The *Chlamydomonas reinhardtii* ODA3 Gene Encodes a Protein of the Outer Dynein Arm Docking Complex

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Abstract. We have used an insertional mutagenesis/gene tagging technique to generate new *Chlamydomonas reinhardtii* mutants that are defective in assembly of the outer dynein arm. Among 39 insertional *oda* mutants characterized, two are alleles of the previously uncloned *ODA3* gene, one is an allele of the uncloned *ODA10* gene, and one represents a novel *ODA* gene (termed *ODA12*). *ODA3* is of particular interest because it is essential for assembly of both the outer dynein arm and the outer dynein arm docking complex (*ODA-DC*) onto flagellar doublet microtubules (Takada, S., and R. Kamiya. 1994. *J. Cell Biol.* 126:737–745). Beginning with the inserted DNA as a tag, the *ODA3* gene and a full-length cDNA were cloned. The cloned gene rescues the phenotype of *oda3* mutants. The cDNA sequence predicts a novel 83.4-kD protein with extensive coiled-coil domains. The ODA-DC contains three polypeptides; direct amino acid sequencing indicates that the largest of these polypeptides corresponds to ODA3. This protein is likely to have an important role in the precise positioning of the outer dynein arms on the flagellar axoneme.

Dyneins are ubiquitous molecular motors that generate force against microtubules to produce many different types of cellular movements, including vesicle transport, localization of the Golgi apparatus, nuclear migration, spindle formation and orientation, possibly some types of chromosome movements, and beating of cilia and flagella (Holzbauer and Vallee, 1994). Both the cytoplasm (Vaisberg et al., 1996) and ciliary/flagellar axonemes (Witman et al., 1994) contain multiple isoforms of dynein, and the maintenance of many essential cell functions undoubtedly depends on the correct attachment of a specific dynein isoform to a specific cell structure. Hence, the mechanisms by which dyneins are targeted to and bind cell organelles are of considerable interest (Vallee and Sheetz, 1996).

An ideal system for studying targeting of dynein isoforms to specific attachment sites is the flagellum of the green alga *Chlamydomonas reinhardtii*, which is amenable to genetic (classical and molecular), biochemical, and physiological approaches. In *C. reinhardtii*, as in most other organisms, the flagellar dyneins make up biochemically distinct structures known as the outer dynein arms and the inner dynein arms, each of which is anchored to a specialized site on the A-tubule of the doublet microtubule (see Fig. 1). Both types of dynein arms interact transiently with the B-tubule of the apposing doublet microtubule to generate interdoublet sliding that is the basis for flagellar bending (for review see Witman, 1990). The outer dynein arms, which repeat with a 24-nm spacing along the doublet microtubules, are believed to contribute as much as four-fifths of this sliding force (Brokaw, 1994).

The outer dynein arm of *C. reinhardtii* has been extensively investigated and consists of at least 15 polypeptides, including three dynein heavy chains (DHCs: DHCα, DHCβ, DHCγ), two intermediate chains (ICs: IC69, IC78), and 10 light chains (LCs) (Huang et al., 1979; Piperno and Luck, 1979; Pfister et al., 1982; King and Witman, 1989) (see Fig. 1A). Each DHC consists of a globular head, containing one or more ATP-hydrolytic sites, and a flexible stem, which extends toward the base of the dynein (Witman et al., 1983). The two ICs are associated with each other and with several of the LCs to form an IC/LC complex (Mitchell and Rosenbaum, 1986; King et al., 1991) that is located at the base of the DHC stems (King and Witman, 1990). One component of this complex, IC78, is in direct contact with α-tubulin in the axoneme (King et al., 1991) and probably plays an important role in binding the outer dynein arm to the A-tubule of the doublet microtubule.

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It is composed of three polypeptides of that of wild-type cells (Table I). A mutation at an 11th locus encoded structural components of the outer dynein arm. (Mitchell and Kang, 1991), Witman, manuscript in preparation). Moreover, in both outer arm and cytoplasmic dynein, the ICs interact directly with the structure to which the dynein attaches (King et al., 1991, 1995; Vaughan and Vallee, 1995). As a result, the outer dynein arm of \textit{C. reinhardtii} has been a very useful model for studying how dyneins in general are targeted to specific attachment sites (Paschal et al., 1992; King et al., 1995).

Since the outer dynein arm attaches to a precisely defined site on the doublet microtubule, it is important to understand what is structurally or biochemically unique about that site. Recently, Takada and Kamiya (1994) demonstrated the existence of a factor that assembles onto the outer dynein arm binding site in the absence of arms in vivo, and that promotes functional reconstitution of outer dynein arms onto armless axonemes in vitro. This factor therefore has the properties expected for an outer dynein arm docking complex (ODA-DC). The putative ODA-DC is visible in certain outer arm–less mutants as a small projection at the site where the outer dynein arm normally would be attached to the doublet microtubule (Takada and Kamiya, 1994; compare Fig. 1 C to 1 D in this report). It is composed of three polypeptides of \textasciitilde 105, 62.5, and \textasciitilde 25 kD (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation).

Mutational loss of the outer dynein arm in \textit{C. reinhardtii} usually results in a slow, jerky swimming phenotype (Kamiya, 1988). Such cells are viable and easily detected in mutant screens, so the outer dynein arm is readily studied by genetic methods (for a recent review see Kamiya, 1995). Currently, mutations at 10 independent loci (\textit{ODA1}–\textit{ODA10})\textsuperscript{2} are known that result in loss of the outer dynein arm and jerky swimming at a rate approximately one-third that of wild-type cells (Table I). A mutation at an 11th locus (\textit{ODA11}) results in loss of part of the outer arm and a reduction in swimming speed to approximately two-thirds the wild-type rate (Sakakibara et al., 1991). Mutations at two other loci (\textit{PF13} and \textit{PF22}) result in loss of outer arms and paralyzed flagella (Huang et al., 1979); the reason for the complete loss of motility in these mutants is not yet understood. Several of the \textit{ODA} loci have been shown to encode structural components of the outer dynein arm. \textit{ODA2} encodes DHC\gamma (Wilkerson et al., 1994), \textit{ODA4} encodes DHC\beta (Sakakibara et al., 1993), \textit{ODA6} encodes IC69 (Mitchell and Kang, 1991), \textit{ODA9} encodes IC78 (Wilkerson et al., 1995), and \textit{ODA11} encodes DHC\alpha (Sakakibara et al., 1991). Therefore, with the apparent exception of \textit{ODA1} and \textit{ODA2}, defects in any one of the outer arm DHCs or ICs result in loss of the complete outer dynein arm. \textit{oda1} and \textit{oda3} lack the ODA-DC in addition to the outer dynein arm (Takada and Kamiya, 1994); \textit{ODA1} recently has been shown to encode the 62.5-kD polypeptide of the ODA-DC (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). Therefore, defects in an ODA-DC component also can lead to loss of the outer dynein arm. The bases for the loss of outer arms in the other \textit{oda} mutants and in \textit{pf13} and \textit{pf22} are not yet known.

The development of techniques for the efficient transformation of the nuclear genome in \textit{C. reinhardtii} (Kindle, 1990) now makes it possible to use insertional mutagenesis (Tam and Lefebvre, 1993; Gumpel and Purton, 1994) to study the polypeptides necessary for outer dynein arm assembly and binding to the doublet microtubule. When \textit{C. reinhardtii} is transformed, the transforming DNA usually is inserted into the genome by nonhomologous recombination, causing disruption or deletion of any gene at the site of insertion. For molecular genetic studies, this has two major benefits. First, if a gene already has been cloned, insertional mutants of that gene can be identified easily by restriction fragment length polymorphism (RFLP) analysis; this technique was used previously to identify mutants with defects in the \textit{IC78} gene (Wilkerson et al., 1995). Second, for insertional mutants defective in genes that have not yet been cloned, it is possible to use the inserted DNA as a tag to clone host DNA near the site of insertion, and then use that DNA to select wild-type genomic DNA clones containing the gene of interest. That the correct gene has been cloned can be confirmed by rescuing the mutant by transformation with the cloned wild-type DNA.

Here we report the use of these two powerful approaches to generate and identify insertional alleles for previously known but still uncharacterized \textit{ODA} loci, as well as to identify a new \textit{ODA} locus. One of the insertional mutants was defective in the \textit{ODA3} gene and hence was of particular interest because that gene is necessary for assembly of the ODA-DC (Takada and Kamiya, 1994). Start-

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|}
\hline
Mutant & Protein affected & Reference\textsuperscript{1} \\
\hline
\textit{oda1} & 62.5 kD & 1, 2 \\
\hline
\textit{oda2} (pf 28) & DHC\gamma & 1, 3, 4 \\
\hline
\textit{oda3} & 105 kD & 1, this study \\
\hline
\textit{oda4} (\textit{oda4-s7, supzf1}) & DHC\beta & 1, 5, 6, 7, 8 \\
\hline
\textit{oda5} & unknown & 1 \\
\hline
\textit{oda6} & IC69 & 1, 9 \\
\hline
\textit{oda7} & unknown & 1 \\
\hline
\textit{oda8} & unknown & 1 \\
\hline
\textit{oda9} & IC78 & 1, 10 \\
\hline
\textit{oda10} & unknown & 1 \\
\hline
\textit{oda11} & DHC\alpha & 11 \\
\hline
\textit{oda12} & unknown this study & 12 \\
\hline
pf13 & unknown & 12 \\
\hline
pf22 & unknown & 12 \\
\hline
\end{tabular}
\caption{\textit{Chlamydomonas} Outer Dynein Arm Mutants* \textsuperscript{1}}
\end{table}

\textsuperscript{1}Table updated from Kamiya (1988, 1995).

2. To conform to recommendations for a standard generic nomenclature in \textit{Chlamydomonas} (Dutcher, 1995), generic loci and the wild-type allele for those loci are denoted using uppercase italic letters, followed by a number when there is more than one locus with the same name (e.g., \textit{ODA3}); mutant alleles are designated in lowercase italic (e.g., \textit{oda3}); different mutant alleles at the same locus are distinguished by a number separated from the locus designation by a hyphen (e.g., \textit{oda3-s4}); phenotypes are indicated by an uppercase letter followed by lowercase letters and a plus or minus (e.g., \textit{Oda} –), and gene products are designated using uppercase roman letters (e.g., \textit{ODA3}).

ing with the inserted DNA as a tag, we cloned wild-type genomic DNA containing the ODA3 gene and showed that it could rescue both the new oda3 insertional mutant and a preexisting oda3 mutant. An ODA3 CDNA clone was isolated and sequenced; the sequence is predicted to encode a novel 83.4-kD protein with three long coiled-coil domains. Protein sequencing and in vitro translation experiments independently demonstrated that the ODA3 gene product is the ~105,000-M$_r$ polypeptide of the ODA-DC. Since ODA3 is predicted to have coiled-coil domains of similar length to those of the 62.5-kD ODA-DC polypeptide (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation), it is possible that these two proteins interact to form a coiled-coil structure that precisely positions outer dynein arms in the axoneme.

**Materials and Methods**

**Strains**

*Chlamydomonas reinhardtii* strains studied in this work (and their relevant genotypes) include: g1 (nit1, NIT2, agt1, mt+), and 1330.1 (nit1, NIT2, ac14, agt1, mt–), both selected for ease of transformation; B214 (nit1, NIT2, ac17, agt1, mt–) (Pazour et al., 1995); CC2229 (oda1, nlt1, nlt2, AC17, mt–); CC2321 (oda2); CC2233 (oda3-1, nlt1, nlt2, AC17, mt–); CC2235 (oda4; CC2237 (oda5); CC2239 (oda6); CC2242 (oda8); CC2245. CC2249); CC2247 (oda10); CC2073 (oda13); CCI030 (p/f1); CCI382 (p/g2); CCI24 (nlt1, nlt2); 137C; 209A (oda3-1, nlt1, NIT2) and 210A (oda3-1, nlt2, NIT2), both derived from a g1 × CC2233 cross. A *Chlamydomonas smithii* strain (CCI373) that is interfertile with *C. reinhardtii* also was used. Strains with “C” numbers and 137C are from the *Chlamydomonas Genetics Center*, Department of Botany, Duke University, Durham, NC. Insertional mutants generated by transformation of g1 and 1330.1 were assigned “V” and “F” numbers, respectively. One of these (V40) had a disruption of the ODA3 gene and was termed oda3-4 (oda3-4::pUC119-NIT1); this strain was crossed to B214 and CC124 to create strains 27D (oda3-4::pUC119-NIT1; NIT2, ac17, mt+) and 98A (oda3-4::pUC119-NIT1; nlt2), respectively.

**Growth Media**

*Chlamydomonas* strains were grown in the following media: M (Sager and Granick [1953] medium I modified to contain 0.0022 M KH$_2$PO$_4$ and 0.00171 M K$_2$HPO$_4$), R (M medium supplemented with 0.0075 M sodium acetate), M–N (M medium without nitrogen), M–N/5 (M–N medium dilituted 5× except for K$_2$HPO$_4$, which was increased 2×), M–N/5+SA (M–N/5 medium supplemented with 0.1% sodium acetate), M–N/KNO$_3$ (M–N medium supplemented with 0.0375% KNO$_3$), SGII/N03 (Sager and Granick [1953] medium II modified to contain 0.003 M KNO$_3$ as the sole nitrogen supply), and SGII/N03 (Sager and Granick [1953] medium II modified to contain 0.0065 M NaNO$_3$ as the sole nitrogen supply). DNA probes of cloned ODA genes. DNA probes used included inserts of pycy (ODA2) (Wilkerson et al., 1994), pEB1.2 (ODA4-5 end) and pBS4-2a (ODA4-3 end) (Mitchell and Brown, 1994), pGB70-16 (ODA6) (Mitchell and Kang, 1991), p78k (ODA9) (Wilkerson et al., 1995), and pBA1.3a (ODA11–5' end) and pBA4.7 (ODA11–3' end) (Mitchell and Brown, 1994).

**DNA and RNA Isolation**

*Chlamydomonas* miniprep DNA was obtained by digesting ~0.3 ml of packed cells with 0.5 ml of 1 mg/ml proteinase K (Boehringer Mannheim, Mannheim, Germany) in 5% sodium lauryl sulfate, 20 mM EDTA, and 20 mM Tris, pH 7.5, at 50°C overnight. NH$_4$ acetate was added to 1.5 M, and the DNA was extracted once with phenol/chloroform (1:1) and once with chloroform, precipitated with ethanol, and then resuspended in 1 mM EDTA and 10 mM Tris, pH 8.0 (Pazour et al., 1995). *Chlamydomonas* mRNA was obtained from wild-type (g1) cells before deflagellation (control) and 30 min after deflagellation. Deflagellation was carried out by pH shock (Witman et al., 1972) to induce transcripts of flagellar genes (Lefebvre and Rosenbaum, 1986). Total RNA was obtained and poly A+ mRNA was isolated using oligo(DT)-cellulose spin columns (mRNA Separator Kit; Clontech, Palo Alto, CA). Gel electrophoresis, Southern blotting, and Northern blotting were performed using standard procedures (Sambrook et al., 1987).

**Genetic Analysis**

Mating and tetrad analysis were performed using techniques described in Levine and Ebersold (1960) and Harris (1989). Meiotic products were separated using a glass needle held by a micromanipulator (1-axis, 36S Tilt Platform; Newport Corp., Irvine, CA). Products were scored for motility (Oda++/+ by light microscopy; for NIT1 by comparing growth on solid SGII/N03 and M; for NIT2 by comparing growth on solid SGII/N03, SGII/N03, and M; for AC17 by comparing growth on solid M and R; and for the presence of pUC119 sequences by Southern blot analysis).

**Complementation in Stable Diploids**

Stable diploids can be selected in *Chlamydomonas* by mating complementing auxotrophs and by plating on selective medium. In this study known oda mutants unable to grow on NO$_3$ as the sole nitrogen source were mated to new oda mutants that could use NO$_3$ but required acetate for growth (ac17 or ac14 mutants). Mated cells were plated on medium without acetate but with NO$_3$ as the sole nitrogen source. This selects for stable diploids and haploid progeny that have recombined to become Ntr+++. To distinguish diploids from haploids, selected lines were examined by RFLP analysis using a probe that detects mating type (Ferris and Goodenough, 1994). Haploids have one of two bands while diploids have both bands.

**Cloning and Sequencing of ODA3**

To generate a ~ phage library of genomic oda3-4 DNA, genomic DNA was partially digested with Sau3AI, and 14–20-kb fragments were isolated by sucrose gradient centrifugation. Fragments were ligated into ADASH II and packaged using Gigapack II extract (Stratagene, La Jolla, CA). To obtain sequences flanking the site of pGP505 insertion, phage clones containing pUC119 were isolated from the above library. These clones were digested with a variety of restriction enzymes, and Southern blots were probed with pGP505 to identify DNA fragments that did not contain either pUC119 or NIT1. These DNA fragments potentially represented genomic DNA closely linked to the inserted DNA. Two such fragments were obtained but neither showed an RFLP between oda3-4 and its wild-type parent (g1). To determine if they were nevertheless linked to the site of insertion, they were used as RFLP probes to score progeny of a cross between *C. reinhardtii* (oda3-4) and *C. smithii*. One fragment (termed ODA3 probe) detected a BamHI RFLP that segregated with the oda3-4 motility defect (12 parental, 0 nonparental), indicating that it was tightly linked to oda3-4. The ODA3 probe was then used to isolate wild-type genomic cosmid clones that spanned the ODA3 locus. These clones were from an unamplified cosmids library constructed from genomic DNA.

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that was size fractionated after partial digestion with Sau3A (Pazour, G.J., and G.B. Witman, unpublished data).

Rescue of oda3 was achieved by cotransforming strains 98A, 209A, and 210A with the cosmid or plasmid clone being tested and a selectable marker. Cotransformation was performed using the glass bead method as described above. Transformants were scored for Oda+ phenotype by light microscopy and, in selected cases, for outer dynein arms by EM.

To obtain DNA clones, a 1-kb HindIII fragment (see Fig. 3) was used to probe a ZAPII DNA library (Wilkerson et al., 1994). Positive DNA clones were sequenced using the 7-deaza Sequenase kit (United States Biochemical Corp., Cleveland, OH); double-stranded DNA was denatured with NaOH, while single-stranded DNA was produced using VCSM13 helper phage. Primers used for sequencing were selected using the PRIMER3 program (S. Rozen and H.J. Skaltsky, 1996. Primer3. http://www-genome.wi.mit.edu/genome_software/other/primer3.html).

Computational Analysis

The Genetics Computer Group suite of programs (Devereux et al., 1984) was used for sequence assembly and protein structure predictions; NEWCOILS (Lupas et al., 1991; Lupas, 1996a) was used to predict coiled-coil regions; and BLAST (Altschul et al., 1990) was used to search for related sequences. The ExpASY molecular biology server (http://expasy.hgcuge.cn/) was used to obtain the theoretical isoelectric point and mass of the ODA3 gene product (Bellqvist et al., 1993). The PROSITE database was used to determine possible sites for posttranslational modifications (Bairoch et al., 1995).

In Vitro Transcription and Translation

T3 RNA polymerase was used to generate synthetic RNA from XhoI-digested pk101. RNA was translated in vitro using a reticulocyte lysate system (Promega, Madison, WI) including [35S]methionine. In vitro transcription products were separated by SDS-PAGE (7.5% separating gel) and detected by autoradiography.

Isolation and Direct Protein Sequencing of the ~105-kD ODA-DC Protein

Axonemes were isolated from CC2239 (oda6) as described in Witman (1986), preextracted with 0.5 M potassium acetate, and then further extracted with 0.6 M KCl (Nakamura et al., 1997). The latter extract was subjected to 5–20% sucrose density gradient centrifugation under Mg2+-free conditions (Takada et al., 1992). The 7S fraction (Takada and Kamiya, 1994) was collected and the ~105-kD protein of the ODA-DC was separated from other proteins by electrophoresis in a 5–20% gradient polyacrylamide gel (Takada, S., C.B. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). The gel was blotted to a polyvinylidene difluoride membrane and stained with Ponceau S. The portion containing the ~105-kD protein was excised and digested with trypsin. The resultant peptides were separated by HPLC; selected peptides were then directly sequenced in an amino acid sequencer (model 477a; Applied Biosystems, Foster City, CA).

Results

Generation of oda3 Mutants by Insertional Mutagenesis

To obtain motility mutants by insertional mutagenesis, nit1 cell lines of C. reinhardtii (g1 or 1330.1) were transformed with the plasmid pGP505 (Pazour et al., 1995), which contains the cloned NIT1 gene in the pUC119 vector (Fernandez et al., 1989). Transformants were picked into liquid selective medium and examined for motility defects (see Materials and Methods). Motility mutants were obtained that exhibited a wide range of phenotypes, including slow-jerky swimmers, slow-smooth swimmers, unflagellate cells, aflagellate cells, cells with paralyzed flagella, and cells with long flagella. The slow-jerky swimmers were of particular interest for this project, as this phenotype (termed Oda−) is indicative of a defect in the outer dynein arm (Kamiya, 1991).

Insertional mutagenesis in C. reinhardtii occurs via non-homologous recombination and usually results in deletions, insertions, and/or rearrangements (Tam and Lefebvre, 1993), all of which should be detectable by RFLP analysis using appropriate probes. Therefore, to determine if any of the insertional Oda− mutants had defects in genes encoding the previously cloned outer dynein arm heavy or intermediate chains, Southern blots of the Oda− mutants were probed with DNA clones encoding portions of DHCγ (ODA2), DHCβ (ODA4), IC69 (ODA6), IC78 (ODA9), and DHCα (ODA11). The results indicated that 12 of the Oda− mutants were defective in ODA4, while 15 were defective in ODA9 (data not shown; see, e.g., Fig. 8 in Wilkerson et al., 1995).

The remaining Oda− mutants did not exhibit RFLPs when probed with DNA encoding the outer dynein arm heavy and intermediate chains. These mutants therefore are potentially defective in uncloned genes necessary for outer arm assembly or function. To determine if these mutants were actually missing their outer arms, they were examined by EM. The vast majority had specific defects in this structure. Most completely lacked outer arms (Fig. 1, C and D) and therefore are “complete” oda mutants. In seven cell lines, the arms were missing on only some of the doublet microtubules (data not shown); these are termed “partial” oda mutants. A few cell lines had apparently intact outer arms and are not currently considered to be oda mutants.

Taken together, the data from the RFLP analysis and EM indicated that 18 oda mutants (five of which were partial) were obtained from 2,978 transformants examined in one transformation experiment (V series), and 21 oda mutants (two partial) were obtained from 6,903 transformants examined in another transformation experiment (F series) (Table II).

Identification of an oda3 Insertional Mutant

The five complete oda insertional mutants that did not show an RFLP when probed with DNA encoding cloned outer dynein arm heavy or intermediate chain genes were tested to determine if they would complement those existing oda and related pf mutants whose defective gene products are unknown. Stable diploids were generated by mating haploid insertional mutant cell lines with oda1, oda3, oda5, oda7, oda8, and oda10 cell lines; F series mutants also were mated to pf13 and pf22. RFLP analysis using a probe that is closely linked to the mating-type locus (Ferris and Goodenough, 1994) confirmed that colonies selected for further analysis were true stable diploids containing both mating-type loci (e.g., Fig. 2 A). One cell line (V40) did not complement the motility defect of oda3 (Fig. 2 A). Complementation studies with another cell line (F60) were not complete when the ODA3 probe became available (see below); RFLP analysis using this probe revealed that F60 had a defect in the ODA3 gene (results not shown). Thus, V40 and F60 represent new alleles of ODA3; since there already were three mutant alleles of ODA3 (Kamiya, 1988), V40 and F60 have been named oda3-4 and oda3-5, respectively. A third mutant (V87.2) did not complement oda10 and therefore appears to be a new ODA10 allele.

The remaining two complete oda insertional mutants
dynein arms are putative ODA-DCs (al., 1995). Tiny projections at sites normally occupied by outer oda 9 insertional mutant lacking outer dynein arms (Wilkerson et al., 1995). (B–E) Electron micrographs of cross-sections of C. reinhardtii flagella. (B) Wild-type (g1). Note outer dynein arms (arrows) and inner dynein arms (arrowheads). The arms occupy precise positions on the A-tubule of the doublet microtubule. (C) oda9 insertional mutant lacking outer dynein arms (Wilkerson et al., 1995). Tiny projections at sites normally occupied by outer dynein arms are putative ODA-DCs (arrow). (D) oda3-4 insertional mutant (V40) lacking outer dynein arms. The profiles of the doublet microtubules at the sites normally occupied by the outer dynein arms are rounder than in oda9, indicating that this mutant lacks the ODA-DC (Takada and Kamiya, 1994). (E) oda3-4 insertional mutant rescued by transformation with the cosmid pK001. The outer dynein arms are completely restored.

complemented all of the uncharacterized preexisting oda cell lines, suggesting that these insertional mutants represented novel oda genes. However, it also was possible that the RFLP analysis failed to reveal defects in outer dynein arm heavy and intermediate chain genes because the defect is caused by a small disruption not detectable by RFLP analysis. To distinguish between these possibilities, these two mutants were tested further to determine if they would complement oda2, oda4, oda6, oda9, and oda11 in stable diploids. One of the insertional mutants (F56) complemented all of these strains, indicating that it represents a new ODA gene; the mutant allele has been named oda12. The other mutant complemented all alleles tested except for oda4, indicating that it is an ODA4 allele that had been missed by the RFLP analysis. The seven partial oda mutants have not yet been completely characterized by complementation analysis; some of these may represent mutant alleles of additional new ODA genes.

Defects in ODA3 result in loss of both the outer dynein arm and the ODA-DC (Takada and Kamiya, 1994; see Fig. 1, C and D). Further work therefore concentrated on the oda3-4 insertional mutant, as it provided an opportunity to learn more about a component necessary for assembly of the ODA-DC.

oda3-4 Is Tagged with pUC119

An advantage of insertional mutagenesis is that the mutated gene frequently is tagged with the inserted DNA, facilitating the cloning of the disrupted gene. To determine if the oda3-4 allele is tagged, the mutant was crossed to a wild-type cell line (B214), and meiotic progeny were scored for motility (Oda+/−), ability to grow on medium containing nitrate as the sole nitrogen source (Nit+/−), and the presence of pUC119 sequences in the genome (Fig. 2 B). In all cases, the Oda− phenotype segregated with both the Nit+ phenotype and pUC119, indicating that the oda3-4 allele is tagged with both Nit1 and pUC119.

Cloning of the ODA3 Gene and Rescue of oda3 Mutants

To obtain DNA at or near the ODA3 locus, a phage library was constructed using genomic oda3-4 DNA, and clones containing pUC119 sequences were selected. Using RFLP analysis (see Materials and Methods), a DNA frag-

<table>
<thead>
<tr>
<th>Locus</th>
<th>V series*</th>
<th>F series†</th>
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<tbody>
<tr>
<td>ODA1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>ODA2</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>ODA3</td>
<td>1 (V40)</td>
<td>1 (F60)</td>
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<td>Unknown†</td>
<td>5</td>
<td>2</td>
</tr>
<tr>
<td>Total</td>
<td>18</td>
<td>21</td>
</tr>
</tbody>
</table>

*From 2,978 Nit + transformants.
†From 6,902 Nit + transformants.
§Seven insertional mutants, all partial odas, have yet to be categorized. It has not been determined whether any of these are alleles of oda5, oda7, oda8, and oda12; question marks have therefore been placed at appropriate positions in the table.
of the insertional mutant V40, was mated to
mt
oda
1
stable diploids demonstrates that V40 is an
Figure 2.
mt
oda
1
oda
2
mutants to generate stable diploids D93 and D98,
probed in Southern blots with a 1.6-kb XbaI fragment of
from the indicated cell lines was digested with BamHI and
cell lines D93 and D98 were true stable diploids, genomic DNA
QK7 that is tightly linked to the
oda
3-4, indicating that the
oda
3-4. The insertional mutant
oda
3-4 (V40) (oda
−
and Nit
−
phenotypes) was crossed to B214
(oda
+ and Nit
−
phenotypes). Progeny were scored for motility
(oda
+/
−
) and ability to grow on nitrate as sole nitrogen source
oda
3-4 and a
oda
3-4, indicating that these cell lines contain both mating
ypes, indicat-
oda
3-4, and from the untransformed parental cell line (g1).
The DNA was digested with Pst1 and probed in Southern blots with
pUC119, which contains only the plasmid sequence used for
transformation. The pUC119 sequence (3.5-kb fragment) is
present in
oda
3-4 but not g1; in the progeny, the plasmid se-
genomic DNA was isolated from the progeny, from
oda
3-4 (V40), and from the untransformed parental cell line (g1).
DNA from products of a cross between
oda
3-4 and a rescued strain (T278) showed that the pK001 DNA segre-
gated with the
oda
+ phenotype (Fig. 2 C), confirming that the rescued phenotype was linked to the inserted
cosmid.

To locate the
oda
3 gene on pK001, the cosmid was di-
gested with several restriction enzymes and the resultant
fragments were inserted into plasmid vectors (Fig. 3). These subclones were then assayed in cotransformation experiments (with
NIT1 as the selectable marker) for the ability to rescue the motility phenotype of strains 209A and 210A, which contain the original
oda
3 allele (Kamiya, 1988) in a
nit1, NIT2 background. The smallest frag-
ment that was capable of rescue (pK144; Fig. 3) was ~10 kb long. EM of several of the rescued transformants confir-
mated that the outer dynein arms were restored (Fig. 1 E), indicat-
ing that pK001 contained the
oda
3 gene. RFLP analysis of DNA from products of a cross between
oda
3-4 and a rescued strain (T278) showed that the pK001 DNA segre-
gated with the
oda
+ phenotype (Fig. 2 C), confirming that the rescued phenotype was linked to the inserted
cosmid.

Isolation of
oda
3 cDNA

A 1-kb HindIII fragment (Fig. 3) that spans the central
Xhol site in pK144 was used to probe Northern blots of total
C. reinhardtii mRNA isolated from wild-type cells that
were not deflagellated, and from cells that had been de-
flagellated and allowed to regenerate their flagella for 30
min to induce mRNAs encoding flagellar proteins. The probe hybridized with an ~3.1-kb mRNA that was
induced after deflagellation (Fig. 4 A). As the
oda
3 mRNA was expected to be one of a relatively small subset of transcripts induced during flagellar regeneration, this further
suggested that the 1-kb HindIII fragment encoded part of the
oda
3 gene. The 1-kb HindIII fragment therefore was
used to probe
C. reinhardtii cDNA libraries, yielding
cDNA clones of ~3.0 and ~2.4 kb. Sequence analysis indi-
cated that these clones were the same at their 3’ ends.

oda
3 cDNA Sequence

The ~3.0-kb clone was sequenced in both directions and
found to contain a 1,797-bp open reading frame. The puta-
tive initiation codon was preceded by two in-frame stop
codons located 45 and 99 nucleotides upstream (Fig. 5). The open reading frame ended with an amber stop codon

Figure 2. Genetic analysis of
oda
3. (A) Complementation in
stable diploids demonstrates that V40 is an
oda
3 allele (referred to as
oda
3-4). Cell line 27D, a plus mating type
(mt+)
derivative of the insertional mutant V40, was mated to
oda
1 (mt−) and
oda
3-1 (mt−) mutants to generate stable diploids D93 and D98,
respectively. Complementation (oda
+ phenotype) was observed in D93 but not in D98, indicating that the
oda
1 gene is functional in V40, while the
oda
3 gene is defective. To confirm that
cell lines D93 and D98 were true stable diploids, genomic DNA
from the indicated cell lines was digested with BamHI and
probed in Southern blots with a 1.6-kb XbaI fragment of
phage
QK7 that is tightly linked to the
mt locus. mt+
and mt− correlate
with hybridization to ~9- and ~5-kb fragments, respectively, when
using this probe. Hybridization to both ~9- and ~5-kb fragments in D93 and D98 indicates that these cell lines contain both mating
types and thus are stable diploids. (B) Linkage between inserted
DNA and
oda
− phenotype in
oda
3-4. The insertional mutant
oda
3-4 (V40) (oda
−
and Nit
+ phenotypes) was crossed to B214
(oda
+ and Nit
−
phenotypes). Progeny were scored for motility
(oda
+/
−
) and ability to grow on nitrate as sole nitrogen source
(Nit
+/
−
). Genomic DNA was isolated from the progeny, from
oda
3-4 (V40), and from the untransformed parental cell line (g1).
The DNA was digested with Pst1 and probed in Southern blots with
pUC119, which contains only the plasmid sequence used for
transformation. The pUC119 sequence (3.5-kb fragment) is
present in
oda
3-4 but not g1; in the progeny, the plasmid se-
genomic DNA was isolated from the progeny, from
oda
3-4 (V40), and from the untransformed parental cell line (g1).
The DNA was digested with Pst1 and probed in Southern blots with
pUC119, which contains only the plasmid sequence used for
transformation. The pUC119 sequence (3.5-kb fragment) is
present in
oda
3-4 but not g1; in the progeny, the plasmid se-
sequence segregates with the
oda
− and Nit
+ phenotypes, indicating that the
oda
− phenotype is due to the inserted DNA. Data are shown for four products (274–27D, bracketed) from one tet-
rad and individual products (28.1–33.1) from five other tetrads.
(C) Linkage between rescuing DNA (pK001) and restored
oda
+ phenotype. Cell line 98A (a
nit2
derivative of
oda
3-4) was co-
transformed with pMN68 (containing
NIT2) and the cosmid pK001. One of the rescued transformants, T278, was crossed to
oda
3-4, and progeny were scored for motility
(oda
+/
−
). Genomic DNA from
98A, T278, and the progeny was digested with PvuII and
probed in Southern blots with the
oda
+ probe, which flanked the
original insertion in
oda
3-4 and is contained in pK001. The probe hybridizes to an ~3.2-kb endogenous fragment, as well as to
new fragments of ~3.2 and ~1.8 kb in T278. The new bands
cosegregate with the
oda
+ phenotype in the progeny from the
cross of T278 and
oda
3-4, indicating that the
oda
+ phenotype is linked to stably inserted pK001 DNA. Data are shown for 14 mei-
osis of DNA from products of a cross between
oda
3-4 and a rescued strain (T278) showed that the pK001 DNA segre-
gated with the
oda
+ phenotype (Fig. 2 C), confirming that the rescued phenotype was linked to the inserted
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Isolation of
oda
3 cDNA

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flagellated and allowed to regenerate their flagella for 30
min to induce mRNAs encoding flagellar proteins. The probe hybridized with an ~3.1-kb mRNA that was
induced after deflagellation (Fig. 4 A). As the
oda
3 mRNA was expected to be one of a relatively small subset of transcripts induced during flagellar regeneration, this further
suggested that the 1-kb HindIII fragment encoded part of the
oda
3 gene. The 1-kb HindIII fragment therefore was
used to probe
C. reinhardtii cDNA libraries, yielding
cDNA clones of ~3.0 and ~2.4 kb. Sequence analysis indi-
cated that these clones were the same at their 3’ ends.

oda
3 cDNA Sequence

The ~3.0-kb clone was sequenced in both directions and
found to contain a 1,797-bp open reading frame. The puta-
tive initiation codon was preceded by two in-frame stop
codons located 45 and 99 nucleotides upstream (Fig. 5). The open reading frame ended with an amber stop codon

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The predicted amino acid sequence was compared to the nonredundant PDB + Swiss-Prot + SPupdate + PIR + GenPept + Genupdate databases (13.6%; the average in the NCBI database is 6.3%).

The nucleotide sequence of the ODA3 cDNA predicts a protein of 749 amino acids (Fig. 5) with a mass of 83,376 daltons and an isoelectric point of 5.72. Based on secondary structure predicted using the NEWCOILS program (Lupas et al., 1991; Lupas, 1996a), the ODA3 gene product can be divided into three domains (Fig. 6): an NH2-terminal domain (residues 1–85), a central domain (residues 86–462), and a COOH-terminal domain (residues 463–749).

The central domain contains four predicted coiled-coil regions (A, B, C, and D) that are interrupted by three short non-coil regions (Fig. 6). Regions A, B, and D contain a total of 226 (120+/70+/36) amino acid residues and are very likely to form coiled-coils (probability ~1), whereas region C (28 amino acids) is slightly less likely to form a coiled-coil (probability ~0.67). An imperfect 11–amino acid tandem repeat [\textsuperscript{560}KOLERKRMRE\textsubscript{569} and \textsuperscript{360}KOLERKQRE\textsubscript{369}] is located at the predicted junction of the COOH end of region B and the adjacent non-coil region. The COOH-terminal portion of the molecule contains both a highly basic region (\textsuperscript{560}KRKKGGK\textsubscript{567}) and a highly acidic region (\textsuperscript{360}DVEEEPEEEE\textsubscript{371}). Overall, the protein has an unusually high percentage of glutamic acid residues (13.6%; the average in the NCBI database is 6.3%).

The predicted amino acid sequence was compared to the protein sequences in the nonredundant PDB + Swiss-Prot + SPupdate + PIR + GenPept + Genupdate database.

Figure 3. Restriction maps of cosmids pK001 insert and plasmid inserts, and ability of the inserts to rescue motility in oda3. A 1-kb HindIII fragment was used to isolate cDNA clones. (Asterisk) A plasmid (pK143) was constructed that contained an insert with an inverted 6-kb XhoI fragment (arrow pointing right). In cotransformation experiments using pK143, one cell line was generated in which the rate of swimming was restored to near wild-type levels. EM indicated that outer dynein arms also were restored. However, in crosses to g1, mixed motility phenotypes were recovered, and the results could not be interpreted in terms of Mendelian segregation of a single gene (data not shown). Further study will be necessary to understand the basis for apparent rescue in this transformant.

Figure 4. Expression of ODA3. (A) The ODA3 gene is induced after deflagellation. Northern blot of C. reinhardtii wild-type (g1) mRNA isolated from nondelflagellated cells (Control) and from cells 30 min after deflagellation (Deflag). The blot was probed on two different occasions and the resultant autoradiographs were superimposed. The 3.1-kb band represents mRNA hybridizing to the 1-kb HindIII fragment of pK001 (Fig. 3), which contains part of the ODA3 sequence; this mRNA is strongly induced in the deflagellated cells. The 1.5-kb band represents mRNA hybridizing to a PTX2 cDNA clone (pG557 insert, provided by G.J. Pazour); PTX2 is not induced by deflagellation (Pazour, G.J., and G.B. Witman, manuscript in preparation) and is used here to demonstrate that similar amounts of mRNA were loaded into each lane. (B) The ODA3 gene occurs once in the C. reinhardtii genome. Southern blot of C. reinhardtii wild-type (g1) genomic DNA digested with five different restriction enzymes and probed with a full-length ODA3 cDNA clone (pK101 insert). A single band was detected in each lane. (C) ODA3 produced by translation in vitro migrates as an ~105,000-M<sub>r</sub> protein. Synthetic mRNA was prepared from ODA3 cDNA (pK101) and translated in a rabbit reticulocyte lysate system. An autoradiograph of an SDS-polyacrylamide gel of the product is shown. M<sub>r</sub> standards were β-galactosidase (116), phosphatase b (97), and BSA (66).
base using the BLAST program (Altschul et al., 1990). The hypothetical coiled-coil regions of the ODA3 gene product were similar to the coiled-coil regions of other proteins, but no convincing similarities were found when non-coiled-coil regions were used to search the database. ODA3 therefore represents a novel protein. Interestingly, a BLAST search using only the tandem repeat of ODA3 revealed that the motif is very similar to the extensive repeats present in the mammalian intermediate filament-associated protein trichohyalin (Fietz et al., 1993):

**ODA3 repeat**

<table>
<thead>
<tr>
<th>KQLERERQERE</th>
<th>ovine trichohyalin</th>
</tr>
</thead>
<tbody>
<tr>
<td>RKFREEEQLQRERQERE</td>
<td>VRRKREEQERE/</td>
</tr>
</tbody>
</table>

The ODA3 Gene Encodes a Component of the ODA-DC

*oda3* mutants lack the ODA-DC (Takada and Kamiya, 1994; Fig. 1 D), which contains polypeptides of ~105, 62.5, and ~25 kD (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). A cDNA encoding the 62.5-kD protein recently has been cloned and shown to correspond to the ODA1 gene (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). To investigate the possibility that the ODA3 gene product corresponds to the ~105-kD protein, the latter was partially purified and four tryptic fragments were sequenced directly (see Materials and Methods). Three of these (VEAMEHSWNR, TISGADTPEEVLAYWEG, and LMIALEEIHPDQLR) matched the ODA3 sequence exactly (Fig. 5, underlines) (the fourth fragment did not match and presumably was from a contaminant). This provides direct evidence that the ODA3 gene product is the largest component of the ODA-DC. The size of this protein previously was estimated to ~105 kD by SDS-PAGE (Takada and Kamiya, 1994). To understand the discrepancy between this value and the mass of 83.4 kD predicted from the sequence, we produced synthetic mRNA from pK101 and translated it in an in vitro reticulocyte lysate translation system. The major translational product migrated with an apparent mass of ~105 kD (Fig. 4 C). Therefore, the ODA3 gene product either runs anomalously in SDS-PAGE, or undergoes significant posttranslational modifications that also occur in the reticulocyte lysate system (see Discussion).

**Discussion**

Utility of Insertional Mutagenesis in Chlamydomonas

*oda1-oda11* mutants lack the ODA-DC (Takada and Kamiya, 1994; Fig. 1 D), which contains polypeptides of ~105, 62.5, and ~25 kD (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). A cDNA encoding the 62.5-kD protein recently has been cloned and shown to correspond to the ODA1 gene (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). To investigate the possibility that the ODA3 gene product corresponds to the ~105-kD protein, the latter was partially purified and four tryptic fragments were sequenced directly (see Materials and Methods). Three of these (VEAMEHSWNR, TISGADTPEEVLAYWEG, and LMIALEEIHPDQLR) matched the ODA3 sequence exactly (Fig. 5, underlines) (the fourth fragment did not match and presumably was from a contaminant). This provides direct evidence that the ODA3 gene product is the largest component of the ODA-DC. The size of this protein previously was estimated to be ~105 kD by SDS-PAGE (Takada and Kamiya, 1994). To understand the discrepancy between this value and the mass of 83.4 kD predicted from the sequence, we produced synthetic mRNA from pK101 and translated it in an in vitro reticulocyte lysate translation system. The major translational product migrated with an apparent mass of ~105 kD (Fig. 4 C). Therefore, the ODA3 gene product either runs anomalously in SDS-PAGE, or undergoes significant posttranslational modifications that also occur in the reticulocyte lysate system (see Discussion).
tagenesis (Tam and Lefebvre, 1993; Gumpel and Purton, 1994; Pazour et al., 1995), we have begun to identify insertional mutants defective in ODA genes that are still uncloned. Since the defective genes in these insertional mutants are likely to be tagged by exogenous DNA, it should be relatively straightforward to clone and characterize the genes that are affected.

In two separate transformations, 39 oda mutants were identified in a total of 9,881 transformants screened. 32 of these represent “complete” oda mutants that totally lack outer dynein arms as assayed by thin-section EM, whereas seven are “partial” oda mutants that have decreased numbers of outer arms. By combining RFLP analysis and complementation studies in stable diploids, we determined that 13 of the 32 complete oda mutants are alleles of ODA4 (cloned), 15 are alleles of ODA9 (cloned), two are alleles of ODA3 (cloning reported here), one is an allele of ODA10 (uncloned), and one represents an allele of a novel ODA gene (assigned oda12, uncloned) (Table II). The seven partial oda mutants have yet to be completely characterized and may represent additional novel ODA genes.

Mutant alleles of the cloned ODA11 gene were not found in this study. This result was expected as oda11 mutants swim much faster than the other oda mutants (Sakakibara et al., 1991), and therefore probably would not have been picked in our screen for slow, jerky swimming mutants. More surprising is that we did not find any mutant alleles of ODA1, ODA2, ODA5, ODA6, ODA7, or ODA8, as mutations at these loci cause loss of the outer arm and slow, jerky swimming. In contrast, 28 of the 39 oda insertional mutants represented alleles of ODA4 and ODA9. Taken together, these results indicate that the insertion of DNA into the C. reinhardtii genome in our transformations was not a random event, and that there may be hot spots for the incorporation of foreign DNA. On the other hand, Mitchell and colleagues, using a similar mutagenesis and screening procedure, found alleles of ODA1, ODA2, ODA3, ODA4, ODA8, and ODA9 among 30 new Oda–insertional mutants (Mitchell, D., personal communication). Therefore, the distribution of insertionally mutated sites may depend on some still undetermined factor such as the cell line used for transformation, the state of the cells during transformation, or the presence in the transforming DNA of unidentified sequence that is homologous to sequence at the apparent hot spots.

In addition to the oda mutants, insertional mutants with a number of other motility phenotypes were obtained using our screening procedure. These mutants should be very useful for the identification and characterization of the genes involved in processes such as inner dynein arm assembly (slow smooth swimming mutants [Kamiya et al., 1991]), flagellar length control (long flagella mutants), and basal body maturation (unflagellated mutants). The breadth of phenotypes obtained confirms the utility of insertional mutagenesis for studying motility phenotypes in C. reinhardtii (Tam and Lefebvre, 1993; Pazour et al., 1995; Smith and Lefebvre, 1996).

The ODA3 Gene Encodes the ~105-kD Protein of the ODA-DC

Beginning with the pUC119 sequence inserted into oda3-4 as a gene tag, we isolated a wild-type genomic cosmid clone and from that a 10-kb wild-type genomic fragment that could rescue the Oda phenotype of both the oda3-4 insertional mutant and the original oda3-1 mutant. A 1-kb subfragment from a part of the 10-kb fragment that appeared to contain the ODA3 gene hybridized in Northern blots to a 3.1-kb mRNA that was induced by deflagellation. The 1-kb fragment was used to isolate ODA3 cDNA clones. Sequence analysis of the cDNA clones revealed that the ODA3 gene encodes a 749-amino acid protein with a predicted molecular mass of 83.4 kD. Southern blot analysis indicated that the C. reinhardtii genome contains a single copy of the ODA3 gene.

oda1 and oda3 mutants differ from oda2, oda4, oda5, oda6, and oda9 mutants in that they lack the ODA-DC in addition to the outer dynein arms (Takada and Kamiya, 1994; Fig. 1, C and D). The ODA-DC is composed of proteins of ~105 kD, 62.5 kD and ~25 kD (Takada and Kamiya, 1994; Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). During the course of this work, a full-length cDNA encoding the 62.5-kD protein was cloned and found to correspond to the ODA1 gene (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). This raised the possibility that the ODA3 gene might encode the ~105-kD protein, notwithstanding the difference in size between the predicted mass of the ODA3 gene product and the apparent mass of the ~105-kD protein as estimated by SDS-PAGE. To investigate this possibility, the ~105-kD protein was isolated and four tryptic fragments were sequenced. The amino acid sequence of three out of four of the fragments exactly matched sequence predicted for the ODA3 gene product. Therefore, ODA3 does encode the largest polypeptide of the ODA-DC.

To understand the difference between the predicted and estimated masses for the ODA3 gene product, we generated synthetic mRNA from an ODA3 cDNA clone and translated it in vitro in a reticulocyte lysate system. The
major translational product migrated in SDS-PAGE with an \( M_r \) of \( \sim105,000 \). This provided additional evidence that the cDNA clone was full length, and indicated that the protein either runs anomalously in SDS-PAGE, or undergoes significant posttranslational modification that also can occur in the in vitro translation system. A ProSite search (Bairoch et al., 1995) revealed that the ODA3 gene product contains 35 possible phosphorylation sites, which is in agreement with the observation that the protein is heavily phosphorylated in vivo (King, S.M., personal communication). Phosphorylation can occur in reticulocyte lysates (Hiremath et al., 1989; Schubert et al., 1994), and the presence of phosphate groups can cause a shift in the relative mobility of a protein in SDS-PAGE far in excess of the added mass (Tang et al., 1993). However, in preliminary experiments, treatment of the in vitro translated protein with alkaline phosphatase caused its relative mobility to shift only slightly. Therefore, the basis for the unexpectedly slow mobility of the ODA3 gene product remains unknown. A similar behavior has been reported for another microtubule-associated protein, E-MAP-115, which has a predicted mass of 84,051 daltons but migrates with an \( M_r = 115,000 \) in SDS-PAGE (Masson and Kreis, 1993).

### Coiled-Coil Domains

The secondary structure of the ODA3 gene product is predicted to contain regions of extended \( \alpha \) helix with a strong propensity to form coiled-coil structures. Coiled-coils have a variety of functions in proteins (Adamson et al., 1993; Lupas, 1996b). Most commonly, they mediate homodimer and heterodimer formation. Therefore, it may be relevant that the ODA1 gene product, with which the ODA3 gene product is associated in vivo, also is predicted to contain extensive \( \alpha \)-helical domains that have a very high probability of forming a coiled-coil. ODA3 contains three regions (\( \sim120, \sim70, \) and \( \sim36 \) residues) with a total of \( \sim226 \) residues that are highly likely to form coiled-coil structures; similarly, ODA1 has three regions (\( \sim96, \sim48, \) and \( \sim34 \) residues) with a total of \( \sim178 \) residues that are strongly predicted to form coiled-coils (Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation). Thus, these two proteins may interact via one or more coiled-coils to form a heterodimer. In this case, the dynein docking function of the ODA-DC might be fulfilled if the residues on one side of the exterior of the coiled-coil interacted with tubulin or some other protein on the doublet microtubule, while residues on the other side of the coiled-coil interacted with one or more outer dynein arm polypeptides, much as the coiled-coil of vertebrate striated muscle tropomyosin interacts with actin and troponin (Phillips et al., 1986).

It also is possible that the coiled-coil domains of the ODA-DC serve as a molecular ruler, either longitudinally along or circumferentially around the doublet microtubule. Although the ODA-DC is important for outer dynein arm binding to the A-tubule, it is unlikely that it is solely responsible for establishing the 24-nm repeat of the arms, because purified sea urchin sperm outer arm dynein has been shown to assemble onto brain microtubules with a 24-nm periodicity in the apparent absence of the ODA-DC (Moss et al., 1992). However, inasmuch as the ODA-DC can assemble onto the doublet microtubule in the absence of the outer dynein arm (Takada and Kamiya, 1994), it may establish its own periodicity, perhaps by binding end-to-end along the microtubule. The length of a coiled-coil region can be estimated by assuming 1.5 Å per residue (Fraser and MacRae, 1973). If all residues in the extended \( \alpha \)-helical regions of ODA1 formed a heterodimeric coiled-coil with an equivalent number of residues in ODA3, this would form a structure \( \sim26.7 \) nm long. Interestingly, if only regions A (120 residues) and D (36 residues) of ODA3 formed a coiled-coil with ODA1, the resulting structure would be \( \sim23.4 \) nm long, almost exactly the same as the outer dynein arm repeat. Alternatively, the ODA-DC, or portions of it, may interact with inner dynein arm polypeptides, or with some other topographical feature of the doublet microtubule, to place the outer dynein arm the correct distance from, and in the correct phase relative to, the inner dynein arm. In either case, the nonhelical portions between coiled-coil regions probably would form discontinuities that extended from the coil, as has been proposed for intermediate filament proteins (Steinert et al., 1983). These and other non–coiled-coil regions could form linkages to the 25-kD ODA-DC polypeptide or to proteins outside of the ODA-DC (see below).

### Charged Domains

The central portion of ODA3 contains two short, highly charged repeats that are nearly identical to a portion of a sequence repeated many times in mammalian trichohyalin, an intermediate filament–associated protein of the hair follicle. In trichohyalin, the repeats are predicted to form an elongated, single-stranded, \( \alpha \)-helical rod structure (Fietz et al., 1993). Trichohyalin is a substrate for transglutaminase, which cross-links glutamine and lysine residues, and for peptidylarginine deiminase, which converts arginine to citrulline; it has been proposed that the repeats provide an ordered array of glutamine, lysine, and arginine for catalysis by these enzymes. Further investigation will be necessary to determine if the repeats of ODA3 are similarly modified posttranslationally. In any case, the presence of these repeats in *Chlamydomonas* raises the possibility that the motif arose early in evolution and is now widespread.

The COOH-terminal end of ODA3 contains one region that is highly basic (\( \sim70 \) DVEEEEPESEEETEEEE716); these regions may be important for interactions between the ODA-DC and the proteins to which it binds. The outer dynein arm together with the ODA-DC are released from the axoneme by treatment with high salt (0.6 M KCl) (Takada and Kamiya, 1994; Takada, S., C.G. Wilkerson, R. Kamiya, and G.B. Witman, manuscript in preparation), indicating that the interaction between the axonemal microtubule and the outer dynein arm/ODA-DC is ionic in nature. Both \( \alpha \) and \( \beta \) tubulins of *C. reinhardtii* have highly acidic COOH termini (Silflow et al., 1985; Younghblom et al., 1984), which may interact with the basic domain of ODA3 to create the salt-sensitive bond. The COOH-terminal region of tubulin is exposed on the surface of the microtubule and has been implicated in binding of other microtu-
bule-associated proteins (Littauer et al., 1986; Paschal et al., 1989).

The outer dynein arm polypeptides that bind to the ODA-DC have not yet been identified, but IC78, which is located at the base of the dynein (King and Witman, 1990), has both acidic and basic domains (King et al., 1995). These could interact with oppositely charged domains on ODA3 to anchor the outer arm to the ODA-DC. The availability of the ODA3 gene and sequence will greatly facilitate identification of the polypeptides that are in direct contact with ODA3, and will permit analysis of the functional domains responsible for the interactions between ODA3 and its binding partners.

**The ODA-DC as a Paradigm for Dynein Targeting**

The results presented here provide definitive evidence that loss of ODA3 results in loss of the ODA-DC, with concomitant failure of the outer dynein arm to assemble onto the doublet microtubule. Therefore, ODA3 is essential for assembly of the ODA-DC, and the ODA-DC is necessary for attachment of the outer dynein arm to the doublet microtubule. These findings suggest that the ODA-DC is responsible for the specific targeting of the outer dynein arm to its correct binding site on the flagellar axoneme. Inasmuch as the inner dynein arms also are targeted to specific binding sites on the doublet microtubules (Smith and Sale, 1992), it is likely that inner dynein arm docking proteins also exist. The insertional mutagenesis technique used here should be helpful in identifying and characterizing such polypeptides. Indeed, because slow, smooth swimming is characteristic of inner armless mutants (Kamiya et al., 1991), some of the slow-smooth swimming insertional mutants that we have isolated may be defective in the protein(s) responsible for inner arm docking.

It has been proposed that the ~20S dynactin complex (Gill et al., 1991) may anchor cytoplasmic dynein to the kinetochore and various membranous structures (Vallee and Sheetz, 1996). If so, dynactin and the ODA-DC would have analogous roles in targeting of cytoplasmic dynein and outer arm dynein to their respective organelles. Although the sequences of all the polypeptides of both complexes have not been reported, the data to date do not reveal any obvious homology between dynactin and the ODA-DC. Therefore, despite their close evolutionary relationship, these two dyneins appear to bind to distinctly different docking complexes. This may turn out to be the most biologically significant difference between these dyneins. On the other hand, cytoplasmic dynein interacts with several different cell organelles at different times in the cell cycle, and it is possible that a cytoplasmic homolog of the ODA-DC will yet be found to mediate binding of cytoplasmic dynein to a specific subset of cellular sites.

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