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Early signaling dynamics of the epidermal growth factor receptor

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Despite extensive study of the EGFR receptor (EGFR) signaling network, the immediate posttranslational changes that occur in response to growth factor stimulation remain poorly characterized; as a result, the biological mechanisms underlying signaling initiation remain obscured. To address this deficiency, we have used a mass spectrometry-based approach to measure system-wide phosphorylation changes throughout the network with 10-s resolution in the 80 s after stimulation in response to a range of eight growth factor concentrations. Significant changes were observed on proteins far downstream in the network as early as 10 s after stimulation, indicating a system capable of transmitting information quickly. Meanwhile, canonical members of the EGFR signaling network fall into clusters with distinct activation patterns. Src homology 2 domain containing transforming protein (Shc) and phosphoinositol 3-kinase (PI3K) phosphorylation levels increase rapidly, but equilibrate within 20 s, whereas proteins such as Grb2-associated binder-1 (Gab1) and SH2-containing tyrosine phosphatase (SHP2) show slower, sustained increases. Proximity ligation assays reveal that Shc and Gab1 phosphorylation patterns are representative of separate timescales for physical association with the receptor. Inhibition of phosphatases with vanadate reveals site-specific regulatory mechanisms and also uncovers primed activating components in the network, including Src family kinases, whose inhibition affects only a subset of proteins within the network. The results presented highlight the complexity of signaling initiation and provide a window into exploring mechanistic hypotheses about receptor tyrosine kinase (RTK) biology.

Significance

To date, poor temporal resolution of response measurement has obscured the complex initiation of receptor tyrosine kinase (RTK) signaling that governs cellular response to stimulation. To address this deficiency, we have performed a systems-level characterization of the phosphorylation changes that occur in the immediate period after growth factor stimulation with 10-s resolution. We treated MCF-10A cells with EGFR and measured tyrosine phosphorylation levels from 0 to 80 s on hundreds of sites in the cell. Examining phosphorylation dynamics on this timescale reveals patterns that were not observable with slower sampling rates. We further explore the roles of negative and positive feedback, providing further insight into systems-level behaviors of the EGFR receptor (EGFR) signaling network.


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Fig. 1. Schematic of experimental workflow. (A) MCF-10A cells were serum-starved for 24 h and treated with 0.2, 0.4, 1, 2.5, 5, 10, 20, or 100 nM EGF for 0, 10, 20, 30, 40, 50, 60, 70, and 80 s. Pooled normalization lysate was stimulated with 20 nM EGF for 60 s. Proteins were processed to peptides, desalted, enriched for phosphotyrosine, and analyzed on a QExactive mass spectrometer. Sample secondary fragmentation spectrum used for (B) sequence assignment and (C) reporter ion quantification.

of 8 M urea (Fig. 1). To assess the phosphorylation changes that may occur during freezing or lysis, control, nonstimulated cells and cells treated with 20 nM EGF for 60 s were left on liquid nitrogen for 1.5, or 15 min before lysis with urea or frozen on liquid nitrogen for 1 min, followed by incubation in urea lysis buffer for 15 or 30 min. Phosphoproteomic analysis of these samples demonstrate no significant change in phosphorylation during these steps, indicating that phosphorylation is rapidly and irreversibly stopped by our freezing and lysis procedure (Fig. S2). Lysates were then processed to peptides, labeled with stable isotope mass tags, and combined. One of the 10 channels was reserved for a fraction of a pooled lysate common to all runs, which provided a normalization value to facilitate comparison between multiple runs. Tyrosine-phosphorylated peptides were enriched by sequential immunoprecipitation using three antiphosphotyrosine antibodies and immobilized metal affinity chromatography before analysis by data-dependent liquid chromatography tandem mass spectrometry (LC-MS/MS). Secondary fragmentation spectra provided peptide sequence information and precisely localized phosphorylation modifications on each peptide, while reporter ion intensities, representing the relative amount of peptide in each of the 10 channels, were extracted to quantify temporal dynamics of phosphorylation (Fig. 1B and C). This approach yielded data on several hundred phosphorylation sites that occur on tryptic peptides amenable to LC-MS/MS analysis. Although not a complete list of all phosphorylation events in the network, they make up a representative fraction of many proteins implicated in EGFR signaling. Heat maps were used to visualize phosphorylation dynamics of peptides seen in multiple replicates for all eight of the ligand concentrations (Fig. 2 and Dataset S1).

Self-Organizing Map Clustering Reveals Distinct Signaling Modules. To identify sites with similar response patterns in the network, self-organizing maps (SOMs) were used. SOMs cluster self-similar patterns in complex data, including protein phosphorylation profiles (12). To correct for variability within single conditions while maximizing the number of phosphorylation sites seen across multiple conditions, phosphorytic dynamics from the 10-, 20-, and 100-nM stimulation conditions were concatenated to create an input vector with 27 features (0–80 s for three conditions). These profiles were clustered by Pearson correlation 10,000 times, and the frequency with which peptides appeared in the same cluster provided a quantitative metric of similarity (Fig. S3A). One prominent feature of this analysis is the tight integration of four EGFR autophosphorylation sites (pY1045, pY1068, pY1148, and pY1173), which all appeared in the same cluster >85% of the time, without once clustering with any other peptide (Fig. S3D) (13, 14). Meanwhile, EGFR pY974, which has been shown to be distinctly regulated from the autophosphorylation sites, clustered separately (15).

Two additional groups that emerged contained many phosphorylation sites canonically implicated in the EGFR network. The first sites contained on the adaptor proteins Src homology 2 domain containing transforming protein (Shc), Grb2 associated and regulator of MAPK protein (GAREM), and Src homology 2 domain containing adaptor protein B (Shb), which serve to recruit additional proteins including growth-factor-receptor-bound protein-2 (Grb2), a site on p85 alpha (PIK3R1), whose phosphorylation is thought to relieve inhibition of phosphoinositide 3-kinase (PI3K), and the nucleotide exchange factor Arhgef5 (Fig. S3C) (16). The abundance of these phosphopeptides is strongly increased at 10 s after stimulation, but plateau by ~30 s, and remains relatively constant thereafter. The second cluster included Grb2-associated binder-1 (Gab1), Y627 and Y659, SH2-containing tyrosine phosphatase (SHP2) Y584, tensin 3 Y802, and Y455 on ANKS1A, a regulator of EGFR recycling (Fig. S3D) (17). When phosphorylated, the two sites observed on Gab1 form a binding motif for SHP2, necessary for extracellular signal-regulated kinase (ERK) activation (18). We observe minimal delay between the appearance of this binding motif and phosphorylation of SHP2 on Y584, a site associated with activation (19). Compared with the other clusters, these peptides show a more gradual change in phosphorylation at the earliest times measured, but continue increasing until they reach similar relative levels as EGFR and Shc by 60 s (Fig. S3E).

Measurement of Receptor-Adaptor Complex Formation. One candidate mechanism for the slower phosphorylation kinetics of Gab1 versus Shc is a difference in recruitment to the receptor. To test this hypothesis, proximity ligation assays (PLAs) were used to verify selected immediate early phosphorylation events in situ and quantify receptor–adaptor interactions with low-nanometer resolution intact cells (20, 21). To measure early association events, cells seeded on glass coverslips were stimulated with 20 nM EGF for 0, 10, or 60 s; frozen on liquid nitrogen; and fixed in 4% paraformaldehyde for analysis. Orthogonal measurement of phosphorylation using PLA corroborates the previous observations of near-immediate phosphorylation of EGFR at 10 s, whereas ERK phosphorylation shows no increase at 10 s after stimulation. Meanwhile, both show strong activation at 60 s (Fig. S4). Imaging also confirmed uniform receptor dimerization patterns across the dish. Endogenous pairwise interactions between EGFR–Shc and EGFR–Gab1 were then measured at 0, 10, and 60 s after 20-nM EGF stimulation (Fig. 3A). By 10 s, EGFR–Shc interactions had reached 74% of the level observed at 60 s, whereas EGFR–Gab1 complexes had reached a corresponding level of 34%. Intriguingly, receptor–adaptor proximity almost identically matched relative phosphorylation dynamics, suggesting recruitment mechanisms may be the dominant component governing early phosphorylation behaviors (Fig. 3B).
Role of Phosphatases in Early Signaling. An intriguing feature observed at the phosphopeptide and protein complex level was the faster dynamics of Shc phosphorylation compared with EGFR phosphorylation, which was a counterintuitive result, as receptor phosphorylation would presumably precede recruitment and phosphorylation of adaptor proteins. One explanation for these patterns may be negative regulation by phosphatases, which may preferentially dephosphorylate receptor binding sites over adaptors, allowing more rapid accumulation of phosphorylation on Shc than EGFR.

To investigate the role of phosphatases in early signaling, cells were treated with 1 mM activated sodium orthovanadate (Na$_3$VO$_4$), a pan-tyrosine phosphatase inhibitor, for 15 min before 20 nM EGF stimulation. Several hundred phosphorylation sites observed demonstrated significant changes in phosphorylation response (Dataset S1). On EGFR, basal levels of phosphorylation were unaffected by phosphatase inhibition, but dynamics on stimulation showed significant differences within 10 s after stimulation (Fig. 4A). Although the four autophosphorylation sites have similar temporal profiles in untreated cells, relative levels of pY1148 in the stimulated, vanadate-treated condition are much greater than levels on other sites, indicating specific phosphatase activity against this site. The relative maxima reached in the vanadate condition were $\sim$fivefold higher compared with the untreated condition, suggesting the highest levels in the 20 nM condition represent at most $\sim$20% of the receptor in the cell. In addition to changes on the receptor, Gab1 also shows site-specific phosphatase regulation. Although Y406, Y627, and Y659 all show similar magnitude changes with vanadate treatment, Y373 shows a much larger relative increase (Fig. 4B).

In contrast to proteins near the top of the network, levels of ERK phosphorylation were significantly higher in the vanadate-treated, unstimulated cells. Ligand-independent ERK activation could either be a result of basal ERK signaling that is continually repressed by phosphatases or vanadate-mediated activation of an alternative pathway for ERK phosphorylation. To this latter possibility, one candidate mechanism involves Src family kinases (SFKs), which are both positively and negatively regulated by phosphorylation (22). In vanadate-treated, unstimulated cells, phosphorylation levels on many SFK sites were elevated, along with several canonical Src substrates, such as p130Cas and delta catenin (Fig. 4C) (23, 24). These data suggest that SFKs are active in the absence of ligand, but are continually repressed by phosphatase activity in unstimulated cells.

Role of Src Family Kinases in Early Signaling. To explore the potential for Src kinase activity as a primed activating component of immediate EGFR signaling, MCF-10A cells were treated with dasatinib, a SFK inhibitor, at 100 nM for 15 min before 20 nM EGF stimulation. Phosphorylation of Src substrates was significantly decreased by inhibitor treatment, but receptor phosphorylation dynamics on...
network at 10-s resolution in adherent epithelial cells in response to eight different ligand concentrations. This method builds on previous mass spectrometric analyses by incorporating a flash-freezing step to precisely terminate signaling with no residual phosphorylation changes.

In addition to rapid responses seen on EGFR and proximal adaptor proteins, we observed phosphorylation changes within 10 s on proteins such as cortactin, plakophilin, and tensin, all proteins not known to be directly regulated by the receptor (Dataset S1). Early phosphorylation on these cytoskeletal components may explain the rapid phenotypic responses observed in membrane protrusion. Rapid downstream phosphorylation changes challenge the preconceived timing of signaling, where receptors and adaptor proteins are phosphorylated and initiate signaling cascades that occur on the timeframe of minutes, and instead indicate a system capable of disseminating information almost immediately across many regions of the network, with specific regulatory mechanisms that tightly control the dynamics of protein phosphorylation.

Fig. 3. PLA measurements of EGFR-adaptor complex formation. (A) Early dynamics of EGFR-adaptor complex formation monitored with in situ PLA after 20 nM EGF treatment. Cells were identified with nuclear stain (DNA), and individual complexes were detected as distinct spots (green). Images are representative of replicate experiments. (Scale bars, 5 μm.) (B) Quantification of complex formation expressed as a fraction of 60-s measurement (solid), plotted with relative phosphorylation levels (dashed) for Gab1 (red) and Shc (black). Complex values represent mean ± SEM.

Y1045, Y1068, Y1148, and Y1173 were not significantly changed, reinforcing their status as primarily autophosphorylation sites (Fig. S5 A and B). Shc pY317, thought to be primarily controlled by direct interaction with EGFR, also showed no significant changes in phosphorylation (Fig. S5C) (25).

Despite apparently intact signaling through Shc, PI3KR1 phosphorylation is significantly impaired as early as 10 s by Src kinase inhibition (Fig. S4). Likewise, phosphorylation on Gab1 and SHP2 are impaired in the early period after growth factor stimulation (Fig. 5 B and C). In addition to these upstream changes, ERK activation downstream is also impaired without SFK activity (Fig. 5D). These data indicate a necessary role for SFKs in activating select components of the EGFR signaling network immediately after ligand binding.

Discussion

Because of its implication in human disease and broad control over cellular behavior, the EGFR signaling network has been one of the most extensively studied systems in biology. Recently, systems biology has provided tools that offer increasing coverage of network behavior. However, much of these data are projected onto a framework derived from traditional biochemical studies. Although knowledge of binary interactions between network proteins has been useful, they often obscure the emergent features that arise when the network is examined as a whole. To shed light on the functional characteristics of the network, we report the first systems-level measurement, to our knowledge, of the EGFR signaling network at 10-s resolution in adherent epithelial cells in response to eight different ligand concentrations. This method builds on previous mass spectrometric analyses by incorporating a flash-freezing step to precisely terminate signaling with no residual phosphorylation changes.

In addition to rapid responses seen on EGFR and proximal adaptor proteins, we observed phosphorylation changes within 10 s on proteins such as cortactin, plakophilin, and tensin, all proteins not known to be directly regulated by the receptor (Dataset S1). Early phosphorylation on these cytoskeletal components may explain the rapid phenotypic responses observed in membrane protrusion. Rapid downstream phosphorylation changes challenge the preconceived timing of signaling, where receptors and adaptor proteins are phosphorylated and initiate signaling cascades that occur on the timeframe of minutes, and instead indicate a system capable of disseminating information almost immediately across many regions of the network, with specific regulatory mechanisms that tightly control the dynamics of protein phosphorylation.

Fig. 4. Tyrosine phosphatase inhibition alters many components of the network. Selected phosphorylation patterns in response to 20 nM EGF stimulation with (dashed) and without (solid) 15 min pretreatment with 1 mM NaVO₃ for (A) EGFR pY1045, pY1068, pY1148, and pY1173 and (B) Gab1 pY259, pY373, pY659, and pY695. (C) Phosphorylation levels for several peptides were significantly altered in the pretreated (white) condition compared with untreated (black) in the absence of EGF.
One of these mechanisms, initially uncovered through clustering via SOMs, appears to be recruitment to the receptor. Through a combination of temporal phosphorylation profiling and PLA, we have shown that Gab1 and Shc phosphorylation dynamics are nearly identical to the speed with which these proteins become proximal to the receptor, implying that the rate-limiting step in adaptor phosphorylation is physical association with the receptor. This observation also may imply that the EGFR kinase domain does not have significant substrate-selective catalytic activity. One of the proteins observed to behave similarly to Shc is GAREM, an adaptor protein recruited to EGFR through interaction with the SH3 domain of Grb2 (26). Although Gab1 is also capable of being recruited to EGFR through interaction with Grb2, GAREM phosphorylation occurs more rapidly than Gab1, suggesting that Grb2 recruitment to the receptor may not be the limiting factor in Gab1 phosphorylation. Instead, an alternative mechanism, PIP₃ interaction with the PH domain of Gab1, may be the primary mode of recruiting Gab1 to the membrane, where it can interact with the receptor (27).

Recruitment is not necessarily the rate-limiting step for phosphorylation of all proteins in the network. For instance, temporal phosphorylation profiles for pY627 and pY659 on Gab1 and pYS84 on SHP2 were identical, despite previous reports suggesting that Gab1 phosphorylation on these sites was required for SHP2 recruitment to the receptor complex.

Our data indicate a prominent and site-selective role for tyrosine phosphatases in mediating the immediate–early signaling events after receptor activation. Although inhibition of tyrosine phosphatase activity has minimal effect on the basal levels of phosphorylation on EGFR and Gab1, the temporal response to stimulation of selected sites in the network was significantly affected by vanadate treatment. Maximum phosphorylation levels of EGFR pY1148 increased by fivefold in stimulated, vanadate-treated cells relative to stimulated control cells, whereas other EGFR sites only showed an ∼twofold increase. Coupling the vanadate results with previous stoichiometry data, which have shown pY1148 to be ∼threefold more phosphorylated compared with other sites on the receptor, indicates that phosphorylation of other sites, such as EGFR pY1173, represents no more than 10% of the total receptor population, even under saturating ligand conditions (28).

Gab1 pY373 appears to be more strongly targeted by phosphatase activity compared with other Gab1 sites. It is also the...
only observed Gab1 site that contains the consensus binding motif for RasGAPs (pY-X-X-P), which oppose ERK activation. Previous work has shown specific phosphatase activity against such binding sites, which may explain the observed changes (29).

Coupling our high-temporal-resolution measurements with inhibition of Src-family kinases with dasatinib uncovered a mechanistic connection between SFKs and ERK activation. In cells treated with dasatinib, Gab1 phosphorylation response to stimulation was significantly decreased, suggesting its recruitment may be affected by SFK activity. As mentioned previously, Gab1 interaction with EGFRF may be limited by PI3P-mediated recruitment to the membrane. Correspondingly, we observe decreased phosphorylation of PI3K in stimulated, dasatinib-treated cells. Together, these data highlight a pathway in which EGFR activation leads to increased SFK activity, resulting in phosphorylation of PI3K within 10 s. Increased PI3K activity increases PI3P levels, recruiting Gab1 to the membrane, where it can interact with EGFR and be phosphorylated. SFK inhibition leads to loss of this pathway, decreased phosphorylation of pY627 and pY659, and SHP2 pY584. Without SHP2 activity to eliminate RasGAP binding sites, ERK shows significantly impaired activation at 80 s in dasatinib-treated, stimulated cells (Fig. 6). We propose that both Shc and SFK-mediated, PI3K-PI3P-associated Gab1 recruitment are necessary for full activation of the ERK MAPKs.

These proposed mechanisms for signaling are derived from combining improved temporal resolution with select network perturbations. The method outlined here allows for the examination of specific biochemical hypotheses in the context of the full signaling network, which will provide a more comprehensive, functional understanding of network behavior. The platform is applicable for a variety of cell types, including those that express multiple members of the ErbB family, and can also be used to query the mechanistic effects of small molecules and biologics. Combining the methods described here with new genetic engineering techniques will further allow for the in vivo characterization of specific phosphorylation sites in the whole network. With improved tools for making measurements that clarify complex biology, it is our hope that this process may improve our understanding of signaling systems.

Methods
MCF-10A cells were maintained in Complete Media, as in ref. 28. Cells were serum-starved for 24 h before stimulation with EGF dissolved in water for 0, 10, 20, 30, 40, 50, 60, 70, or 80 s. The common pooled normalization channel was treated with 20 nM EGF for 60 s. lysates were processed to peptides, labeled with TMT 10-plex, combined, and immunoprecipitated as described previously (28). After elution from immunoprecipitation, a second round of enrichment was performed by immobilized metal affinity chromatography. Peptides eluted from immobilized metal affinity chromatography were analyzed by ESI LC-MS/MS on a QExactive Mass Spectrometer (Thermo Scientific) operated in information-dependent acquisition mode, as described in ref. 12. Raw mass spectral data files were searched against SwissProt database containing Homo sapiens protein sequences using Mascot version 2.4. MS/MS spectra of phosphorylated peptides observed in multiple biological replicates were validated using computer-assisted manual validation (CAMV) software to confirm phosphorylation assignment and isolation purity (30). From accepted scans, reporter ion quantification was extracted, isoCote corrected, and normalized based on median relative protein quantification ratios.

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