Exon Skipping in the RET Gene Encodes Novel Isoforms That Differentially Regulate RET Protein Signal Transduction*

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Received for publication, December 11, 2015, and in revised form, May 9, 2016. Published, JBC Papers in Press, May 23, 2016, DOI 10.1074/jbc.M115.709675

Rearranged during transfection (RET), a receptor tyrosine kinase that is activated by the glial cell line-derived neurotrophic factor family ligands (GFLs), plays a crucial role in the development and function of the nervous system and additionally is required for kidney development and spermatogenesis. RET encodes a transmembrane receptor that is 20 exons long and produces two known protein isoforms differing in C-terminal amino acid composition, referred to as RET9 and RET51. Studies of human pheochromocytomas identified two additional novel transcripts involving the skipping of exon 3 or exons 3, 4, and 5 and are referred to as RET∆E3 and RET∆E345, respectively. Here we report the presence of RET∆E3 and RET∆E345 in zebrafish, mice, and rats and show that these transcripts are dynamically expressed throughout development of the CNS, peripheral nervous system, and kidneys. We further explore the biochemical properties of these isoforms, demonstrating that, like full-length RET, RET∆E3 and RET∆E345 are trafficked to the cell surface, interact with all four GFRα co-receptors, and have the ability to heterodimerize with full-length RET. Signaling experiments indicate that RETΔE3 is phosphorylated in a similar manner to full-length RET. RET∆E345, in contrast, displays higher baseline autophosphorylation, specifically on the catalytic tyrosine, Tyr1065, and also on one of the most important signaling residues, Tyr1062. These data provide the first evidence for a physiologic role of these isoforms in RET pathway function.

RET is a receptor tyrosine kinase that is critical for kidney morphogenesis, spermatogenesis, and development of the nervous system (1–4). RET is activated by a family of four growth factors known as the glial cell line-derived neurotrophic factor (GDNF) family ligands (GFLs), which includes GDNF, neurturin, artemin, and persephin (1). Each GFL binds to one of four cognate glycosylphosphatidylinositol-anchored co-receptors known as the GDNF family receptor αs (GFRαs) (2, 5). The GFL-GFRα complex binds to RET, inducing RET dimerization and subsequent autophosphorylation on multiple tyrosine residues within the intracellular tyrosine kinase domain. This enhances tyrosine kinase activity and initiates the association of adaptor proteins and enzymes that trigger multiple second messenger cascades (6).

The presence of two major Ret isoforms, RET9 and RET51, has been extensively described in the literature, and a third isoform, RET43, has also been observed in humans (7–9). Ret is 20 exons long, and RET9 and RET51 transcripts differ in alternative splicing of intron 19. Intron 19 in the RET51 transcript is excised properly, whereas, in the RET9 transcript, the intron is retained, changing the reading frame and inserting a premature stop codon into the amino acid sequence. This creates a unique nine-amino acid C-terminal sequence for RET9 and a unique 51-amino acid C-terminal sequence for RET51. Interestingly, these two different isoforms display marked differences in their degradation and function (7, 10, 11).

Three additional Ret transcripts have been reported in various tumor sources as well as adult human tissues (12). These novel transcripts are a product of exon skipping in the 5′ region of RET, which encodes for the extracellular domain of the protein. Skipping of exons 3 (RET∆E3) or exons 3, 4 and 5 (RET∆E345) gives rise to transcripts that encode for full-length Ret proteins but with deletions in the extracellular domain, specifically cadherin-like domain 1 (CLD1) or CLD1–3, respectively. These deletions are hypothesized to change the extracellular domain structure as well as binding to GFL-GFRα complexes and may impact the overall stability of the proteins. The skipping of exons 3 and 4 results in a frameshift that culminates in a premature stop. This protein encodes only a small portion of the RET extracellular domain with no transmembrane or intracellular domain. Because this transcript is likely to be subjected to nonsense-mediated decay, it is unlikely to be involved in RET signal transduction.

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Here we show that \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) transcripts are conserved in vertebrates and that the mRNA and proteins of these splice variants are expressed throughout the nervous system in mice. We also show that these isoforms are trafficked to the cell surface and that both isoforms interact with all four GFRs. Additionally, we find that \( \text{RET}^{\Delta E/3} \) is phosphorylated to a similar level as full-length \( \text{RET} \) and is activated in a GDNF-dependent manner. However, \( \text{RET}^{\Delta E/345} \) displays higher baseline autophosphorylation, specifically on the catalytic tyrosine Tyr\(^{905}\), and also on an additional signaling tyrosine, Tyr\(^{1062}\). Interestingly, \( \text{RET}^{\Delta E/345} \) is not activated in a GDNF-dependent manner. Taken together, these isoforms may have unique and important unidentified roles in the development and maintenance of the nervous system and kidneys as well as in the pathophysiology of neuroendocrine gland diseases.

Results

Exons 3, 4, and 5 of Ret Are Not Highly Conserved, but \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) Transcripts Are Observed in Several Organisms—It has been described previously that the intracellular domain of RET, which contains a tyrosine kinase domain, is more highly conserved than the extracellular domain (13). To better understand the level of conservation of amino acids encoded by each exon, we compared sequences from zebrafish (\textit{Danio rerio}), mouse (\textit{Mus musculus}), rat (\textit{Rattus norvegicus}), and human (\textit{Homo sapiens}). We found that the entire intracellular domain was more than 75% conserved at the amino acid level. This was not surprising because mutations in many of these highly conserved regions give rise to loss of function (\textit{e.g.} Hirschsprung disease) or gain of function (\textit{e.g.} multiple endocrine neoplasia type 2A (MEN2A) and 2B (MEN2B)) phenotypes (Fig. 1A). Interestingly, exons 3, 4, and 5 are less than 50% conserved between these species. Although this lack of conservation is suggestive that these regions of RET are not as functionally important, it could also be interpreted as allowing for amino acid flexibility between the different RET isoforms.

Our initial sequence analysis suggested that \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) transcripts could be expressed in additional vertebrates other than \( H. \text{sapiens} \). These transcripts were originally identified in human kidney and substantia nigra fetal tissues (12). Sequence analysis demonstrated that exon skipping also encodes potential full-length transcripts in zebrafish, mice, and rats (Fig. 1A). To determine whether these novel transcripts were expressed in these organisms, species-specific primers were created to identify each of the Ret transcripts: full-length Ret, \( \text{Ret}^{\Delta E/3} \), and \( \text{Ret}^{\Delta E/345} \). RT-PCR analysis identified the presence of \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) from 48 hours post fertilization zebrafish embryos, E19.5 rat dorsal root ganglia, and E18.5 mouse spinal cord (Fig. 1B). Sequence analysis confirmed these amplicons to be \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) (data not shown). This is the first time these transcripts have been identified in vertebrates other than \( H. \text{sapiens} \). Taken together, these data indicate that 5’ exon skipping is conserved and results in the expression of \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) transcripts.

Exon Skipping Creates Large Deletions in the Extracellular Domain of RET—To understand the impact deletions in the extracellular domain would have on RET, we analyzed the deletions in terms of a CLD1–4 model derived from electron microscopy and small angle x-ray scattering analyses (14). The main consequence of deletion of exon 3 is the removal of half of CLD1 (residues 113–150) and half of CLD2 (residues 154–208). Each domain adopts a \( \beta \) sandwich cadherin fold comprised of two \( \beta \) sheets separated by numerous buried hydrophobic residues. Elimination of exon 3 removes one sheet from CLD1 and several \( \beta \) strands from both sheets of CLD2 (Fig. 1C). Therefore, it is likely that both CLD domains are substantially perturbed leaving \( \beta \) strands without the cadherin fold structure and exposing some hydrophobic residues. Exon 3 removal also eliminates a highly constrained Cis-Pro loop (Cys\(^{137}\)–Cys\(^{142}\)) unique to higher vertebrate RET sequences. The portion of CLD1–2 removed coincides with regions stabilizing the clamb shell arrangement of CLD1–2 (15). This would result in a more open and extended arrangement for the remaining parts of CLD1 and CLD2 with much greater flexibility between the domains. An odd number of cysteine residues (three in total) are deleted by removing exon 3, including the Cys\(^{157}\)–Cys\(^{197}\) disulfide. Cys\(^{166}\) is also removed, leaving its partner Cys\(^{243}\) unpaired but buried. The two unpaired cysteines, Cys\(^{197}\) and Cys\(^{216}\), which constitute a known folding bottleneck in wild-type RET, are left untouched. We expect \( \text{Ret}^{\Delta E/3} \) to be a less stable transmembrane protein with a short half-life that retains at least some ligand-binding properties.

For \( \text{RET}^{\Delta E/345} \), removal of exons 3, 4, and 5 effectively eliminates much of CLD1–3 but leaves CLD4-cysteine-rich domain (CRD) intact (Fig. 1C). This region by itself was shown to be insufficient to bind ligand but retains at least one important binding epitope (\textit{i.e.} necessary for ligand binding but not sufficient) (14). Many receptor tyrosine kinases have extracellular domains that have autoinhibitory functions in the absence of ligand. Deletion of such regions can lead to autoactivation in a ligand-independent manner. By analogy with these other systems, removal of CLD1–3 could leave a truncated form of RET that would be constitutively activated in the absence of ligand and likely would be unresponsive to the GFLs.

\( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) Are Co-expressed in Several Tissues with Ret throughout Development—To determine where and when \( \text{Ret}^{\Delta E/3} \) and \( \text{Ret}^{\Delta E/345} \) transcripts are expressed and whether their expression levels change over time, we analyzed tissues in which full-length Ret is expressed at several developmental time points. Within the central nervous system, we analyzed the brain, as signaling via Ret is important for the survival of central noradrenergic neurons and dopaminergic neurons in the ventral midbrain, as well as the spinal cord, where GDNF/ GFRA1/Ret signaling is critical for axon guidance into the hind limbs and survival of \( \gamma \) motor neurons (16–21). In the peripheral nervous system, we analyzed sensory neurons of the dorsal root ganglia (DRG), where neurturin/GFRA2/Ret signaling is important for the survival and maintenance of mechanoreceptors during development (22, 23). Additionally, shortly after birth, a subclass of nociceptors transition from being TRKA-positive to expressing GFRA1/Ret to support the development and survival of nonpeptidergic nociceptors (24–26). Sympathetic neurons of the superior cervical ganglia (SCG) were also analyzed, as it has been well established that artemin/GFRA3/Ret signaling is necessary for proper sympathetic chain migra-
tion and axon guidance during development (27–29). Last, the kidney was analyzed because GDNF/GFRα1/Ret signaling is crucial for ureteric bud branching and morphogenesis of the kidney (30–34). Using qPCR, we examined the relative expression of Ret, RetΔE3, and RetΔE345 in each of these tissues in mice and found that they were detected in all five tissues. The relative expression of Ret, RetΔE3, and RetΔE345 was lower in the brain, spinal cord, and kidney (Fig. 2A) compared with the DRG and SCG (Fig. 2B). Interestingly, in the spinal cord, DRG, and SCG, there was a significant increase of RetΔE3 expression at E19.5 that preceded a significant increase of Ret at P3. The expression of all three transcripts was highest in the DRG, particularly at E19.5 and P3, when the IB4+ subpopulation of nociceptors is emerging.

5’ and 3’ Alternative Splicing of Ret transcripts Are Not Mutually Exclusive—Alternative splicing of intron 19 in Ret allows for the translation of two isoforms of Ret, RET9 and RET51. We sought to determine whether alternative splicing could occur simultaneously, both 5’ and 3’ in Ret transcripts, allowing for increased diversity of encoded RET proteins. cDNA was isolated from E19.5 mouse DRGs, as we found this tissue to have the highest relative expression of Ret, RetΔE3, and RetΔE345 (Fig. 2B). Forward primers specific for each 5’ splicing event were paired with reverse primers that would detect either Ret9 or Ret51 alternative splicing. Full-length Ret9 and Ret51 transcripts were detected, as expected, at 3.33 kb and 4.38 kb, respectively. We observed RetΔE3 and RetΔE345 with intron 19 retention, thus encoding RetΔE3 and RetΔE345 transcripts with

FIGURE 1. Removal of exons 3 or 3, 4, and 5 of RET creates proteins with large deletions in the extracellular domain, and transcripts encoding these receptors are expressed in multiple organisms. A, comparison of the amino acid sequences of RET between zebrafish, mice, rats, and humans was performed, and the conservation within each exon is indicated by color. Interestingly, the amino acids encoded in exons 3, 4, and 5 are the least conserved. Additionally, the impact of exon skipping on the extracellular domain within the protein is shown. B, RetΔE3 and RetΔE345 transcripts are detectable in zebrafish (D. rerio), mice (M. musculus), and rats (R. norvegicus) by qPCR. Like human RETΔE3 and RETΔE345, alternative splicing of exon 3 or exons 3, 4, and 5 in these vertebrates are predicted to encode full-length, transmembrane RET proteins with deletions in the extracellular domain. C, deleted regions are shown in red, projected onto a structural model for RET cadherin domains 1–4 derived from PDB code 4UX8.
amplicons of 3.05 kb and 2.61 kb, respectively. However, we were only able to detect \textit{Ret}/H9004 E345 with a properly excised intron 19 at an amplicon size of 3.66 kb, encoding a \textit{Ret51}/H9004 E345 transcript. Although we were unable to detect a \textit{Ret51}/H9004 E3 transcript (4.10 kb), this is likely due to the generally low expression levels of \textit{Ret}/H9004 E3.

Taken together, these data indicate that the \textit{Ret} locus encodes for at least three previously unidentified transcripts and at least five total Ret isoforms: \textit{Ret9}, \textit{Ret9}/H9004 E3, \textit{Ret9}/H9004 E345, \textit{Ret51}, and \textit{Ret51}/H9004 E345.

\textbf{RET Proteins of the Same Molecular Weight as RET\textsuperscript{E345} Are Detected in Vivo}—Although we can detect \textit{Ret\textsuperscript{AE3}} and \textit{Ret\textsuperscript{AE345}} mRNA transcripts, which are likely being processed for translation, we sought to detect the presence of these isoforms at the protein level. Plasmids encoding human \textit{RET51}, \textit{RET51}/H9004 E3, and \textit{RET51}/H9004 E345 were transfected into NIH/3T3 cells, and lysates were run as size standards to aid in detecting Ret isoforms of the appropriate molecular weights from E18.5 mouse brain. The molecular weight of native \textit{RET51} protein is 120 kDa, but

\begin{figure}[h]
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\caption{\textit{Ret\textsuperscript{AE3}} and \textit{Ret\textsuperscript{AE345}} are dynamically expressed throughout development, and RET proteins with similar molecular weights as \textit{RET\textsuperscript{AE345}} are detect \textit{in vivo}. A, expression of \textit{Ret}, \textit{Ret\textsuperscript{AE3}}, and \textit{Ret\textsuperscript{AE345}} in mouse brain (E15.5, E19.5, P3), spinal cord (E15.5, E19.5, P3, adult), and kidney (E15.5, E19.5, P3, adult) by qPCR is shown as relative expression compared with actin. B, expression of \textit{Ret}, \textit{Ret\textsuperscript{AE3}}, and \textit{Ret\textsuperscript{AE345}} in mouse sympathetic (E19.5, P3, adult) and DRG sensory neurons (E15.5, E19.5, P3, adult) by qPCR is graphed as in A. C, cDNA from E19.5 DRGs was synthesized from poly(A) RNA, and RT-PCR was performed to determine the presence of \textit{Ret} transcripts with both 5' and 3' alternative splicing. \textit{Ret9}, \textit{Ret9}/H9004 E3, \textit{Ret9}/H9004 E345, \textit{Ret51}, and \textit{Ret51}/H9004 E345 transcripts were identified. D, E18.5 mouse brains were lysed, and immunoprecipitations (IP) for \textit{RET9} or \textit{RET51} were performed. Immunoblotting for \textit{RET} detected full-length \textit{RET9} and \textit{RET51}, as expected. The presence of \textit{RET9} and \textit{RET51} proteins at the same molecular weight as \textit{RET\textsuperscript{AE345}} suggested the existence of \textit{RET\textsuperscript{AE345}} and \textit{RET\textsuperscript{AE345}} isoforms.}
\end{figure}

\begin{enumerate}
\item \textbf{A}, expression of \textit{Ret}, \textit{Ret\textsuperscript{AE3}}, and \textit{Ret\textsuperscript{AE345}} in mouse brain (E15.5, E19.5, P3), spinal cord (E15.5, E19.5, P3, adult), and kidney (E15.5, E19.5, P3, adult) by qPCR is shown as relative expression compared with actin.
\item \textbf{B}, expression of \textit{Ret}, \textit{Ret\textsuperscript{AE3}}, and \textit{Ret\textsuperscript{AE345}} in mouse sympathetic (E19.5, P3, adult) and DRG sensory neurons (E15.5, E19.5, P3, adult) by qPCR is graphed as in A.
\item \textbf{C}, cDNA from E19.5 DRGs was synthesized from poly(A) RNA, and RT-PCR was performed to determine the presence of \textit{Ret} transcripts with both 5' and 3' alternative splicing. \textit{Ret9}, \textit{Ret9}/H9004 E3, \textit{Ret9}/H9004 E345, \textit{Ret51}, and \textit{Ret51}/H9004 E345 transcripts were identified.
\item \textbf{D}, E18.5 mouse brains were lysed, and immunoprecipitations (IP) for \textit{RET9} or \textit{RET51} were performed. Immunoblotting for \textit{RET} detected full-length \textit{RET9} and \textit{RET51}, as expected.}
\end{enumerate}
Novel RET Isoforms Differentially Regulate RET Signaling

because of posttranslational modifications, specifically glycosylation of the extracellular domain, RET51 has a molecular weight of ~180 kDa. The native molecular weights of RET\(\Delta E3\) and RET\(\Delta E345\) were predicted to be 109 kDa and 93 kDa, respectively. In transfected cells, we consistently observe the mature, processed proteins to have molecular weights of ~150 kDa (RET\(\Delta E3\)) and ~125 kDa (RET\(\Delta E345\)) (Fig. 2D).

Whole brains from Ret\(^{+/+}\) or Ret\(^{-/-}\) E18.5 littermate mice were lysed and divided in two, and immunoprecipitations were performed to isolate either RET9 or RET51 proteins. Although the qPCR data suggested that Ret transcripts have an overall lower relative expression in the brain compared with PNS tissues, the brain was chosen for analysis because of the abundant amount of protein that could be isolated. Ret\(^{-/-}\) mice were used as a control to demonstrate the specificity of RET immunoblotting and to show the potential presence of RET proteins at the same molecular weight as RET\(\Delta E3\) and RET\(\Delta E345\). Because we previously observed more highly expressed Ret9\(\Delta E345\) and Ret51\(\Delta E345\) transcripts compared with those of Ret\(\Delta E3\), we hypothesized that we would be most able to detect both RET9\(\Delta E345\) and RET51\(\Delta E345\) proteins. By immunoblotting for RET, we were able to detect full-length RET9 and RET51 proteins, as expected (Fig. 2D). Additionally, we observed RET9 and RET51 bands with similar molecular weights to RET\(\Delta E345\), suggesting the presence of RET9\(\Delta E345\) and RET51\(\Delta E345\) proteins in mouse brain (Fig. 2D). Because the antibodies used for immunoprecipitations of RET9 and RET51 are to the C terminus, and the RET antibody for immunoblotting is to the common N-terminal region, this RET protein is unlikely to be a degradation product. However, we cannot exclude the possibility that this is an immature, non-glycosylated, full-length RET protein.

RET\(\Delta E3\) and RET\(\Delta E345\) Are Trafficked to the Cell Surface—To examine the biochemical properties of RET\(\Delta E3\) and RET\(\Delta E345\), we transitioned to an in vitro system for these experiments. Although RET\(\Delta E3\) and RET\(\Delta E345\) encode proteins that may be capable of signaling, this does not guarantee that they will be correctly trafficked to the cell surface for activation. To determine whether RET\(\Delta E3\) and RET\(\Delta E345\) are trafficked to the plasma membrane, NIH/3T3 cells were transfected with C-terminally epitope-tagged Ret constructs that also contain an IRES-GFP to serve as an indicator for transfected cells. Taking advantage of a RET antibody specific for the extracellular domain and using an HA antibody to label the C terminus of RET, we were able to selectively visualize RET located on the plasma membrane. Full-length RET was detected on the cell surface of transfected NIH/3T3 cells, as expected (Fig. 3A). Permeabilization revealed the C-terminal HA tag, confirming the integrity of the plasma membrane (Fig. 3P). Similar to full-length RET, both RET\(\Delta E3\) and RET\(\Delta E345\) were trafficked to the cell surface, as determined with the extracellular RET antibody in non-permeabilized cells (Fig. 3, S and A). Transfection of GFP alone showed no labeling with the RET and HA antibodies, regardless of permeabilization, confirming the specificity of our immunostaining (Fig. 3, A–H). A cell surface biotinylation experiment was performed as a secondary measure to confirm these results, and we again observed trafficking of full-length RET, RET\(\Delta E3\), and RET\(\Delta E345\) to the cell surface (data not shown). Thus, RET\(\Delta E3\) and RET\(\Delta E345\) are trafficked properly to the cell surface, where they can participate in GFL-mediated activation.

RET\(\Delta E3\) and RET\(\Delta E345\) Bind to GFRα1, GFRα2, GFRα3, and GFRα4—Previous structural analysis (Fig. 1C) suggested that interactions between RET\(\Delta E3\) and the GFRαs may be unstable and that RET\(\Delta E345\) would be unlikely to bind to the co-receptors because of deletions of important binding residues in CLD1 and CLD3. To confirm these results, we co-transfected HEK293T cells with RET\(\Delta E3\) or RET\(\Delta E345\) and HA epitope-tagged GFRα constructs (HA::GFRα1 (mouse), HA::GFRα2 (rat), HA::GFRα3 (mouse), and HA::GFRα4 (chicken)). Immunoprecipitations were performed using a stringent immunoprecipitation buffer (modified radioimmune precipitation assay buffer) to eliminate any weak, nonspecific interactions. Surprisingly, we found that both RET\(\Delta E3\) and RET\(\Delta E345\) bind to each of the four GFRαs (Fig. 4). Co-transfection of the GFRαs with TrkA, another receptor tyrosine kinase, was performed as a negative control to show selectivity of binding between the GFRαs and RET (Fig. 4). As expected, TrkA did not associate with any of the GFRαs. There were no apparent differences between the four different GFRαs in regard to the extent of their association with RET\(\Delta E3\) and RET\(\Delta E345\), suggesting that these splice variants could potentially be activated by all four GFLs (Fig. 4).

RET\(\Delta E3\) and RET\(\Delta E345\) Heterodimerize with Full-length RET—In addition to being potential signaling molecules on their own, we observed that these transcripts are normally expressed in tissues where full-length RET is also expressed (Fig. 2). Although these isoforms may be present in different cells within these tissues, these observations raise the question of whether RET\(\Delta E3\) and RET\(\Delta E345\) could heterodimerize with full-length RET and affect its function. To test this possibility, HEK293T cells were co-transfected with HA epitope-tagged RET51\(\Delta E345\) and HA epitope-tagged RET9\(\Delta E3\) and RET51\(\Delta E3\) associated with full-length RET51 and that RET\(\Delta E3\) and RET51\(\Delta E345\) could also bind to one another (Fig. 5B). We found that both RET51\(\Delta E3\) and RET51\(\Delta E345\) associated with full-length RET51 and that RET\(\Delta E3\) and RET51\(\Delta E345\) associated with full-length RET51 and that RET\(\Delta E3\) and RET51\(\Delta E345\) could also bind to one another (Fig. 5B). This is consistent with former experiments suggesting that overexpression of RET proteins allows for ligand-independent dimerization via the transmembrane domain (35).

RET\(\Delta E345\) Has Increased Ligand-independent Activation Compared with RET and RET\(\Delta E3\)—To determine levels of activation of RET\(\Delta E3\) and RET\(\Delta E345\), the basal phosphorylation of tyrosine residues of the RET isoforms were compared with full-length RET phosphorylation. Overexpression of the RET isoforms in vitro, either in the presence or absence of a GFRα, allows for ligand-independent dimerization and activation of RET. The level of activation can be assessed by determining the summed level of all of the phosphorylated tyrosines using a pan-phosphotyrosine antibody, or individual tyrosine residues can be analyzed using residue-specific RET phosphotyrosine antibodies. We evaluated the level of total phosphotyrosine...
FIGURE 3. RET^{AE3} and RET^{AE345} are trafficked to the cell surface. NIH/3T3 cells were transfected with GFP (A–H), Ret51::HA (I–P), Ret51^{AE3}::HA (Q–X), or Ret51^{AE345}::HA (Y–F). Constructs for the RET51 isoforms were bicistronic, allowing for expression of GFP as an indicator for positive transfection, which was encoded by an IRES-GFP following RET cDNA sequences. Cells were washed in 1× PBS and fixed briefly in 4% paraformaldehyde 24 h post-transfection. Cells were then blocked in immunofluorescence blocking solution with or without Triton X-100 to allow for cell membrane permeabilization. In the absence of Triton X-100, antibodies specific for the RET extracellular domain were able to bind to Ret51::HA (K), Ret51^{AE3}::HA (S), and Ret51^{AE345}::HA (A). However, using an antibody against the HA epitope that would detect the C terminus located intracellularly for Ret51::HA (L), Ret51^{AE3}::HA (T), and Ret51^{AE345}::HA (B') showed no binding/fluorescence in the absence of Triton X-100. When immunofluorescence blocking solution with Triton X-100 was used, fluorescence was detected using the anti-HA antibody for Ret51::HA (P), Ret51^{AE3}::HA (X), and Ret51^{AE345}::HA (F'). These data show that Ret51^{AE3} and Ret51^{AE345} are trafficked to the cell surface similarly as full-length RET51. Scale bars represent 20 μm.
between the RET isoforms and individually evaluated phosphorylation at two tyrosine residues of RET, Tyr905 and Tyr1062. Tyrosine 905 in the RET kinase domain is an autocatalytic tyrosine that is conserved in many receptor tyrosine kinases and is a binding site for GRB7 and GRB10 (36–38). Additionally, mutation of tyrosine 905 to phenylalanine (Y905F) impairs the kinase activity of RET (39). Tyrosine 1062 is a binding site for SHC, Dok4/5, IRS-1, and FRS-2 when phosphorylated and is a binding site for Enigma in a phosphorylation-independent manner (36, 40–47). Interaction between Tyr1062 and these adaptor proteins leads to activation of the Ras/ERK and PI3K/AKT pathways (41, 42, 45, 46, 48).

To evaluate RET phosphorylation, NIH/3T3 cells were transfected with RET51, RET51/E3, or RET51/E345 constructs in the presence or absence of GFRα1, and the total levels of phosphorylated RET or the level of phosphorylation at Tyr905 and Tyr1062 were assessed. To analyze total Tyr(P) levels of RET, immunoprecipitations of RET51 were performed, followed by immuno-

**FIGURE 4. RET**<sup>153</sup> and RET**<sup>1E345</sup> bind to GFRα1, GFRα2, GFRα3, and GFRα4. RET51, RET51<sup>ΔE3</sup>, and RET51<sup>ΔE345</sup> were co-transfected with HA epitope-tagged GFRα1 (A), GFRα2 (B), GFRα3 (C), or GFRα4 (D). Immunoprecipitations (IP) were performed using an HA antibody to select for GFRαs. Immunoblotting (IB) for RET showed interaction of RET51<sup>ΔE3</sup> and RET51<sup>ΔE345</sup> with all four GFRα proteins. Co-transfection of HA::GFRα1, HA::GFRα2, HA::GFRα3, or HA::GFRα4 with TrkA was performed as a negative control, and, as expected, TrkA did not associate with any of the GFRαs.
Novel RET Isoforms Differentially Regulate RET Signaling

Blotting for Tyr(P) (Fig. 6A). An increase in the level of RET\(^{-\Delta E3}\) phosphorylation compared with full-length RET was observed, but this was not statistically significant (\(p = 0.0837\)). We did detect, however, a significant increase in the Tyr(P) level of RET\(^{-\Delta E3}\) co-expressed with GFR\(\alpha1\) compared with full-length RET co-expressed with GFR\(\alpha1\) (\(p = 0.0366\)). The phosphorylation of RET\(^{-\Delta E3}\) co-expressed with GFR\(\alpha1\) or expressed alone was not significantly different from full-length RET (\(p = 0.6231\)).

To determine levels of Tyr(P)\(^{905}\) and Tyr(P)\(^{1062}\), pre-immunoprecipitation lysates were collected from samples in the previous experiments, and immunoblotting was performed using antibodies specific for these two phosphorylated tyrosine residues (Fig. 6B). Levels of Tyr(P)\(^{905}\) and Tyr(P)\(^{1062}\) were significantly elevated for RET\(^{-\Delta E345}\) compared with RET (\(p = 0.0439\) and \(p = 0.0411\), respectively) and also for RET\(^{-\Delta E345}\) co-expressed with GFR\(\alpha1\) compared with full-length RET co-expressed with GFR\(\alpha1\) (\(p = 0.0106\) and \(p = 0.0096\), respectively). However, levels of Tyr(P)\(^{905}\) and Tyr(P)\(^{1062}\) were unchanged for RET\(^{-\Delta E3}\) compared with RET and also for RET\(^{-\Delta E3}\) co-expressed with GFR\(\alpha1\) compared with full-length RET co-expressed with GFR\(\alpha1\). Taken together, these data suggest that RET\(^{-\Delta E3}\) has a similar activation level as full-length RET, whereas RET\(^{-\Delta E345}\) has an elevated activation level compared with RET in a ligand-independent manner. Additionally, the lower total phosphoryl-tyrosine level for RET\(^{-\Delta E345}\) compared with TYR\(^{905}\) and TYR\(^{1062}\) suggests that one or more of the other tyrosine residues in RET is not phosphorylated as highly as TYR\(^{905}\) and TYR\(^{1062}\) compared with full-length RET.

Because we consistently observed an increased level of tyrosine phosphorylation of the RET\(^{-\Delta E345}\) isoform compared with full-length RET, we wondered whether this level of receptor activation was similar to oncogenic RET mutations, particularly those found in MEN2. Human RET point mutations known to be causal for either MEN2A or MEN2B were introduced into the RET\(^{51}\) expression plasmid, specifically C634R (RET\(^{MEN2A}\)) and M918T (RET\(^{MEN2B}\)). NIH/3T3 cells were transfected with RET\(^{S}\), RET\(^{S,\Delta E345}\), RET\(^{S,\Delta E345}\), RET\(^{S,\Delta E345}\), RET\(^{S,\Delta E345}\), or RET\(^{S,\Delta E345}\) constructs. Immunoprecipitations for RET\(^{51}\) were performed to isolate RET proteins, followed by immunoblotting for Tyr(P) as described previously. We observed a significant increase in the level of phosphorylation of both RET\(^{MEN2A}\) and RET\(^{MEN2B}\) oncogenic proteins compared with full-length RET (\(p = 0.0282\) and \(p = 0.0097\), respectively) (Fig. 6F). The levels of phosphorylation of RET\(^{MEN2A}\) and RET\(^{MEN2B}\) co-expressed with RET\(^{-\Delta E345}\) showed no significant difference (\(p = 0.4898\) and \(p = 0.1712\), respectively). These data indicate that the RET\(^{-\Delta E345}\) isoform is phosphorylated at similar levels as oncogenic RET mutations.

RET\(^{-\Delta E3}\) Is Activated in a GDNF-dependent Manner but RET\(^{-\Delta E345}\) Is Not—Although differences are observed in the level of ligand-independent activation for RET\(^{-\Delta E3}\) and RET\(^{-\Delta E345}\), we wondered whether these two receptors could be activated in a GDNF-dependent manner. To evaluate this, Neuro2A cells were co-transfected with HA epitope-tagged RET\(^{S}\), RET\(^{S,\Delta E3}\), or RET\(^{S,\Delta E345}\) and with untagged GFR\(\alpha1\). Twenty-four hours post-transfection, the culture medium was removed, and cells were stimulated with 200 ng/ml recombinant mouse GDNF for 30 min. Cells were detergent-extracted, and immunoprecipitations for HA were performed to isolate the exogenous RET isoforms. Immunoblotting for GFR\(\alpha1\) was performed to determine the level of GDNF-dependent association of GFR\(\alpha1\) to RET (Fig. 7A). For our positive control, where GFR\(\alpha1\) and RET\(^{S}\) were co-expressed, we observed a significant GDNF-dependent association of these proteins after 30 min of stimulation (\(p = 0.0373\)) (Fig. 7B). Under conditions
where GFRα1 and RET51ΔE3 or RET51ΔE345 were co-expressed, we observed a trend of GDNF-dependent association of GFRα1 to both RET isoforms, but this ligand-dependent interaction was not statistically significant ($p = 0.1750$ and $p = 0.3269$, respectively).

To determine whether the RET isoforms were activated in a GDNF-dependent manner, immunoblotting for poly-ubiquitin was performed. We have shown previously that, upon ligand activation and autophosphorylation, RET is rapidly poly-ubiquitinated and targeted for degradation (49). Because ectopic

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**FIGURE 6.** RETΔE345 displays increased ligand-independent activation compared with full-length RET. A, RET51 was co-transfected with HA epitope-tagged GFRα1 in NIH/3T3 cells. Immunoprecipitations (IP) were performed with a RET51 antibody to select for the RET isoforms. Immunoblotting (IB) for phospho-tyrosine and RET was performed, and integrated density values were calculated for each. B, preimmunoprecipitation lysates from the previous experiment were subjected to SDS-PAGE, and Western blotting was performed to determine the levels of residue-specific tyrosine phosphorylation of RET. Immunoblotting was performed for phospho-RET (Tyr905), phospho-RET (Tyr1062), and RET, and integrated density values were calculated for each. Ratios of phospho-RET/RET (C), phospho-RET (Tyr905)/RET (D), and phospho-RET (Tyr1062)/RET (E) are reported, with values normalized to RET51. F, expression plasmids encoding oncogenic RET proteins, RET51MEN2A and RET51MEN2B, were transfected into NIH/3T3 cells, and integrated density values were calculated and analyzed for phospho-RET/RET for all RET isoforms. *, $p < 0.05$; **, $p < 0.01$; n.s., not significant.
expression of RET results in high levels of autophosphorylation in the absence of ligand, we were not able to use phosphotyrosine immunoblotting as a measure of RET activation upon GDNF stimulation. We observed a significant increase in poly-ubiquitinated RET51 ($p = 0.0370$) following GDNF stimulation, as expected, and also observed a significant increase in poly-ubiquitinated RET51/E3 ($p = 0.0493$), suggesting that this isoform is activated in a GDNF-dependent manner. Interestingly, we did not observe a difference in the level of poly-ubiquitinated RET51/E345 following GDNF stimulation ($p = 0.0696$). These data suggest that, like full-length RET, RET/E3 is activated in a GDNF-dependent manner but RET/E345 is not.

Discussion

Alternative splicing of precursor mRNA is one of many processes that mediates gene regulation in metazoans, and the frequency of alternative splicing increases with species complexity (50, 51). For example, of the ∼25,000 genes encoded by the human genome, ∼95% produce transcripts that are alternatively spliced (50, 51). By expanding the proteome through the synthesis of various protein isoforms, alternative splicing allows for increased protein diversity, with isoforms performing different biological functions.

Here we provide evidence for two alternative splicing events in Ret that, in combination with Ret9 or Ret51 alternative splicing, give rise to at least five Ret transcripts (Fig. 2C). The splicing events for RET/E3 and RET/E345 have only been described previously in human tissues (12). Because an additional Ret transcript, RET43, is only expressed in humans, we examined the expression of RET/E3 and RET/E345 in additional vertebrate tissues to determine whether these might also be human-specific transcripts (8, 52). To this end, we discovered that RET/E3 and RET/E345 are not human-specific Ret transcripts but are also expressed in normal developing tissues of zebrafish, mice, and rats (Fig. 1C). Functionally important transcripts are transcriptionally conserved between species, arguing that RET/E3 and RET/E345 have physiologically important functions.

Analysis of an experimentally validated structural model for the RET cadherin-like domains 1–4 was made to evaluate the
Novel RET Isoforms Differentially Regulate RET Signaling

consequence of exon deletions of RET alternative splicing on the ligand binding and activation characteristics of RETΔE3 and RETΔE345 (Fig. 1B). Previous experiments indicate that the GDNF-GFrα1 complex binds directly to CLD4 and the cysteine-rich domain (53). This suggested that both RETΔE3 and RETΔE345 should be able to bind to the GDNF-GFrα1 complex, and indeed this interaction was observed not only for GFrα1 but also for GFrα2, GFrα3, and GFrα4 (Fig. 4). It should be noted that, in our experiments, the chicken GFrα4 construct contained all three cysteine-rich domains, unlike mammalian GFrα4, which lacks domain 1. Although it was originally thought that CLD1–3 of RET was involved in direct binding from results obtained through mutagenesis studies, a direct interaction between CLD1–3 and the GDNF-GFrα1 complex has not been observed (53, 54). Instead, it has been proposed that CLD1–3 contributes to the stability of the tertiary structure of the GDNF-GFrα1-RET complex by forming a secondary dimerization site to trap the GDNF-GFrα1 binary complex (14). These studies predict that RETΔE3, which forms a fused CLD1-CLD2 region, likely gives rise to a less stable tertiary GDNF-GFrα1-RET complex compared with full-length RET. This may result in a receptor variant with differential ligand binding specificities. FGF receptors are a subfamily of four receptor tyrosine kinases that are receptors for FGFs, of which there are 18 ligands (55). Alternative splicing of the FGF receptors in the region coding for the extracellular ligand-binding domain of the receptors allows for differential ligand binding. It is possible that the alternative splicing responsible for RETΔE3 similarly modifies GFL-GFrαo affinity. Alternatively, the truncated extracellular domain of RETΔE3 may cause alterations in the conformational changes that occur upon GFL-GFrα binding that induces receptor autophosphorylation. This would cause different autophosphorylation kinetics in RETΔE3 compared with full-length RET and, therefore, differential activation of downstream second messenger cascades. Biochemical experiments using an in vitro system that allows for the monitoring of autophosphorylation upon direct ligand activation of RETΔE3 or full-length RET in isolation will be necessary to distinguish between these possibilities.

The large deletion in the extracellular domain of RETΔE345, including the Ca2+ binding domain, which is important for RET folding and ligand-dependent signaling, is predicted to have dramatic effects on RETΔE345 activation. Indeed, it is likely that RETΔE345 functions as a constitutively active form of RET, consistent with the mechanistic data shown in Fig. 6. Autophosphorylation of RETΔE345 is increased compared with full-length RET in the presence and absence of GFrα1 in vitro (Fig. 6B). Constitutively active receptor tyrosine kinases have not been found in normal tissues and usually arise from germline and somatic mutations, where they are most commonly found to cause neoplasia. Activating mutations in RET have been shown to be responsible for multiple endocrine neoplasia, type 2, including neoplasias such as medullary thyroid carcinoma and pheochromocytomas (56). Unlike other constitutively active forms of RET, RETΔE345 may be post transcriptionally or translationally regulated in vivo in a manner that is not recapitulated in the in vitro system used here. In this way, the signaling of RETΔE345 may be under tight regulation for this isoform to perform specific functions in vivo.

To interrogate the functions of RETΔE3 and RETΔE345 in primary neurons and tissues, we attempted to generate isoform-specific antibodies. Likewise, we also tried to create siRNAs to selectively knock down RetΔE3 and RetΔE345 transcripts. Neither of these experimental approaches were successful at targeting RETΔE3 and RETΔE345 without also affecting full-length RET. Therefore, to determine the physiologic functions of RETΔE3 and RETΔE345, a transgenic approach in a vertebrate will be required to selectively remove these individual transcripts. This could be accomplished by deleting specific introns and fusing the flanking exons in-frame to inhibit exon skipping (e.g. remove intron 3 and fuse exons 2 and 3 together) to inhibit alternative splicing for RetΔE3. How this approach would affect RET expression and other splicing events, however, is unknown. In addition, because of the invariably coincident expression of RetΔE3 and RetΔE345 in the same tissues as Ret, it is possible that full-length RET may compensate for the functions of RETΔE3 and RETΔE345. As an alternative approach, production of knockin animals in which only RetΔE3 or RetΔE3 is expressed would determine whether these isoforms are capable of supporting the normal development of the nervous system and kidneys.

Last, to gain a mechanistic understanding of the factors responsible for the alternative splicing of Ret, it would be interesting to explore RNA motifs encoding exon splicing enhancers within the locus that impact the formation of RetΔE3 and RetΔE345. Understanding the spliceosome machinery involved in this process may help us understand why the expression of these transcripts is elevated in tumors, such as in pheochromocytomas (12). Overall, it may be necessary to return to a disease model, such as neuroendocrine gland tumors, where these RET transcripts were originally identified and use the pathobiology to understand the signaling capacities of RETΔE3 and RETΔE345 to identify their functions in normal development.

Experimental Procedures

Molecular Modeling—This analysis was performed as described previously (14). Models of cadherin-like domains 1–4 were taken from PDB code 4UX8, defined by a combination of EM and small angle x-ray scattering analyses. The figure was made using PyMOL software (Schrödinger).

RNA Isolation, RT-PCR, and qPCR—Total RNA was isolated from zebrafish, mouse, or rat tissues using TRIzol reagent (Thermo Fisher Scientific) according to the instructions of the manufacturer. cDNA synthesis was performed from 1 μg of total RNA to generate poly(A) first-strand cDNA using the SuperScript III First-Strand Synthesis System for RT-PCR kit (Life Technologies). For RT-PCR reactions with amplicons of less than 200 bp, GoTaq Green Master Mix (Promega) and species-specific oligonucleotide primers for Ret, RetΔE3, and RetΔE345 were used. RT-PCR reactions with amplicons greater than 200 bp were performed using the Phusion high-fidelity DNA polymerase kit (New England Biolabs) following the instructions of the manufacturer. qPCR was performed using FastStart Universal SYBR Green Master Mix (Roche) and run on a Quant-
Novel RET Isoforms Differentially Regulate RET Signaling

Quantitative analyses were performed by calculating the ΔCt (Ct of transcript divided by Ct of the housekeeping gene actin) for each gene analyzed and transforming that value to log₂. These values are reported as the relative expression.

**Primer Sequences**—The primers used for this study were as follows: ZF.RetFL.F, 5'-TCG TAG TTT ACG CAG CCG CCC TTG TTA AGT TG T-3'; ZF.RetFL.R, 5'-CAG TAG TTT ACG ATC TTC TTG ACC G-3'; ZF.Ret3.F, 5'-CAG TAG TTT ACG ATC TTC TTG ACC G-3'; ZF.Ret3.R, 5'-TTC AGT GTC ATG CCC GTT GA-3'; ZF.Ret345.F, 5'-CAG TAG TTT ACA GCT GAA ACT CAG TC-3'; ZF.Ret345.R, 5'-TGA CAT TGG AGA AGC GAA TTG-3'. Mouse.Ret9.R, 5'-AGC ATC CGC AGG GAT AGT AGG A-3'; Mouse.Ret3.R, 5'-ACA CTG TCA CTG GGA AGG G-3'; Mouse.Ret3.F, 5'-AGC ATC CGC AGG GAT ATG CT-3'; Mouse.Ret3.R, 5'-TCC AGT GAG TAG TAG GC-3'; Rat.RetFL.F, 5'-AGC ATC CGC AAT GGC GGC TT-3'; Rat.RetFL.R, 5'-TAG CAT CGC GAA CTG GTA GA-3'; Rat.Ret3.F, 5'-AGC ATC CGC AAT GGC GGC TT-3'; Rat.Ret3.R, 5'-CCG CTT AAA CTC CAC AG-3'; Rat.Ret345.F, 5'-AGC ATC CGC AAT GGC GGC TT-3'; Rat.Ret345.R, 5'-TAG GCC ATG GTG AGG TTT AGC-3'; and Mouse.Ret9.R, 5'-ATT TAC TGT CCA TTG CAA GCC-3'; and Mouse.Ret51.R, 5'-CCT ACT AGT GCT TTA AGT CTG AG-3'.

Mice—Wild-type C57BL/6J mice were purchased from The Jackson Laboratories (Bar Harbor, ME). Ret⁻/- mice have been described previously and were maintained in a mixed genetic background (57). For timed matings, noon of the day on which a vaginal plug was detected was considered E0.5. All housing and procedures performed on mice were approved by the University of Michigan Animal Care and Use Committee.

**Detergent Extraction and Preclearing of Whole Tissues**—Tissues were harvested, placed in a 2-mL tube with 250 μL of immunoprecipitation buffer lacking Nonidet P-40 (TBS (pH 7.4), 10% glycerol, 500 μM sodium vanadate, and protease inhibitors) along with a steel grinding ball (5 mm, 69989, Qiagen, Valencia, CA) and mechanically homogenized using a TissueLyser II (Qiagen) for 2 min at a frequency of 20 Hz. Following this, 250 μL of 2% Nonidet P-40-containing immunoprecipitation buffer was added to homogenates and incubated for 1 h at 4 °C under gentle agitation. Homogenates were then centrifuged for 10 min at 16,000 × g and subjected to an initial preclearing step with protein A and protein G alone at 4 °C for 2 h under gentle agitation. A second preclearing step was performed with protein A, protein G, and a species-matched nonspecific control IgG for 2 h under gentle agitation.

**Immunoprecipitations and Immunoblotting**—Plates were placed on ice, gently washed twice with 1× PBS (pH 7.4), and lysed with modified radiolabeled immunoprecipitation assay buffer (TBS (pH 7.4), 10% glycerol, 1% Triton X-100, 0.1% SDS, 500 μM sodium vanadate, and protease inhibitors). Proteins were immunoprecipitated using anti-RET9 (goat, Santa Cruz Biotechnology), anti-RET51 (goat, Santa Cruz Biotechnology), anti-HA (mouse, Millipore), or anti-FLAG (rabbit, Cell Signaling Technology) selective antibodies. Immunoprecipitates were subjected to SDS-PAGE, followed by electrophoresis onto PVDF membranes (Immobilon P, Millipore). Immunoblotting was performed on blots using antibodies selective for RET (1:500, goat, R&D Systems), HA (1:5000, rabbit, Cell Signaling Technology), phospho-tyrosine (1:1000, mouse, Millipore), phospho-RET ( Tyr905) (1:500, rabbit), phospho-RET (Tyr1062) (1:500, rabbit), GFRα1 (1:500, goat, R&D Systems), or poly-ubiquitin (1:500, mouse, Enzo Life Sciences). Phospho-RET (Tyr905) and phospho-RET (Tyr1062) have been described previously (58). Lysates collected prior to immunoprecipitations served as loading controls to assess protein expression levels and were also subjected to immunoblotting for actin (1:1000, goat, Santa Cruz Biotechnology). Each biochemical experiment was performed three to four times with similar results.

**Mammalian Cell Culture and Transfections**—NIH/3T3 (ATCC), HEK293T (ATCC), and Neuro2A cells (ATCC) were maintained at 37 °C with 5% CO₂ in DMEM supplemented with FBS (10% by volume) and a penicillin/streptomycin/glutamine mixture. Cells were plated into 6-well tissue culture plates and allowed to proliferate until cells were 70% confluent. Transfections of cells were performed using Lipofectamine 2000 according to the instructions of the manufacturer (Invitrogen). For experiments testing the interaction of proteins, a total of 4 μg of plasmid DNA was added per well using a GFP plasmid as a control for transfecting equal amounts of DNA. For in vitro ligand-independent activation experiments, 2 μg of HA::GFRα1 and 1 μg of one of the RET constructs were transfected for a total of 3 μg of DNA per condition, with a GFP plasmid used as a control for transfecting equal amounts of DNA. RET51AE3 and RET51ΔE345 constructs were subcloned from RET51 constructs into pcDNA3.1 (Thermo Fisher Scientific). A Ret51 construct encoding mouse RET51 fused at the C terminus with an HA epitope tag (RET51:HA) was acquired from Dr. Ben Allen. This plasmid was subcloned to create similarly tagged RET51AE3 and RET51ΔE345 constructs, which were subcloned into pLentilox-IRES-GFP (University of Michigan Vector Core). For in vitro ligand-dependent activation experiments, 3 μg of HA epitope-tagged Ret51, Ret51AE3, or Ret51ΔE345 was co-transfected with 3 μg of untagged GFRα1. Twenty-four hours post-transfection, the medium was aspirated, and cells were treated with 200 ng/ml recombinant mouse GDNF (Peprotech) in fresh cell culture medium for 30 min. The HA::GFRα1 (mouse) plasmid was generously donated by Dr. Ben Allen. The HA::GFRα2 (rat), HA::GFRα3 (mouse), and HA::GFRα4 (chicken) plasmids were generously donated by Dr. Carlos Ibáñez. The TrkA (rat) plasmid was generously donated by Dr. Cristin Carter-Su. Constructs encoding human RET51 harboring MEN2A (C634R) and MEN2B (M918T) point mutations in RET were created using the QuickChange II XL site-directed mutagenesis kit (Agilent).

**Quantification of Immunobots**—Scanned images of x-ray films were imported into ImageJ (National Institutes of Health) and processed using the gel analysis tool. Integrated density values obtained from immunoblotting were reported as mean ± S.E. with arbitrary units on the vertical axis. Values were normalized to the level of RET51 phosphorylation in each experiment. Statistical analyses were performed using
Student’s t test for each experiment, and all biochemical experiments were performed at least three times with similar results.

**Acknowledgments**—We thank Dr. Benjamin Allen for reagents and critical scientific comments on the manuscript. Additionally, we thank Dr. Carlos Ibáñez, Dr. Christin Carter-Su, and Dr. Frank Costantini for reagents. We also thank the Pierchala laboratory for helpfuful scientific discussions.

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J. Biol. Chem. 2016, 291:16249-16262.
doi: 10.1074/jbc.M115.709675 originally published online May 23, 2016

Access the most updated version of this article at doi: 10.1074/jbc.M115.709675

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