



A live-cell assay for the detection of pre-microRNA–protein interactions†

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Recent efforts in genome-wide sequencing and proteomics have revealed the fundamental roles that RNA-binding proteins (RBPs) play in the life cycle and function of coding and non-coding RNAs. While these methodologies provide a systems-level view of the networking of RNA and proteins, approaches to enable the cellular validation of discovered interactions are lacking. Leveraging the power of bioorthogonal chemistry- and split-luciferase-based assay technologies, we have devised a conceptually new assay for the live-cell detection of RNA–protein interactions (RPIs), RNA interaction with Protein-mediated Complementation Assay, or RiPCA. As proof-of-concept, we utilized the interaction of the pre-microRNA, pre-let-7, with its binding partner, Lin28. Using this system, we have demonstrated the selective detection of the pre-let-7-Lin28 RPI in both the cytoplasm and nucleus. Furthermore, we determined that this technology can be used to discern relative affinities for specific sequences as well as of individual RNA binding domains. Thus, RiPCA has the potential to serve as a useful tool in supporting the investigation of cellular RPIs.

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Introduction

Recent studies have shown that RNAs are invariably bound to and often modified by RNA-binding proteins (RBPs).^{1–6} Thus, it is no surprise that RBPs have been found to play key roles in regulating many aspects of coding and non-coding RNA biology, including RNA processing, nuclear export, cellular transport, function, localization, and stability.^{1,5,7,8} These efforts are carried out by >1500 unique RBPs that utilize a variety of RNA-binding domains (RBDs) to achieve oftentimes specific and high affinity interactions with target transcripts.^{1,2,7–9} Accordingly, disruption of this complex network of RNA–protein interactions (RPIs) has been implicated in a number of human diseases including neurodegenerative, neuromuscular, neurodevelopmental and cardiovascular diseases, encephalo-, ribosomo- and neuropathies, autoimmune disorders, and many types of cancer.^{7,10–13} Thus, the targeting of RBPs and RPIs has arisen as a new frontier in RNA-targeted drug discovery.¹⁴

Much of our knowledge regarding RPIs has been generated *via* large-scale sequencing and proteomics efforts.^{15–17} Methods

such as RNA-immunoprecipitation followed by sequencing (RIP-seq) and crosslinking followed by immunoprecipitation and sequencing (CLIP-seq) have enabled the protein-centric identification of the RNA targets of a select RBP.^{18,19} On the other hand, RNA-centric approaches, which typically utilize terminally-tagged RNA substrates for affinity enrichment,²⁰ have employed quantitative mass spectrometry for the proteomic discovery of proteins bound to a specific RNA.^{21–24} While both of these methods have proved overlapping, yet non-redundant,²³ each provides a systems biology view of the networking between RNA and protein, warranting the need for downstream technologies for validating discovered interactions.

Traditionally, the validation of RPIs has been carried out *via in vitro* biochemical and biophysical methodologies.^{25–28} Although useful, these techniques are low-throughput and quite laborious as they require the expression and purification of RBPs. Moreover, as these experiments are performed outside of cells, a high rate of false hits may be observed: false negatives due to the lack of required RBP binding partners or RBP post-translational modifications, and false positives due to formation of non-physiological RPIs in solution.^{8,28–30} Early cellular assays of RPIs relied on yeast three-hybrid systems for the high-throughput screening of RNA sequences that bind to a specific RBP; yet, like the *in vitro* methods, this approach is performed outside of human cells and relies solely on the physical properties of the RNA and protein, not their natural biological activities.^{31–33} Mammalian cell-based assays have been reported and primarily rely on fluorescence imaging-based approaches for the cellular detection of RPIs.^{34–39} These methods are also not without drawbacks as tools for

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validating RPIs, particularly for interactions with poorly characterized RBPs, due to reliance on antibodies for RPI detection. Additionally, several of these techniques require that the RNA-of-interest is labeled with a protein-binding RNA affinity tag, such as the MS2 hairpin, to enable the design of multicomponent RNA–protein complementation assays to detect cellular RPIs *via* fluorescence and, more recently, chemiluminescence.^{39–41} While these methods have been shown to successfully detect interactions between proteins and long RNAs like mRNAs and lncRNAs, as well as enable the discovery of new RPIs,⁴⁰ the addition of a large hairpin RNA tag precludes it from use with small RNA species and could interfere with RPIs by altering local RNA structure and dynamics and affect RNA processing and localization.⁴²

Over the past few years, the Garner laboratory has developed high-throughput screening technology for the discovery of small molecule modulators of RNA biology, including RPIs.^{43–46} Most recently, we have developed a click chemistry-mediated complementation assay, a homogeneous platform in which signal is dependent upon RPI-driven protein complementation of a split luciferase engineered from NanoLuc (NanoLuc Binary Technology; NanoBiT), allowing for catalytic signal amplification upon detection of full-length RPIs.⁴⁷ As our previously developed assay platforms were biochemical in nature, to further advance our ability to study and manipulate RNA biology, we became interested in the development of a live-cell assay for RPIs. Herein, we report our efforts toward the development of a conceptually new approach for detecting RPIs in live cells.

Results and discussion

Development of a live-cell assay for detecting RPIs

Leveraging a related bioorthogonal chemistry-based strategy previously utilized by our group,⁴⁷ and making use of Promega's HaloTag (HT)⁴⁸ and NanoBiT⁴⁹ systems, we devised an approach to directly monitor RPIs in living cells called RNA interaction with Protein-mediated Complementation Assay, or RiPCA (Fig. 1). In brief, cells stably expressing SmBiT-HT (step 1) are transiently transfected with an RBP-LgBiT plasmid and RNA probe which contains a PEGylated chloroalkane motif (step 2). In the cell, the RNA probe is first labeled with SmBiT through a covalent reaction between HT and the chloroalkane handle on the RNA (step 3). Successful interaction between the RNA and RBP is then monitored *via* chemiluminescence (step 5), which is produced upon SmBiT-LgBiT reassembly, forming the functional NanoLuc protein (step 4) (Fig. 1).

RiPCA differs from the previously reported complementation assays for RPIs^{41,50–53} in several important ways: (1) the weak affinity of SmBiT and LgBiT (K_d of 190 μM)⁴⁹ ensures that signal generation is driven by the RPI, thereby enabling quantitative and accurate analysis. (2) Because our technology does not rely on the use of a protein-binding RNA affinity tag (*e.g.* MS2 hairpin) for RPI detection,^{39–41,54,55} we have the capability to detect RPIs beyond those involving mRNAs, including interactions with small non-coding RNAs such as miRNAs. (3) By using a NanoLuc-based format, enhanced sensitivity will be observed due to catalytic

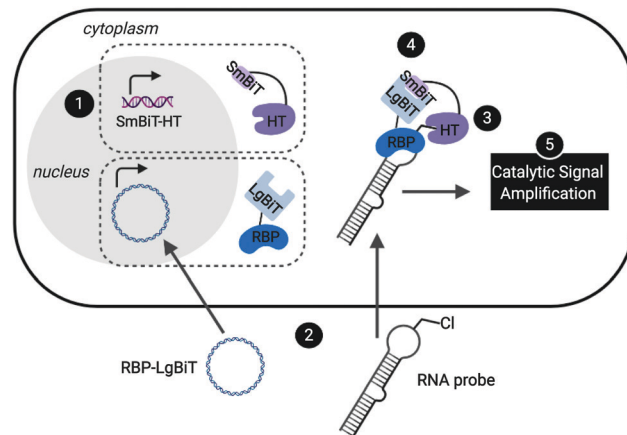


Fig. 1 Live-cell assay for the detection of RPIs. In RiPCA, cells stably expressing SmBiT-HT (step 1) are transfected with an RBP-LgBiT plasmid and chloroalkane labelled-pre-miRNA probe (step 2). HT covalently labels the RNA probe with SmBiT (step 3) and upon interaction of the RBP and RNA-HT-SmBiT, the BiTs reassemble forming functional NanoLuc (step 4), which when treated with a luciferase substrate results in chemiluminescence (step 5). Created with BioRender.com.

generation of chemiluminescence signal to promote favorable assay statistics. (4) The association of SmBiT and LgBiT is reversible, demonstrating potential for the manipulation of the RPI using small molecule inhibitors or other cellular stimuli.⁴⁹ Thus, we envisioned that the development of RiPCA would serve as a useful tool for the validation and study of cellular RPIs.

Based on our previous work in investigating the RPI between the pre-miRNA, pre-let-7, and RBP, Lin28,^{46,47} we used this as a model for assay development. By binding to the terminal loop of primary or precursor forms of select let-7 family members, Lin28 inhibits maturation, and in some cases induces degradation, reducing levels of mature let-7.^{56,57} Crystal structures of Lin28 in complex with the pre-element of pre-let-7 have been reported,^{24,57,58} which revealed that the protein uses three RNA-binding domains to interact with the hairpin loop of pre-let-7: an N-terminal cold shock domain (CSD) and two CCHC zinc knuckle domains (ZKD).⁵⁸ The CSD and tandem ZKDs are connected by a flexible linker, which allows Lin28 to bind to the diverse let-7 family members, as well as its mRNA targets.⁵⁸ To generate an RNA substrate, we used a chemically synthesized, 5' biotinylated pre-let-7d containing an aminohexylacrylamino uridine substitution at U35 in the loop region, in line with our previous work,^{43–45} that was subsequently converted into a PEGylated chloroalkane, herein referred to as pre-let-7d-Cl (Fig. S1, ESI†). We were confident that these modifications to the RNA would not disrupt interaction with Lin28 since our laboratory has previously developed biochemical assays using similar RNA probes containing a *trans*-cyclooctene (TCO) modification at the U35 position⁴⁷ or at the 5' end of the pre-let-7d hairpin.⁴⁶

To minimize the number of reagents that require transfection, a stable cell line expressing SmBiT-HT was generated from Flp-In HEK 293 cells. The 293-SmBiT-HT cells were then transiently transfected with pre-let-7d-Cl and LgBiT-tagged Lin28A constructs and grown in a 96-well plate. Both N- and C-terminally



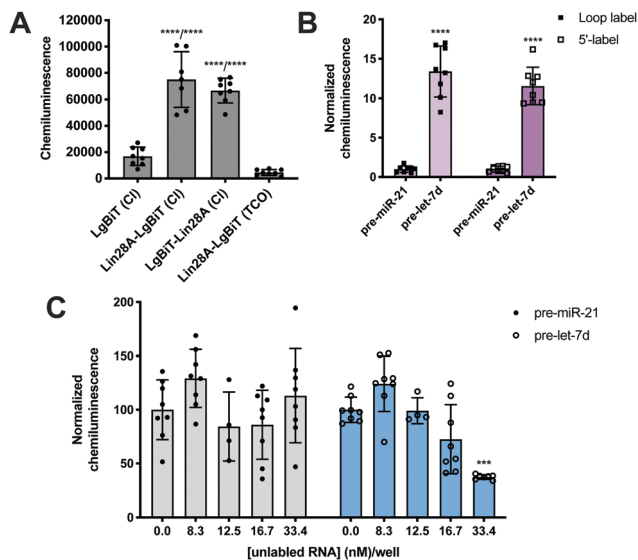


Fig. 2 Proof-of-concept. (A) Selective detection of the Lin28A-pre-let-7d RPI using RiPCA. (B) The effect of varying the location of the chloroalkane linker. (C) Selective inhibition of pre-let-7d signal in RiPCA by titrating in increasing concentrations of an unlabeled pre-miRNA probe. See Tables S3–S5 (ESI[†]) for *p* values for each set of experiments.

tagged Lin28A constructs were tested to determine if the location of the LgBiT tag would affect Lin28A binding to pre-let-7d-Cl. After incubation for 24 h, the media was changed, the cells were treated with NanoGlo Live Cell reagent and chemiluminescence resulting from the RPI was measured. Excitingly, as shown in Fig. 2A, our proof-of-concept was successful and the assay selectively reported the interaction of pre-let-7d-Cl with both LgBiT-labeled Lin28A constructs over a LgBiT only control (Fig. 2A). We also demonstrated the necessity of the chloroalkane modification, as minimal signal was detected with the corresponding non-reactive RNA probe, which bears a TCO handle in the terminal loop (Fig. 2A).⁴⁵ Similar levels of chemiluminescence signal enhancement were observed with both Lin28A-LgBiT constructs, and we chose to use the C-terminally tagged protein in further experiments.

We next turned our attention to the RNA substrate. To confirm that signal was dependent upon the selective interaction of Lin28A with pre-let-7d, we included a negative control sequence, pre-miR-21, that is not known to interact with Lin28A.²⁴ Notably, signal produced by pre-miR-21 was 12–14-fold less than that of pre-let-7d, indicating that RiPCA can selectively detect the pre-let-7d-Lin28A RPI (Fig. 2B). As such, we used pre-miR-21 treatment as a measure of assay background in subsequent experiments. Wondering whether the location of the chloroalkane modification might impact signal detection, we generated additional pre-miRNA probes that placed the chloroalkane modification at the 5' terminus of the hairpin. As shown in Fig. 2B, the location of the modification did not result in a notable difference in raw signal for pre-let-7d, nor a significant difference in signal enhancement measured as signal-to-background (S/B) for pre-let-7d compared to pre-miR-21 (Fig. S5, ESI[†]); negligible signal was again observed with pre-miR-21 demonstrating assay specificity. We further

explored the effect of modifications to our RNA probes by altering the length of the chloroalkane linker and observed that shorter linkers (PEG2) produced higher S/B compared to longer linkers (PEG4 and PEG6) (Fig. S6A, ESI[†]). Of note, neither differences in linker length nor uridine modifications altered the signal produced by pre-miR-21 probes (Fig. S6B, ESI[†]). As pre-miRNA probes containing a PEG4 linker produced more consistent signal in the assay (Fig. S6A, ESI[†]), we opted to proceed with the PEG4-modified RNA probes. Finally, we confirmed that the 5' biotin does not inhibit the pre-let-7d-Lin28A RPI in RiPCA (Fig. S6C, ESI[†]). It is prudent to note that our RNA probes contain biotin due to their utility in other applications in the lab and is not explicitly required for RiPCA.

With our ideal Lin28A and pre-let-7d substrates in hand, we examined the dynamic range of the assay by adjusting the amount of RBP-LgBiT plasmid and pre-let-7d-Cl transfected per well. Interestingly, as the amount of RBP-LgBiT plasmid transfected per well was increased, the S/B observed between pre-let-7d and pre-miR-21 decreased (Fig. S7A, ESI[†]). This is likely a result of increased instances of non-specific interactions between SmBiT and LgBiT due to elevated cellular concentration of Lin28A-LgBiT. In contrast, increasing the amount of pre-miRNA-Cl transfected per well resulted in a concentration-dependent increase in S/B (Fig. S7B, ESI[†]). From these results, and with the goal of maximizing S/B while minimizing the amount of transfection reagent used in each experiment, we proceeded with loop labeled pre-miRNA probes and transfected minimal quantities of the RNA and RBP reagents.

Finally, to demonstrate that RiPCA signal is produced as a result of ternary complex formation between pre-let-7d-Cl, SmBiT-HT, and Lin28A-LgBiT, cells were treated with increasing amounts of an unlabeled pre-miRNA, pre-let-7d or pre-miR-21. Importantly, we observed that unlabeled pre-let-7d, but not pre-miR-21, inhibited signal generated by treatment with the corresponding chloroalkane-labeled RNA probe (Fig. 2C). These data indicate that labeling of RNA with SmBiT does not contribute to non-specific interactions, further demonstrating that the specificity and strength of the signal in the assay is a direct measure of Lin28A binding to its target pre-miRNA.

Encouraged by these promising data, we next set out to determine whether our read-out reflects Lin28 binding specificity by performing RiPCA with a variety of pre-miRNA probes (Fig. 3A). Again, we observed no signal in the presence of pre-miR-21, as well as pre-miR-34a, another sequence that has not been shown to interact with Lin28A (Fig. 3B).²⁴ Additionally, we examined the interaction between Lin28A and other pre-let-7 isoforms, pre-let-7a-1 and pre-let-7g, which belong to different sub-classes of pre-let-7s. Analysis of let-7 sequences enriched in recent CLIP experiments revealed differential binding of distinct pre-let-7s by Lin28.⁵⁶ This study found that Lin28 more strongly interacts with and regulates pre-let-7s that contain both a CSD and ZKD binding site (CSD⁺) compared to those that only contain a ZKD binding site (CSD⁻) (Fig. 3A).^{56,57} This finding was corroborated in a proteomics study that identified RBPs enriched with specific pre-miRNA sequences, which demonstrated that Lin28 interacted primarily with CSD⁺ pre-let-7s.²⁴



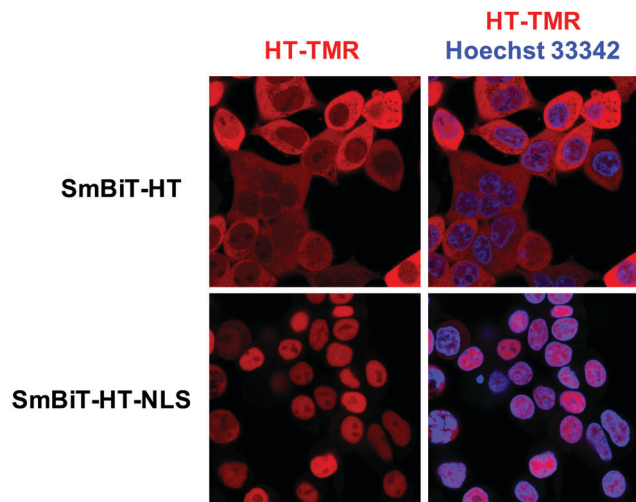


Fig. 5 Localization of SmBiT-HT. Detection of SmBiT-HT and SmBiT-HT-NLS in Flp-In HEK293 cells *via* confocal microscopy. SmBiTs were detected following conjugation with a tetramethylrhodamine (TMR)-labelled HT ligand (red). Nuclei were stained with Hoechst 33342 (blue). Images on the left are of the SmBiT-HTs alone; those on the right are overlaid with the nuclear stain.

Because Lin28A and Lin28B have been shown to function as let-7 repressors both within the cytoplasm and nucleus,^{59,60} we again used this as a model. Using the SmBiT-HT-NLS expressing cell line, we detected the production of chemiluminescence signal indicative of the interaction of Lin28A-LgBiT, as well as LgBiT-Lin28A, with pre-let-7d (Fig. S9, ESI[†]). Relative to cytoplasmic RiPCA, we observed a 2.7- and 2.2-fold reduction in raw chemiluminescence signal produced upon interaction of pre-let-7d with Lin28A-LgBiT and LgBiT-Lin28A, respectively (Fig. S9, ESI[†]).

Curious to evaluate the signal produced by RNA probes generated with variable PEG linkers, we performed nuc-RiPCA with each set of probes. While much less difference was seen in the nucleus compared to the cytoplasm, shorter linkers (PEG2) produced slightly higher signal than longer linkers (PEG4 and PEG6) (Fig. S6D, ESI[†]). The consistent low-level signal detected with the various modified pre-miR-21 probes was recapitulated in nuc-RiPCA (Fig. S6E, ESI[†]). We were then interested in exploring whether we could tune the signal in nuc-RiPCA by adjusting the amount of DNA and pre-miRNA-Cl transfected. In contrast to cytoplasmic RiPCA, increasing the amount of DNA transfected resulted in greater signal relative to the pre-miR-21 control (Fig. S10A, ESI[†]). With respect to RNA transfection, nuc-RiPCA yielded the same trend as in the cytoplasm, and S/B increased with increasing pre-miRNA-Cl (Fig. S10B, ESI[†]).

Following this successful result, we tested nuc-RiPCA against other pre-miRNAs with both Lin28A-LgBiT and Lin28B-LgBiT. In accordance with our results obtained using the cytoplasmic assay, we observed signal only with the two CSD⁺ sequences, pre-let-7d and pre-let-7g, with both Lin28A- and Lin28B-LgBiT (Fig. 6). In addition to observing lower overall signal in the nucleus than in the cytoplasm, we also found that CSD⁺ sequences produced S/B that was between 1.8- and 2.7-fold lower

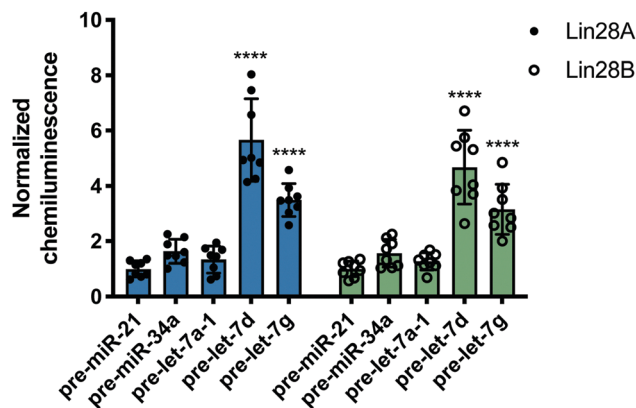


Fig. 6 Nuc-RiPCA. Selectivity of Lin28A-LgBiT (left) and Lin28B-LgBiT (right) against various pre-miRNAs. See Table S8 (ESI[†]) for *p* values.

than in the cytoplasm (Fig. 3 and Fig. 6). We hypothesize that the lower signal and S/B, as well as increased tolerance of LgBiT expression, could be the result of short residence time of the LgBiT-tagged protein in the nucleus, competition with endogenous Lin28B in the nucleus,^{59,60} or decreased pre-miRNA probe uptake into the nucleus. With respect to the latter, this is unlikely to be problematic, as molecules <40 kDa have been shown to readily traverse through the nuclear pore complex.⁶¹ We anticipate that the ability of RiPCA to enable organelle-specific RPI detection will open the door to its future application in the study of disease-relevant nuclear RBPs critical in human health.

Conclusions

RBPs and RPIs play important, but currently understudied roles in the maintenance of human health. While the advent of sequencing and quantitative mass spectrometry has dramatically enhanced our ability to globally profile these interactions, revealing intricate and complicated networks of RPIs, experimental validation of these data sets remains a challenge. Using chemical biology- and bioorthogonal chemistry-based strategies, we have developed an innovative assay for the live-cell detection of RPIs, RiPCA. Through this approach, we have selectively detected the interaction of the Lin28 proteins and their pre-let-7 targets in both the cytoplasm and the nucleus. We also demonstrate that RiPCA is capable of discerning between high and low affinity RPIs, using the individual Lin28 RBDs as a model. Combined, these data provide encouraging proof-of-concept for this emerging technology. Efforts towards its adaptation to validate putative RBP-pre-miRNA interactions discovered *via* proteomics,²⁴ as well as RPIs outside of those with pre-miRNAs, are currently being explored and will be reported in due course.

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

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