

Cryo-EM structure of catalytic ribonucleoprotein complex RNase MRP

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Abstract:

RNase MRP is an essential eukaryotic ribonucleoprotein complex involved in the maturation of rRNA and the regulation of the cell cycle. RNase MRP is related to the ribozyme-based RNase P, but it has evolved to have distinct cellular roles. We report a cryo-EM structure of the *S. cerevisiae* RNase MRP holoenzyme solved to 3.0 Å. We describe the structure of this 450 kDa complex, interactions between its components, and the organization of its catalytic RNA. We show that while the catalytic center of RNase MRP is inherited from the ancestral enzyme RNase P, the substrate binding pocket of RNase MRP is significantly altered by the addition of unique RNA and protein elements, as well as by RNA-driven protein remodeling.

One Sentence Summary:

Changes in peripheral RNA elements and RNA-driven protein remodeling result in diversification of related catalytic RNPs

Main Text:

Ribonuclease (RNase) MRP, a site-specific endoribonuclease, is a ribonucleoprotein complex (RNP) comprising a catalytic RNA moiety and multiple (ten in *S. cerevisiae*) protein components (1-4). RNase MRP is an essential eukaryotic enzyme that has been found in practically all eukaryotes analyzed (5). RNase MRP is localized to the nucleolus and, transiently, to the cytoplasm (3). Known RNase MRP functions include its participation in the maturation of rRNA and in the metabolism of specific mRNAs involved in the regulation of the cell cycle (6-13). Defects in RNase MRP result in a range of pleiotropic developmental disorders in humans (14).

RNase MRP is evolutionarily related to RNase P, a ribozyme-based RNP primarily involved in the maturation of tRNA (15-17). RNase MRP appears to have split from the RNase P lineage early in the evolution of eukaryotes, acquiring distinct substrate specificity and cellular functions (5, 18, 19).

The catalytic (C-) domain of RNase MRP RNA (Fig. 1A) has the secondary structure closely resembling that of the C-domain of RNase P (Figs. 1B) and includes elements forming a highly conserved catalytic core (3-5). The specificity (S-) domain of RNase MRP RNA does not have any apparent similarities with the S-domain of RNase P (Figs 1A, B). Crosslinking studies (20) indicate the involvement of the RNase MRP S-domain in substrate recognition.

Most of the RNase MRP protein components are also found in eukaryotic RNase P (2). *S. cerevisiae* RNase MRP and RNase P share 8 proteins (Pop1, Pop3, Pop4, Pop5, Pop6, Pop7, Pop8, and Rpp1 (2 copies); RNase MRP protein Snm1 has a homologue in RNase P (Rpr2), while Rmp1

is found only in RNase MRP. Shared proteins bind to both C- and S-domains (21-23). RNase MRP proteins Pop1, Pop6, Pop7 are also an essential part of yeast telomerase, where they form a structural module that stabilizes the binding of telomerase components Est1 and Est2 (24).

Here, we report a cryo-EM structure of the *S. cerevisiae* RNase MRP holoenzyme solved to the nominal resolution of 3.0 Å. We reveal the overall architecture of the RNP, the structural organization of its catalytic RNA moiety, the substrate binding pocket of the enzyme, and interactions between RNase MRP components. Further, we compare the structure of RNase MRP to the structure of the progenitor RNP, eukaryotic RNase P. We show that several of the proteins shared by RNase MRP and RNase P undergo RNA-driven structural remodeling, allowing the same proteins to function in distinct structural contexts. We demonstrate that while the structure of the catalytic center of RNase MRP is practically identical to that of RNase P, the topology of the substrate binding pocket of RNase MRP diverges from that of RNase P due to the presence of auxiliary RNA elements positioned in the immediate vicinity of the conserved catalytic center, due to the binding of RNase MRP protein Rmp1 near the catalytic center, as well as due to RNA-driven protein remodeling.

Overall structure of RNase MRP

RNase MRP holoenzyme used in the final 3D reconstruction was isolated from *S. cerevisiae* as a 1:1 mix with RNase P using a Tap-tag approach with the purification handle fused to protein Pop4 (25). The isolate had all expected proteins and RNA components; RNase MRP was active (Fig. S1). RNase MRP particles were separated from RNase P during data processing using 3D classification (Supplementary Material).

The resultant electron density map had the overall resolution of 3.0 Å, with the central regions as good as 2.5 Å (Figs. S2, S3). The final model included all known components of RNase MRP, except for a peripheral protein Pop3. The electron density corresponding to Pop3 was clearly present, but the map quality in this region was not sufficient for a reliable atomic reconstruction of the Pop3 structure; for illustrative purposes Pop3 was modelled into the RNase MRP map using its structure in yeast RNase P (23).

RNase MRP structure (Fig. 2) is dominated by proteins. The proteins are forming a clamp-like structure embedding RNA and protecting most of it from the solvent, consistent with the available biochemical data (25). The basic patches of RNase MRP proteins largely face the RNA component (Fig. S4). The RNA elements corresponding to the catalytic center in RNase P and forming the catalytic center in RNase MRP protrude into the central part of the protein clamp opening and are exposed to the solvent.

Unlike bacterial RNase P RNA (26), RNase MRP RNA is missing elements that can serve to stabilize its 3-dimensional organization and, similar to eukaryotic RNase P (23, 27), uses protein components as the main structural scaffold (below).

Structure of RNase MRP RNA

RNase MRP RNA forms an essentially single-layered structure dominated by coaxially stacked helical regions (Fig. 1C). In the C-domain, helical stem P1 forms a coaxial stack with stem P4, while stems P19, P2, and the proximal part of P3 form a semi-continuous helix connected to the

P1/P4 stack via a 4-way junction. The fold of the C-domain of RNase MRP RNA is remarkably similar to the structurally conserved fold of the C-domains found in RNases P from all domains of life (23, 27-31), including yeast RNase P (Fig. 1D).

The key areas forming the catalytic center in RNase P (regions CR-IV, CR-Va, stems P4 and P2), as well as P1 and the proximal part of P3 are folded virtually identically in RNase MRP, even in the regions that are somewhat divergent in their sequences. Specifically, stem P4 (which, based on secondary structure predictions, was expected to be shorter than that in RNase P (3)), is terminated by a non-canonical C93-U307 base pair, bringing the length of the stem to 8 base pairs, same as in RNase P. Additionally, the sequence of the CR-IV region differs in RNase MRP and RNase P; however, the structural differences are localized to the U265-U266 RNase MRP section, which substitutes for a longer U311-A314 stretch found in RNase P, while the rest of CR-IV adopts practically identical conformations in the two enzymes. The linkers connecting stem P3 to P2 and P4 differ in RNase MRP and RNase P. However, the position and orientation of the proximal part of stem P3 are defined by its coaxial stacking with P2 and are not affected by differences in the linkers; similarly, the 4-nucleotide A318-C321 loop does not affect the orientation of the distal part of stem P1. Thus, the sequence differences found in the central regions of the C-domains of RNases MRP and RNase P do not lead to substantial structural differences.

The main structural difference between the C-domains of RNase MRP and RNase P is the orientation of the peripheral P15 stems, which changes by $\sim 45^\circ$ (Figs. 1C, D).

The A70G mutation in human RNase MRP (corresponding to A84G in *S. cerevisiae*) leads to the developmental disorder Cartilage Hair Hypoplasia (CHH) (14). A84 forms a reversed Hoogsteen base pair with U314, which is not isosteric to any of the possible GU base pairs; thus the change in the geometry caused by the transition to GU is the likely cause of the RNase MRP functional deficiency leading to CHH.

The S-domains of RNase MRP and RNase P are structurally different, consistent with the divergent substrate specificities of the two enzymes. The central feature of the RNase MRP S-domain is a coaxial stack of helices P5 and P8 (Fig. 1C). The distal part of P8 hosts a salient phylogenetically conserved feature of RNase MRP: the 5'-GA(G/A)A(G/A)-3' element (nucleotides 109-113) (32). This element folds into a canonical GNRA tetraloop capping the P8 stem, with the fifth nucleotide (A113) bulging out; A113 is observed adopting both anti- and syn- conformations (Figs. 3A, S5). This region is involved in interactions with the protein component Pop1; a stacking interaction with the phylogenetically conserved Arg107 appears to stabilize the syn- conformation of A113 (Fig. 3A). Mutations that disrupt the equivalent of the P8 stem in human RNase MRP lead to developmental disorders (14).

Helix P5 is closed by a 29 nucleotide-long loop dubbed here L5 (Fig. 1A). The proximal part of the loop folds into a well-ordered complex structure devoid of canonical base pairs and stabilized by extensive interactions with proteins Pop4 and Snm1 (Figs. 3B-E). The distal part of L5 (U132 to U143) is disordered in the structure.

The P8/P5 helical stack of RNase MRP is docked against the P1/P4 helical stack of the C-domain in a manner closely resembling the docking of the P8/P9 helical stack against P1/P4 in yeast RNase P (Fig. S6): RNase MRP P5 stem occupies a position matching the position of the P9 stem in the S-domain of RNase P, while the position of the RNase MRP P8 stem closely matches the position of the P8 stem in RNase P. Thus, the structural similarity between RNase MRP and RNase P RNAs extends beyond the C-domains, albeit the similarities in the S-domains are somewhat superficial as the loops closing the matching stems are different in RNase MRP and RNase P, as is the connectivity of the elements (Fig. 1).

Long helical stems P6 and P7 are located in the peripheral part of the S-domain (Fig. 1C). The helical elements of the S-domain are connected by linker regions dubbed here J8/9, J6/7, and J7/9 (Figs. 1A, C). RNase MRP S-domain is connected to the C-domain by a short helical stem dubbed here P9; it should be noted that RNase MRP P9 is not a structural equivalent of the stem P7 in RNase P.

Proteins in RNase MRP

Pop1, the largest protein component of RNase MRP, forms the main structural brace in the RNP (Figs. 4A, S7A). The C-terminal part of the protein engages the C-domain RNA, as it does in yeast RNase P (Fig. S7B) (23). The structures of the C-domain RNAs in RNase MRP and RNase P are similar (above), and the C-terminal part of Pop1 (residues 203-875) folds and interacts with RNA essentially the same way in the two enzymes (RMSD = 2.6 Å).

At the same time, the N-terminal part of Pop1 engages mostly the S-domain RNA, which is different in RNase MRP and RNase P. The N-terminal part of Pop1 undergoes RNA-driven structural remodeling and adopts distinct conformations in the two RNPs (Figs. 4B, C). RNA-driven structural remodeling of Pop1 can explain the ability of this protein to stabilize diverse RNPs (17, 24), even when the engaged RNA components are structurally distinct. It should be noted that a short stretch of the N-terminal part of Pop1 (residues 91-99) interacts with an area of the C-domain RNA (parts of P4 stem and CR-IV), which adopts the same conformation in RNase MRP and RNase P. In this region the structures of Pop1 in RNase MRP and RNase P converge, only to diverge away from this short stretch (Fig. S7C).

Proteins Pop6 and Pop7 form a heterodimer that helps to engage Pop1 in both RNase MRP (Fig. 4A) and RNase P (22, 23, 33). Pop6 and Pop7 interact with P3 RNA regions that adopt similar folds in RNase MRP and RNase P and are folded similarly in the two enzymes (RMSD = 1.8 Å and 1.3 Å, respectively).

Protein Pop4 bridges together stem P1 of the C-domain RNA and the L5 loop of the S-domain (Figs. 4D, S8A). This structural role is similar to the role played by Pop4 in RNase P, where it bridges together stems P1 and P9 (23) (Figs. 4E, S8B). The structures of the S-domains of RNase MRP and RNase P differ (above), and the part of Pop4 that engages the S-domain RNA undergo RNA-driven structural remodeling (Figs. 4D, E, S8C): the long alpha helix $\alpha 3$ found in RNase MRP Pop4 is disrupted in RNase P. In RNase MRP, the N-terminal part of $\alpha 3$ is involved in extensive interactions with the L5 RNA loop, stabilizing its structure (Figs. 3B-E, 4D), whereas in RNase P the corresponding fragment of Pop4 is rotated $\sim 135^\circ$ and engages stem P9 (Fig. 4E). It

should be noted that the extended version of the $\alpha 3$ helix found in RNase MRP would clash with the CR-III element of the RNase P S-domain (Figs. 4D, E). The differences in the S-domains of RNase MRP and RNase P RNAs necessitate the structural remodeling of Pop4; however, the similarity in the positioning of P5/L5 in RNase MRP and stem P9 in RNase P (Figs. 4D, E, S6) allows for the structural divergence to be localized to only the N-terminal region of Pop4, while the rest of the protein adopts essentially the same fold in the two enzymes (RMSD = 2.3 Å for residues 85-279).

Proteins Pop5, Pop8, and two copies of Rpp1 form a heterotetramer that interacts with the C-domain (Fig. S9A). The structure of the engaged RNA region is very similar in RNase MRP and RNase P; accordingly, these proteins interact with RNA the same way in the two RNPs and have essentially the same structure (RMSD = 1.7 Å for the heterotetramer), with one notable exception. In RNase P, Pop5 interacts with the N-terminal alpha helix $\alpha 1$ of Pop1 (Fig. S9C). The fold of the N-terminal part of Pop1 in RNase MRP differs from that in RNase P (above), and Pop1 helix $\alpha 1$ does not reach Pop5, instead engaging protein Rmp1, which is not present in RNase P, and in RNase MRP the structural role of Pop1 helix $\alpha 1$ is played by the helix $\alpha 6$ of Pop5 itself (Fig. S9B); $\alpha 6$ appears to be disordered in RNase P (23). Thus, the RNA-driven structural remodeling of Pop1 leads to the remodeling of another protein component, Pop5.

RNase MRP protein Snm1 has a homologue in yeast RNase P, Rpr2. Both Snm1 and Rpr2 are nestled between Pop4 and Pop3 (Figs. S8A, B), interact with Pop4 in a similar fashion, and adopt generally similar folds (Figs. S8D, E). Snm1 binds to the S-domain of the RNase MRP RNA and is involved in the stabilization of the structure of the L5 RNA loop (Figs. 3B-E).

RNase MRP protein Rmp1 is a core RNase MRP protein involved in extensive interactions with the RNA component, binding primarily to the foundation of the P15 stem, as well as to the J7/9 region and stem P9 (Figs. 5A-D, S10). Rmp1 binds in the vicinity of the catalytic center of the enzyme and is involved in the formation of the substrate binding pocket of RNase MRP (below).

The comparison of the protein components of RNase MRP and RNase P demonstrates that RNA-binding protein components shared by several RNPs can cope with the divergence in RNA by adopting distinct conformations. This RNA-driven protein remodeling allows for the binding of the same protein to different RNAs that are not necessarily similar at the protein binding sites.

Substrate binding pocket of RNase MRP

Photoreactive RNase MRP substrates form UV-induced crosslinks to nucleotides A113, U225 in the S-domain and nucleotides G87-U88, G264-U266 in the C-domain of RNase MRP RNA, as well as to proteins Pop1, Pop4, Pop5, and Rpp1 (20). These elements define the substrate binding pocket of RNase MRP (Fig. 6A); the RNase MRP analogue of the RNase P catalytic center (regions CR-IV, CR-Va, stems P4 and P2 (23, 27-31)) is also localized to this pocket.

RNase MRP RNA elements lining the substrate binding pocket include S-domain stem P8 with its terminal loop containing the phylogenetically conserved 5'-GA(G/A)A(G/A)-3' element (32) (which crosslinks to substrates (20)), as well as the S-domain stem/loop P5/L5 and S-domain junction element J6/7 (which also crosslinks to substrates (20)). The elements of the RNA C-domain forming the catalytic center in RNase P (CR-IV, CR-Va, stems P4 and P2 (23, 27-31)),

are also exposed to the substrate binding pocket of RNase MRP. RNase MRP protein parts contributing to the formation of the substrate binding pocket (Fig. 6A) include the N-terminal segment of Pop1, proteins Pop4, Pop5, Rmp1, and one of the copies of Rpp1.

Despite the virtually identical folds of the RNA catalytic centers, the overall topologies of the larger substrate binding pockets in RNase MRP and RNase P are different, in keeping with the divergent specificities of the two related enzymes (Figs. 6A, B).

The CRII/III element and P10/11 stem that contribute to the recognition of the pre-tRNA substrate in RNase P (23, 27-31, 34) are missing in RNase MRP. RNase MRP S-domain elements P5/L5, P8, J6/7 define a pocket that is tighter than that in RNase P. It should be noted that tRNA is not expected to fit into the substrate binding pocket of RNase MRP, mainly due to the steric clash between the “elbow” part of pre-tRNA and the L5 loop of the S-domain (Fig. 6).

In addition to the difference in the S-domains of RNase MRP and RNase P RNAs, the presence of a unique RNase MRP protein Rmp1 near the catalytic center, as well as RNA-driven remodeling of the N-terminal part of protein Pop1 (above) are also major contributors to the divergence of the substrate binding pocket of RNase MRP from that of RNase P and, by inference, to differences in the specificities of the two enzymes.

Our results demonstrate that RNase MRP has acquired its distinct specificity without significant changes in the structural organization of the catalytic center inherited from the progenitor RNase P RNP. Instead, unique auxiliary RNA elements are introduced into the immediate vicinity of the

conserved catalytic center and, acting together with RNA-driven remodeling of proteins shared with RNase P, as well as changes in the protein composition, substantially alter the substrate binding pocket of this RNA-based enzyme.

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Author contributions: AP designed experiments, isolated and verified RNPs, vitrified samples, and built the structure; DL and HL built the structure; CB prepared grids, vitrified samples, and collected cryo-EM data; IB isolated RNPs; SLH designed experiments and contributed to the implementation of the research; ASK designed experiments, collected and processed cryo-EM data, built the structure, and wrote the paper.

Competing interests: The authors declare no competing interests.

Data and materials availability: The accession numbers for the reported structure are: PDB 6W6V and EMDB EMD-21564.

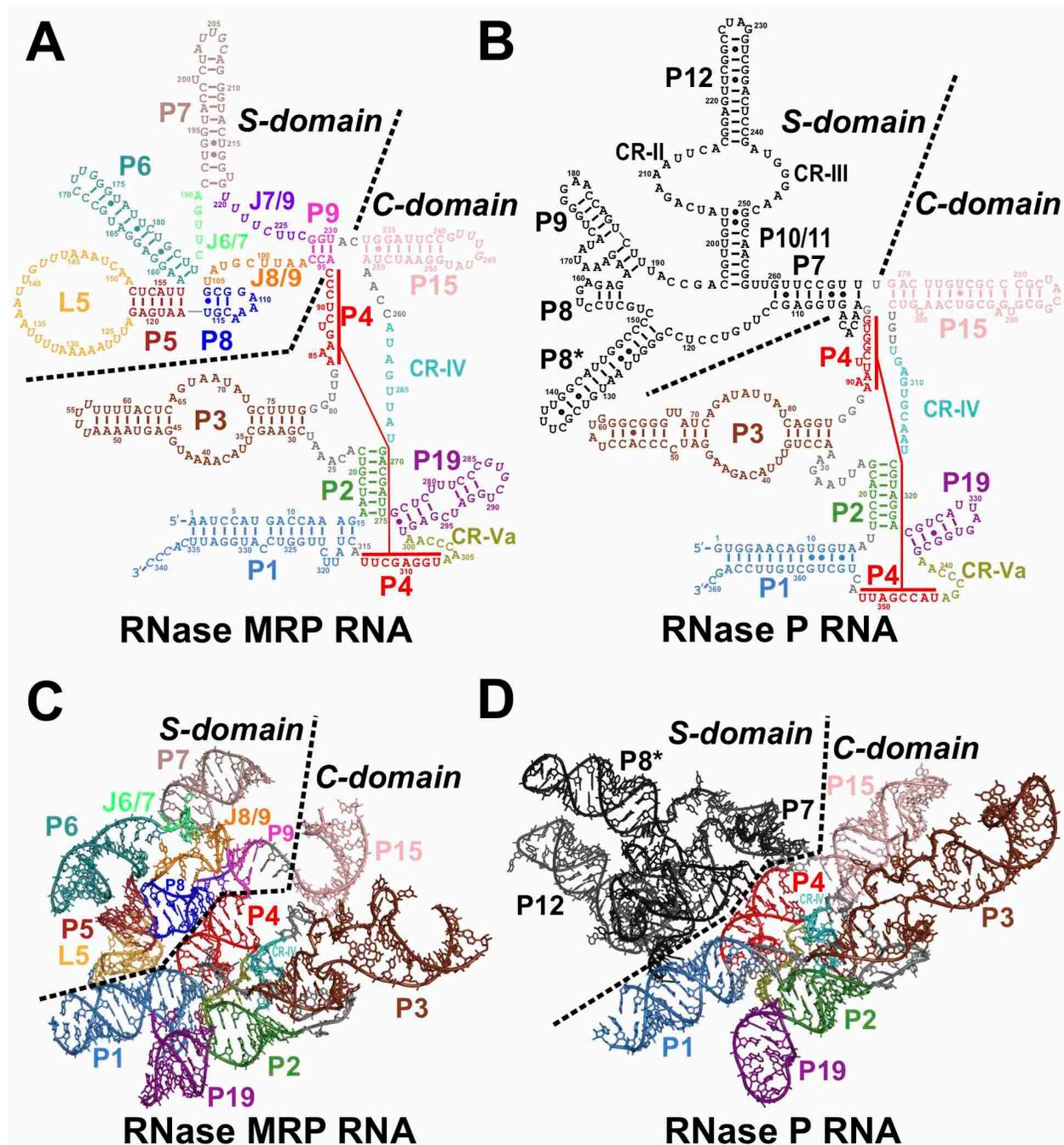


Fig. 1. RNA components of RNase MRP and RNase P.

(A, B) Secondary structure diagrams of the RNase MRP and RNase P RNAs, respectively.

(C, D) Folding of the RNase MRP and RNase P RNAs, respectively, color-coded as in (A, B).

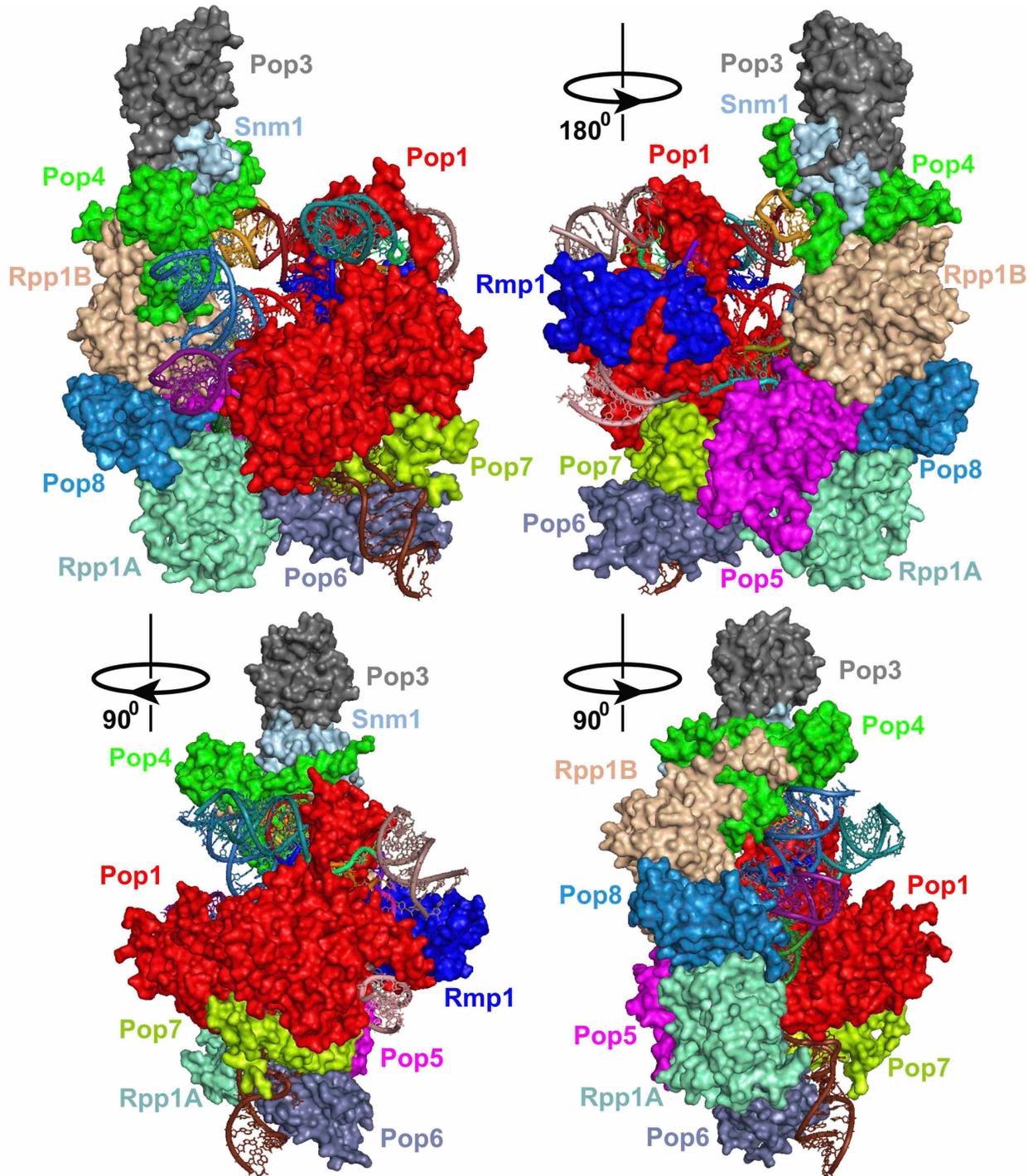


Fig. 2. Structure of the RNase MRP holoenzyme. Protein components (shown as surfaces) are color-coded as marked; the RNA elements (shown as a cartoon) are color-coded according to Fig. 1A.

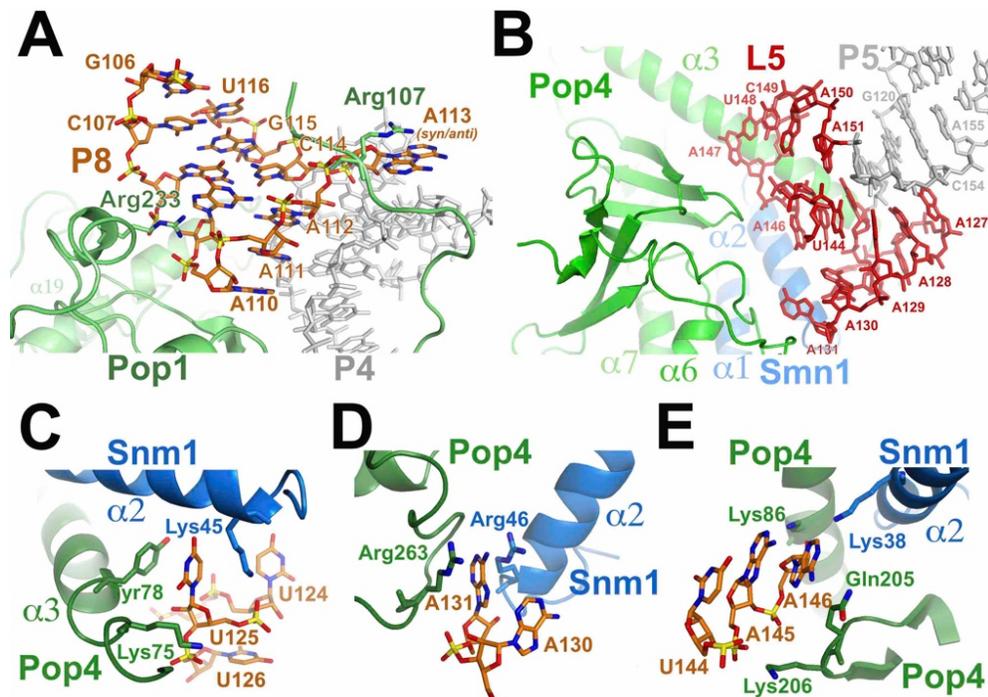


Fig. 3. RNA-protein interactions in RNase MRP.

(A) The structure of the conserved element 5'-GA(G/A)A(G/A)-3' (nucleotides 109-113) and its interactions with protein Pop1.

(B) The structure of the L5 loop and its interactions with proteins Pop4 and Snm1.

(C-E) Proteins Pop4 and Snm1 stabilize the L5 loop.

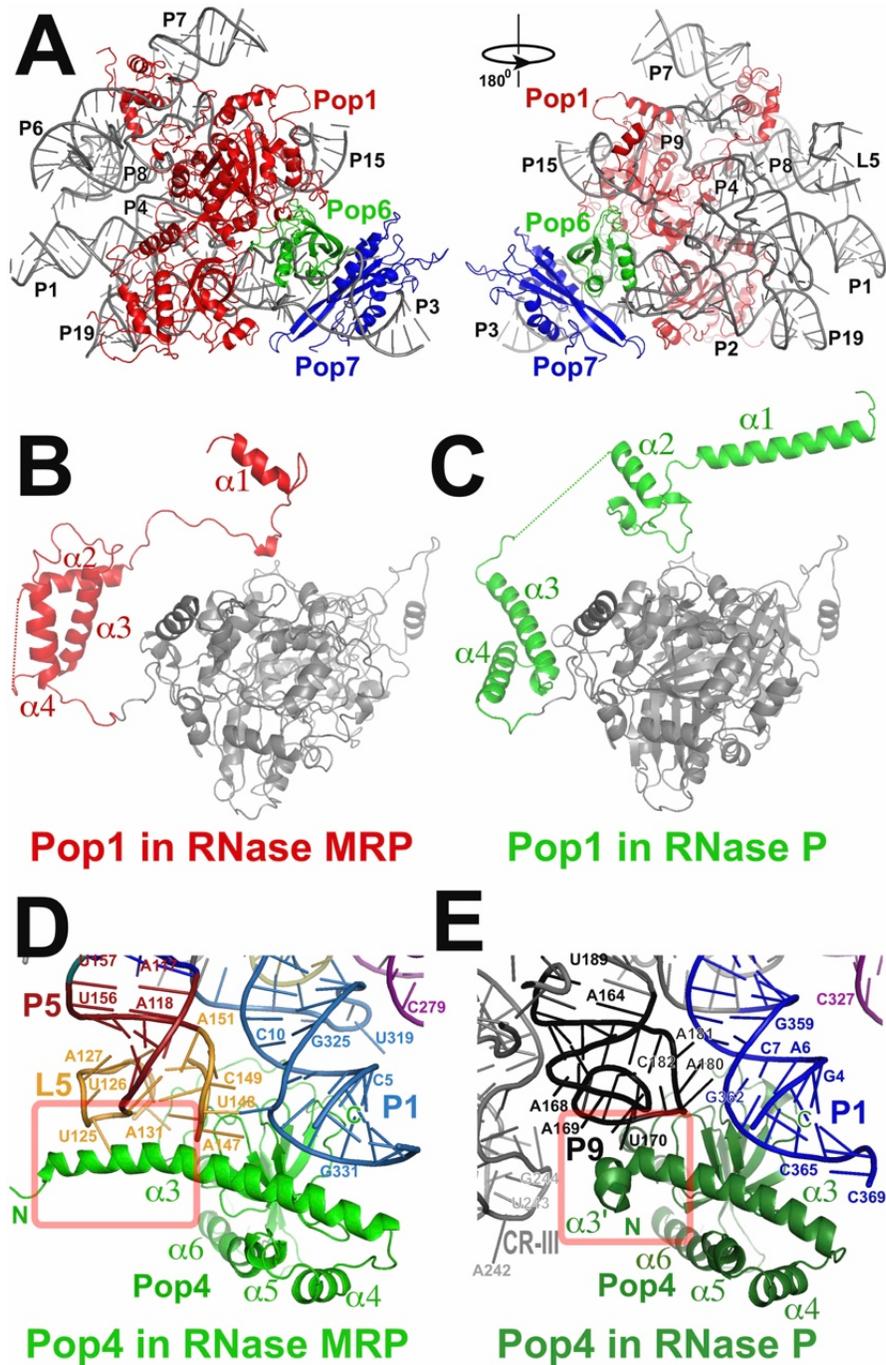


Fig. 4. Proteins Pop1 and Pop4 stabilize the overall RNase MRP RNA structure and undergo RNA-driven remodeling.

(A) Interactions of Pop1 (red), Pop6 (green), and Pop7 (blue) with RNase MRP RNA (grey).

(B, C) The fold of the N-terminal part of Pop1 in RNase MRP (B) differs from that in RNase P (C). The N-terminal parts of Pop1 are shown in red (for RNase MRP) and green (for RNase P); the similarly folded C-terminal parts of Pop1 are shown in grey.

(D, E) Interactions of Pop4 with the RNA components of RNase MRP (D) and RNase P (E). RNA elements are marked and color-coded according to Fig. 1A, B. The part of the Pop4 $\alpha 3$ helix that folds differently in RNase MRP and RNase P is boxed.

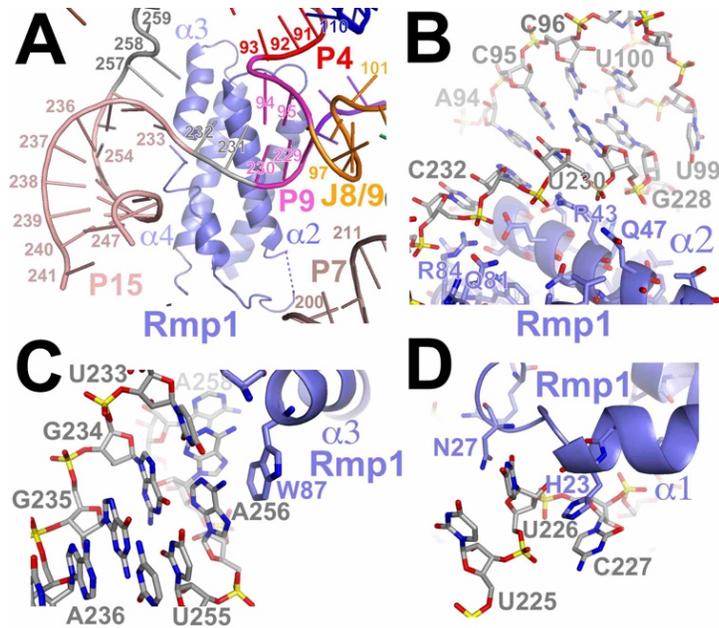


Fig. 5. Interactions of protein Rmp1 with the RNA component of RNase MRP.

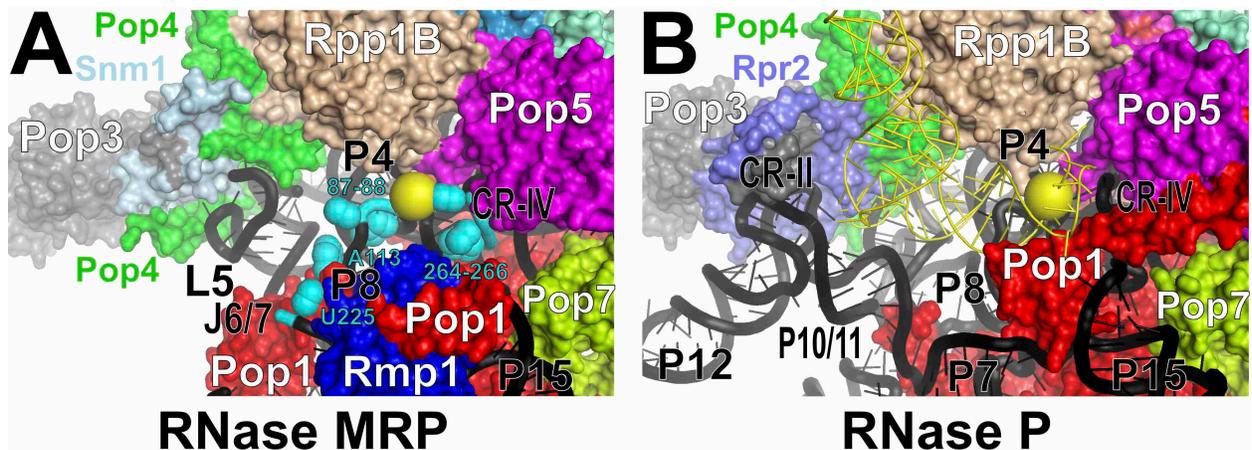


Fig. 6. Substrate binding pockets in RNase MRP and RNase P. Protein components are color-coded according to Fig. 2. RNase MRP and RNase P RNAs are shown as cartoons in black. The substrate pre-tRNA in the substrate binding pocket of RNase P (23) (B) is shown by thin yellow lines. The location of the scissile bonds is shown by yellow spheres (A, B). The locations of the RNase MRP nucleotides that crosslink to substrates (20) are shown as spheres in cyan (A).

Supplementary Materials for

Cryo-EM structure of catalytic ribonucleoprotein complex RNase MRP

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Materials and Methods

Overview

Two attempts to determine the structure of RNase MRP have been made. The first attempt relied on the RNase MRP holoenzyme isolated from *S. cerevisiae* using a purification tag fused to RNase MRP protein component Rmp1 as described below. The isolated complex was unstable when stored on ice, which compromised its usefulness for structural studies. Ultimately, the quality of the resulting reconstruction (6.8 Å resolution) was not satisfactory.

The second attempt relied on the samples isolated from *S. cerevisiae* using a purification tag fused to protein Pop4. Since Pop4 is found in both RNase MRP and the related RNase P, the resulting samples contained a mix of the two complexes. RNase MRP and RNase P particles were separated *in silico*, by 3D classification using the low-resolution RNase MRP reconstruction obtained in the first attempt (above) and the published reconstruction of the RNase P holoenzyme (23) as references. This approach yielded a 3.0 Å reconstruction of the RNase MRP particle.

RNase MRP isolation

The RNase MRP holoenzyme was isolated from *S. cerevisiae* strain OE1004 (35). This strain (*MATa RMP1::TAPHIS8::TRP1 sep 1::URA3 pep4::LEU2 nucl::LEU2 ade2-1 trp1-1 his3-11,15 can1-100 ura3-1 leu2-3,112*) has a purification tag (TAPHIS8) fused to the C-terminus of protein Rmp1. The purification tag was similar to the standard TAP tag (36), but with eight histidine residues replacing the calmodulin-binding fragment (35).

The RNase MRP holoenzyme was isolated as previously described (35) with minor modifications. Yeast were grown in 32 liters of YPD media at 30°C to the late logarithmic phase with vigorous aeration. The culture was cooled on ice; the cells (~300 g) were harvested by centrifugation at 4,000 g (4°C), washed with water twice, followed by a wash with 20% glycerol, frozen in liquid nitrogen, and stored at -75°C. Frozen cells were thawed in cold water and resuspended in 200 ml of a buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 5 mM Mg-acetate, 10% glycerol, 1 mM PMSF, and 400 mg of deproteinated total yeast RNA (Sigma). The cells were disrupted using a BeadBeater (Biospec) (15 x 15-sec pulses on ice), then Tween 20 was added to 0.1% (v/v), and the extract was clarified by centrifugation at 17,000 g for 1 hr (4°C) followed by ultracentrifugation at 100,000 g for 3 hrs (4°C). The clarified extract was mixed with 3 ml of rabbit IgG agarose (Sigma) and incubated for 5 hrs at 4°C with light agitation. The IgG agarose was washed six times with 5 volumes of the buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 1 mM Mg-acetate, 10% glycerol, 1 mM PMSF, and 0.1% (v/v) Tween 20 (Buffer A), and resuspended in 2 ml of the same buffer, supplemented with 10 mg of deproteinated total yeast RNA. Then 80 units of tobacco etch virus protease (TEV) were added and the sample was incubated for 12 h at 4°C with light agitation. The resin was pelleted by centrifugation at 500 g for 5 min (4°C), and the supernatant was collected; the resin was additionally washed twice with 5 ml of Buffer A. The three fractions of the supernatant were combined and cleared of the residual resin by the centrifugation at 4,000 g for 3 min. TEV protease was essentially removed by 3 rounds of the sample concentration to the final volume of 0.5 ml using an Amicon-Ultra 15 (100 kDa MWCO) concentrator (Millipore), followed by the dilution to the final volume of 4 ml of the Buffer A. The final volume was adjusted to 3 ml, and the sample was incubated with 2 ml of Ni-NTA Agarose (QIAGEN) in buffer A for 5 hrs at 4°C with light agitation. The resin was washed six times with 10 ml of Buffer A supplemented with 10 mM Na-imidazole (pH 7.4). After the final wash, the resin was resuspended in a buffer containing 10 ml of 400 mM Na-imidazole (pH 7.4), 100 mM KCl, 2.5 mM Mg-acetate, 5% glycerol, 0.5 mM PMSF, and 0.1% (v/v) Tween 20, and RNase MRP was eluted for 25 min at 4°C with light agitation. After elution, the residual resin was removed by centrifugation at 4,000 g for 3 min, and the elution buffer was exchanged for a buffer containing 20 mM Tris-HCl (pH 8.0), 75 mM KCl, 75 mM NaCl, 5 mM Mg-acetate, 1 mM TCEP, 0.1 mM EDTA, 0.1 mM PMSF, 0.5% glycerol, and 0.1% (v/v) Tween 20 (Buffer B), and concentrated using an Amicon-Ultra 4 (100 kDa MWCO) concentrator (Millipore). The enzyme concentration was adjusted to approximately 0.15 pmol/μl as estimated by denaturing gel electrophoresis of the RNA component run alongside with known quantities of reference RNA.

RNase MRP/RNase P mix isolation

A mix of RNase P and RNase MRP was isolated from *S. cerevisiae* strain YSW1 (a generous gift from Mark Schmitt, *MATa POP4::TAPTAG::TRP1ks pep4::LEU2 nuc1::LEU2 sep1::URA3 trp1-1 his3-11,15 can-100 ura3-1 leu2-3,112 (10)*) using a TAP-tag approach with the tag fused to the C-terminus of protein Pop4. Since Pop4 is a component of both RNase MRP and RNase P, the isolation resulted in the mix of RNase MRP, mature RNase P, and RNase P precursor particles; RNase MRP and RNase P particles were found at approximately 1:1 molar ratio as judged by denaturing gel electrophoresis of the RNA moieties (Fig. S1A). The sample contained all expected proteins (Fig. S1B) as verified by mass spectrometry, the RNA components were essentially intact (Fig. S1A), and RNase MRP was active as judged by the well-characterized cleavage of a 148 nt-long fragment of the internal transcribed spacer 1 (ITS1) of the rRNA precursor, containing the A3 site (9) (Fig. S1C).

The purification procedure described in (25) was used with modifications. Yeast were grown in 32 liters of YPD media at 30°C to the late logarithmic phase with vigorous aeration. The culture was cooled on ice; the cells (~300 g) were harvested by centrifugation at 4,000 g (4°C), washed with water twice, followed by a wash with 20% glycerol, frozen in liquid nitrogen and stored at -75°C. Yeast cells were thawed in cold water and resuspended in a buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 10% glycerol, 1 mM PMSF, 1mM EDTA, supplemented with a protease inhibitor (Pierce). The cells were disrupted using a BeadBeater (Biospec) (15 x 15-sec pulses on ice), then Tween 20 was added to 0.1% (v/v), and the extract was clarified by centrifugation at 17,000 g for 60 min (4°C) followed by ultracentrifugation at 100,000 g for 3 h (4°C). The clarified extract was mixed with 3 ml of rabbit IgG agarose (Sigma) and incubated for 5 hrs at 4°C with light agitation. The IgG agarose was washed 6 times with 5 volumes of the buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 10% glycerol, 1 mM PMSF, 1mM EDTA, 0.1% (v/v) Tween 20 (Buffer LB), and resuspended in 5 ml of a buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 10% glycerol, 1 mM PMSF, 0.1mM EDTA, 4 mM CaCl₂, 1 mM Na-imidazole (pH 8.0), 5 mM Mg-acetate, and 0.1% (v/v) Tween 20 (Buffer CB). 80 units of TEV protease and 1,600 units of RNasine RNase inhibitor (Promega) were added, and the sample was incubated for 12 h at 4°C with light agitation. The resin was pelleted by centrifugation at 500 g for 5 min (4°C), and the supernatant was collected; the resin was additionally washed twice with 5 ml of Buffer CB. The three fractions of the supernatant were combined; DTT was added to the final concentration of 1 mM.

The sample was cleared by centrifugation at 4,000 g for 3 min and 2 ml of calmodulin resin (Agilent Technologies) was added to the sample. The sample was incubated with the resin for 3 hrs at 4°C with light agitation. The resin was washed six times with 10 ml of Buffer CB supplemented with 1 mM DTT. After the final wash, the resin was resuspended in 10 ml of a buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 10% glycerol, 1 mM PMSF, 10 mM EGTA, 1 mM Na-imidazole (pH 8.0), 5 mM Mg-acetate, 1 mM DTT, and 0.1% (v/v) Tween 20. RNase MRP/RNase P mix was eluted for 2 hrs at 4°C with light agitation. After the elution, the elution buffer was exchanged for a buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 5 mM Mg-acetate, 1 mM DTT, 0.5% glycerol, 0.1 mM EDTA, 0.1 mM PMSF, and 0.1% (v/v) Tween 20 (Buffer S), and concentrated using an Amicon-Ultra 4 (100 kDa MWCO) concentrator (Millipore). RNase MRP concentration was adjusted to approximately 1 pmol/μl as estimated by denaturing gel electrophoresis of the isolated RNA run alongside with known quantities of reference RNA. The samples used for vitrification were stored on ice overnight, and vitrified the next day.

Preparation of continuous carbon grids

Quantifoil grids (Cu 300 R1.2/1.3 or R2/1, Quantifoil) were coated with a thin continuous layer as follows. 0.12 g of Formvar 15/95 Resin (Electron Microscopy Sciences) was added to 50 ml of 1,2-dichloroethane (Sigma-Aldrich) and allowed to dissolve overnight in a Coplin jar. A 75 mm x 25 mm microscope slide (Ted Pella) was used to gently stir the solution. A new glass slide was dunked in and out of the solution 10 times to evenly coat the surface of the slide. The Formvar solution was allowed to dry on the slide a few seconds until no remaining liquid was visible. A razor blade was used to score the edges of the film and the Formvar film was floated onto the surface of water in a small basin. Quantifoil grids were placed top side down on the floating Formvar film. The grids and film were picked up with a piece of printer paper placed on top of them. With the top side of the Formvar-coated grids facing up, the paper was allowed to dry overnight at room temperature on a filter paper in a glass Petri dish. Coins were used to hold down the edges of the paper to prevent it from curling. The Leica EM ACE600 sputter coater was used to apply a thin carbon coating on the Formvar-coated grids. The Leica EM ACE 600 reported the final measured thickness of the carbon application of 1.7 nm – 2.1 nm, depending on the batch. The paper with the Formvar-coated grids was placed on a filter paper saturated with 1,2-dichloroethane in a covered glass Petri dish for 30 min to dissolve the Formvar. The grids were transferred to another glass Petri dish lined with dry filter paper and allowed to dry before use. The continuous carbon-coated Quantifoil grids were stored in a desiccator and protected from light.

Sample vitrification, cryo-EM data acquisition and processing

6.8 Å (initial) data obtained using isolated RNase MRP holoenzyme

RNase MRP was isolated from yeast using a purification handle fused to RNase MRP protein Rmp1 as described above.

The sample was diluted 3-8-fold in the vitrification buffer containing 20 mM Tris-HCl (pH 8.0), 75 mM KCl, 75 mM NaCl, 5 mM Mg-acetate, 1 mM TCEP, 0.1 mM EDTA, and 0.1% (v/v) Tween 20, and centrifuged at 18,000 g for 5 min at 4°C. Immediately following the centrifugation, 3.5 µl of the solution was applied to Quantifoil Cu 300 R1.2/1.3 continuous carbon-coated grids prepared as described above and vitrified using Vitrobot Mark IV (FEI) operating at 5°C and 95% humidity. After the sample loading, the grids were incubated for 15 sec, blotted for 3-4 sec, and then plunged into liquid ethane.

Cryo-EM data was collected using FEI Titan Krios electron microscope (300 kV). The images were recorded by Falcon 3EC direct electron detector operating in the counting mode at the nominal magnification of 75,000 (pixel size 0.886 Å). EPU software (FEI) was used for the automated data collection. One image per grid hole was collected at 41 frames per stack. The total exposure rate was 1 e⁻/Å² per frame. The defocus was set to vary in the 1.6-3.2 µm range.

The total of 2,597 micrographs were collected. Frame alignment and integration were performed as implemented in cisTEM (37). Excessive particle drift was commonly observed closer to the end of the stacks; accordingly, only the first 33 frames were used. The contrast transfer function (CTF) parameters were estimated using CTFFIND as implemented in cisTEM (37). Low quality images were discarded, and the remaining 1,875 images were used to extract 236,629 candidate particles using an *Ab initio* algorithm implemented in cisTEM. False-positives as well as small and aggregated particles were removed using 2D classification, leaving 71,917 RNase MRP particles. To generate an initial RNase MRP reconstruction, *ab initio* 3D reconstruction was performed as implemented in cisTEM. The result was used as a starting reference volume in the 3D autorefinement and classification into three 3D classes as implemented in cisTEM. Upon the convergence of the refinement/classification, the single higher resolution class

contained 71.3% of the particles. The resolution of the corresponding electron density map (gold standard Fourier shell correlation (FSC) 0.143) was 6.8 Å. The resulting 3D reconstruction of RNase MRP was used as a reference for *in silico* separation of RNase MRP and RNase P particles, and as a starting reference for the high-resolution refinement as described below.

3.0 Å (final) data obtained using a mix of RNase MRP and RNase P holoenzymes

The mix of RNase MRP and RNase P holoenzyme particles was isolated from yeast using a purification handle fused to protein Pop4 as described above.

For vitrification, the sample was diluted 12-30-fold in the vitrification buffer containing 20 mM Tris-HCl (pH 7.9), 150 mM KCl, 5 mM Mg-acetate, 1 mM DTT, 0.1 mM EDTA, and 0.1% (v/v) Tween 20, and centrifuged at 18,000 g for 5 min at 4°C. Immediately following the centrifugation, 3.5 µl of the solution was applied to Quantifoil Cu 300 R2/1 continuous carbon-coated grids prepared as described above and vitrified using Vitrobot Mark IV (FEI) operating at 5°C and 95% humidity. After the sample loading, the grids were incubated for 30 sec, blotted for 5 sec, and then plunged into liquid ethane.

Cryo-EM data was collected using FEI Titan Krios electron microscope (300 kV). The images were recorded by Falcon 3EC direct electron detector operating in the counting mode at the nominal magnification of 75,000 (pixel size 0.886 Å). EPU software (FEI) was used for the automated data collection. Three non-overlapping images were collected per grid hole at 39 frames per stack. The total exposure rate was 1 e⁻/Å² per frame. The defocus was set to vary in the 1.1-2.7 µm range.

The total of 6,869 micrographs were collected. All 39 frames in the stacks were aligned and integrated using the Unblur algorithm as implemented in cisTEM (37). Dose weighting was applied as implemented in cisTEM (37); the exposure dose for the dose weighting procedure was set to 70% of the actual dose. The CTF parameters were estimated using CTFFIND as implemented in cisTEM (37); 4 movie frames were averaged for the CTF estimation. All images and the corresponding CTF estimations were inspected manually and the images judged to be of low quality were excluded from the further data processing. The remaining 4,781 images were used to extract 1,653,406 candidate

particles using an *Ab initio* algorithm implemented in cisTEM (37). False-positive particles were removed using 2D classification, resulting in 1,086,037 particles, which included both RNase MRP and RNase P.

RNase MRP particles were separated from RNase P particles *in silico* using 3D classification/refinement into two classes as implemented in cisTEM (37); the resolution limit for the classification was set to 20 Å. The published 3D reconstruction of the RNase P holoenzyme particle (23) was used as the initial reference for one of the 3D classes, and the 6.8 Å 3D reconstruction of the RNase MRP particle (above) was used as the initial reference for the other 3D class. After the convergence of the classification, the RNase MRP class contained 545,483 particles, while the RNase P class contained 540,554 particles, consistent with the approximately 1:1 RNase MRP RNA to RNase P RNA molar ratio observed for the isolated sample (Fig. S1A).

RNase MRP particles were subjected to additional 2D classification and low quality/false positive classes were removed from the further processing. The resulting 520,656 particles were subjected to 3D autorefinement/classification into 2 classes as implemented in cisTEM (37). The initial RNase MRP reconstruction (above), subjected to a 20 Å low-pass filtering was used as the starting reference for the 3D refinement. Upon the convergence of the refinement/classification, the low-resolution class was discarded, and the 3D refinement/classification into 2 classes was repeated once more. The resultant selection contained 155,205 RNase MRP particles, which were used in the final 3D refinement. It should be noted that a similarly relatively small fraction of the “good” particles was also observed for the related yeast RNase P (23). The selected RNase MRP particles were used to perform the final 3D refinement as implemented in cisTEM (37) (Fig. S2). The 3D reconstruction algorithm implemented in cisTEM allows for the rejection of a specified fraction of the worst fitting particles. The best map was obtained with the rejection of 25% of particles; this map was used for the model building and refinement. The overall resolution of the map was 3.0 Å based on the gold standard Fourier shell correlation (FSC) of 0.143 criterion (Fig. S2D) and was calculated using Phenix 1.17.1 suite (38).

Model building and refinement

The published structure of the RNase P holoenzyme and the corresponding electron density map were superposed on the RNase MRP electron density map to aid at the initial stage of RNase MRP model building. Further model building

relied on the RNase MRP map. The predicted secondary structure of the RNase MRP RNA component (3), the predicted secondary structures of the protein components, the crystal structure of proteins Pop6/Pop7 in complex with the P3 subdomain of RNase MRP (33), and the results of the crosslinking analysis of RNase MRP (21) were used as additional aids. Overall, the quality of the RNase MRP electron density map (Fig. S3) was sufficient for the unambiguous modeling of most parts of the RNA component, and all protein components, except for Pop3, for which electron density was observed, but the local quality of the map was not sufficient to build a model.

Model building was carried out in Coot 0.8.9 (39). The refinement and validation were performed using Phenix 1.17.1 suite (38); secondary structure restraints as implemented in Phenix were used in the refinement of the helical regions of the RNA component as appropriate. Protein Pop3 was not included in the final model; for illustrative purposes, the cryo-EM structure of Pop3 as it is found in the context of yeast RNase P (23) was fitted into the corresponding part of the RNase MRP map by maximizing the correlation between the map and the model as implemented in Chimera (40).

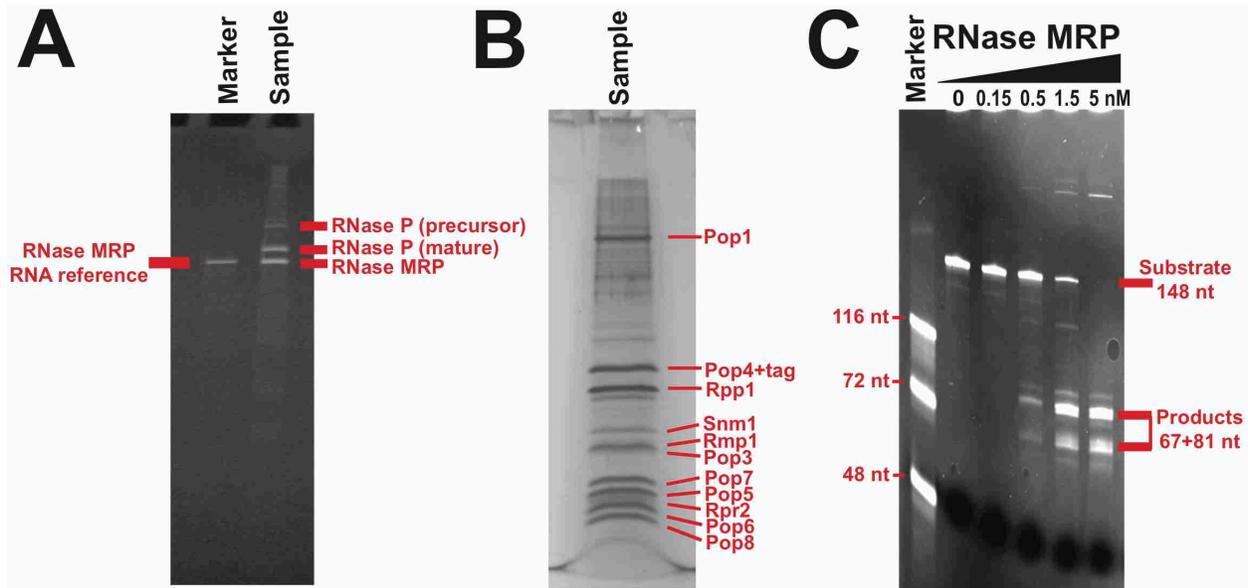


Fig. S1. Isolation of RNase MRP.

(A) RNA components in the isolated sample. Ethidium bromide stained denaturing PAAG.

(B) Protein components as identified by mass spectrometry. Silver stained SDS PAAG.

(C) RNase MRP cleavage of a fragment of the internal transcribed spacer 1 (ITS1) of the rRNA precursor containing the A3 site (9). The substrate (400 nM) was digested by 0; 0.15 nM; 0.5 nM; 1.5 nM; 5.0 nM of RNase MRP (as shown above the gel) for 60 min at 30°C in a buffer containing 20 mM Tris-HCl (pH 7.5), 50 mM NaCl, 10 mM MgCl₂, 1 mM DTT, 50 µg/ml BSA. Ethidium bromide stained denaturing PAAG.

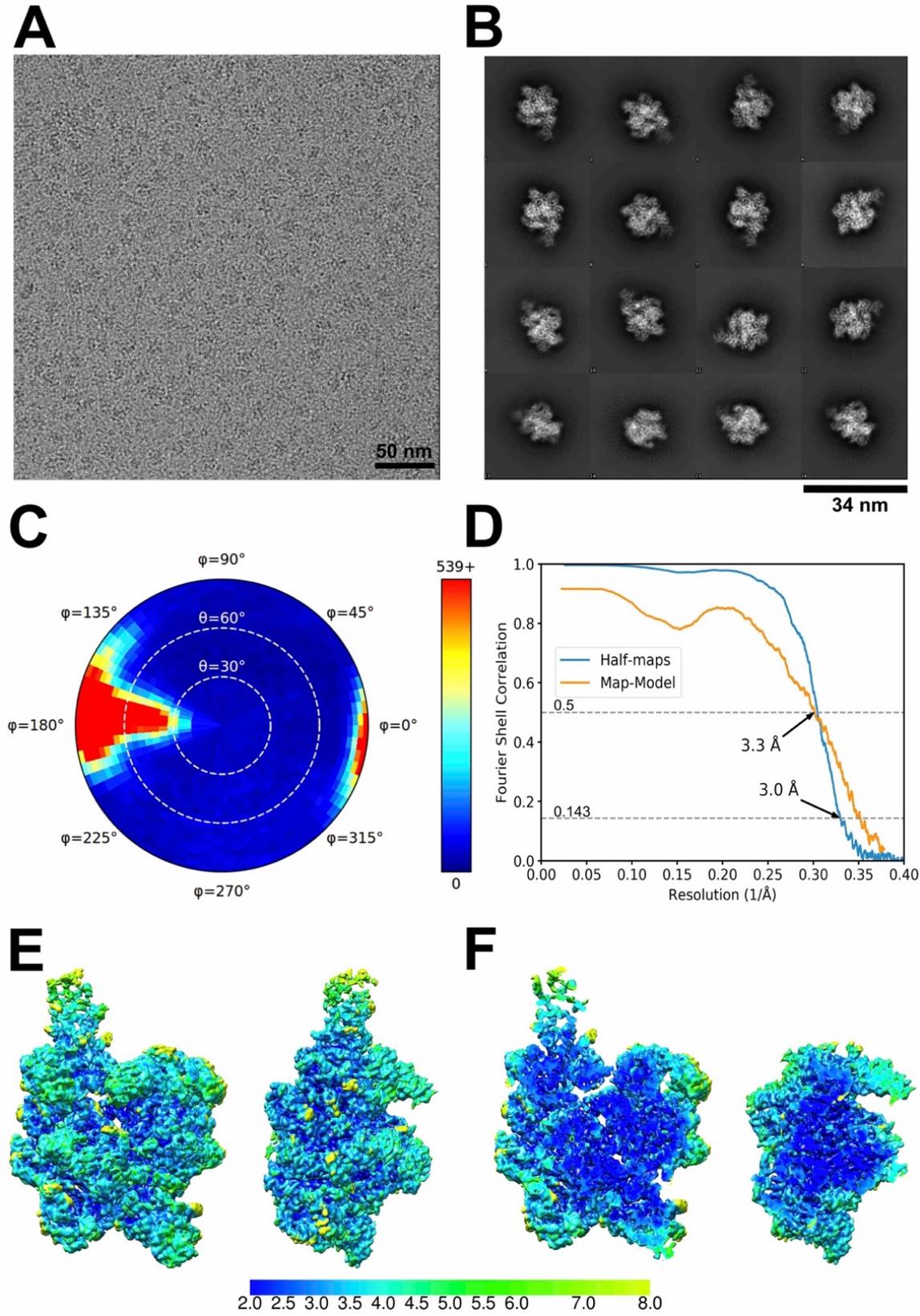


Fig. S2

Fig. S2. Cryo-EM data collection and processing.

(A) A representative cryo-EM micrograph.

(B) Representative 2D class averages for RNase MRP particles.

(C) The distribution of the Euler angles for the aligned particles used in the final RNase MRP reconstruction.

(D) Fourier shell correlation curves (FCS) for the final electron density map and model. The gold-standard FCS curve is shown in blue; the final model to the final map FSC curve is shown in orange.

(E, F) Local resolution map (surface and with the top half removed, correspondingly). The resolution (in Å) is color-coded according to the bar below the map.

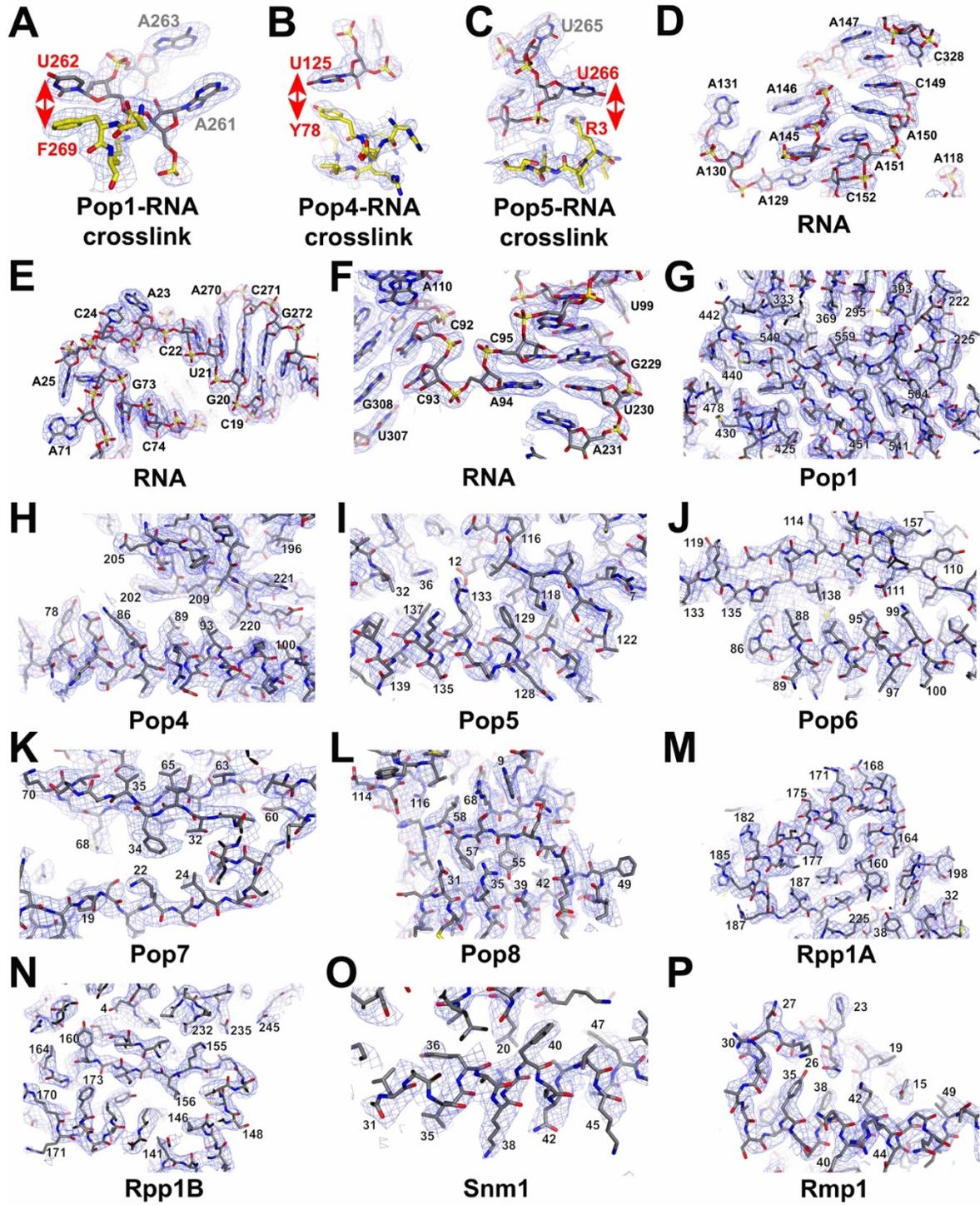


Fig. S3

Fig. S3. Representative fragments of the final electron density map used in the model building and refinement.

(A-C) Fragments corresponding to the previously identified UV-induced RNA-protein crosslinks (shown by red arrows) (21).

(D-F) Representative fragments of the RNase MRP RNA component.

(G-P) Representative fragments of RNase MRP proteins Pop1, Pop4, Pop5, Pop6, Pop7, Pop8, Rpp1 (copies A and B), Snm1, and Rmp1, as marked below the panels.

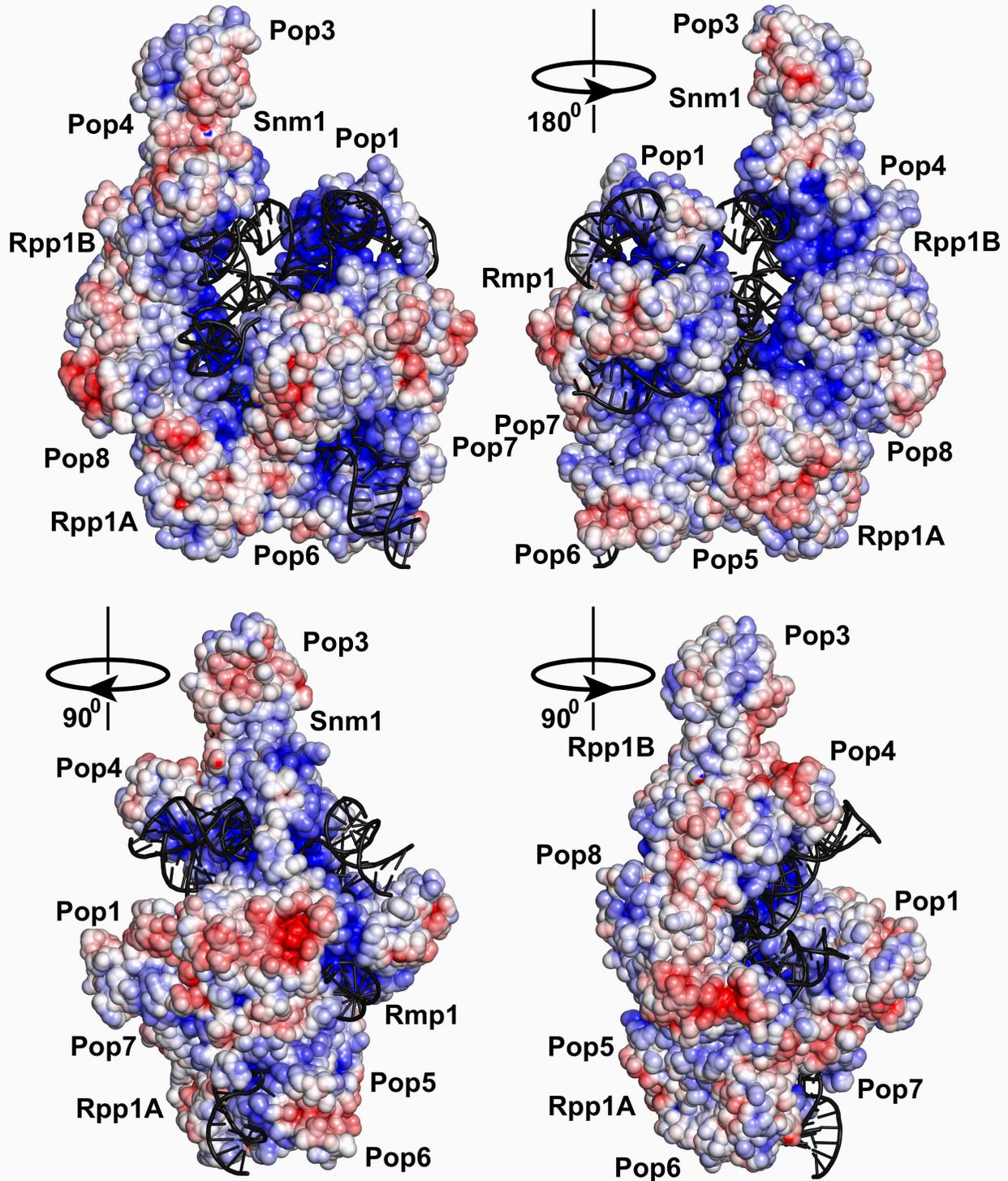


Fig. S4. Surface electrostatic potential distribution for the RNase MRP proteins and the RNA component of RNase MRP.

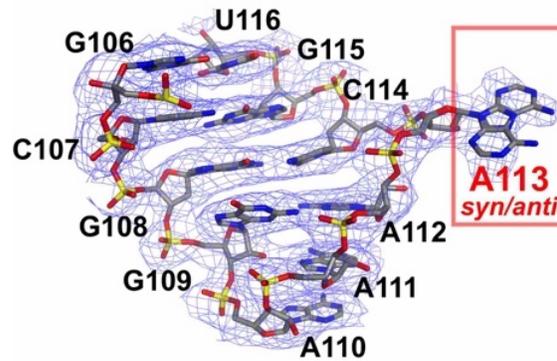


Fig. S5. The structure of the phylogenetically conserved RNase MRP RNA element 5'-GA(G/A)A(G/A)-3' (nucleotides 109-113) and the corresponding electron density. Conserved purine A113 that is observed in both syn- and anti- conformations is shown boxed.

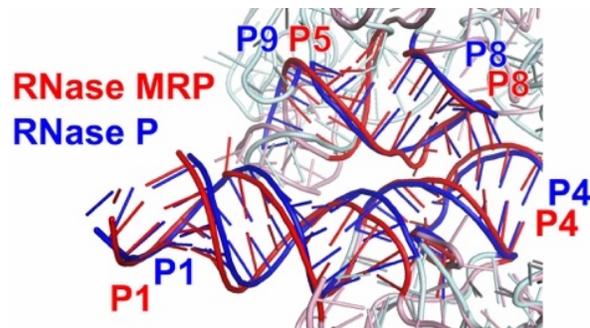


Fig. S6. The docking of the P8/P5 helical stack against the P1/P4 stack in RNase MRP is similar to the docking of the P8/P9 stack against the P1/P4 stack in RNase P. RNase MRP RNA is shown in red; RNase P RNA is shown in blue.

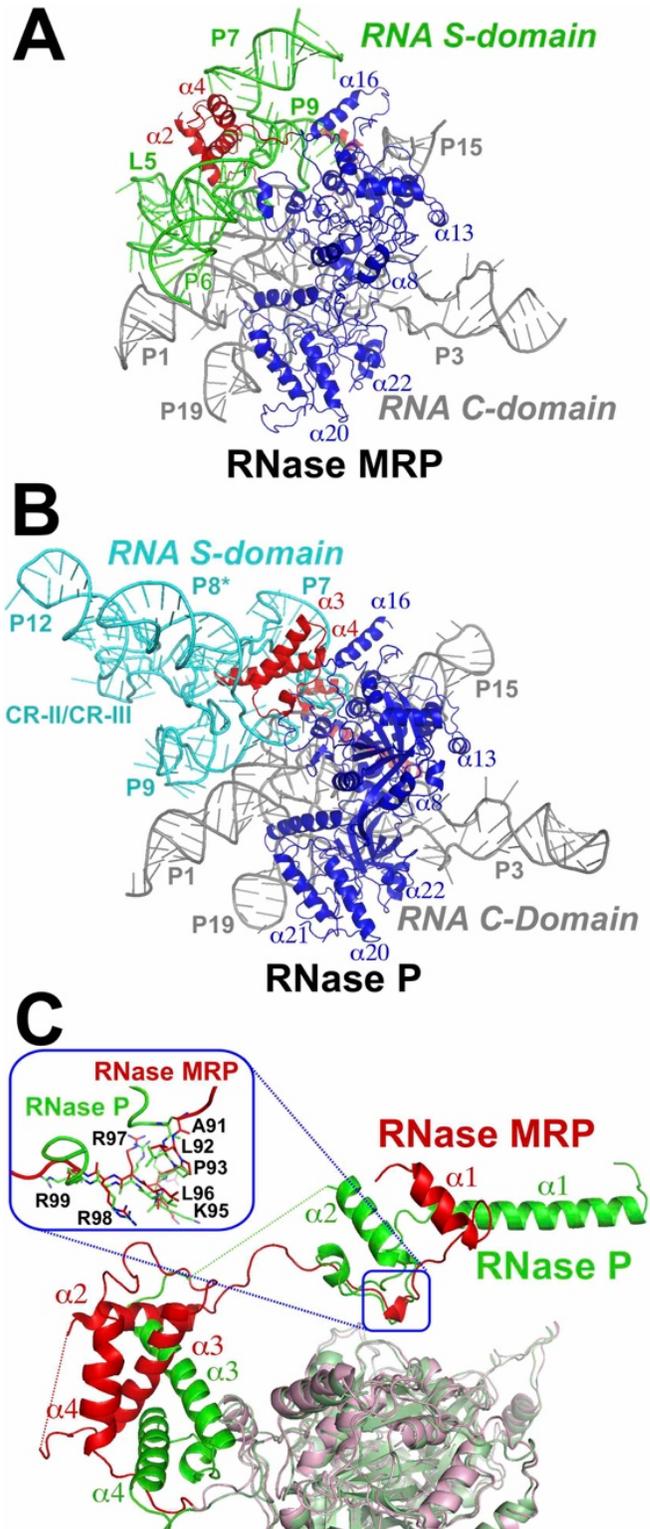


Fig. S7

Fig. S7. Protein Pop1 in RNase MRP and RNase P.

(A, B) The N-terminal part of Pop1 (shown in red) interacts predominantly with the S-domains of RNase MRP (panel A, shown in green) and RNase P (panel B, shown in cyan), whereas the C-terminal part of Pop1 (residues 203-875, shown in blue) interacts with the C-domains of RNase MRP and RNase P (shown in grey).

(C) Superposition of Pop1 in RNase MRP (red and pink) and Pop1 in RNase P (green and light green). The folds of the N-terminal parts (red and green for RNase MRP and RNase P, respectively) differ significantly, with the exception of the residues 91-99, where the two folds converge (insert).

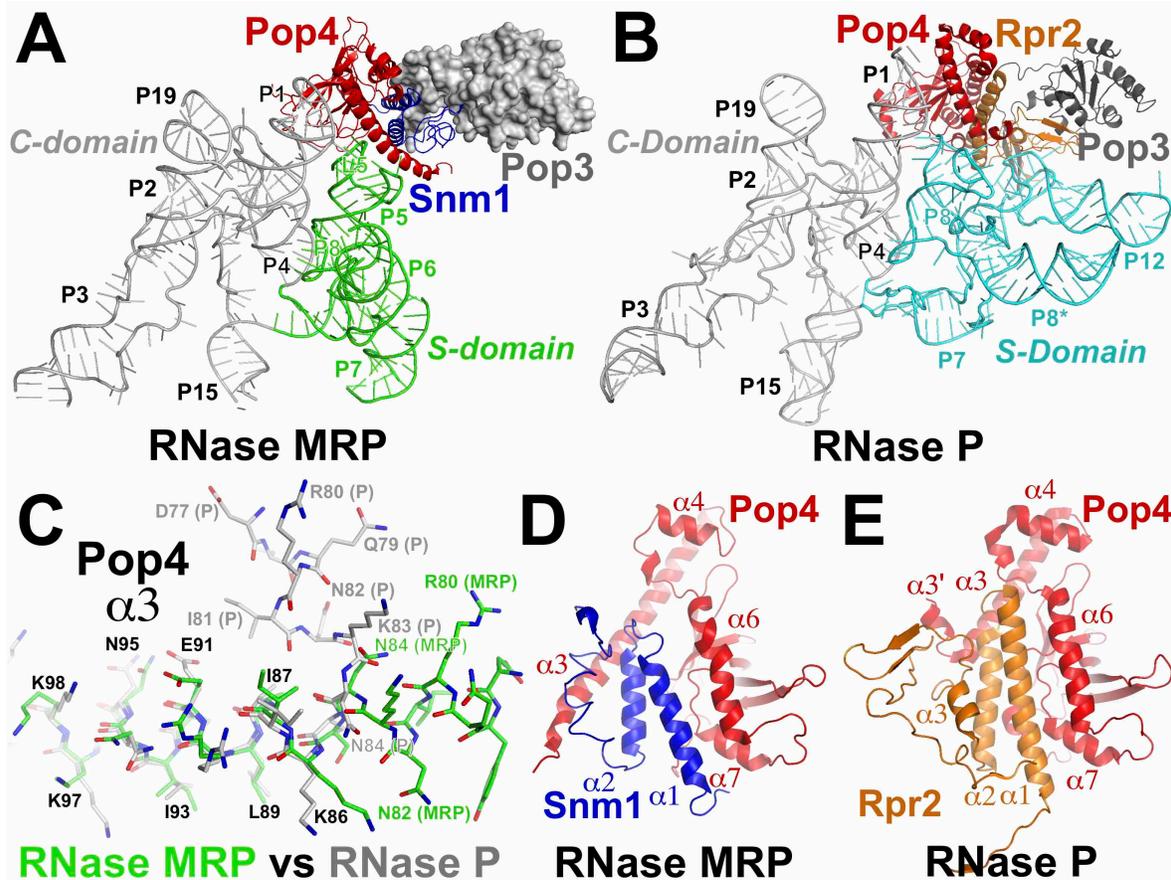


Fig. S8. Proteins Pop4, Snm1, Rpr2, Pop3 in RNase MRP and RNase P.

(A) In RNase MRP, Pop4 (red) bridges stem P1 of the C-domain RNA (grey) and P5/L5 stem/loop of the S-domain RNA (green) and interacts with Snm1 (blue) and Pop3 (grey).

(B) In RNase P, Pop4 (red) bridges stem P1 of the C-domain RNA (grey) and P9 stem of the S-domain RNA (cyan) and interacts with Rpr2 (orange) and Pop3 (grey).

(C) The $\alpha 3$ helix of Pop4 in RNase MRP is disrupted in RNase P. The $\alpha 3$ helix in RNase MRP is shown in green; the corresponding part of Pop4 in RNase P is shown in grey.

(D, E) Interactions of RNase MRP protein Snm1 (blue) with Pop4 (red) in RNase MRP (D) versus interactions of RNase P protein Rpr2 (orange) with Pop4 (red) in RNase P (E).

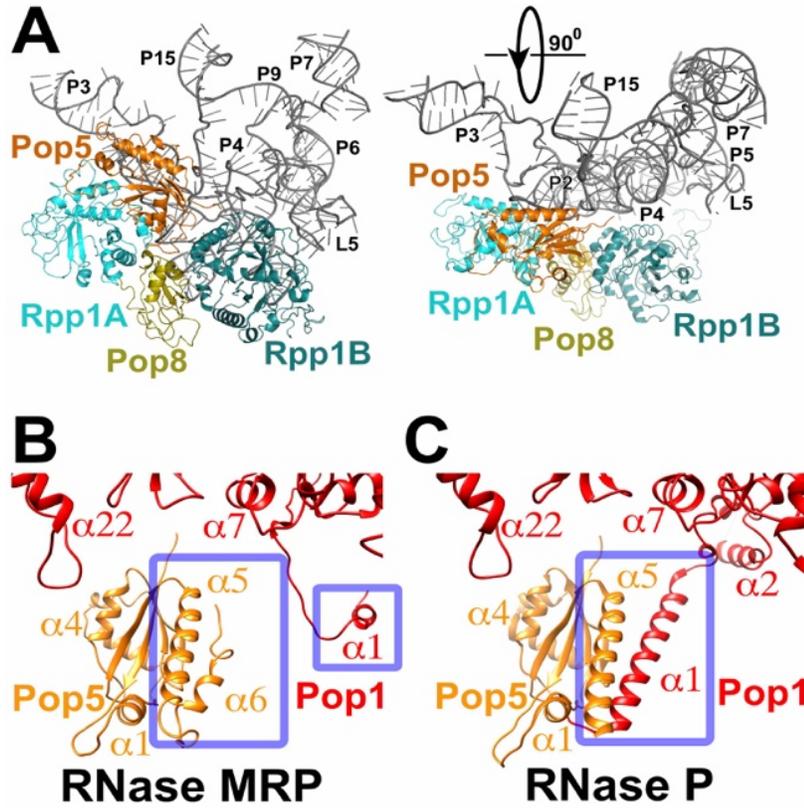


Fig. S9. Proteins Pop5, Pop8, Rpp1 in RNase MRP.

(A) Interactions of the Pop5/Pop8/Rpp1_(A,B) heterotetramer with the RNA component of RNase MRP. Proteins are colored as shown.

(B, C) Differences in the Pop5 fold and interactions with Pop1 in RNase MRP (B) versus RNase P (C). The divergent parts are shown boxed; Pop5 is shown in orange; Pop1 is shown in red.

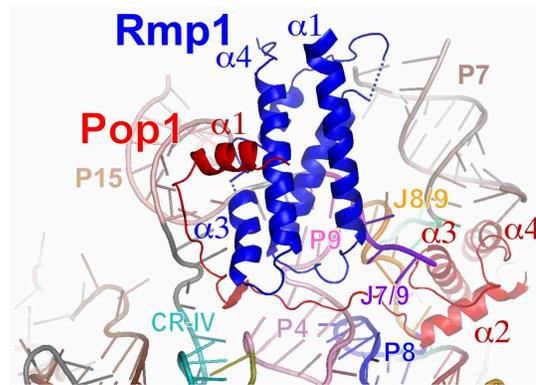


Fig. S10. Protein Rmp1 interacts with the N-terminal elements of Pop1 and RNase MRP RNA. Rmp1 is shown in blue; Pop1 is shown in red; RNase MRP RNA is shown in colors matching Fig. 1A.

Table S1. Cryo-EM data collection, refinement and validation statistics

	RNase MRP (EMD-21564) (PDB 6W6V)
Data collection	
EM equipment	FEI Titan Krios
Detector	Falcon 3EC
Pixel size (Å)	0.886
Electron exposure (e ⁻ /Å ²)	39
Defocus range (µm)	1.1-2.7
Reconstruction	
Software	cisTEM-1.0.0
Number of used particles	155,205
Map resolution (Å)	3.0
FSC threshold	0.143
Model building	
Coot	0.8.9
Refinement	
Software	Phenix 1.17.1
Model composition	
Chains	11
Non-hydrogen atoms	24,850
Protein residues	2306
RNA nucleotides	294
Validation	
R.m.s. deviations	
Bond lengths (Å) (# >4σ)	0.004 (0)
Bond angles (°) (# >4σ)	0.590 (3)
Ramachandran plot	
Favored (%)	88.58
Allowed (%)	11.42
Disallowed (%)	0.0
MolProbity score	1.97
Clashscore	6.89