

The protist, *Monosiga brevicollis*, has a tyrosine kinase signaling network more elaborate and diverse than found in any known metazoan

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Tyrosine kinase signaling has long been considered a hallmark of intercellular communication, unique to multicellular animals. Our genomic analysis of the unicellular choanoflagellate *Monosiga brevicollis* discovers a remarkable count of 128 tyrosine kinases, 38 tyrosine phosphatases, and 123 phosphotyrosine (pTyr)-binding SH2 proteins, all higher counts than seen in any metazoan. This elaborate signaling network shows little orthology to metazoan counterparts yet displays many innovations reminiscent of metazoans. These include extracellular domains structurally related to those of metazoan receptor kinases, alternative methods for membrane anchoring and phosphotyrosine interaction in cytoplasmic kinases, and domain combinations that link kinases to small GTPase signaling and transcription. These proteins also display a wealth of combinations of known signaling domains. This uniquely divergent and elaborate signaling network illuminates the early evolution of pTyr signaling, explores innovative ways to traverse the cellular signaling circuitry, and shows extensive convergent evolution, highlighting pervasive constraints on pTyr signaling.

choanoflagellate | evolution | genome | kinome | phosphotyrosine

Choanoflagellates such as *Monosiga brevicollis* are unicellular aquatic protists and the closest known relatives of multicellular animals (metazoans). The sequencing of the *Monosiga* genome now provides a key evolutionary node between metazoans and fungi, close to the origin of animal multicellularity (1). The role of the tyrosine-specific group of kinases (TKs) in intercellular signaling and their restriction to metazoans suggested that TKs were key to metazoan evolution (2). Plants and unicellular organisms lack TKs, although they have a small number of dual-specificity kinases and associated tyrosine phosphatases (PTPs) and SH2 phosphotyrosine-binding domains generally not involved in intercellular signaling. The surprising discovery of TKs in choanoflagellates (3–5) showed that invention of these key mediators of intercellular signaling preceded their expansion in metazoans. We show here that choanoflagellates have invested hugely in a largely independent pTyr signaling system, yet many of these genes suggest functional convergence between choanoflagellates and metazoans and new combinations of signaling modules, both of which hint at restricted pathways through the signaling network.

Results

Determination and Classification of *Monosiga* Tyrosine Kinases. Our analysis of the draft *Monosiga* genome predicts 128 TKs within a total kinome of ≈ 380 protein kinases (<http://kinase.com/kinbase>). Extensive gene model curation and selected cDNA and genome resequencing allowed us to improve predictions for 102 of these sequences, although several fragments and likely imperfect predictions remain. These constitute the largest known tyrosine kinome and make up over twice the fraction of the proteome than that of any metazoan (6–9), a startling result for a unicellular organism. Sequence analysis of the kinase domain and other regions clusters these TKs into 22 families and 26 singletons (Fig. 1). Their scope is

paralleled by their diversity: when compared with metazoan TKs by pairwise and multiple sequence alignment and family profile–profile alignments, the only clearly identifiable specific homologs were of the Src subgroup kinases (Src, Csk, Abl, and Tec).

Receptor TKs (RTKs). Eighty-eight RTKs are predicted, based on predicted signal peptides and transmembrane (TM) regions, known extracellular domains, and paralogy. Most are typical type I TM proteins, but two are multipass (six to nine adjacent predicted TMs), including one encoding transporter domain [supporting information (SI) Fig. S1]. Seventy-three RTKs belong to 15 families, none of which have obvious metazoan orthologs, although kinase domain profile–profile alignments do show weakly specific similarity between RTKB and RTKC families and the metazoan FGFR and Eph families, respectively. Their domain organization is often similar to that of metazoans, whether due to common origin or convergent evolution (Table S1, Fig. 2). For instance, *Monosiga* lacks the Ig domains found in many metazoan RTKs, but 15 *Monosiga* RTKs have divergent repeats similar to hyalin (HYR) domains, which in turn are predicted to be structurally related to Ig and FN3 domains (10). Similarly, 21 *Monosiga* RTKs have cysteine-rich extracellular repeats and several families of CxxC motifs. These are weakly similar to the TNFR and furin-like domains of some metazoan RTKs. Variant EGF-like domains are also seen (Table S1). Several of these domains are found in other predicted receptor and secreted *Monosiga* proteins. For instance, the *Monosiga*-specific RM1 motif is repeated 8–13 times in three RTKs (SI Text) and in 40 other proteins, most of which are predicted to be secreted.

Cytoplasmic TKs (CTKs). Most CTKs are associated with membrane and pTyr binding and, as in metazoans, are likely to transduce signals from activated receptors, although frequently with distinct domain combinations. Twenty-nine of the 40 CTKs fall into eight families, seven of which also contain SH2 or phosphotyrosine binding (PTB) domains (Fig. 2, Fig. S2). These include homologs of all four Src subgroup families, based on presence of SH2 and SH3 domains and on kinase domain sequence similarity, which averages

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Data deposition: The sequences reported in this paper have been deposited in the kinase.com database, <http://kinase.com/kinbase/FastaFiles> (accession nos. Mb0001–Mb0128).

See Commentary on page 9453.

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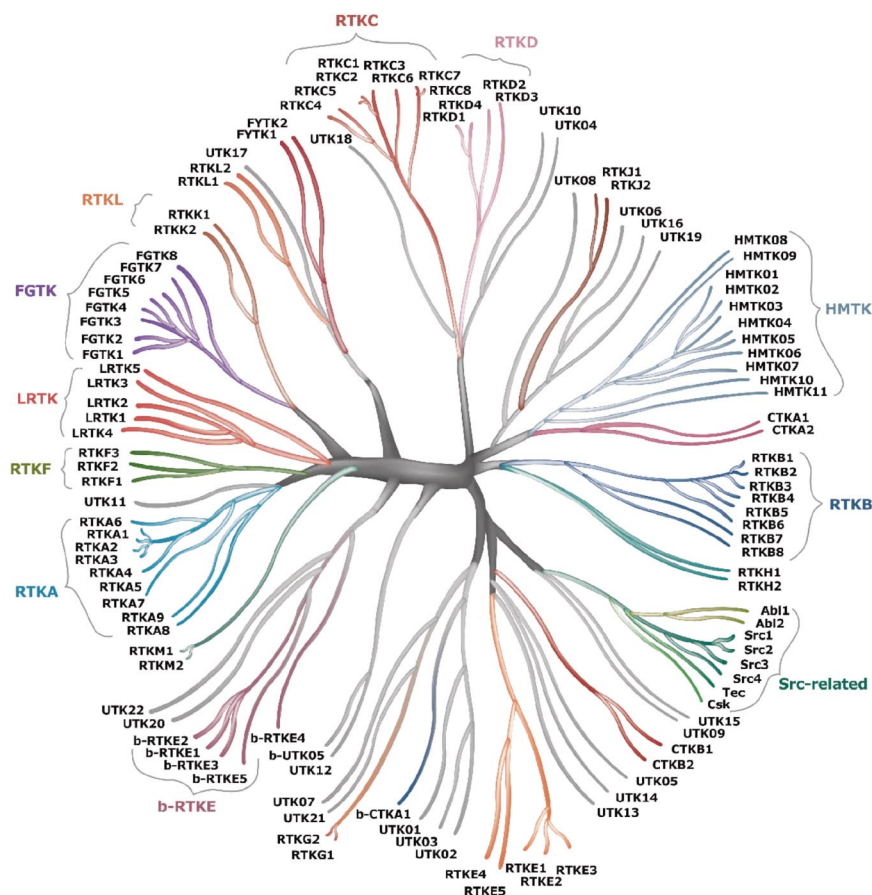


Fig. 1. Phylogenetic tree of *Monosiga* tyrosine kinases, based on alignment of kinase domains, pairwise similarity, and conservation of key residues. Second kinase domains are prefixed by b-. Specific branching patterns between most families are relatively poorly supported.

60% identity to their closest metazoan homologs, compared with <50% for any other *Monosiga* kinase. As in metazoans, all but Csk have an activation loop phosphorylation site, and all four Src family kinases have conserved Csk phosphorylation sites at their C termini. In *Monosiga ovata*, Csk has been shown to phosphorylate and partially repress Src activity through this site (5).

Three of the four Srcs have predicted membrane-anchoring myristoylation sites, indicating that they function proximal to receptors, as with their metazoan counterparts. Curiously, the fourth replaces this with a predicted lipid-binding C2 domain that suggests a novel mechanism of membrane targeting, perhaps similar to the PH domain of Tec kinases (Fig. S2).

Other CTK families also have pTyr-binding domains and may be downstream of RTKs. The two FYTK kinases have SH2 and inositol lipid-binding FYVE domains, one CTKA kinase has SH2 and PH domains, and 10 of the 15 HMTK (HM-motif TK) kinases have PTB domains (Fig. S1). Although FYVE and PTB domains have not previously been seen in TKs, they may function analogously to the membrane targeting (PH, myristoylation) and pTyr-binding (SH2) domains of Src subgroup kinases.

Several RTKs contain predicted Src phosphorylation and SH2-binding sites, most notably at four conserved tyrosines in the RM2 motif within the tail of several RTKB kinases (Fig. S1, SI Text). We tested biochemical activity of *Monosiga* Src1 on peptides generated from two copies of this motif from RTKB2, along with *Monosiga* STAT (a predicted Src substrate) and an optimal vertebrate Src substrate. All showed distinct activity, but the specific activity toward the RTKB2 peptides under these conditions was 6-fold higher (Fig. 3). Kinetic analysis of phosphorylation showed that RTKB2-1 had a k_{cat} of 97.4 min⁻¹ and a K_m of 280 μ M, whereas

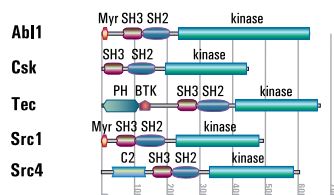
the c-Src optimal peptide gave k_{cat} = 6.5 min⁻¹ and K_m = 90 μ M. Thus, specific recognition of RTKB2-1 by Src1 is driven primarily by a high maximal velocity toward this substrate. These data raise the possibility that the RTKB tail is a Src1 substrate, thus linking RTK and CTK signaling, as in metazoans, and that initial auto-phosphorylation of one of these sites by the RTK recruits Src for further phosphorylation.

Kinase Domain Conservation and Catalytic Activity. Given their ancient divergence, we tested whether *Monosiga* TK domains had unique sequence features. Comparison of all *Monosiga* TK domains to all human, *Drosophila*, and *Caenorhabditis elegans* TK domains by HMM profiles shows a remarkable similarity (Fig. S3), with no clear difference in the conservation profile at any part of the domain. This suggests that TKs in both lineages are under similar constraints, and that their common ancestor had already taken on a “mature” TK structure. Most appear to be activated by phosphorylation, because 103 TKs conserve one or more tyrosines in the phosphorylatable region of the activation loop (Dataset S1).

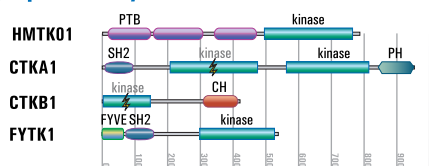
In other species, several RTKs have lost key catalytic residues and are thought to act as scaffolds or coreceptors, including the EGF receptor ErbB3 and several Eph receptors (7). By this measure, 13 *Monosiga* TKs are inactive (Dataset S1). Most belong to the RTKB or RTKM families or are unclassified. Unlike in human, three of the inactive *Monosiga* kinases appear to be cytoplasmic.

Nine kinases have dual catalytic domains, including the six RTKE receptors and the two CTKA cytoplasmic kinases. In all cases, one of the two domains is predicted to be catalytically inactive and is usually very divergent or fragmentary. This situation is analogous to but distinct from metazoan Jak kinases, whose inactive second kinase domains are autoinhibitory (11).

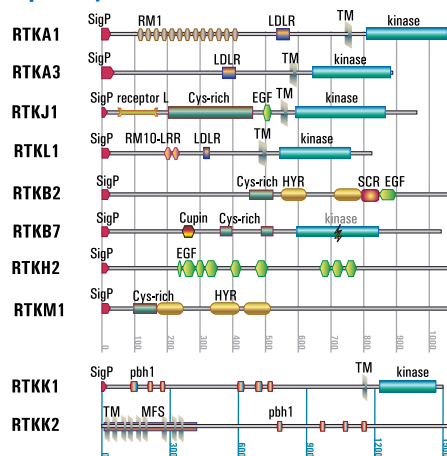
Src-related Kinases



Cytoplasmic Tyrosine Kinases



Receptor Tyrosine Kinases



Unclassified Tyrosine Kinases

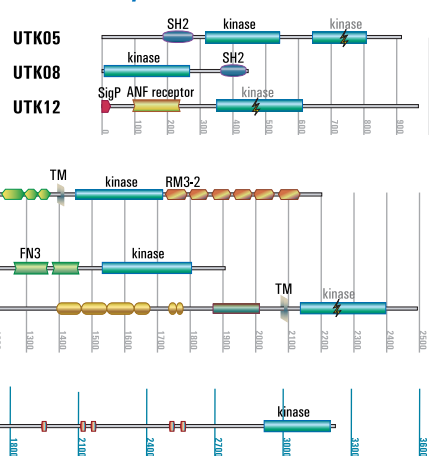


Fig. 2. Domain architecture of representative tyrosine kinases. Predicted inactive kinase domains indicated by lightning bolt, fragments or partial matches to domains indicated by shortened icons. SigP: signal peptide; Myr, myristoylation site; other names from Pfam, SMART, or Table S1. For fuller tyrosine kinome, see Fig. S1 and <http://kinase.com/kinbase>.

Other Phosphotyrosine Signaling Proteins. The richness and diversity of tyrosine kinases are reflected in downstream pTyr-dependent proteins. Conventional tyrosine-specific phosphatases (PTP) and pTyr-binding domains (SH2, PTB) are also greatly expanded in number and domain complexity when compared with yeast, *Dictyostelium*, or *Tetrahymena* and surpass even the human counts for PTP and SH2 proteins (Table 1). As with TKs, we see limited orthology to metazoans, tremendous diversity and several recurrent themes and variations in domain architecture (Fig. 4).

Unlike the few other unicellular PTPs, 4 of the 39 *Monosiga* PTPs have clear human orthologs. These include SHP, PTPN13

(PTP-BAS), PTP23 (HD-PTP), and PTP N3/N4. Curiously, *Drosophila* lacks PTPN13, and both *Drosophila* and *C. elegans* lack PTPN23, so, although ancient, these are not evolutionarily indispensable. Both SHP and PTPN13 have been shown to dephosphorylate Src in mammals (12). As in metazoans, some PTPs appear to be catalytically inactive, and four have lost their HCxxxxR active site motifs (Fig. S2).

By contrast, over one-fifth (26 of 123) of the SH2 proteins have human orthologs covering 15 classes (13) and all 11 major functional categories (Table 2, Fig. 4, Fig. S2). This indicates that much of the cellular pTyr-modulated circuitry was present in the unicellular common ancestor, despite the limited orthology in TKs and PTPs. These shared SH2 proteins mediate pTyr modulation of major signaling pathways, including Ras, Rho, Rac, and Cdc42 small GTPases, phospholipid and calcium signaling, transcription, cytoskeletal interactions, Src subgroup tyrosine kinase and SHP phosphatase signaling, and several adaptors and scaffolds. The remaining 98 SH2 proteins and 35 PTPs lack metazoan orthologs, but many have domain combinations that suggest common themes and the development of new circuits within a constrained set of

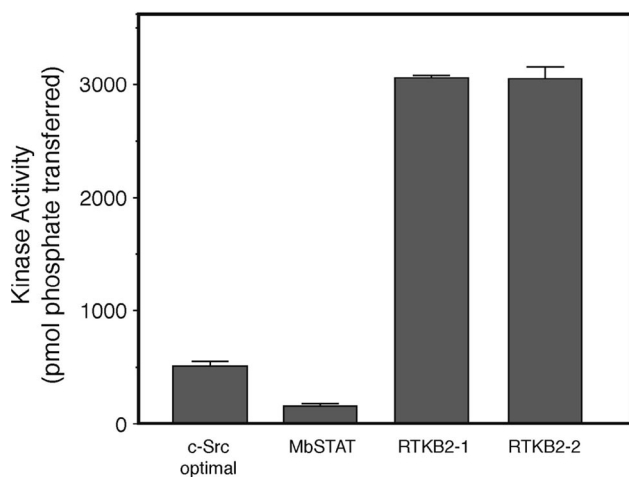


Fig. 3. *Monosiga* Src1 kinase efficiently phosphorylates two RM2 motifs in the cytoplasmic tail of RTKB2 (MbRTK1). The higher efficiency relative to a site on a *Monosiga* STAT homolog or a consensus c-Src substrate suggests that these are specific Src1 phosphorylation sites.

Table 1. Number of proteins with pTyr associated signaling domains in selected genomes

Species	TK	PTP	SH2	PTB
<i>Tetrahymena thermophila</i>	0	3	1	0
<i>Dictyostelium discoideum</i>	0	3	13 (14)	0
<i>S. cerevisiae</i>	0	7	1	0
<i>M. brevicollis</i>	128 (136)	39 (40)	123 (143)	20 (31)
<i>Drosophila melanogaster</i>	33 (34)	16 (23)	28 (34)	10
Human	90 (94)	38 (50)	110 (120)	46 (51)
Human– <i>Monosiga</i> orthologs	4	4–5	19	1

Parentheses indicate domain count due to multidomain proteins. Human counts from RefSeq analysis and published studies (13, 16, 29).



Several PTP and SH2 domains are fused to Class III myosins.

Many more PTP and SH2 proteins are linked to other signaling domains but in unique architectures or with no specific homology to human counterparts (Fig. S2). Partner domains consist mostly of protein, lipid, and calcium-binding adaptor modules, including SH2, SH3, PDZ, SAM, WW, C1, PH, ankyrin, and EF hand domains. *Monosiga* lacks orthologs of the metazoan SH2-RasGEF and SH2-C1-RhoGAP proteins but has unique and possibly analogous

Table 2. Human orthologs of *Monosiga* SH2-containing proteins and their functions

Ortholog	Function
Crk	SH2-SH3 adaptor (RAP/RAC GEFs for adhesion)
Grb2	SH2-SH3 adaptor (SOS, Gab1-MAPK/PI3K)
SHP	PTP phosphatase: Src activator, RTK signaling
STAT	Transcription factor
Cbl	Ubiquitination, receptor trafficking
PIK3R (p85)	Phosphoinositide signaling; PI3K regulatory subunit
PLC γ	Phospholipase: PI3K/Ca signaling adaptor
SHIP2	Phosphoinositide phosphatase
RASA1	Small GTPase: Ras adaptor
Rin	Small GTPase: CDC42 adaptor?
Vav	Small GTPase: Rho adaptor
Src/Abl/Csk/Tec	Src kinase signaling
TNS1	Cytoskeleton
SH2D4	Unknown
Supt6 h	Regulator of chromatin structure. Conserved in yeast, probably non-tyr-binding

RasGEF-SH2-SH2 and RhoGAP-SH2 combinations. Similarly, *Monosiga* and metazoans have several nonorthologous proteins containing SH2 and PH domains, which may share related functions, although lacking obvious common ancestry. A few metazoan proteins have dual SH2 domains, but *Monosiga* has 27 multi-SH2 proteins, with up to six domains seen in a single protein. Many more proteins (13 PTP, 28 SH2) consist of only one recognizable domain and probably include many fragmentary gene predictions.

PTB domains are prevalent in *Monosiga* and metazoans (Table 1) but are absent from lower organisms. Metazoan PTBs can bind peptides, phosphopeptides, or phospholipids (16). The specific ligands in *Monosiga* could not be predicted by sequence analysis, although domain combinations indicate several are associated with pTyr signaling. The PTB-kinase association in the HMTK family is novel, although reminiscent of the pairing of SH2 with the Src subgroup kinases and a likely case of convergent evolution. *Monosiga* has one PTB-SH2 protein that might be a homolog of the Shc adaptor, but other PTB proteins are unique and have no other domains.

Discussion

The *Monosiga* genome has revealed a treasure trove of diverse tyrosine kinases and associated pTyr signaling proteins. These demonstrate an unprecedented diversity relative to all known (metazoan) TK-based signaling yet reveal several common themes that suggest convergent evolution and a limited set of recurring molecular themes favored by signaling pathways. These data also highlight the unresolved puzzle of why a unicellular organism has such an elaborate signaling system based on external cues. Some choanoflagellates such as *Proterospongia* sp. do form colonies, and it may be that such a colonial ancestor drove the evolution of this system, yet it is clear from sequence conservation that pTyr signaling proteins continue to be essential for the current unicellular lifestyle of *M. brevicollis*. Possible functions include response to prey, predators, mates, and the abiotic environment.

Only the four Src-subgroup kinase families have detectable metazoan orthologs, although possible divergent homologs of FGFR and Eph RTKs may also be present. By contrast, most metazoan TK families are clearly visible in sponges (EGFR, FGFR, Eph, InsR, Ret, Musk, Sev, DDR RTKs and Jak, Syk, and Fer CTKs) (17, 18). This suggests that the choanoflagellate-metazoan common ancestor had a mature Src signaling and some RTKs, but that most metazoan TK families were established closer to the base of metazoans. The story is similar with PTPs, but the common

ancestor apparently had a very extensive set of SH2 proteins, with many more classes invented within the choanoflagellate lineage. Extracellular domains evolve rapidly and can swap between families, such as in the RM1 domains found in both RTKA and RTKG1 kinases. Even more remarkably, one, or possibly two, of the receptor SH2 proteins have extracellular domains that are highly similar (>90% sequence identity) to RTKB3 and RTKB5, indicating these are recent fusions and suggesting a kinase-SH2 interaction by receptor heterodimerization. *Monosiga*-specific extracellular motifs are also seen in many other receptor proteins, including rPTPs and rSH2s, and secreted proteins, suggesting they have common ligands or may interact homotypically.

Intracellular domains are more evolutionarily stable and are dominated by pTyr, lipid, and protein interaction domains, as seen in metazoans. In addition, *Monosiga* pTyr-associated proteins have a strong association with the cytoskeleton, as evidenced by an unusual abundance of myosin, CAP-Gly, and calponin homology domains, and with a variety of small GTPase GAPs and GEFs.

Common themes and possible convergent evolution are seen in the domain structures of many pTyr signaling proteins. These include the swapping of a myristoylation site for a putative lipid-binding C2 domain in Src4, the common occurrence of HYR domains reminiscent of Ig domains, the use of cysteine-rich motifs in extracellular regions, and the use of PTB domains as membrane or phosphopeptide anchors that may be analogous to SH2 and myristoylation domains in Src kinases. The development of dual-domain and catalytically inactive kinases are also probably independent innovations in both lineages. In other cases, *Monosiga* proteins are associated with signaling domains not found in metazoans or found in a different architecture, indicating it has successfully explored new paths within signaling space.

Many of the combinatorial aspects of metazoan pTyr signaling are also found in *Monosiga*, including the widespread occurrence of activation loop phosphorylation sites, the likely phosphorylation of RTKs by Src and of Srcs by Csk, the predicted membrane localization of most CTKs, the conservation of most major classes of SH2 domain proteins, and the occurrence of many multi-SH2 proteins that may link distinct pTyr signals. Conversely, the absence of many metazoan components may allow experimental investigation of pathway alternatives, such as the likely specific activation of STAT by Src kinases in the absence of JAKs or the possible link between RTKs and MAPK signaling given the absence of any Raf kinase in *Monosiga* (1).

Future Prospects. This analysis of the draft *Monosiga* genome is surely just an exploratory step in understanding this elaborate and divergent network. The sequence divergence in *Monosiga* and the presence of many short exons hamper gene prediction. We manually improved 102 the kinase sequences over the genome predictions, but several are still clearly incomplete. Impending large-scale choanoflagellate EST and genome sequencing, including those for *Proterospongia* sp. and *Monosiga ovata*, will greatly improve our predictions and provide an evolutionary context. New proteomic technologies to identify TK substrate sites and signaling protein interactions (19–21) could quickly fill in much of the signaling network and allow large-scale comparisons to other systems. A greater understanding of pTyr signaling in choanoflagellates promises to reveal both variations on an important biological theme and commonalities that indicate common origin or convergent evolution.

Materials and Methods

Gene Identification. Protein sequences were predicted from release 1.0 of the *Monosiga* genome (1). Protein kinases, PTP, SH2, and PTB-containing proteins were identified by profile HMM searches against genomic, EST, and predicted gene sequences, using HMMer (<http://hmm.janelia.org>), GeneWise (www.ebi.ac.uk/Wise2), and Gene Detective (a hardware-accelerated implementation of

GeneWise). Individual hits were merged by sequence comparison and mapping to genomic sequence using Blat (22).

TKs were identified by their characteristic HrD[IVLM]AaRN motif [uppercase letters are invariant; Ser/Thr kinases (STKs) are typically HrDIKPEN] and by scoring against kinase group-specific HMMs. These TKs also strongly conserved the [KR]Wm[as]PE motif ([KR]YM[AS]PE in STKs) (Dataset S1).

All sequences were extensively curated using ESTs, sequence similarity to *Monosiga* and published proteins and to Pfam/SMART domains in surrounding genomic regions. Seven questionable cases were improved by targeted cDNA or genomic sequencing. Kinase domains were compared with metazoan kinase families by multiple alignment and tree building, by pairwise blast analysis, and by comparison of profile HMMs built from metazoan and *Monosiga* families using PRC (<http://supfam.org/PRC>).

Domain Profiles. HMM searches on Pfam, SMART, TIGR, and in-house HMM (SI Text, Figs. S3 and S4) with Global and Glocal models were performed with a hardware-accelerated DeCypher system (Active Motif). E value cutoffs of $e = 10$ were used to pick up repeated elements whose individual scores were very low. Sequence-level scores of $e > 0.001$ were discarded and scores of $e > 1e-8$ inspected manually. Overlapping domains from different profile families were merged. Cysteine-rich regions were identified by multiple overlapping hits to the Pfam and SMART profiles GCC2, GCC3, TNFR3, TNFRc6, NCD3G, and to internal models for RM5, RM6, RM9, RM15, and RM15t. Adjacent cysteine-rich regions were merged when separated by < 10 residues.

Custom HMM profiles were built for several unique conserved regions, found by manual inspection and the MEME motif-finder (23), followed by HMM searches of *Monosiga* and GenBank protein and EST sequences to diversify the motifs found and occasional merging of adjacent motifs into gapped profiles. The Vav PH domain and Supt6 CSZ domain were detected by alignment to proteins with these domains but did not score significantly on the HMMs.

Signal peptides and TM segments were predicted by SignalP (24) and TM-HMM (25). TMs that overlapped kinase domains or signal peptides were eliminated. Likely receptors that lacked either signal were subjected to gene prediction and evaluated in part on the basis of these motifs; this may have led to some overprediction of such motifs. Myristoylation sites were predicted with NMT (<http://mendel.imp.ac.at/myristate>) (26).

Kinase Domain Conservation. The alignment of *Monosiga* and metazoan TK domain HMMs was built from a hand-edited alignment of all *Monosiga* TK kinase domains (Dataset S2) and an alignment of published TK domains of *C. elegans*, *Drosophila*, and human (6, 7, 9). The logo was generated with Logomat-M (27).

Other Genomic Searches. Sequence files used for profile searches included *Dictyostelium*: "dicty.predicted.proteins" (<http://dictybase.org>, June 2007 download) *Saccharomyces cerevisiae*: "SGD1.01.45.known.pep" (www.ensembl.org/); *Drosophila* BDGP.4.3.46 "all.pep" (www.ensembl.org/); *Tetrahymena thermophila* "gene.prediction" (<http://ciliate.org>, May 2005 download), and human RefSeq proteins from GenBank, June 2007 download.

Sequencing. Resequencing used either a *M. brevicollis* cDNA library (3) or cDNA. To generate cDNA, *M. brevicollis* (American Type Culture Collection 50154) was cocultured with *Enterobacter aerogenes* at 25°C in natural seawater infused with cereal grass (5 g/liter) in 150- × 25-mm polystyrene dishes (Falcon). Total RNA was extracted and DNase treated with RNeasy Midi-prep kit (Qiagen). This was reverse-transcribed with an oligo(dT) primer (Invitrogen) and amplified using gene specific primers. PCR amplicons were cloned into pCR-Blunt II-TOPO (Invitrogen) and sequenced by using the vector specific primers M13F (5'-TGTAACGACGACGCGCCAGT-3') and M13R (5'-AACAGCTATGACCATG-3') or the gene-specific PCR primers.

Src1 Phosphorylation Assay. Src1 was expressed and purified from insect cells (30). Phosphorylation assays were carried out in total volumes of 25 μ l at 30°C, containing 50 mM Tris-HCl (pH 7.4), 10 mM MgCl₂, 1 mg/ml BSA, 400 μ M [γ -³²P]ATP (15 cpm/pmol), and 750 μ M peptide. Peptide sequences were: c-Src optimal, AEEIYGEFEAKKKK; MbStat, KKKASGYVMADIA; RTKB2-1, SEEVYGA-VDDKKK; RTKB2-2, AEEVYEAIDKKK. Reactions were initiated by addition of purified Src1 to 1.5 μ M and terminated with 45 μ l of cold 10% trichloroacetic acid at 20 min. This time point was within the linear range of the enzyme assay. Samples were centrifuged for 1 min, and 35- μ l aliquots of the supernatants were spotted onto 2.1-cm phosphocellulose paper circles (27). The circles were washed three times with cold 0.5% phosphoric acid and once with acetone, dried, and counted dry in a liquid scintillation counter to measure incorporation of ³²P into peptide. Reactions were carried out in duplicate and are presented \pm standard deviation. For kinetic measurements, reactions were carried out with varying concentrations of peptide substrates (5–1,000 μ M). Kinetic parameters were calculated by fitting data to the Michaelis–Menten equation.

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