Quantifying the strength of heterointeractions among receptor tyrosine kinases from different subfamilies: Implications for cell signaling

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Receptor tyrosine kinases (RTKs) are single-pass membrane proteins that control vital cell processes such as cell growth, survival, and differentiation. There is a growing body of evidence that RTKs from different subfamilies can interact and that these diverse interactions can have important biological consequences. However, these heterointeractions are often ignored, and their strengths are unknown. In this work, we studied the heterointeractions of nine RTK pairs, epidermal growth factor receptor (EGFR)–EPH receptor A2 (EPHA2), EGFR–vascular endothelial growth factor receptor 2 (VEGFR2), EPHA2–VEGFR2, EPHA2–fibroblast growth factor receptor 1 (FGFR1), EPHA2–FGFR2, EPHA2–FGFR3, VEGFR2–FGFR1, VEGFR2–FGFR2, and VEGFR2–FGFR3, using a FRET-based method. Surprisingly, we found that RTK heterodimerization and homodimerization strengths can be similar, underscoring the significance of RTK heterointeractions in signaling. We discuss how these heterointeractions can contribute to the complexity of RTK signal transduction, and we highlight the utility of quantitative FRET for probing multiple interactions in the plasma membrane.

The 58 human receptor tyrosine kinases (RTKs) are grouped into 20 subfamilies based on the homology of their N-terminal extracellular (EC) domains (1–4). They are single-pass membrane receptors, which are activated upon dimerization and phosphorylation of tyrosines in their C-terminal intracellular (IC) kinase domains (3, 5–8). Adaptor proteins bind to the phosphorylated tyrosines, leading to the initiation of downstream signaling cascades that control cellular processes such as growth, motility, survival, and differentiation (9–14). Moreover, RTKs are often dysregulated in many growth disorders and diseases, such as cancer (15–20).

Whereas RTKs are best known for forming signaling homodimers, they are also known to interact with many other binding partners, including other RTKs from the same subfamily, G protein–coupled receptors, and cell adhesion molecules (21–26). Furthermore, it is becoming clear that they are capable of interacting with RTKs from unrelated subfamilies. A recent review of the literature identified nearly 100 studies that support the idea that RTKs from different subfamilies can interact (27). Despite these studies, heterodimerization across subfamilies has largely been ignored, both in conceptual models of RTK signaling and in the interpretation of RTK signaling data. However, these heterodimers have been shown to play important and diverse roles, such as amplifying or inhibiting downstream signaling (28–34), increasing the signal diversity (35–38), or providing signaling back-up (27).

It is also becoming increasingly apparent that unliganded RTK dimers can form in physiological contexts and can have important effects, especially in disease (39–41). RTK overexpression has been linked to many cancers, as it promotes RTK interactions even in the absence of ligand (17, 18). Interestingly, some of these cancers are associated with not just increased RTK expression, but loss of ligand as well. It can be expected that such conditions (high RTK expression and no ligand) also favor the formation of heterodimers of unrelated RTKs, as these RTKs typically do not share ligands. However, it is currently not clear how likely cross-subfamily heterointeractions are to occur, and it is thus difficult to say how significant a role they play in RTK signaling. Indeed, heterointeraction strengths for RTKs from different subfamilies are unknown, as prior work has had limited quantification capabilities.

In this work, we study the heterointeractions of nine RTK pairs (EGFR–EPHA2, EGFR–VEGFR2, EPHA2–VEGFR2, EPHA2–FGFR1, EPHA2–FGFR2, EPHA2–FGFR3, VEGFR2–FGFR1, VEGFR2–FGFR2, and VEGFR2–FGFR3) using a fluorescence-based method: fully quantified spectral imaging Förster resonance energy transfer (FSI-FRET). The FSI-FRET method reports on the occurrence of specific interactions and yields the strength of the interactions (1). We advance this FRET methodology to also enable the determination of the oligomer size (e.g. dimer, trimer, tetramer) of the heterointeraction. For three of the RTK pairs, we demonstrate that they form dimers in the absence of ligand, and we quantify their heterodimerization strengths. Surprisingly, we find that the heterodimerization and homodimerization strengths are similar. This suggests that the heterodimers are likely to form under physiological and pathological conditions. Furthermore, we show that the presence of ligands modulates the abundance of the heterodimers. This work underscores the significance of heterointeractions in RTK signaling and highlights a method that can be used to study a multitude of interactions in the plasma membrane of live cells.

Results

Heterointeraction models

In most cases, RTKs are thought to signal as homodimers (2, 3). Accordingly, if two different RTKs, X and Y, were to

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interact, the obvious assumption is that they would form a heterodimer, XY. However, there is no experimental evidence that excludes the possibility of higher-order heterooligomers (e.g. trimers, tetramers) forming. For example, as each RTK explores a monomer-dimer equilibrium, it is possible that a dimer, YY, and a monomer, X, interact to form a trimer, XYY. It is also possible that two dimers interact to form a tetramer, XXYY. These cases are illustrated in Fig. 1. Here, we develop physical-chemical models for these cases, which can be used to identify the model that best describes experimental data.

**Heterodimers (Fig. 1A)**

The heterodimer case has been considered previously by Del Piccolo et al. (42). Even in this simplest case, three coupled reactions are needed to describe the fact that both X and Y form homodimers, XX and YY, in addition to the heterodimer, XY.

\[
\begin{align*}
    [X] + [X] & \rightleftharpoons K_X [XX] \\
    [Y] + [Y] & \rightleftharpoons K_Y [YY] \\
    [X] + [Y] & \rightleftharpoons K_{XY} [XY]
\end{align*}
\]

Reactions 1 – 3

Here, \( K_X \) and \( K_Y \) are the homodimer association constants, and \( K_{XY} \) is the heterodimer association constant. These three association constants can be written in terms of the concentrations of the monomers and dimers.

\[
\begin{align*}
    K_X &= \frac{[XX]}{[X]^2} \\
    K_Y &= \frac{[YY]}{[Y]^2} \\
    K_{XY} &= \frac{[XY]}{[X][Y]}
\end{align*}
\]

Assuming that the total concentration of each RTK is constant—an assumption that is valid for the conditions under which the FRET experiments are performed (43)—then simple equations for mass conservation can be written as follows.

\[ [X_{total}] = [X] + 2[XX] + [XY] \]  
\[ [Y_{total}] = [Y] + 2[YY] + [XY] \]  

By substituting in the values of \( K_X \) and \( K_Y \) from Equations 1–2, the total concentrations can be written in terms of the monomer and heterodimer concentrations as follows.

\[ [X_{total}] = [X] + 2K_X[X]^2 + [XY] \]  
\[ [Y_{total}] = [Y] + 2K_Y[Y]^2 + [XY] \]
These are quadratic equations which can be solved for in terms of the monomer concentration.

\[
[X] = \frac{-1 + \sqrt{1 - 8K_X ([XY] - [X_{total}])}}{4K_X} \quad \text{(Eq. 8)}
\]

\[
[Y] = \frac{-1 + \sqrt{1 - 8K_Y ([XY] - [Y_{total}])}}{4K_Y} \quad \text{(Eq. 9)}
\]

Last, the concentration of the heterodimer can be written in terms of the monomer concentrations and the heterodimer association constant by rearranging Equation 3 as follows.

\[
[XY] = K_{XY} [X] [Y] \quad \text{(Eq. 10)}
\]

By substituting the values of [X] and [Y] from Equations 8 and 9 into Equation 10, we arrive at an equation where \(K_{XY}\) and [XY] are defined in terms of \([X_{total}]\), \([Y_{total}]\), \(K_X\), and \(K_Y\). As described previously (1, 6, 42, 44), quantitative FRET experiments can directly measure \([X_{total}]\) and \([Y_{total}]\), and the FRET efficiency. FRET efficiency is a function of [XY], as shown in Equation 32 (see “Materials and methods”). Accordingly, if \(K_X\) and \(K_Y\) have been measured previously, then FRET experiments can be used to determine the best-fit \(K_{XY}\). For a detailed description of this fitting process, see Ref. 42.

**Heterotrimers (Fig. 1B)**

There are two possible heterotrimers, XYY and XYX; however, these are equivalent, as the assignment of X and Y for the two RTKs can be switched. Accordingly, only the case of XYY will be considered here. We begin with defining the three coupled reactions needed to fully describe the formation of the homodimers, XX and YY, and the heterotrimer, XYY:

\[
\begin{align*}
[X] + [X] &\rightleftharpoons K_X [XX] \\
[Y] + [Y] &\rightleftharpoons K_Y [YY] \\
[X] + [YY] &\rightleftharpoons K_{XY} [XYY]
\end{align*}
\]

Reactions 4 – 6

where \(K_{XY}\) is the heterotrimer association constant. These three association constants can be written in terms of the concentrations of the monomers, homodimers, and heterotrimers:

\[
K_X = \frac{[XX]}{[X]^2} \quad \text{(Eq. 11)}
\]

\[
K_Y = \frac{[YY]}{[Y]^2} \quad \text{(Eq. 12)}
\]

\[
K_{XY} = \frac{[XYY]}{[X][YY]} \quad \text{(Eq. 13)}
\]

The equations for mass conservation can be written as follows.

\[
[X_{total}] = [X] + 2[XX] + [XYY] \quad \text{(Eq. 14)}
\]

\[
[Y_{total}] = [Y] + 2[YY] + 2[XYY] \quad \text{(Eq. 15)}
\]

and these can be rewritten in terms of the association constants.

\[
[X_{total}] = [X] + 2K_X [X]^2 + [XYY] \quad \text{(Eq. 16)}
\]

\[
[Y_{total}] = [Y] + 2K_Y [Y]^2 + 2[XYY] \quad \text{(Eq. 17)}
\]

These are quadratic equations which can be solved for in terms of [X] and [Y].

\[
[X] = \frac{-1 + \sqrt{1 - 8K_X ([XYY] - [X_{total}])}}{4K_X} \quad \text{(Eq. 18)}
\]

\[
[Y] = \frac{-1 + \sqrt{1 - 8K_Y (2[XYY] - [Y_{total}])}}{4K_Y} \quad \text{(Eq. 19)}
\]

Finally, the concentration of the heterotrimer can be written in terms of the monomer concentrations and the heterotrimer association constant by rearranging Equation 13.

\[
[XYY] = K_{XY} [X] [Y] = K_{XY} [X] [K_Y] [Y]^2 \quad \text{(Eq. 20)}
\]

By substituting the values of [X] and [Y] from Equations 18 and 19 into Equation 20, we arrive at an equation where \(K_{XY}\) and [XY] are defined in terms of \([X_{total}]\), \([Y_{total}]\), \(K_X\), and \(K_Y\), as is the heterotrimer association constant. These three association constants can be written in terms of the concentrations of the monomers, dimers, and tetramers as follows.

**Heterotetramers (Fig. 1C)**

Here, we consider two dimers, XX and YY, forming a tetramer, XYY. We begin with defining the three coupled reactions needed to fully describe the formation of the homodimers and the heterotetramer,

\[
\begin{align*}
[X] + [X] &\rightleftharpoons K_X [XX] \\
[Y] + [Y] &\rightleftharpoons K_Y [YY] \\
[XX] + [YY] &\rightleftharpoons K_{XXYY} [XYY]
\end{align*}
\]

Reactions 7 – 9

where \(K_{XXYY}\) is the heterotetramer association constant. These three association constants can be written in terms of the concentrations of the monomers, dimers, and tetramers as follows.

\[
K_X = \frac{[XX]}{[X]^2} \quad \text{(Eq. 11)}
\]

\[
K_Y = \frac{[YY]}{[Y]^2} \quad \text{(Eq. 12)}
\]

\[
K_{XXYY} = \frac{[XYY]}{[X][YY]} \quad \text{(Eq. 13)}
\]
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\[
K_X = \frac{[XX]}{[X]^2} \quad \text{(Eq. 21)}
\]

\[
K_Y = \frac{[YY]}{[Y]^2} \quad \text{(Eq. 22)}
\]

\[
K_{XYY} = \frac{[XXYY]}{XX][YY] \quad \text{(Eq. 23)}
\]

The equations for mass conservation are written as follows,

\[
[X]_{\text{total}} = [X] + 2[XX] + 2[XXYY] \quad \text{(Eq. 24)}
\]

\[
[Y]_{\text{total}} = [Y] + 2[YY] + 2[XXYY] \quad \text{(Eq. 25)}
\]

and these can be rewritten in terms of the association constants.

\[
[X]_{\text{total}} = [X] + 2K_X[X]^2 + 2[XXYY] \quad \text{(Eq. 26)}
\]

\[
[Y]_{\text{total}} = [Y] + 2K_Y[Y]^2 + 2[XXYY] \quad \text{(Eq. 27)}
\]

The monomer concentration can be determined by solving these quadratic equations.

\[
[X] = \frac{-1 + \sqrt{1 - 8K_X(2[XXYY] - [X]_{\text{total}})}}{4K_X} \quad \text{(Eq. 28)}
\]

\[
[Y] = \frac{-1 + \sqrt{1 - 8K_Y(2[XXYY] - [Y]_{\text{total}})}}{4K_Y} \quad \text{(Eq. 29)}
\]

Last, the concentration of the heterotetramer can be written in terms of the monomer concentrations and the heterotetramer association constant by rearranging Equation 23 as follows.

\[
[XXYY] = K_{XYY}[XX][YY] = K_{XYY}K_X[X]^2K_Y[Y]^2 \quad \text{(Eq. 30)}
\]

By substituting the values of [X] and [Y] from Equations 28 and 29 into Equation 30, we arrive at an equation where \(K_{XYY}\) and [XXYY] are defined in terms of \([X]_{\text{total}}, [Y]_{\text{total}}, K_X, \) and \(K_Y\). As with the heterodimer and heterotramer cases, FRET can be used to determine \(K_{XYY}\) and hence heterotetramer stability.

**Experimental assessments of heterointeractions**

The first RTK pair that we studied was EGFR and EPHA2. EGFR is a member of the ErbB family and is important for cell growth, division, migration, and survival \((45–47)\). It is known to be overexpressed or mutated in many cancers \((48, 49)\). EPHA2 is a member of the EPH family, and it plays a role in neuronal development, cell proliferation, and axon guidance \((50–52)\). Overexpression and mutations of the EPH receptor are generally oncogenic \((16, 53)\), but unliganded EPHA2 dimers have been shown to suppress tumorigenic signaling \((54)\). Both EGFR and EPHA2 can be expressed in the same cells \((55–57)\), and they have been reported to interact, based on fluorescence colocalization and coimmunoprecipitation \((58)\). Moreover, there is a report that increased EPHA2 levels in colorectal cancer correlates with a poor response to the EGFR inhibitor cetuximab \((59)\). Taken together, these studies indicate that the EGFR–EPHA2 interaction could be affecting signaling and could be a potential target in therapeutic strategies. Therefore, we asked whether we could detect the heterointeraction between EGFR and EPHA2 using FRET.

To study heterointeractions using FRET, one RTK must be labeled with a donor fluorophore, and the other RTK must be labeled with an acceptor fluorophore. If the receptors interact, the donor will be brought close to the acceptor, and FRET will occur. In these FRET experiments, HEK 293T cells were transiently co-transfected with plasmids that encode for full-length EGFR labeled with mTurquoise (MT, a FRET donor) and full-length EPHA2 labeled with eYFP (a FRET acceptor). The cells were grown for at least 24 h after transfection and were serum-starved for at least 12 h prior to imaging to ensure that no ligands were present.

FRET experiments were performed using the FSI method, which has been described in detail previously \((1)\). In brief, the cells were imaged using a two-photon microscope. Two scans were taken of each cell: one scan to maximally excite the donor and one scan to maximally excite the acceptor. Fluorescence intensities were converted into concentrations using purified solutions of the fluorophores of known concentrations. The cells were analyzed using MATLAB to determine the 2D concentration of the donor, the 2D concentration of the acceptor, and the apparent FRET efficiency of selected regions of the plasma membrane.

We encountered difficulties in these experiments: whereas the co-transfection was performed with a 2:1 or 3:1 EGFR–MT/EPHA2–YFP ratio of plasmid DNA, all cells had very low EGFR expression. It appeared that EPHA2 expression suppressed EGFR expression, such that only a small percentage of the cells that we imaged had quantifiable concentrations of both EGFR and EPHA2 in the plasma membrane. A representative image of such a cell that was selected for analysis is shown in Fig. 2A. For this cellular subset, the experimentally measured \((i.e.\) apparent) FRET efficiencies \((E_{\text{app}})\) for the EGFR–EPHA2 pair are shown in Fig. 2B as a function of acceptor concentration (EPHA2–YFP). Each black star represents the measurement for a 2–3-μm stretch of plasma membrane; two such membrane regions were selected per cell. The blue line in Fig. 2B is a previously modeled monomer proximity line \((60, 61)\), which represents the maximum proximity FRET that can occur at a given acceptor concentration. Proximity FRET increases with increasing fluorophore concentration, as it occurs when a donor and acceptor are randomly close to each in the absence of specific interactions.

We sought to experimentally validate the predicted monomer proximity FRET using a monomer control protein. To do this, we performed FRET experiments with linker for activation of T cells (LAT). LAT is a single-pass membrane protein that plays a key role in T-cell receptor downstream signaling \((62–64)\), and it has been used as a monomer control in the literature \((65)\). HEK 293T cells were transfected with versions of LAT that have been labeled by either MT or eYFP on the C terminus via a flexible linker. As seen in the plot of apparent FRET versus acceptor concentration in Fig. 2C, the LAT FRET...
effectively straddles the monomer proximity line, confirming that the modeled monomer proximity line accurately describes the proximity FRET of a monomer. Further evidence is seen in Fig. 2D, where the FRET data in Fig. 2C have been corrected for the monomer proximity FRET contribution, histogrammed (blue bars), and fit to a Gaussian (red line). As expected, the result is effectively random noise (i.e., a Gaussian centered around zero).

As can be seen in Fig. 2A, almost all of the FRET data for the EGFR–EPHA2 pair lie above the modeled proximity line. This is indicative of a specific interaction occurring between EGFR and EPHA2, consistent with the literature (58, 59). Thus, we were able to observe the heterointeraction between full-length EGFR and EPHA2 using FRET. However, we were not able to calculate the heteroassociation constant, as a much larger data set is needed to fit the data with the models shown in Fig. 1 (A–C).

There is a published report that the deletion of amino acids in the EGFR post-kinase tail may improve EGFR expression (66). Accordingly, we created an EGFRΔTail construct where the post-kinase tail (~220 amino acids) was deleted and replaced with a fluorescent protein (MT). We also used an EPHA2ΔSAM construct where the SAM domain was replaced with a fluorophore (eYFP), such that the fluorescent proteins were attached to the kinase domains in both constructs (see Fig. 1D, middle). This design should ensure that the FRET pair will be in close proximity if heterointeractions occur, allowing us to measure FRET.

The overall expression of EGFR was improved upon the deletion of the post-kinase tail, but most of it did not traffic to the plasma membrane, and thus the concentration of EGFR in the plasma membrane was still low. Only a small fraction of the imaged cells had a quantifiable amount of both receptors. For these cells, Fig. 3A shows the FRET efficiency versus acceptor concentration for the EGFRΔTail–EPHA2ΔSAM pair. Again, almost all of the data lie above the modeled proximity line, indicating that a specific interaction is occurring between EGFR and EPHA2. Accordingly, the post-kinase tail of EGFR and the SAM domain of EPHA2 are not required for the interaction to occur.

In Fig. 3B, we show FRET data reporting on the interactions between VEGFR2 and EPHA2. Specifically, we studied the interactions between VEGFR2ΔTail–MT and EPHA2ΔSAM–YFP, as the full-length VEGFR2 exhibited virtually no plasma membrane expression in HEK 293T cells. VEGFR2 is critically important for angiogenesis. It plays a role in organ development
RTK heterointeractions

Figure 3 Heterointeractions between truncated EGFR and EPHA2 (A) and truncated VEGFR2 and EPHA2 (B). In the case of EPHA2, the SAM domain was replaced with eYFP, and in the case of EGFR and VEGFR2, the post-kinase tail was replaced with MT. The cells were transfected with varying donor/acceptor ratios, with a total of 0.5–3 μg of EPHA2–eYFP DNA and either 1–5 μg of EGFR–MT DNA (A) or 3–8 μg of VEGFR–MT DNA (B). Each black star represents the measurement for a 2–3-μm stretch of plasma membrane in a cell. Two regions were selected per cell, for a total of 56 and 50 data points in A and B, respectively. The solid blue line is the modeled monomer FRET proximity as a function of acceptor concentration (60, 61). In both cases, almost all data points lie above the monomer proximity line, demonstrating specific interactions.

We next performed FRET experiments with ECTM EGFR and ECTM EPHA2, following the same imaging protocol as in the full-length and post-kinase tail deletion cases. Fig. 4A shows the apparent FRET for the ECTM EGFR–ECTM EPHA2 pair as a function of acceptor concentration. These data combine the results of two types of experiments, one where EGFR was labeled with MT while EPHA2 was labeled with eYFP (orange) and one where EGFR was labeled with eYFP while EPHA2 was labeled with MT (green). Fig. 4B shows the concentration of EGFR versus the concentration of EPHA2 for the entire data set. We see that the FRET data in Fig. 4A lie above the proximity line, which indicates that a specific interaction is occurring between ECTM EGFR and ECTM EPHA2.

Next, we asked which of the different interaction models (dimer, trimer, or tetramer) in Fig. 1 (A–C) best describe these data. To answer this question and to determine the heteroassociation constant, we need to know the homodimer association constants, $K_X$ and $K_Y$, for both ECTM EGFR and ECTM EPHA2. The homodimer association constant $K_{EGFR}$ has been measured previously (77), but $K_{EPHA2}$ has not been. To measure it, we performed FRET homodimerization experiments in HEK 293T cells that were co-transfected with plasmids for ECTM EPHA2–MT and ECTM EPHA2–eYFP, and we followed a well-established homodimer data analysis protocol (1, 81). Fig. 4C shows the apparent FRET efficiency versus the total concentration of receptors, and Fig. 4D shows the donor versus acceptor concentration. The ECTM EPHA2 homodimer association constant is determined by fitting a monomer-dimer model to the FRET data (1, 81). The best-fit value for $K_{EPHA2}$ is 0.0012 ± 0.0001 μm$^2$/receptor, and the corresponding dissociation constant ($K_{EPHA2}^{-1}$) is 860 ± 50 receptors/μm$^2$. Using $ΔG = RT ln K$—where $R$ is the ideal gas constant and $T$ is temperature—and a reference state of 1 nm$^2$/receptor gives $ΔG_{EPHA2}$ of −4.0 ± 0.1 kcal/mol.

With $K_{EGFR}$ and $K_{EPHA2}$ known, the heterointeraction FRET data in Fig. 4 (A and B) were fit to predictions derived from the three heterodimerization models in Fig. 1 (A–C) as well as a...
monomer model where the proximity line is regarded as the best fit \((60, 61)\). For these four models, the best-fit theoretical FRET was compared with the experimental FRET, and mean squared errors (MSEs) were calculated for each of the four models, as shown in Fig. 4E. The lowest MSE in Fig. 4E is observed for the heterodimer, suggesting that EGFR and
**Table 1**

Homo- and heterodimerization strengths for three of the studied RTKs

| RTKs             | $K_d$ (receptors/µm²) | $\Delta G$ (kcal/mol) |
|------------------|-----------------------|-----------------------|
| EGF              | 2800 ± 200            | −3.5 ± 0.1            |
| EPHA2            | 860 ± 50              | −4.0 ± 0.1            |
| VEGFR2           | 2700 ± 450            | −3.5 ± 0.1            |
| EGF–EPHA2        | 5000 ± 1000           | −3.1 ± 0.2            |
| EGF–VEGFR2       | 5200 ± 1200           | −3.1 ± 0.2            |
| EPHA2–VEGFR2     | 2900 ± 900            | −3.4 ± 0.2            |

RTK heterointeractions

EPHA2 likely form a heterodimer and not higher-order oligomers. Fig. 4F shows the dimeric fractions, which depend on both EGF and EPHA2 concentrations. The experimentally derived fractions are shown as purple circles, and the best-fit surface is shown in cyan. The best-fit $K_{\text{EPHA2–VEGFR2}}$ for the heterodimer is 5000 ± 1000 receptors/µm², corresponding to a heterodimer stability of $\Delta G_{\text{EPHA2–VEGFR2}} = −3.1 ± 0.2$ kcal/mol. In Table 1, these values are compared with the previously determined homodimer dissociation constants and $\Delta G$ values for ECTM EPHA2 and EPHA2 homodimers. We see that the heterointeraction strength is very similar to the homointeraction strengths.

**ECTM EGF and ECTM VEGFR2 form a heterodimer**

Next, we studied the heterointeraction between ECTM EGF and ECTM VEGFR2 using the same FRET imaging protocol. Fig. 5A shows the apparent FRET efficiencies versus acceptor concentration data for the ECTM EGF–ECTM VEGFR2 pair. Again, experiments were performed with two different labeling schemes: ECTM EGF–MT, ECTM VEGFR2–eYFP (orange) and ECTM EGF–eYFP, ECTM VEGFR2–MT (green). Fig. 5B shows the expression of VEGFR2 versus EGF expression. Virtually all of the data in Fig. 5A lie above the proximity line, which indicates that a specific interaction is occurring between ECTM EGF and ECTM VEGFR2.

The data in Fig. 5 (A and B) were fit to the models shown in Fig. 1 (A–C) and the monomer model using the previously measured values of $K_{\text{EGF}}$ (77) and $K_{\text{VEGFR2}}$ (1). An MSE analysis for the different heterointeraction models is shown in Fig. 5C, and it suggests that EGF and VEGFR2 form a heterodimer. Fig. 5D shows the dimeric fraction as a function of EGF and VEGFR2 concentrations, where the best-fit surface is in cyan and the data are the purple circles. The best-fit $K_{\text{EGF–VEGFR2}}$ for the heterodimer is 5200 ± 1200 receptors/µm², which corresponds to heterodimer stability of $\Delta G_{\text{EGF–VEGFR2}} = −3.1 ± 0.2$ kcal/mol. These values are comparable with the previously determined homodimerization constants and $\Delta G$ values for the ECTM EGF and ECTM VEGFR2 homodimers, as shown in Table 1.

**ECTM EPHA2 and ECTM VEGFR2 both engage in heterointeractions with ECTM FGFR1, ECTM FGFR2, and ECTM FGFR3**

We performed FRET experiments with FGFR1, FGFR2, and FGFR3, with the goal to study their heterointeractions with EPHA2 and VEGFR2. The FGFRs are important for musculoskeletal development (82, 83), and their misfunction has been linked to many growth disorders (20, 84). These three FGFRs have been shown to interact with each other (42), and immunoprecipitation studies suggest that they can also interact with many RTKs from other subfamilies (27). To the best of our knowledge, no interactions have been reported for FGFRs and EPHA2 or for FGFRs and VEGFR2. However, all FGFRs have been shown to interact with EPHA4 (85, 86). We therefore hypothesized that there may be FGFR interactions, in particular FGF-EPHA2 interactions, that have not yet been identified.

In the experiments, HEK 293T cells were co-transfected with either ECTM VEGFR2–MT or ECTM EPHA2–MT and one of ECTM FGFR1–eYFP, ECTM FGFR2–eYFP, or ECTM FGFR3–eYFP. The apparent FRET efficiencies for all six pairs are shown as a function of acceptor concentration in Fig. 7, with the EPHA2 data in Fig. 7A and the VEGFR2 data in Fig. 7B. In all cases, virtually all of the data lie above the proximity line, which indicates that EPHA2–FGFR1, EPHA2–FGFR2, EPHA2–FGFR3, VEGFR2–FGFR1, VEGFR2–FGFR2, and VEGFR2–FGFR3 all engage in specific heterointeractions.

**EGF and mEphrin-A1 Fc ligands decrease the heterointeractions**

The data presented above suggest that RTK homo- and heterodimers are approximately equally likely to form in the absence of ligand; accordingly, their abundance will depend on the expression of the RTKs. If ligands are present, however, homodimers that are bound to their cognate ligand will be stabilized, and thus the concentration of monomers will decrease, which is expected to lead to a depletion of the heterodimer.
population. In other words, if the ligands do not bind the heterodimers, the heterodimer population will decrease as the ligand concentration is increased. Thermodynamic predictions have shown that when the ligand concentration substantially exceeds the ligand dissociation constant (27), the heterodimer population becomes near zero.

With these considerations in mind, we quantified the effect of the EGF ligand on the ECTM EGFR–ECTM VEGFR2 heterodimer population. We used 10 nmol of EGF, which is about an order of magnitude higher than the reported EGF-EGFR apparent dissociation constant (87–90). Cells were prepared identically to the EGFR–VEGFR2 experiments, except 0.1% (w/v) BSA was added to the starvation and imaging media to prevent ligand binding to the imaging dish, and the ligand was added 10 min prior to imaging. In Fig. 8A, the apparent FRET versus acceptor concentration is shown for cells in the presence of 10 nmol of EGF; EGF data are shown in red, and the no-ligand data (see Fig. 5A) are in black. We observe a marked decrease in the FRET efficiency in the presence of EGF, as the FRET efficiency now appears to be straddling the monomer proximity line. Thus, a substantial decrease in the heterointeraction occurred, as expected.

Under the conditions of this experiment, over 90% of the EGF present is expected to be ligand-bound. We therefore considered these data in the context of the models in Fig. 1 (A–C), where one receptor is EGF-bound EGFR, and the other receptor is unliganded VEGFR2. The homodimerization constant for EGF-bound EGFR has been measured previously as 0.053 μm²/receptor (77). The MSE analysis, shown in Fig. 8B, reveals that the monomer model exhibits the lowest MSE, supporting the fact that EGF largely abolished the heterointeraction.

We further analyzed the FRET data by first correcting for the monomer proximity FRET contribution and then histogramming the corrected FRET data. As seen in Fig. 8C, the data for EGFR–VEGFR2 in the presence of EGF is effectively a Gaussian centered around zero, similar to the case of the monomer.
control LAT (Fig. 2D) but very different from the case of EGFR–VEGFR2 in the absence of ligand. In agreement with this assessment, an analysis of variance indicates that the LAT and EGF data are not significantly different, but the no-ligand data and EGF data are very significantly different (p < 0.0001). These results provide more evidence that EGF substantially decreases the EGFR–VEGFR heterointeraction, consistent with expectations.

Next, we performed a similar experiment, adding the ligand ephrin-A1-Fc to ECTM EGFR–ECTM EPHA2 heterodimers. This is a ligand with subnanomolar affinity for EPHA2, which induces EPHA2 cluster formation (91, 92). It is dimeric, as it is fused to the Fc fragment of an antibody that dimerizes constitutively. As shown in Fig. 8D, the presence of 50 nM ephrin-A1-Fc caused a substantial decrease in FRET. Of note, the majority of the measured FRET efficiencies are very low and now lie below the monomer FRET proximity line. This can be explained by the fact that the liganded EPHA2 forms clusters, and thus the monomer proximity FRET model is now inaccurate. Indeed, it has been shown previously that the contribution of proximity FRET decreases as the cluster size grows (60, 61). Thus, the data in Fig. 8D are consistent with the idea that ephrin-A1-Fc sequesters EPHA2 into clusters and decreases its heterointeractions with EGFR.

**Discussion**

**FRET as a tool to study heterointeractions**

Here, we demonstrate that RTKs from unrelated subfamilies can interact with each other in the absence of ligand using a quantitative FRET technique that reports on the heterointeraction strength in live cells. We expand the previously published methodology (42) to determine the type of heterooligomer that forms when two different RTKs interact. In particular, we
develop models that describe heterodimers, heterotrimmers, and heterotetramers, and then we fit these models to FRET data to determine which of these models best describe the data. This method requires the acquisition of FRET binding curves over a broad range of concentrations. We were able to collect such binding curves for ECTM versions of several RTKs and to determine the oligomer size and the stability of the oligomer. Constructs that incorporated the kinase domains had poor plasma membrane expression, such that only a small fraction of the imaged cells had both receptors in their plasma membranes. Thus, whereas we showed that these RTKs interact, we were not able to quantitatively assess the interactions. In all cases where we were able to determine the oligomer size, we showed that the RTKs interact as heterodimers.

Although such heterointeractions have been studied previously, almost all of the research has been done with qualitative techniques. The most common method used in the literature to assess heterointeractions is co-immunoprecipitation (co-IP), followed by Western blotting (93, 94). One of the RTKs is immunoprecipitated, and the other RTK is blotted against. In this technique, DNA probes are attached to primary or secondary antibodies against the proteins of interest. When the probes come in close proximity, a rolling circular amplification reaction occurs, and this DNA product can be hybridized with fluorescently labeled oligonucleotides. This assay is performed in cells rather than lysates, preserving the native environment, although fixation and permeabilization are generally required. However, this assay is not able to provide information about interaction energies or stoichiometries, as it simply reports on the close proximity of the proteins of interest.

Another method that has been used to assess propensities for interaction is chemical cross-linking, where a chemical cross-linking agent is added before Western blotting is performed under non-denaturing conditions (44, 95, 96). This helps to stabilize weak or environment-dependent interactions that are often lost in co-IP. However, in addition to having similar issues as co-IP and not being able to provide interaction free energies, it has other shortcomings. First, the cross-linker is nonspecific, meaning that it cross-links all proteins in close proximity. As a result, bands of cross-linked proteins on gels are smeared and generally cannot be accurately quantified. Second, the cross-linking propensities depend not only on close proximity, but also on structure. Third, the cross-linker itself is expected to drive the equilibrium toward the oligomeric state, although this effect has not been investigated in detail.

Although less common than the previous two methods, another method that is used to study heterointeractions relatively frequently is the proximity ligation assay (97–100). In this technique, DNA probes are attached to primary or secondary antibodies against the proteins of interest. When the probes come in close proximity, a rolling circular amplification reaction occurs, and this DNA product can be hybridized with fluorescently labeled oligonucleotides. This assay is performed in cells rather than lysates, preserving the native environment, although fixation and permeabilization are generally required. However, this assay is not able to provide information about interaction energies or stoichiometries, as it simply reports on the close proximity of the proteins of interest.

Whereas these methods will continue to have utility in the future, the FRET method that we use here can be used in conjunction with these methods to yield a more comprehensive description of the interactions. We are thus hopeful that this method will have a broad applicability in RTK research.
Here we find that in the absence of ligand, EPHA2 interacts with FGFR1, FGFR2, and FGFR3; VEGFR2 interacts with FGFR1, FGFR2, and FGFR3. We further show that EGFR–VEGFR2, EGFR–EPHA2, and EPHA2–VEGFR2 all form heterodimers. Surprisingly, we find that the stabilities of the heterodimers are quite similar to the stabilities of the homodimers in the absence of ligand. This is an unexpected result, and it suggests that these heterointeractions could have a significant effect in regulating RTK function. As the strengths of interaction are similar, both homodimers and heterodimers are expected to form under physiological conditions and to depend primarily on the relative RTK expression levels, in accordance with the law of mass action. In cancer, RTKs are often overexpressed, and ligand levels can be reduced (17, 18). Under these conditions, unliganded RTK heterodimers are expected to form with high probabilities.

**Open structural and mechanistic questions**

An open question is what physical contacts between the RTKs are enabling these heterointeractions. One possibility is that the TM domains play a key role, as it has been argued in the literature that the interactions between RTK TM domains are promiscuous, forming many weak, nonspecific interactions (15, 101, 102). It is also possible that interactions between the kinase domains contribute, as the kinase domain is highly conserved across families (79, 80). Previous FRET studies of RTK homodimers have shown that the deletion of their intracellular domains usually leads to dimer destabilization, both in the presence and absence of ligand (6, 78).
However, the intracellular domain is not required, as the ECTM versions of all RTKs that were studied here engaged in heterointeractions. It is also possible that there is no universal mechanism for the heterointeraction, but rather, each pair interacts differently.

High-resolution structural information about RTK heterodimers is currently not available, and the steps leading to their phosphorylation and signaling are unknown. The mechanism of activation of the different RTK homodimers is believed to be different. For instance, EGFR is believed to be activated through a distinct allosteric mechanism that involves the formation of asymmetric dimers (103). Questions therefore arise as to how the EGFR kinase and the other kinases will function within heterodimers. Further work is needed to understand the mechanistic details of RTK heterodimer formation and activation.

**Interactions with ligands and the RTK interactome**

We investigated the effect of ligand on heterointeractions by adding the cognate ligands of EGFR, EGF, and EPHA2, ephrinA1-Fc. We showed that the presence of saturating quantities of these ligands abolished the EGFR–VEGFR2 and EGFR–EPHA2 interactions, respectively. This means that the presence of ligand does not just shift the population of RTKs from monomers to dimers, but also decreases the concentration of heterodimers. One study found that an interaction between ROR1 and ErbB3 resulted in a unique tyrosine on ErbB3 being phosphorylated, and this resulted in a novel modulation of the Hippo–YAP pathway (38). Thus, the ligand can act as a signaling modulator that alters the extent to which a signaling pathway is being activated, by altering the equilibrium between heterodimers and homodimers.

It is possible that all RTKs interact with each other and form an interaction network, which we called the “RTK interactome” (27). In this view, each RTK can influence numerous interconnected signaling pathways via direct interactions with RTK partners in the plasma membrane. There exists a complex equilibrium between multiple RTKs and their ligands, which is controlled by the expression of all interaction partners and the association constants that govern every interaction. This greatly increases the degree of signaling complexity that RTKs are able to achieve and makes the signaling highly responsive to environmental changes that alter RTK and ligand concentrations. The data presented here provide experimental support for this view and add a quantitative perspective to our understanding of the RTK interactome.

**Implications**

The RTK interactome concept may provide an explanation for differences in experimental data acquired for an RTK in different cell lines (27). Typically, the expression of unrelated RTKs is not taken into account, and the possibility of RTK heterodimer formation is ignored. However, it is conceivable that the presence of these heterodimers greatly influences the cell signaling outcomes. The data in this study are consistent with this view.

Heterointeractions, such as the ones quantified here, need to be considered in the design of RTK inhibitors for cancers and other diseases or disorders (27). Even if a drug successfully inhibits the signaling of an RTK homodimer, the drug will not have the desired effect if signaling-competent heterodimers are also present. Therefore, the RTK expression pattern in a cell type may be a critical factor in determining the performance of the inhibitor. Because an inhibitor can affect the functions of both homodimers and heterodimers—or possibly even affect the entire RTK interactome—it can lead to many unanticipated consequences. Accordingly, the understanding of RTK heterointeractions may help in understanding drug resistance in cancer therapy and in designing more effective treatments. Significant work is still needed to quantify all homo- and heterointeractions and to arrive at a comprehensive understanding of RTK signaling and its biological functions. The use of quantitative experimental methodologies and a theoretical framework will help guide such investigations.

**Materials and methods**

**Plasmids**

Plasmids encoding for different constructs of EGFR, EPHA2, VEGFR2, FGFR1, FGFR2, FGFR3, and LAT were used. In almost all cases, the plasmids was in the pcDNA3.1+ vector, and the fluorescent protein (either MT or eYFP, which together form a FRET pair) was positioned at the C terminus of the RTK constructs, attached via a flexible (GGS)3 linker to enable free rotation (104). The one exception is the full-length EGFR, which is in the pSSX vector and has a GGS linker. The plasmids encoding for ECTM EGFR, ECTM VEGFR2, ECTM FGFR1, ECTM FGFR2, ECTM FGFR3, EPHA2ΔSAM, full-length EGFR, and full-length EPHA2 have been described in detail in previous studies (1, 77, 78, 81, 92, 105, 106). The plasmids encoding for ECTM EPHA2, EGFRΔTail, VEGFR2ΔTail, and LAT were cloned using Gibson assembly. In all four cases, the receptor constructs were cloned into the pcDNA3.1+ vector using the NEBuilder HiFi DNA assembly kit (New England Biolabs, E5520S), and the primers were designed using the online NEBuilder assembly tool. For ECTM EPHA2, the last amino acid in the TM domain before the hydrophilic flexible (GGS)3 linker was Ile-558; for EGFRΔTail, the last amino acid was Leu-979; and for VEGFR2ΔTail, the last amino acid was Asn-1162; numbering is based on the sequences for isoform 1 in UniProt.

**Cell culture and transient transfection**

HEK 293T cells were purchased from ATCC. The cells were cultured in Dulbecco’s modified Eagle’s medium (Thermo Fisher Scientific, 31600034) supplemented with 3.5 g/liter glucose, 1.5 g/liter sodium bicarbonate, and 10% fetal bovine serum (HyClone, SH30070.03). The cells were maintained at 37 °C with 5% CO2 and passed every third day.

For the FRET experiments, HEK 293T cells were seeded at a density of 2.0 × 105 cells/dish in collagen-coated, glass-bottom Petri dishes (MatTek, P35GCOL–1.5–14–C). Approximately 24 h later, the cells were transfected with the desired plasmids using Lipofectamine 3000 (Invitrogen, L3000008) according to
the manufacturer’s protocol. Twelve hours after transfection, cells were serum-starved by replacing the medium with fetal bovine serum–free Dulbecco’s modified Eagle’s medium that also lacked phenol red (MilliporeSigma, D2902) to ensure that endogenous ligands were not present during imaging. To remove all traces of phenol red, the imaging dishes were washed twice with the phenol red–free medium. When using the VEGFR2 plasmids, 10 nM sodium butyrate was added 6 h after transfection and after serum starvation to increase expression. In the case of ligand experiments, 0.1% (w/v) BSA was added to the imaging dish during starvation to prevent ligands from binding to the imaging dish.

**FRET imaging and data analysis**

FRET experiments were performed using the FSI method described previously (1). The medium in the imaging dish was replaced with hypotonic swelling medium (1:9 serum-free media/distilled H2O, 25 mM HEPES), as this reversibly “unwrinkled” the plasma membrane (107), enabling accurate concentration measurements. If ligands—either 10 nM EGF (Cell Signaling Technology, 89165C) or 50 nM dimeric Ephrin-A1 Fc (R&D Systems, 602-A1-200)—were used, 0.1% (w/v) BSA was added to the swelling medium, and then the ligands were added. Each dish was imaged for no more than 2 h. Imaging was done using a two-photon microscope equipped with the OptiMiS True Line Spectral Imaging system (Aurora Spectral Technologies, Shorewood, WI, USA) (108, 109). A Mai Tai laser (Spectra-Physics, Santa Clara, CA, USA) was used to generate femtosecond mode-locked pulses, and two images were acquired per cell: one at 840 nm to primarily excite the donor and one at 960 nm to primarily excite the acceptor. Cells singly transfected with a construct labeled with MT as well as cells singly transfected with a construct labeled with eYFP were also imaged to obtain the fluorescent spectra of the fluorophores. Pixel-level fluorescent intensities were converted into 2D concentrations using purified MT and eYFP solutions (110) to create calibration curves as described (1).

Cells were analyzed in MATLAB as described previously (1, 42). For each cell, two 2-3-μm stretches of plasma membrane were analyzed. The 2D concentration of the RTK labeled with the donor ([donor]), the 2D concentration of the RTK labeled with the acceptor ([acceptor]), and the apparent FRET efficiency ($E_{app}$) were determined in each membrane region. Proximity FRET ($E_{prox}$) (111, 60) was corrected for using the following,

$$E_{oligo} = \frac{E_{app} - E_{prox}}{1 - 2E_{prox} + 2E_{app}E_{prox}} \quad \text{(Eq. 31)}$$

where $E_{oligo}$ is the FRET due to sequence-specific interactions (61).

Furthermore,

$$E_{oligo} = \frac{\text{[heterooligomer]} E}{[\text{donor}]} \quad \text{(Eq. 32)}$$

where $\text{[heterooligomer]}$ is the concentration of either heterodimers, heterotrimers, or heterotetramers and $E$ is the intrinsic FRET, a structural parameter depending on the distance between the two fluorophores and their orientation (112). By combining this with the equations under “Heterointeraction models,” a two-parameter fit for the heteroassociation constant, $K_{het}$ (e.g. $K_{XX}$, $K_{XY}$), and $E$ was performed using the experimentally measured [donor], [acceptor], and $E_{app}$. Data fitting was performed as described previously using MATLAB (1, 42). Statistical significance was determined using one-way analysis of variance in PRISM.

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**Data availability**

All raw FRET data are available free of charge upon request from Kalina Hristova (kh@jhu.edu). All other data are contained within the article.

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**Abbreviations**—The abbreviations used are: RTK, receptor tyrosine kinase; FSI, fully quantified spectral imaging; MT, mTurquoise; 2D, two-dimensional; EC, extracellular; TM, transmembrane; IC, intracellular; EGF, epidermal growth factor; EGFR, EGF receptor; VEGFR2, vascular endothelial growth factor receptor 2; FGFR, fibroblast growth factor receptor; EPHA2, EPH receptor A2; YFP, yellow fluorescent protein; eYFP, enhanced YFP; LAT, linker for activation of T cells; MSE, mean squared error; co-IP, co-immunoprecipitation.

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plays an important role in the proliferation of embryonic neurons.