antibodies were applied at 1:500 dilutions, nuclei were stained with Hoechst-33258, and slides were mounted in Cityfluor (London). Preparations were observed using a Zeiss Axioplan 2 plus fluorescence microscope equipped with a Roper CoolSnap-CF intensified camera with GFP filters. Images were processed with Metamorph software (Universal Imaging).

To the protocol of the manufacturer (Qiagen, Courtaboeuf, France). The sequences of Paramecium genes were retrieved from ParameciumDB (2) (http://paramecium.cgm.cnrs-gif.fr) and verified by rescoring after cloning.

Transfection. (i) HeLa cells. HeLa cells were maintained in Dulbecco’s mod-
ified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (FBS) and 1% penicillin-streptomycin (all from Gibco) at 37°C in 5% CO2. The day prior to transfection, HeLa cells (1 × 10^5) were plated on a 24-well plate containing prestertilized coverslips in DMEM supplemented with 10% FBS. The cells were transfected with 0.5 µg of Hs-Bug22-GFP using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s directions.

(ii) Paramecium transformation. ndt-1 cells were transformed by microinjec
tion into their macronuclei of filtered and concentrated plasmid DNA (5 µg/µl) containing a mixture at a ratio of 1:1 of the plasmids of interest (GFP fusion genes and Hs-Bug22 expression) and a plasmid directing the expression of the ND7 gene (31). Microinjection was done under an inverted Nikon phase-contrast microscope, using a Narishige micromanipulation device and an Eppendorf air pressure microinjector. Successfully transfected cells were screened for the ability to discharge trichocysts on picric acid stimulation and further analyzed.

Antibodies. The polyclonal GTL3 anti-Bug22p antibody was purchased from Aviva Systems Biology (San Diego, CA). The mononuclear antibody 2H5, raised against Odhomydomonas centrin (29), was used to label centrioles in HeLa cells. The monoclonal antibody IDS (38), which essentially labels basal bodies and a few microtubule arrays (10), was used to visualize the basal-body pattern. The polyclonal anti-GFP antibody was purchased from Interchim (Montluçon, France). The polyclonal anti-Paramecium ciliary tubulin (9) and anti-ciliary root
ettlet (34) antibodies were homemade antibodies.

The secondary antibodies used in immunofluorescence were Alexa Fluor 488 or 594 (Molecular Probes) or from Invitrogen, the antibody used in Western blot experiments was an anti-rabbit IgG–alkaline phosphatase (AP) conjugate (Promega), and the antibodies used in immunoelectron microscopy were colloidal-gold-conjugated anti-rabbit immunoglobulins (Gar GS and Gar G10; Aurion).

Protein extraction, SDS gels, and Western blots. Paramecium protein extracts were prepared from a pellet of 2 × 10^5 cells homogenized in a potter with trichocyst crude. 10 mM Tris, 1 mM EDTA, 1% SDS, 50 mM NaCl, 2 mg/ml protease inhibitor cocktail (Set1; Calbiochem). Low-speed pellets and superna
tants were separated by centrifugation at 10,000 × g. Total protein extracts of bacteria were obtained by sonication of a bacterial pellet in lysis buffer (50 mM HEPES, pH 7.5, 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, 0.1% SDS, 0.1% Na deoxycholate). Gel electrophoresis was performed using Criterion XT gels at neutral pH and 12% Bis-Tris (Bio-Rad) according to the manufacturer’s recommendations. The primary antibodies used were anti-GTL3 (1:500) and anti-GFP (1:750). The secondary antibody was anti-rabbit–AP (1:500). Western blots were revealed using the 5-bromo-4-chloro-3-indolylphosphate (BCIP)-ni troblue tetrazolium (NBT) liquid substrate system from Sigma.

Immunofluorescence microscopy. (i) HeLa cells. After 24 h of culture, cells were washed with phosphate-buffered saline (PBS), fixed for 6 min at –20°C in ethanol, and blocks of at least 10^5 cells were incubated in 4% paraformaldehyde, 0.1% sodium thiocyanate, 0.1% Tween 20 in PBS for 30 min. The cells were incubated at room temperature with either the 2H5 (1:1,000) or the GTL3 (1:200) antibody for 1 h and then washed three times in 1% BSA and 0.1% Tween 20 and incubated for 1 h with the secondary antibody before three additional washes.

(ii) Paramecium. Immunostaining of cells was carried out as previously de
dscribed (28) using the anti-ciliary tubulin (1:500), the monoclonal ID5 (1:350), or the anti-ciliary rootlet (1:400) antibody. For both HeLa cells and paramecia, appropriate fluorescent secondary antibodies were applied at 1:500 dilutions, nuclei were stained with Hoechst-33258, and slides were mounted in Cityfluor (London). Preparations were observed under a Zeiss Axioplan 2 plus fluorescence microscope equipped with a Roper CoolSnap-CF intensified camera with GFP filters. Images were processed with Metamorph software (Universal Imag

Electron microscopy. For morphological observations, cell pellets were pro-
cessed as previously described (5) using 1% or 2% glutaraldehyde for fixation. For postembedding immunolocalization, cell pellets were processed as described previously (18), using either 2% paraformaldehyde, 0.15% glutaraldehyde, or 3% paraformaldehyde and 0.5% glutaraldehyde for fixation, and embedded in LR White (London Resin). Thin sections were collected on nickel grids and treated with double-diluted water or 0.1 M NH4Cl in 0.1 M PBS and then saturated and processed with 3% BSA in PBS. Primary polyclonal antibodies were diluted 1:50 to 1:75 (anti-GTL3) and 1:10, 1:20, or 1:500 (anti-GFP). After being washed, the sections were incubated with, respectively, 1:40, 1:75, or 1:100 dilutions of 5-nm (GTL3 labeling) or 10-nm (GFP labeling) gold particles. The sections were examined with a Philips CM10 or a Jeol transmission electron microscope.
RNAi by feeding. RNAi gene knockdown was performed according to the method of Galvani and Sperling (13). HT115, an RNAIII-deficient strain of *Escherichia coli* with an isopropyl-β-D-thiogalactopyranoside (IPTG)-inducible T7 polymerase, was transformed by the desired constructs into Littmus or L4440 plasmids. Wild-type paramecia were incubated in double-stranded-RNA (dsRNA)-expressing bacteria and were transferred daily into fresh feeding medium. Control cells were fed with bacteria carrying the complete coding region of the *ND7* gene.

Swimming analysis. Two methods were used for swimming analysis. First, we measured the swimming speed of cells under dark-field microscopy using a Zeiss Axioskop 2-plus microscope with a 10× objective by recording tracks during 650-ms pauses. Second, we used a Zeiss Stemi 2000-C dissecting microscope with 1-min time-lapse acquisitions at 7 frames per second with a Roper CoolSnap-CP camera and Metamorph software (Universal Imaging). The speed means and standard errors were calculated by measuring the lengths of the tracks using ImageJ and subtracting a mean length of the paramecia for the first method and using automated analysis of the swimming tracks currently under development for the second method (J. Ferracci, O. Arnauz, and J. Cohen, unpublished data).

High-speed video microscopy. All of the observations were made under an inverted microscope (Axiovert 200; Carl Zeiss S.A.S., Le Pecq, France) using a 100× objective on 5 μl of a suspension of paramecia immobilized between a cover glass (diameter, 50 mm; thickness, 0.13 to 0.16 mm) from Menzel-Glaser and a 22- by 22-mm coverslip, with a few hundred calibrated 10×, 15×, or 20-μm polystyrene beads (Fluka), during overall periods shorter than 5 min to avoid changes in ciliary activity. A final dilution of ca. 1/100 of India ink was added to the medium to be visualized as particles that materialized the fluid flow during the acquisitions. Cilium movements were digitally recorded with a camera (PixeLink A741; Ottawa, Canada) at a rate of 355 images per second. A movie was constituted of 1,800 images with a definition of 256 by 192 pixels. The size of a pixel was 0.13 by 0.13 μm². In each sequence, we selected a cilium that we were able to follow during an entire cycle of beating. For this, the video sequences were played back frame by frame using Streampix software (Norpix Inc., Montreal, Canada). The angle of beating was calculated by trigonometric calculations in the triangle made by the anchoring point and the two extreme positions of the cilium extremities during the power stroke.

Swimming-behavior assays. Cells to be assayed were first equilibrated for a few minutes in resting solution (2 mM Na citrate, 1 mM Na2HPO4, 1 mM Na2HPO4, 1.5 mM CaCl₂) and then individually transferred to test solutions that consisted of either 30 mM KCl or 5 mM tetrathylammonium, 10 mM NaCl in resting solution (19). Observation of immediate swimming behavior was done during the first minute after transfer under the dissecting microscope.

RESULTS

Conservation of Bug22p in eukaryotes. Bug22p, a small basic protein of 22 kDa, was one of the abundant proteins identified in a high-throughput proteomics analysis of *Paramecium* cilia (9 distinct peptides), and its mRNA level in cells regenerating cilia was increased by a 2- to 3-fold induction (reference 1 and unpublished data). A BLAST search for similar proteins in other species for which whole-genome sequence data are available revealed that it is widespread in eukaryotes harboring centrioles/cilia at some stage of their life cycles. Among eukaryotes devoid of centrioles/cilia, plants possess *BUG22* genes whereas fungi do not. Interestingly, fungi, such as *Batrachochytrium dendrobatidis*, which form flagellated metates also have a *BUG22* gene. The protein sequence of *Bug22p* is highly conserved throughout evolution (Fig. 1), except for the presence of C-terminal extensions in *Trypanosoma* and *Caenorhabditis*. Such lack of sequence variation during evolution suggests a conserved role in different organisms, including but not restricted to ciliary function, since the sequence conservation is also high in plants. Despite high overall amino acid sequence conservation, diagnostic positions in the sequence allow us to distinguish *BUG22* genes of acenotriolar organisms (higher plants) from the others, as shown in a global alignment (Fig. 1).

The whole *Bug22p* protein constitutes by itself the Interpro domain DUF667 (for domain of unknown function) that occurs in some proteins other than *Bug22p*. However, the sequence conservation of the DUF667 domain in these proteins is much lower than within the *Bug22p* subfamily, suggesting that the functions of the less conserved DUF667 domains may be divergent with respect to the *Bug22p* domain.

Four *Bug22* genes for a single *Bug22p* protein in *Paramecium*. Four paralogous *BUG22* genes encoding four 100% identical proteins are present in the *Paramecium* genome. These paralogous genes arose from the last two whole-genome duplications (WGDs) that occurred in this organism (3). The four genes, with the identifiers (IDs) GSPATG00007012001, GSPATG00016658001, GSPATG00012810001, and GSPATG00013199001, were, respectively, called *BUG22a*, -b, -c, and -d. The DNA sequences are almost identical (96.1% identity between *BUG22a* and -b and 96.9% between *BUG22c* and -d [both pairs are paralogs from the most recent WGD]) and 83.2% to 83.8% between the pair *BUG22a* plus -b and the pair *BUG22c* plus -d. Most of the divergence occurs in intron sequences, as visualized by multiple alignments (see Fig. S1 in the supplemental material). The few differences in exons are all silent.

Localization of *Bug22p* in HeLa cells and *Paramecium*. HeLa cells transfected with the GFP*-HisBug22* cDNA displayed labeling of two dots close to the nucleus, which colocalized with centrin, a centriolar marker (11), as revealed by the 20H5 monoclonal antibody (see Fig. S2A in the supplemental material), indicating that Hs*Bug22p* is present at the centrosome. This observation is similar to images published by Keller et al. (16), who transfected U2OS cells with a C-terminal GFP*-Bug22p* fusion protein in a systematic study of centriolar proteins. Our results thus confirm the centriolar localization of *Bug22p*.

Both the *HsBUG22* and the *PbBUG22* GFP fusion genes were used to transform *Paramecium* by DNA microinjection into the macronucleus, and the fluorescence was observed in the clonal descendants of the injected cells. Labeling of basal bodies and the proximal two-thirds of cilia was observed with both GFP fusions (Fig. 2), showing that, in agreement with the proteomics of cilia in several species, *Bug22p* localizes to basal bodies and cilia and that the human ortholog Hs*Bug22p* is targeted to these organelles, as well.

GTL3 is a commercial antibody raised against the C-terminal portion of the human *Bug22p* protein. As the primary sequence is highly conserved between *Paramecium* and humans (89% identity in the amino acid sequences) (Fig. 1), we tested the GTL3 antibody by Western blotting (Fig. 3) and found that it specifically recognized a 22-kDa peptide in *Paramecium* extracts (Fig. 3A), as well as the GFP*-HisBug22* fusion expressed in *E. coli* (Fig. 3B and C). In HeLa cells, the GTL3 labeling coincided with the localization of centrin (see Fig. S2B in the supplemental material) and GFP*-HisBug22* (see Fig. S2C in the supplemental material), confirming its *Bug22p* specificity.

In immunofluorescence, no labeling of *Paramecium* by GTL3 was observed. This negative result may be due to the solubility of *Bug22p* molecules in the medium after detergent permeabilization or to lack of accessibility to the antigen under the fixation and labeling procedures used. The protein local-
ization was then examined by postembedding immuno-electron microscopy, both with the GTL3 antibody on wild-type cells and with an anti-GFP antibody on GFP-Bug22p-expressing cells (Fig. 4). Examination of 142 sections of cilia and basal bodies using the GTL3 antibody and 115 sections with the anti-GFP antibody indicated that both antibodies gave the same labeling in the basal bodies and cilia: gold particles oc-
curred preferentially in the terminal plate of the basal body and the transition zone to the cilia, as well as in the cilia, in the vicinity of the outer doublets of the axoneme, and often between the axoneme and the membrane. In GFP-expressing cells, additional labeling was often found within the basal body, close to its proximal part (Fig. 4G). In GFP-expressing cells, 34 longitudinal sections of cilia revealed an apparent periodicity...
of the labeling as a regular disposition of the gold particles at 500- to 550-nm intervals (Fig. 4J and K) and, less often, at 250- and 800-nm intervals.

**Growth and swimming defects provoked by Bug22p depletion in Paramecium** can be complemented by the human ortholog. The function of Bug22p was investigated in Paramecium using the RNAi feeding method. Since the four Paramecium BUG22 genes encode the same protein, we would expect not qualitative but quantitative differences by inactivating each paralog of the gene family. We used BUG22a and BUG22c sequences for double-stranded-RNA production. The two dsRNAs were expected to silence all four paralogs, based on sequence conservation (see Fig. S1 in the supplemental material). Double silencing using the two sequences was needed to improve the knockdown efficiency on all four genes.

We first evaluated the efficiency of the RNAi procedure on fluorescent Paramecium cells from a clone transformed with the GFP-PtBug22a fusion gene: the fluorescence was completely abolished by the double BUG22a-BUG22c knockdown, showing that the RNAi efficiency depleted Bug22p (data not shown). Phenotypic analyses obtained in further RNAi experiments were performed on nontransformed wild-type cells.

Knocking down each pair of BUG22a-BUG22b or BUG22c-BUG22d genes resulted in a growth rate of 1 or 2 fissions per day compared to 4 or 5 in control cells and led to lethality in 2 to 3 days. When all 4 genes products were knocked down, cells divided 1 or 2 times on the first day and 0 or 1 time on the second day and died within 3 days. Lethality might be the result of starvation owing to impaired ciliary beating (see below) that prevented food capture in the oral apparatus, as observed by defective take-up of India ink (not shown).

Swimming speed was measured by recording swimming tracks under dark-field microscopy, which showed that, after 1 day of silencing, knocking down each pair of gene products decreased the rate of motility by 75%, while knocking down all four gene products decreased the rate of motility by 90% (Fig. 5).

To test whether a human Bug22 protein could complement RNAi depletion of Paramecium orthologs, we transformed Paramecium for heterolog expression of the human BUG22 cDNA and submitted the cells to RNAi against the BUG22a and BUG22c Paramecium sequences, which do not share nucleotide identity stretches sufficient to induce silencing of the human gene, and reciprocally to RNAi against the human BUG22 cDNA, which did not cross-react with Paramecium sequences (see Fig. S1 in the supplemental material). A control with double human and Paramecium BUG22 RNAi was also performed. We observed that neither HsBUG22 nor PtBug22 RNAi alone could alter the growth rate or the swimming speed, whereas the double Hs/Pt RNAi could (Fig. 6 and FIG. 2. Ciliary and basal-body localization of Paramecium and human Bug22p GFP fusions in Paramecium. (A and B) Wild-type paramecia immunofluorescently labeled with the ID5 monoclonal antibody to visualize basal bodies (A) and with the anti-ciliary tubulin to show cilia (B). (C) Paramecium expressing GFP-PtBug22p showing clear labeling of basal bodies and cilia. (D) Same as panel C for a Paramecium expressing GFP-HsBug22p. Bar = 10 μm.

FIG. 3. Western blot evidence that Bug22p is the antigen of a commercial anti-GTL3 antibody. Lanes: A1, molecular mass markers for lane A2; A2, anti-GTL3 labeling of a 10,000 × g supernatant of whole homogenized paramecia showing a positive band between 20 and 25 kDa; B1 and C1, total extracts of E. coli expressing GFP in the pPXV plasmid normally used for expression in Paramecium but which allows low expression in the bacteria; B2 and C2, E. coli expressing the GFP-Bug22p fusion; C3, molecular mass markers for lanes B1 to C2. The B series lanes were revealed with the anti-GTL3 antibody and the C series with the anti-GFP antibody. The asterisk indicates the level of the fusion protein that is recognized by both GTL3 and GFP antibodies.
its legend provide details). This demonstrates the ability of the human protein to rescue depletion of *Paramecium* Bug22p.

**Bug22p depletion affects swimming behavior.** When carefully observed under the dissecting microscope, live Bug22p-depleted cells seem to undergo less frequent backward swimming than control cells. Since these events are rare under control conditions, we tested swimming behavior in ion solutions, as described for the identification of Mendelian mutants (19). Control cells immersed in 30 mM KCl or in 5 mM tetraethylammonium, 10 mM NaCl (see Materials and Methods) immediately underwent continuous backward swimming for more than 20 s and then whirled for a few seconds and resumed normal forward swimming. In contrast, Bug22p-depleted cells showed a very much weakened reaction, with very short or no backward swimming and whirling for less than 5 s. This suggests abnormal ciliary-channel activity or abnormal response of the axonemal motors to ion currents, induced by RNAi knockdown.

**Bug22p depletion disturbs basal-body organization in the cortex of Paramecium.** Since the Bug22p protein is present in basal bodies, in addition to cilia, we wondered whether its depletion causes basal body defects. We therefore examined Bug22p-depleted cells using antibody labeling that revealed, not only the basal bodies themselves (antibody ID5), but also the ciliary rootlets (anti-KD), which provide good markers of the orientation of individual basal bodies in the global pattern of the polarized cell cortex. By the second day under BUG22 RNAi conditions, a sporadic disorganization started to appear, with ciliary rootlets showing diverse erratic orientations (see Fig. S3 in the supplemental material). This effect on basal-body organization and polarity could be a direct consequence of Bug22p depletion or an indirect effect secondary to alteration of ciliary activity.

**Bug22p depletion alters ciliary beating efficiency and stroke shape in Paramecium.** To investigate the cause of the decrease in swimming speed induced by RNAi depletion of Bug22p, we first checked the presence of cilia by immunofluorescence using an anti-tubulin antibody. Cilia appeared unaffected in number, density, shape, or length, in contrast to what is obtained by silencing IFT172, a component of the intraflagellar-transport machinery involved in ciliary growth and maintenance (26), used as a control (see Fig. S4 in the supplemental material). In

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**FIG. 4. Localization of Bug22p in Paramecium by postembedding immunoelectron microscopy.** (A to F) Labeling of wild-type cells with the GTL3 antibody using 5-nm gold particles. (G to N) Labeling of GFP-Bug22p-expressing cells with an anti-GFP antibody using 10-nm gold particles. The arrowheads point to gold particles. (A and H) Labeling of the terminal plates of basal bodies. (B, D, and I) Labeling of the transition zone to the cilium. (F and J to N) Labeling in the vicinity of the outer doublets of the axoneme. (C, E, and J) Labeling between the axoneme and the membrane. (G) Labeling within the basal body close to its proximal part. (J and K) Examples of regular disposition of the gold particles along the axoneme. ci, cilium; bb, basal body; tp, terminal plate; tz, transition zone. Bar = 250 nm.
living cells observed under phase-contrast microscopy at high magnification, the cilia appeared to beat vigorously, in apparent contradiction of the strong decrease in the cell swimming speed. This indicates that other factors than beating frequency can affect swimming speed. We therefore studied the ciliary beating of Bug22p-depleted cells by high-speed video microscopy (Fig. 7; see Videos S1 and S2 in the supplemental material). The recorded images revealed that although the cilia beat at an even higher frequency than normal (42 Hz versus 14 Hz), they beat with reduced efficiency, since (i) the angle of beating was narrower (55° versus 97°) and (ii) the apparent distinction between power and recovery strokes, as well as the coordination of beating over the cell surface, was lost. In addition, cilia appeared "floppy" when they moved, as though they were less rigid, which could explain their altered beating. The global impairment in the beating efficiency of fluid movement could be visualized by particles in the medium moving in a disorganized manner, more or less randomly, compared to the control, where particles were rapidly swept in a dominant direction.

Examination of Bug22p-depleted cells under transmission electron microscopy did not reveal particular defects in the general organization of the basal body or cilium (9-fold symmetry, a central pair of microtubules, and radial spokes in the axoneme). However, the overall shape of the cilia was strongly altered in 30% of the longitudinal sections observed, with sharp bending that was never observed in controls, accompanied by shrinking of the axoneme diameter in the region of the bend (Fig. 8). This indicated that the natural rigidity of cilia was greatly impaired by the depletion of Bug22p, in good agreement with the observation of the shape of beating cilia in live cells.

**DISCUSSION**

**Bug22p, a novel ciliary protein.** The present study of Bug22p was stimulated by finding the protein in proteomic analysis of *Paramecium* cilia (1) and by the fact that we could extract it by mining the results of high-throughput studies of various species. In organisms with centrioles and cilia, high-throughput studies provided strong evidence that Bug22p has a localization/function related to these organelles. In *Chlamydomonas*, Bug22p has been detected in proteomic analysis of flagella (25) and basal bodies (17), and *BUG22* mRNA is upregulated during flagellar biogenesis (17, 36). The protein has also been found in *Trypanosoma* flagella (8), in *Tetrathyamina* cilia (32), in human cilia (23), and in mouse photosensory receptors (21). The promoter of the *BUG22* gene in *Caenorhabditis* displays an X box (6), characteristic of genes encoding proteins related to ciliary biogenesis and regulated by the daf19 transcription factor. The mouse ortholog of *BUG22, GTL3*, has been identified in a gene trap screen (transcript T10-2A2) as expressed in many tissues, but preferentially in the testis (27), an organ where flagellated sperm cells develop. A large-scale cDNA screen by *in situ* hybridization of zebra fish embryos (35; http://zfain.org/) localized major expression of *BUG22* in tissues where cilia are known to play an important role in embryonic development—Kupffers vesicle (equivalent to the embryonic node of mammals), neural tube, spinal cord, and pronephric duct—which is compatible with preferential expression in ciliated tissues. *In situ* hybridization in *Drosophila* (Berkeley *Drosophila* Genome Project [http://www.fruitfly.org/]) revealed preferential expression of *BUG22* in sensory complexes during embryogenesis, again compatible with expression in ciliary tissues. The high sequence conservation and apparently ubiquitous presence of Bug22p in centrosomes, cilia, and flagella stimulated its further study.

**Potential roles of Bug22p in motile cilia.** Cilia and flagella are organelles composed of hundreds of proteins whose assembly and function, apart from main components of the backbone of the axoneme, are not well understood. Even the most abundant proteins found in high-throughput proteomics studies of cilia remain to be studied, and we focused on one of them, Bug22p.

In *Paramecium*, four genes issuing from two successive whole-genome duplications encode four identical proteins, so that only one Bug22p protein is synthesized in the cell although the nucleotide sequences had time to diverge at silent posi-
The localization experiments described here used a GFP-PtBug22p fusion in vivo, or using anti-Bug22p or anti-GFP antibodies on wild-type or GFP-PtBug22p-expressing cells in immunoelectron microscopy, confirmed the basal-body and ciliary localization of the protein in Paramecium. More precisely, the labeling showed a preferential concentration of the gold particles close to the triplet (basal bodies) or doublet (cilia) microtubules, with frequent labeling between the axoneme and the ciliary membrane. Bug22p could therefore be a protein more or less stably associated with microtubules. Its basic isoelectric point (~10) is compatible with possible interaction with tubulin, although this remains to be demonstrated biochemically.

The localization of Bug22p in the transition zone between the basal body and cilium, as well as in foci along the cilium, suggests a role in IFT. Such a view could be supported by the facts that high-throughput 2-hybrid screens revealed CLUAP1/qilin/dyf-3 as a potential interactant of Bug22p in Drosophila melanogaster (15) and that dyf-3 has been shown to belong to the IFT in Caenorhabditis elegans (12, 22). More precisely, dyf-3 is a component of the IFTB particle that is involved in the anterograde movements in cilia that control ciliary growth and turnover. However, knocking down Bug22p in Paramecium does not yield lack-of-cilia phenotypes found in IFTB knockdowns in other species, and in Paramecium, as verified by IFT172 RNAi experiments (see Fig. S4 in the supplemental material). The question of an interaction of Bug22p with dyf-3 in Paramecium, or its potential role in the IFT itself, or as IFT cargo, is still open.

The most obvious effect of Bug22p depletion is a reduction in cell swimming speed resulting from inefficient ciliary beating (Fig. 7; see Videos S1 and S2 in the supplemental material) and apparent loss of ciliary rigidity (Fig. 8). Little is known about the factors that rigidify membrane-bound microtubular complexes. Ciliary bending results from multiple interactions within the axoneme, where the nine microtubular doublets slide over neighboring doublets through dynein-tubulin interactions coordinated by the central pair and radial spokes (30). Interestingly, it has recently been shown that hydin, a central-pair-associated protein, is critical for regulation of ciliary bending during beating strokes and of beat frequency in a manner opposite to our observations for Bug22p. Indeed, hydin depletion in mutants produces excessive ciliary rigidity and reduces beat frequency (20), in contrast to Bug22p-depleted cilia, which display excessive flexibility and beating frequency.
Functional conservation of Bug22p between Paramecium and humans. Bug22p sequence conservation from Paramecium to humans suggests conservation of its biological function, tested here by functional complementation. The expression of the human HsBug22p protein can rescue the depletion of the PtBug22p protein under RNAi conditions. Moreover, a balance can be found in which the expression level of the HsBug22p protein is in a physiological range, rescuing the PtBUG22 RNAi phenotype, while being susceptible to RNAi silencing itself. This is the first example of functional complementation by a human gene in a ciliate, which opens the use of Paramecium as a model to study human proteins by comparing the rescue properties of wild-type and mutated versions associated with diseases.

Different statuses for BUG22 genes through evolution and potential roles of the proteins. The Bug22p protein is a highly conserved eukaryotic protein; however, it is not found in all eukaryotes. All species that display centrioles and cilia, be they motile or not, at some stages of their life cycle have one or several BUG22 genes. With a few exceptions, plants and fungi do not have centrioles and cilia. Fungi do possess a centro-

some-like structure, the spindle pole body, while plants have MTOC structures to organize the microtubules that govern chromosome separation at mitosis. Interestingly, only plants, not fungi, have BUG22 genes in their genomes. Examination of the aligned sequences reveals a consensus at critical positions in Bug22p proteins of species with centriolar/ciliary function (Fig. 1). Plant Bug22p proteins display another consensus at the same positions and not only a lack of the ciliary consensus, suggesting that there is selective pressure to prevent divergence from this consensus. Plants that do possess centrioles and cilia, such as Physcomitrella and Selaginella, have several Bug22p proteins encoded in their genomes that display different combinations of ciliary and plant consensus residues at the critical positions, indicating that both putative centriolar/ciliary and plant functions of Bug22p are needed in these plants.

In the present study of the motile cilia of Paramecium, we have found a striking role of Bug22p in ciliary rigidity and beating efficiency, as well as a possible role in the cortical organization of basal bodies. Bug22p proteins seem to exist in other types of cilia, sensory and primary cilia. Indeed, a BUG22...
gene with a cilium-specific X box in its promoter is found in *C. elegans* (6), which possesses only sensory cilia. Bug22p peptides have also been detected in the primary cilium-derived photosensory rod outer segments of mouse retina (21). Although the role of Bug22p in sensory and primary cilia is unknown, we can infer from its localization in the motile cilia of *Paramecium* that Bug22p plays a role associated with axonemal doublets and the ciliary membrane, potentially important for sensory and signaling functions.

The presence of *BUG22* genes in higher plants devoid of centrioles and cilia, with such sequence conservation, suggests that functions other than those associated with centrioles and cilia also exist for Bug22p. However, although higher plants have lost centrioles, they have kept microtubule-organizing centers, as well as proteins with conserved domains characteristic of centrosomes, such as TOF, LisH, and PLL motif-containing proteins (*TONNEAU1* in plants, FOP and OFD in animals) and centrin 2 (4). The prominent interactions of microtubules with the plasma membrane in these cells could be a clue to the function of Bug22p in plants, by extrapolation of what we have found in motile cilia.

Interestingly, coevolution of the centrin 2 and Bug22p proteins seems to have occurred in eukaryotes. Among the different centrin genes that exist in eukaryotes, the centrin 2 form is found, not only in species with centrioles and cilia, but also, as a derived form called centrin 2', in higher plants lacking centrioles and cilia (7). No Bug22p or centrin 2 sequences are present in fungi.

FIG. 8. Ciliary defects in Bug22p-depleted paramecia observed in electron microscopy. (A to C) Observation of RNAi after 24 h. Magnification, ×22,000. (D and E) Observations of RNAi after 48 h. Magnification, ×16,000. (A and D) Control ND7 RNAi. (B, C, and D) BUG22 RNAi. Note the unusual bending of cilia in Bug22p-depleted cells, as viewed in the longitudinal axis of cilia in panels B and E and with contiguous longitudinal and cross sections of the same cilium in panel C. Images of exaggerated ciliary bending were never observed earlier and were detected for the first time in Bug22p-depleted cells. Among 140 longitudinal sections of cilia from 15 different cells, 43 cilia displayed such exaggerated bending, whereas none were observed in a similar number of sections in the control.
devoid of centrioles and cilia, which rely on centrin 3 isoforms only for centrosomal-like function of the spindle pole body. If we call BUG22 the form found with the ciliary consensus and BUG22’ the form with the plant consensus, a good parallel thus appears between the evolution of the centrin 2 and BUG22 families. A functional relationship between Bug22 and centrin 2 proteins may exist and should now be investigated to understand the essential role of Bug22p.

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