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Exosome Substrate Targeting: The Long and Short of It

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Keywords: Dis3, exosome, ribonuclease, rRNA-processing protein 44 (Rrp44), rRNA-processing protein 6 (Rrp6), Ski, Trf4-Air2-Mtr4 polyadenylation complex (TRAMP complex).

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Exosome substrate targeting: the long and short of it

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Abstract

The exosome ribonuclease complex functions in both the limited trimming of the 3' ends of nuclear substrates during RNA processing events and in the complete destruction of nuclear and cytoplasmic RNAs. The two RNases of the eukaryotic exosome, Rrp44 and Rrp6, are bound at either end of a catalytically inert cylindrical core. RNA substrates are threaded through the internal channel of the core to Rrp44 by RNA helicase components of the nuclear TRAMP or cytoplasmic Ski complexes. Recent studies reveal that Rrp44 can also associate directly with substrates via channel-independent routes. Although the substrates of the exosome are known, it is not clear whether specific substrates are restricted to one or other pathway. Data currently available support the model that processed substrates are targeted directly to the catalytic subunits, whereas at least some substrates that are directed towards discard pathways must be threaded through the exosome core.

Functions of the Exosome

The exosome ribonuclease complex acts on all types of RNA at some point during their lifetime, functioning in (i) the 3' end maturation of stable RNAs (and some mRNAs); (ii) the complete degradation of fragments such as the 5' ETS (5' external transcribed spacer) that are released as a result of RNA processing reactions; (iii) quality control of all classes of nuclear and cytoplasmic RNAs; (iv) the constitutive degradation of unstable nuclear transcripts such as CUTs (cryptic instable transcripts), and (v) the ultimate degradation of cytoplasmic mRNAs^[1,2].

The exosome complex is conserved throughout eukarya and most archaea, and genetic depletion of any subunit except Rrp6 causes a block in mitotic growth in yeast^[3,4]. The importance of exosome function in human biology is reflected by the

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Abbreviations: CSD, cold-shock domain; CUT, cryptic unstable transcript; Dis3L, DIS3-like exonuclease; 5'-ETS, 5'-external transcribed spacer; HRDC, helicase and RNase D C-terminal; hRRP6, human Rrp6 protein; KH, K homology; PIN, PiIT N-terminus; PNPase, polynucleotide phosphorylase; Rrp, rRNA-processing protein; Ski, superkiller; snoRNA, small nucleolar RNA; snoRNP, small nucleolar ribonucleoprotein; TRAMP complex, Trf4-Air2-Mtr4 polyadenylation complex.

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observations that active site mutations in Rrp44 correlate with the incidence of multiple myeloma and relapsed acute myeloid leukaemia^[5,6], while mutations in the related enzyme Dis3L2 are linked to Perlman Syndrome and Wilm's tumour predisposition^[7]. Genetic complementation analyses underscore the functional conservation of exosome subunits through fungi, plants and metazoa^[3,8-10].

Structure of the Exosome

The eukaryotic exosome has a highly stable "core", consisting of an asymmetric double-layered ring. The lower tier comprises six subunits (Rrp41, Rrp42, Rrp43, Rrp45, Rrp46 and Mtr3) that are structurally related to RNase PH. This is overlaid with an upper tier of three "cap" subunits (Rrp4, Rrp40 and Csl4) that have S1 or KH RNA binding domains^[11,12] (see Figure 1). A structurally homologous complex is found in archaeobacteria^[13], which is itself similar in architecture to the bacterial PNPase (polynucleotide phosphorylase) enzyme^[14]. Catalytically, however, there are clear differences between these complexes. While the archaeal complex contains multiple copies of a fewer number of subunits, the subunits of the eukaryotic exosome have diverged through evolution. The eukaryotic exosome core has lost catalytic activity but the substrate RNAs traverse the same pathway taken through the central channel of the archaeal complex^[12,13]. The eukaryotic complex is directly associated with the eukaryotic-specific ribonucleases Rrp44 (also known as Dis3) and Rrp6^[3,4]. Rrp44 is bound to the core at the "bottom" of the channel near the RNA exit site^[15,16], while Rrp6 is bound to the "top" of the core^[12,17] (Figure 1). Rrp44

and Rrp6 are hydrolytic ribonucleases [18,19], whereas the archaeal complex and PNPase are phosphorolytic enzymes.

The exosome RNase Rrp44

Rrp44 is related to the RNase II/R family of 3'-5' exoribonucleases [20]. The catalytic RNB exonuclease domain is flanked at the N-terminal end by two tandem, cold-shock domains (CSD1 and CSD2) and at the C-terminal end by an S1 domain (Figure 1). The cold shock and S1 domains contribute to RNA binding. In addition to its exonuclease activity, Rrp44 also has a distinct endonuclease activity that is associated with its N-terminal PIN (PiIT N-terminus) domain [21-23]. Full-length Rrp44 protein nevertheless degrades RNA in a 3'-5' direction [3,24], indicating that the endo- and exonuclease activities of Rrp44 are functionally coupled. The PIN domain also mediates the interaction between Rrp44 and the exosome core [23,25]. Both RNase activities of Rrp44 are largely suppressed upon interaction with the exosome through allosteric effects that diminish its ability to bind RNA [24]. In yeast there is a single gene encoding Rrp44 and the protein functions in the nucleolus, the nucleoplasm and the cytoplasm [3,26]. In contrast, three Rrp44 homologues are found in human cells; Dis3 is nucleoplasmic and excluded from the nucleolus, while Dis3L and Dis3L2 are cytoplasmic [27,28]. Dis3L lacks endonuclease activity but is associated with the exosome core, while Dis3L2 lacks the PIN domain and is not associated with the exosome. Dis3L2 plays a role in cytoplasmic mRNA degradation, in both human cells and in the fission yeast *Schizosaccharomyces pombe* [28,29]. Thus, at least some eukaryotic Rrp44 homologues function independently of the exosome core. Indeed, a recent study reported that Rrp44 in budding yeast functions in mitochondria independently of the exosome core [30]. Whether yeast Rrp44 can fully function when separated from the exosome remains to be determined. Notably, expression of the PIN domain alone is able to support slow growth [31].

The exosome RNase Rrp6

Rrp6 belongs to the RNase D family of 3'-5' exoribonucleases that contain a DEDD exonuclease domain juxtaposed to one or more HRDC (helicase and RNase D C-terminal) domains [32]. Rrp6 members of this family also contain an N-terminal PMC2NT domain [33] and a C-terminal exosome interacting domain [12] (see Figure 1). Interaction of Rrp6 with the exosome core has little effect on its exonuclease activity but allosterically stimulates the activity of Rrp44 [24]. Yeast Rrp6 is a nuclear protein [18], while the human Rrp6 protein hRRP6 (also known as PM-Sc100) is restricted to the nucleolus [27]. Three different Rrp6 proteins are expressed in *Arabidopsis thaliana*; Rrp6L1 and Rrp6L2 proteins

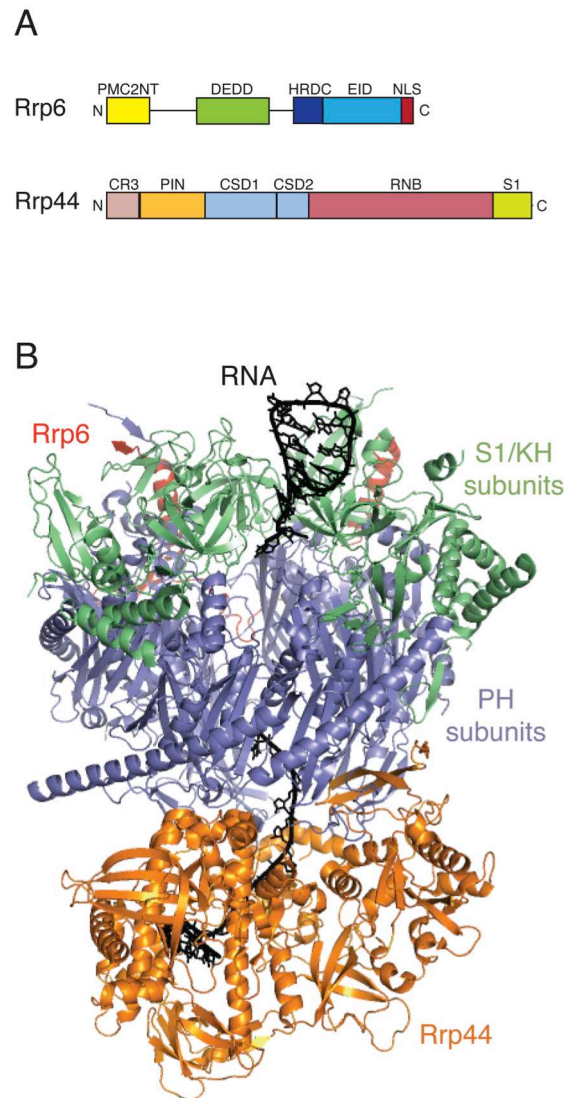


Figure 1. Structure of the exosome. (A) Domain organisation of the catalytic subunits Rrp6 and Rrp44. Rrp6 comprises an N-terminal PMC2NT domain, a DEDD exonuclease domain, an HRDC domain, an Exosome-Interacting domain (EID) and a C-terminal nuclear localization signal (NLS). Rrp44 comprises an N-terminal CR3 motif, a PIN domain, two cold shock domains (CSD), a catalytic RNB domain and an S1 domain. (B) Architecture of the exosome complex bound to RNA. The exosome core consists of a ring of RNase PH-related subunits (in blue) that is overlaid by a second ring of subunits that contain S1 or KH domains (in green). A central channel passes through the core that can bind ~ 30 nucleotides of single-stranded RNA (in black). Rrp44 (in gold) is positioned at the base of the exosome core, with the 3' end of the RNA bound at its exonuclease active site. Rrp6 (the C-terminal exosome-interacting domain is shown in red) is wrapped around the top of the exosome core. The image was generated from PDB accession number 4IFD [12] using MacPyMOL.

are found in the nucleus, while Rrp6L3 is localized to the cytoplasm [9]. Thus, distinct forms of both Rrp44 and Rrp6 have evolved in different systems and presumably carry out specialized functions.

Rrp6 and the small nuclear, RNA binding protein Rrp47 (also known as Lrp1 in yeast, or C1D in human cells) form a heterodimer through an interaction between their N-terminal PMC2NT and Sas10/C1D domains^[34,35]. This interaction provides mutual protein stabilization^[36,37] and limits expression of the Rrp6/Rrp47 complex to the nucleus^[35]. The C-terminal region of Rrp47 mediates interactions with snoRNP components that are critical for the normal 3' end maturation of box C/D snoRNAs^[38]. Thus, the Rrp6/Rrp47 interaction also allows RNA degradation to be coupled to substrate recognition or binding.

Threading substrates through the exosome

Rrp44 alone shields a stretch of 9-12 nucleotides in RNase protection assays, and 31-33 nucleotides when associated with the exosome core^[16]. Substrate binding to Rrp44 requires a single-stranded 3' terminus but, once engaged with the RNA, Rrp44 is a highly processive exoribonuclease that efficiently degrades structured RNA^[11]. The energy released upon hydrolytic activity of Rrp44 is released in bursts, pulling substrates through the complex by a steric occlusion mechanism^[39,40].

The function of the exosome complex *in vivo* is dependent upon additional factors, notably the structurally closely related RNA helicases Mtr4 (also known as Dob1) and Ski2^[41,42]. Yeast Mtr4, together with the noncanonical poly(A) polymerases Trf4 or Trf5 and the RNA-binding proteins Air1 or Air2, constitute the TRAMP complexes that stimulate exosome-mediated RNA degradation *in vitro* and *in vivo*^[43-45]. The human protein hMTR4 is found in a homologous TRAMP complex together with hRRP6 in the nucleolus, as well as being a component of the distinct nuclear exosome targeting (NEXT) complex^[46]. This suggests that hMTR4, like yeast Mtr4, has a major role in targeting nuclear RNAs to the exosome and that it promotes the degradation of some substrates independently of their oligoadenylation. Ski2 is a component of the cytoplasmic Ski complex, which contains the scaffold protein Ski3 and two copies of the protein Ski8^[47,48], and is critical for exosome-mediated cytoplasmic mRNA degradation^[42] and mRNA surveillance pathways^[49-51].

RNase protection assays suggest that the Ski complex is positioned on top of the exosome core^[52], as has been proposed for Mtr4^[53,54]. The RNA helicases Mtr4 and Ski2 are thought to unfold the 3' end of substrates and thread the resultant single-stranded RNA 3' end into the exosome core for subsequent degradation by Rrp44. A number of conserved residues are found at the base of the central DExH core within Mtr4 and Ski2 and on the upper surface of the exosome core, suggesting that they might constitute a mutual binding interface. Nevertheless, additional factors are required for a stable interaction between the RNA helicases and

the exosome core. Mtr4 directly interacts with the exosome-associated protein Mpp6^[55], which may promote binding with the exosome core. Furthermore, hRRP6 is required for stable interaction between hMTR4 and the exosome core^[46]. The interaction between the yeast Ski complex and the exosome is dependent upon another factor, Ski7^[56].

Mutations in Rrp41 (also known as Ski6) and/or Rrp45 that alter conserved residues or obstruct the central channel cause a block in growth in yeast^[24,57]. This suggests that the passage of at least some RNAs through the exosome core is essential for cell growth.

A conformational switch in Rrp44

Recent structural analyses resolve two distinct structural conformations of Rrp44 bound to the exosome core complex^[58] (see Figure 2). In the presence of short structured RNAs, Rrp44 that is associated with the exosome core has an extended structure and the substrate can directly access either the exonuclease domain or the PIN domain. In this state, the CSD2 domain of Rrp44 is positioned at the RNA exit site at the bottom of the core, blocking the pathway of the substrate through the core. In the presence of longer RNAs, Rrp44 undergoes a substantial conformational change and becomes more compact. The exonuclease domain is rotated $\sim 120^\circ$, bringing it close to the RNA exit site and displacing the CSD2 domain. In this conformation, Rrp44 is able to digest RNA that is threaded through the channel. These data suggest that Rrp44 bound to the exosome core can switch between an extended conformation that is accessible to direct substrate binding, and a compact conformation that is aligned for interaction with threaded substrates.

The path(s) taken by RNA to reach Rrp6 is less clear. An Rrp6 mutant lacking the C-terminal exosome-interaction domain is functional *in vivo*, indicating that at least some substrates can bind to Rrp6 independently of the exosome complex^[59]. In the context of the exosome complex, the RNA binding activities of Rrp6 and Rrp44 are interconnected. The interaction between Rrp6 and the exosome core causes an increased RNA binding activity of Rrp44. Conversely, the presence of a substrate that is locked into the threaded pathway, by mutation of the Rrp44 exonuclease active site, decreased the RNA binding activity of Rrp6^[24]. It was suggested that Rrp6 substrates are threaded into the central channel of the exosome core and then diverted laterally to Rrp6. However, recent structures of the yeast exosome complex do not support such a pathway^[12]. Alternatively, the interrelated RNA-binding activities of exosome-associated Rrp6 and Rrp44 could reflect an allosteric mechanism that is mediated through the exosome core. Thus, commitment of substrates to

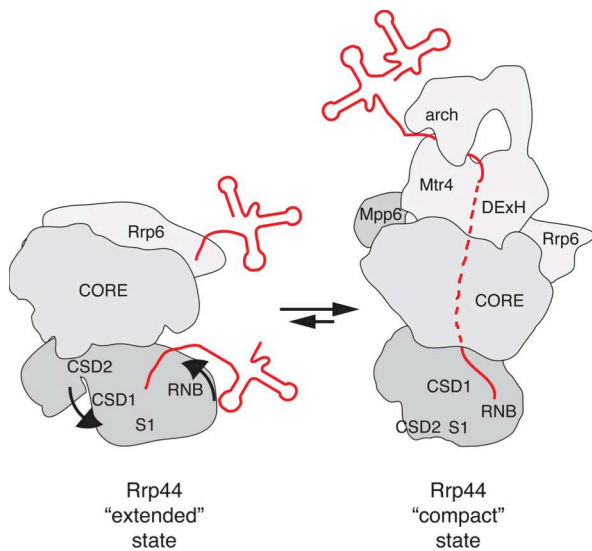


Figure 2. Exosome substrate targeting pathways.

Exosomes are observed in two distinct physical states that differ in the "extended" or "compact" conformational state of Rrp44. In the extended state, RNA substrates (in red) can directly access the active sites of Rrp44 (and presumably that of Rrp6). Upon switching into the "compact" state, the RNB exoribonuclease domain of Rrp44 is rotated into the space directly beneath the core (the movement is indicated by the black arrows). Structured RNAs targeted to discard pathways are threaded into the exosome core by the RNA helicase Mtr4. The portion of the RNA pathway that passes through the inner channel of the DExH domain and the exosome core is indicated by a dashed line. Stable binding of Mtr4 to the exosome core may require either Rrp6 or Mpp6. Adoption of a compact structure by Rrp44 may lead to an allosteric inhibition of Rrp6 RNase activity.

the threaded pathway to Rrp44 might suppress access of substrates to the catalytic domain of Rrp6, thereby ensuring that a single exosome complex can only be engaged with a substrate targeted to the one or other enzyme (see Figure 2).

Long and short pathways for distinct RNAs

Deletion of the exosome interacting domain of Rrp6 does not impair its function in box C/D snoRNA 3' maturation or the trimming of the "5.8S+30" processing intermediate, but does cause a defect in discard pathways that degrade the 5' ETS fragment and truncated degradation intermediates of 5S rRNA^[59]. Furthermore, *in vivo* cross-linking studies have shown that Rrp44 and Rrp6 bind directly to 7S pre-rRNA and snoRNA processing intermediates but very few contacts were observed with exosome core components^[60]. These data suggest that substrates that undergo limited trimming reactions are normally targeted directly to Rrp44 or Rrp6, whereas at least some substrates destined for complete degradation are dependent upon threading through the exosome core. It is unclear whether Rrp6 can efficiently degrade its other substrates, such as pre-mRNAs,

CUTs/SUTs, tRNAs and snRNAs, when uncoupled from the exosome core.

Strikingly, the subset of Rrp6 substrates that accumulate in *rrp6 mpp6* double mutants comprise those that are destined for complete degradation (CUTs, pre-mRNAs, 5' ETS, truncated stable RNAs, snoRNA transcription read-through products) and exclude those that undergo limited 3' end processing reactions (5.8S rRNA, box C/D snoRNAs)^[61,62]. Given that Rrp6 substrates targeted to degradation are dependent upon the interaction between Rrp6 and the exosome core, this suggests that the synthetic lethal relationship between loss of function *rrp6* and *mpp6* alleles^[61] might reflect redundant functions of Rrp6 and Mpp6 in facilitating substrate threading into the exosome core. This model is supported by the direct physical interaction reported between hMPP6 and hMTR4^[55] and the observation that the stable interaction between hMTR4 and the exosome core is dependent upon hRRP6^[46].

Although the exosome was initially characterised through its role in the 3' processing of 5.8S rRNA^[3], the overwhelming majority of exosome substrates are targeted to RNA discard pathways^[60,63]. The exosome is able to degrade extensively structured pre-ribosomal RNP particles in a highly processive manner. It is not clear how substrates destined for limited processing could be effectively differentiated from those that are targeted to discard pathways if they were both engaged in a highly processive mode of degradation. One function proposed for the endonuclease activity of Rrp44 is to release substrates that are stalled in the exosome channel, thereby salvaging blocked exosome complexes^[64]. The endonuclease activity of Rrp44 is not, however, required for exosome-mediated RNA processing events in wild-type cells. Thus, such a role presumably represents a fail-safe recycling function rather than an obligatory step in RNA processing.

Which RNAs might be degraded by Rrp44 using direct, core-independent pathways, and how would these RNAs be targeted without threading through the exosome core? One major class of such substrates might be incorrectly folded tRNAs, which can be targeted directly to Rrp44^[65,66]. Notably, tRNA represents a significant proportion of the RNA subjected to cellular RNA discard pathways^[63]. Other small, structured RNAs targeted to discard pathways may well share this route. Such substrates are likely to require 3' tailing by TRAMP^[65] or another nucleotidyltransferase^[66] to facilitate efficient docking with Rrp44. A major current objective is to define the set of RNAs that are threaded through the exosome core and those that are targeted directly to either Rrp44 or Rrp6.

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