Nanobodies and recombinant binders in cell biology

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Antibodies are key reagents to investigate cellular processes. The development of recombinant antibodies and binders derived from natural protein scaffolds has expanded traditional applications, such as immunofluorescence, binding arrays, and immunoprecipitation. In addition, their small size and high stability in ectopic environments have enabled their use in all areas of cell research, including structural biology, advanced microscopy, and intracellular expression. Understanding these novel reagents as genetic modules that can be integrated into cellular pathways opens up a broad experimental spectrum to monitor and manipulate cellular processes.

Deciphering the inner workings of the cell requires specific molecular probes to measure the spatial and temporal dynamics of cellular structures. One class of probes that has been pivotal to modern cell biology is antibodies. Their ability to bind specifically to antigens such as proteins and even their posttranslational modifications (PTMs) has been exploited extensively to interrogate cellular function. For example, fluorescently labeled antibodies have been crucial staining reagents in molecular imaging techniques to reveal information on subcellular localization, abundance, and molecular interactions of biological antigens of interest. In addition, antibodies are essential to a vast range of biochemical analyses, including classic diagnostic techniques, such as Western blots and ELISA, but also to systems biology methods, such as mass spectrometry and ChIP-Seq (chromatin immunoprecipitation sequencing; Kidder et al., 2011), in which antibodies mediate the initial purification of the biological specimen.

Although conventional full-length antibodies are still the most widely used binding reagents for biochemistry and cell biology applications, their complex structural organization and their tedious manufacturing procedures have urged the development of new, alternative binding reagents. Such binders are either recombinantly generated immunoglobulin derivatives or synthetically designed from very different protein scaffolds. As such, recombinant binders are made to complement antibody-based fields of application or even enable completely new and innovative experiments. Of particular interest are binders that can be robustly expressed in living cells, a feature that is exclusive to small and stable binding molecules and cannot be performed easily with full-length antibodies, as a result of crucial inter- and intramolecular disulphide bridges that do not form in the cytoplasm. Thus, researchers have found a plethora of new applications in which binders have been combined with enzymatic or structural functionalities in living systems.

The development of in vitro screening techniques has been a decisive step for the rise and generation of recombinant binding reagents. These methods include classic phage display but also bacterial and yeast display as well as ribosomal and mRNA display. With such in vitro display techniques at hand, directed evolution strategies and genetic manipulation of binder sequences allow targeted engineering of key features, such as specificity, valence, affinity, and stability, enable derivatization toward smaller and more stable binding entities, and facilitate expression in heterologous hosts. Yet, it is the virtually limitless combinability of binder-mediated target recognition with any other chemical or biological function (viruses, translocation peptides, enzymes, structural proteins, dyes, toxins, and therapeutic agents) that opens up a whole universe of biotechnological innovations. Here, we provide a short overview of the most important recombinant binder formats and how they are generated followed by a few examples of how these valuable reagents can promote innovation and enable new discoveries in various fields of cell research. An in-depth discussion of molecular and structural aspects of some of these formats can be found in recent specialized reviews (Muyldermans, 2013; Plückthun, 2015).

Development of different recombinant binders

Immunoglobulin derivatives. Naturally produced immunoglobulins (IgG, IgM, IgA, IgD, and IgE) are universal weapons against pathogenic threats. The predominant isotype in nature is IgG, a 150-kD multichain/monodomain protein. IgG consists of two heavy chains and two light chains with variable domains (VH and VL; Fig. 1 A). The binding occurs via six (three in VH, three in VL) complementarity determining regions (CDRs). Derivatization of IgG allows the generation of functional Fab (~50 kD), scFv (~25 kD), and single, variable domain Vh or Vl fragments (Fig. 1 A). In Fab and scFv formats, the antigen binding surface is established via specific association of the Vh and the Vl domain, mediated by hydrophobic framework residues (Chothia et al., 1985). The necessity of noncovalent interdomain interactions for functional domain as-
In vitro binder selection with display techniques

Several polypeptide display techniques are available to identify antigen-specific binders in vitro. Although these methods use different biological platforms, they share common features, including the capacity to screen large gene libraries, physical coupling of the encoding DNA sequence with its respective protein, and the possibility to increase binding specificity and affinity by repetitive mutagenesis and selection cycles (panning).

Phage display
Phage display, the most common display technique, involves the display of a recombinant binder library on the surface of bacteriophages upon genetic fusion with a viral coat protein. Individual phages comprise a defined binder on the surface and the respective gene within a phagemid inside the phage particle. Challenging this phage library with an immobilized antigen allows for in vitro selection of specific binders that can be amplified and identified by reination of *Escherichia coli*. Most widely used are M13 filamentous phages.

Bacterial and yeast display
Both techniques rely on fusing the gene of the binder library to respective surface proteins. In contrast to phages, bacteria and yeast can be screened via flow cytometry. Displaying binders on yeast is additionally advantageous because the expression is mediated by an eukaryotic machinery, which has been demonstrated to be more effective in comparison to phage display (Bowley et al., 2007).

Ribosome and mRNA display
Ribosome display involves the in vitro transcription and translation of a DNA library. A genetic trick allows immobilization of the translated, correctly folded protein on the ribosome in a noncovalent complex with the respective mRNA, thereby providing the necessary genotype-phenotype type link. The variable pool of binder-ribosome-mRNA complexes is then incubated with immobilized antigen, and specific binders are amplified by reverse transcription.

mRNA display differs from ribosome display in that the in vitro translated polypeptide is covalently linked to its cognate mRNA via puromycin, a tRNA mimetic. In contrast to other display techniques, ribosome and mRNA display are independent of *E. coli* transformation and the accompanying potential loss of library diversity. In addition, the reverse transcription and PCR amplification steps in between successive selection cycles introduce sporadic mutations that mimic an affinity maturation step.

Nonimmunoglobulin binders
The specific needs of the various fields of application as well as patenting issues motivated the development of several alternative binder formats that are based on defined nonimmunoglobulin protein folds. Here, we describe a few that are well established and suited for cellular research (Fig. 1 B).

There are different strategies to engineer completely new classes of antibody mimetics. On the one hand, researchers have used natural protein folds as universal scaffolds for the generation of recombinant binding reagents. For example, the 10-kD fibronectin protein fold serves as a template for the bioengineered Adnectins/monobodies (Fig. 1 B). They are structured similarly to immunoglobulin domains with seven β sheets and three CDR-like loops located on the top of the barrel-like fold (Koid et al., 1998). Genetic randomization of the CDR-like loops allows the generation of libraries from which to retrieve binders with desired specificities after in vitro display, as was for example demonstrated for a monobody directed against the Ab1 SH2 domain (Wojcik et al., 2010). In contrast to variable Ig domains, the fibronectin structure does not depend on intramolecular disulphide bridges, thus facilitating functional applications in reducing environments such as the cytoplasm of living cells (Gross et al., 2013).

Another example of a synthetic binding reagent is anticalins, which derive from the 20-kD lipocalin fold (Beste et al., 1999). Lipocalins are eight-stranded β-barrel structures that mediate binding and transport of small molecules in their endogenous environment. Upon randomization of the four natural binding loops, the synthetic variant may be designed to specifically recognize antigens of interest with high affinities. The goblet-like binding pocket significantly differs from antibody paratopes (Fig. 1 B). Thus, the anticalin format is considered especially useful to generate binders against small molecules (Kornndörf er et al., 2003) and protruding conformational epitopes (Eggenstein et al., 2014).

Another class of affinity proteins has been designed based on the immunoglobulin binding protein A from *Staphylococcus aureus*. These so-called affibodies comprise three α helices without disulphide bonds and have a molecular mass of ~6.5 kD (Fig. 1 B). Random mutagenesis of defined residues within the two binding helices allows screening for target-specific binding reagents for biotechnological use (Nord et al., 1997).

In terms of size and binding mode, all aforementioned binder formats are more or less restricted to the fold of their natural template. On the other hand, engineering repeat proteins as recombinant antibody mimetics allows more modular design strategies. The basic principle of repeat binders lies within the consecutive arrangement of multiple repeat units (Fig. 1 B). The overall length of the repeat protein may be adjusted with respect to specific target properties. A prominent example for repeat protein binders are designed ankyrin repeat proteins (DARPins). DARPins are based on the ankyrin fold, which naturally mediates protein–protein interactions (PPIs) and has been observed in diverse protein families (Li et al., 2006; Al-Khodir et al., 2010).
The structural subunit of a DARPin consists of a β turn followed by a pair of antiparallel α helices and a loop, typically comprising 33 amino acids. Randomization of defined helix residues enables selection of high-affinity binders (Binz et al., 2003; Forrer et al., 2003; Kohl et al., 2003). In contrast to barrel-like binder folds with antigen-reactive loop structures, DARPins form slightly concave binding surfaces that favor large, conformational epitopes. As a result of the complete absence of disulphide bridges, DARPins are well suited as potential intracellular binders (Parizek et al., 2012) and can be expressed in large amounts in bacteria.

In addition to these established formats, a plethora of novel protein scaffolds has been developed and includes recombinant binders such as avimers (Silverman et al., 2005), affilins (Ebersbach et al., 2007), fynomers (Grabulovski et al., 2007), affitins (Mouratou et al., 2007), knottins (Smith et al., 1998), armadillo repeat proteins (Parramugarni et al., 2008), and the very recently published adhirons, which can be stably produced in large amounts in bacteria (Tiede et al., 2014). As a complement to conventional antibodies, man-made recombinant binders, based on immunoglobulin or nonimmunoglobulin folds, open up new possibilities for the life sciences. The available formats have common and unique properties (Table 1) that can be used to choose the best format for a given application. For all formats, however, the success in retrieving a potent binder with a desired target specificity largely depends on the library size as well as the techniques and conditions used for their screening.

**Applications of recombinant binders in molecular and cellular biology**

**Cellular proteomics.** The analysis of proteomes relies on the availability of high-quality binding reagents. They are necessary for the molecular analysis of specific target proteins and their spatiotemporal cellular abundance using standard detection methods such as microscopy and Western blot analysis. Furthermore, binder-mediated affinity purification in combination with systems biology techniques has contributed to a comprehensive understanding of cellular interactomes in development and disease. Although such experiments were historically performed with conventional antibodies, the use of recombinant binding reagents for proteomic application becomes increasingly popular, and international consortia have been installed with the long-term goal to cover the entire human proteome with renewable binders (Tassig et al., 2007; Colwill and Graßlund, 2011).

Biochemical and proteomic analyses of crude samples include affinity-based purification or depletion of specific com-
ponents using chromatography columns that, however, require large amounts of affinity material. Thus, expensive monoclonal and polyclonal antibodies are less attractive to purify or deplete endogenous proteins, whereas inexpensive immobilized-metal affinity chromatography (IMAC) systems, such as the Ni-NTA (nitrilotriacetic acid)/His-tag system, are widely used but also restricted to artificially tagged proteins. Consequently, recombinant binders that can be produced at low costs in high quantities offer valuable alternatives for immunoaffinity chromatography of biological samples (Blank et al., 2002; Grönwall et al., 2007).

The rise of comprehensive proteomic analyses started with the advent of protein microarrays enabling quantification of cellular protein components (Fig. 2 A). However, array analyses of complex protein samples require highly sensitive and specific detection reagents to measure low-abundant proteins (Fig. 2 A). In terms of recombinant probes, high-affinity scFvs have proven especially useful for protein arrays (Wingren et al., 2007) and allowed for biomarker profiling of cancers (Ingvarsson et al., 2008; Carlsson et al., 2010) and autoimmune diseases (Carlsson et al., 2011). However, the high cost per array is still the limiting factor in protein array technologies.

State-of-the-art proteomics nowadays mostly relies on the use of mass spectrometry to analyze complex protein mixtures (Walther and Mann, 2010). Affinity tools are used in mass spectrometry–based proteomics to purify or enrich defined targets
and subsequently identify PTMs and interacting factors (Fig. 2A). To bypass the time-consuming generation of specific binders, proteins are often targeted with established foreign epitopes. The use of GFP as tag in cellular proteomics allows the experimental link between live cell microscopy and proteome analysis (Cristea et al., 2005) and a high-affinity GFP-binders—are to some extent beneficial effect of auxiliary affinity reagents. The early successes with Fab (Kovari et al., 1995) or Fv (Ostermeier et al., 2009) for crystal contacts (Sennhauser et al., 2007; Bandeiras et al., 2008; Schönfeld et al., 2009; Veesler et al., 2009; Rasmussen et al., 2011a; Koide et al., 2012; Krishnamurthy and Gouaux, 2012).

Especially, β-rich chaperones (nanobodies and monobodies) have been shown to facilitate crystallization of macromolecules (Skrabana et al., 2010; Serrière et al., 2010) and lower the entropic cost of lattice formation. In particular, recombinant biologics are more amenable to the application of chaperones than native macromolecules. The effectiveness of the affinity reagents to assist the crystallization process originates from their shared favorable properties: facile identification and large-scale production of stable and soluble target binders. The DARPins, monobodies, anticalins, and affibodies have an additional advantage over nanobodies, Fabs, and scFvs because they lack disulphide bridges, allowing standard cytoplasmic expression to produce functional chaperones. The different interactions of a DARPin and a nanobody with the complex with human fibronectin ED-B (Extra Domain-B; Gebauer et al., 2013). Finally, affibodies seem to prefer a flat interacting surface (Hoyer et al., 2008). Therefore, all these chaperones should be considered complementary to each other, rather that competitive, to tackle the most difficult proteins to crystallize and to improve the diffraction quality. Moreover, different crystals of the same target protein complexed with alternative chaperones or with the same type of chaperone associated at different epitopes are instrumental to identify various alternative conformations of macromolecules, which might be linked to mechanism of action (Pecqueur et al., 2012; Chaikuad et al., 2014) or to degenerate binding with alternative partners.

Affinity-based biosensors, inhibitors, and imaging tools

Fluorescent biosensors are essential analytical tools to study molecular target structures at the cellular level. Although conventional antibodies are traditionally used to stain antigens in fixed cells, providing endpoint information only, FP-based bio-

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sensors are widely used for the spatiotemporal analysis of target structures in living cells. The genetic tagging of a protein of interest with a fluorescent reporter enables the dynamic visualization of molecular features in living cells. Such studies include the subcellular localization, conformational changes, and PPI that can be measured, e.g., by distance-dependent Förster resonance energy transfer (FRET). However, the fusion with FPs requires genetic manipulation and may compromise the biological function of the protein of interest (Hosein et al., 2003). Moreover, recombinant expression rarely reflects endogenous expression levels, and nonprotein targets cannot be studied with this strategy. For these applications, recombinant affinity binders offer new options to engineer novel intracellular biosensors.

Tracing and tracking. A straightforward application for dynamic intracellular tracking of endogenous target structures consists of the genetic fusion of specific binders with FPs (Fig. 3 A). For these applications, any recombinant binders can be used, as long as they are functionally expressed in living cells. The reported reagents cover diverse aspects of cell biology, including the visualization of cytoskeletal components (Rothbauer et al., 2006; Riedl et al., 2008, 2010), the DNA replication machinery (Burgess et al., 2012), and viral infections (Jones et al., 2010; Helma et al., 2012) as well as reporting on apoptotic progression (Zolghadr et al., 2012) and ubiquitin signaling (Sims et al., 2012). However, it is important to note that live-cell visualization of endogenous structures and their respective cell biology with affinity reagents requires careful monitoring of potential interference upon antigen binding, such as effects on stability or localization of the target. To prevent that genetic fusion of a nanobody impairs its affinity and specificity, fusion partners should be preferentially added at the C terminus of nanobodies, which is the natural connection site for the constant domain and thus distal from the antigen binding site.

Conformation and PTM sensors. As discussed in the structural biology section, affinity reagents can detect and stabilize specific conformational states of proteins. Consequently, refocusing such binders as intracellular biosensors potentially enables the spatiotemporal analysis of specific conformational changes and their biological implications in

Figure 3. Recombinant binders as intracellular biosensors, effectors, and tools for nanoscopy. [A] Categories of affinity-based live-cell biosensors and effectors. (top left) Fluorescently tagged tracers are used to monitor antigen-specific localization patterns. (top right) Conformation-specific and PTM-specific binders are recruited to respective sites of action. (bottom left) Binders that block biologically active target sites or modulate target function upon binding. (bottom right) An F3H assay to investigate PPIs in living cells. A GFP-specific nanobody is anchored to defined subcellular structures such as the artificially introduced LacO array visible as a spot in the nucleus or the endogenous nuclear lamina or centrosomes. In all of these cases, RFP-preys colocalize with GFP-bait fusions in the presence of an interaction. (B) Recombinant binder tools for nanoscopy. Superresolution techniques such as 3D-structured illumination microscopy require repetitive imaging of a single cell, which leads to severe bleaching of FPs. Nanobodies can thus be used to stabilize or enhance FPs and enable high quality image acquisition conditions. Importantly, their small size reduces the linkage error, as the distance between the fluorescent label and the actual specimen is minimized. Bar, 10 µm.
living cells (Fig. 3 A). Such application was shown for a nanobody that binds the activated conformation of the β2-adrenoceptor (Rasmussen et al., 2011a). Fusing this binder with GFP reports on the subcellular localization of activated β2-adrenoceptor and lead to the identification of endosomal membranes as initiation sites of acute G protein–coupled receptor signaling, which has previously been considered to occur exclusively from the plasma membrane (Irannajad et al., 2013).

Similarly, a DARPin that recognizes the activated, phosphorylated conformation of extracellular signal–regulated kinase (ERK; Kummer et al., 2012) was conjugated with a solvatochromatic merocyanine to quantitatively report on active pERK localization in living cells in absence and presence of an inhibitor of the upstream regulatory kinase MEK1/2 (Kummer et al., 2013). Interestingly, this DARPin recognizes a conformational change within the activation loop that is activation-dependent and thus indirectly reports on the primary modification, the ERK phosphorylation. Like conventional antibodies, recombinant binders can be generated against PTMs and nonprotein epitopes. Thus, live-cell application of recombinant binders paves the way toward dynamic analysis of PTMs as was shown with a GFP-tagged scFv that specifically recognizes histone acetylation (H3K9ac) and was used for dynamic live-cell and live-animal monitoring of epigenetic chromatin modulation (Sato et al., 2013).

**PPIs.** Systematic probing of PPIs is often performed in vitro or in yeast. In living cells, PPIs can be tested with FRET, which is technically demanding, or with protein fragment complementation assays, which do not report in real time and are irreversible. In contrast, the recently developed fluorescent 3-hybrid (F3H) assay is based on recombinant immunorecruitment and allows dynamic and reversible monitoring of PPIs in living mammalian cells with a simple optical readout (Fig. 3 A; Herce et al., 2013). GFP fusions with proteins of interest (bait) were recruited to discrete subcellular structures such as artificial LacO DNA arrays by fusing GBP to the Lac repressor. In addition, naturally occurring major satellite repeats, the nuclear periphery, and cytoplasmic structures such as centrosomes can be used as anchor points. Enrichment of red fluorescent prey proteins at the respective anchor sites then indicates specific interaction. The nanobody-based F3H assay is also suited for drug discovery, as small molecule compounds or peptides can be identified that prevent or disrupt PPIs. With this F3H assay, e.g., the disruption of the p53–HDM2 interaction by potential cancer drugs such as Nutlin-3 was monitored in living cells, providing direct information on dose response, kinetics, and bioavailability (Herce et al., 2013).

**Nanoscopy.** In recent years, several novel super-resolution microscopy technologies revolutionized the field of fluorescence microscopy and enabled the analysis of cellular structures at subdiffraction resolution (Schermelleh et al., 2010). However, the higher resolution also imposes new requirements on detection reagents (Fig. 3 B). As a result of the size of primary and secondary antibodies, the attached fluorophore is positioned at a distance from the actual antigen, leading to so-called linkage errors. Thus, using the smallest possible immunofluorescence binding reagents is of utmost importance to unleash the full potential of super-resolution microscopy. In a first exemplary study, the GFP-specific nanobody was used for structured illumination microscopy to reveal the molecular machinery that effects the intercellular abscission during cell division (Guizetti et al., 2011). This gain in resolution was then systematically demonstrated arguing for the use of directly labeled nanobodies for advanced nanoscopy (Ries et al., 2012). Similarly, nanobodies have been used for photothermal single-molecule tracking in living cells with functionalized gold nanoparticles (Leduc et al., 2013).

**Target modulation and validation.** The molecular interaction between binder and antigen potentially alters or inhibits the biological function of the target structure (Fig. 3 A). In combination with intracellular expression, recombinant binders may thus be used for targeted modulation of antigen activity. The conceptual realization of such binder-mediated modulation was demonstrated by targeting the active site of the potato starch branching enzyme A in plant cells (Jobling et al., 2003). In a related approach, estrogen receptor–specific monobodies were used to discriminate ligand-induced conformational changes of estrogen receptor in yeast (Koide et al., 2002). Finally, nanobodies were used to modulate protein conformation and thereby either enhance or minimize fluorescence properties of FPs (Kirchhofer et al., 2010).

Novel recombinant binder formats have been developed as innovative biotherapeutics that bind and block defined disease-related antigens. Many of these potential target antigens, however, reside on the inside of cells, and thus, therapeutic applications depend on efficient techniques for delivery of proteins into affected cells. For research purposes, however, the ectopic expression of inhibitory binders in cells is a promising workaround for the functional validation of potential drug targets and does not require the physical target ablation by knock-out or knockdown. For such purposes, DARPin s were used to selectively inhibit JNK isoforms in human cells (Parizek et al., 2012), a class of enzymes that is critically involved in stress-induced signaling and is discussed as a potential drug target for various indications (Cui et al., 2007). Furthermore, a fibronectin-derived monobody was designed to block an intramolecular interaction of the fusion oncoprotein Bcr-Abl in primary chronic myeloid leukemia cells, revealing a novel interface for therapeutic intervention in chronic myeloid leukemia and demonstrating the general use of recombinant intracellular binders against elusive drug targets (Grebien et al., 2011; Sha et al., 2013).

**Toward synthetic biology.**

**Synthetic library design.** Immunoglobulin-based binders derived from natural postimmunization repertoires reliably deliver high-quality recombinant affinity reagents. However, the natural antibody modularity also allows the rational design of entirely synthetic antibody libraries using advanced molecular biology techniques. Over the last two decades, such synthetic antibody technologies have constantly evolved and are now arguably as good as or even better than natural libraries (Adams and Sidhu, 2014). Design strategies for such synthetic libraries are based on different concepts and include the recreation of natural CDR diversity or, as a next step, the rational implementation of well-known structure–function parameters to create recombinant antibody libraries with predefined, favorable biophysical properties regarding both antigen recognition and overall stability (Hoet et al., 2005; Tiller et al., 2013). Lacking a natural mechanism of diversity, nonimmunoglobulin binder libraries have to be synthetically generated and diversified with continuous improvements in design and performance (Seeger et al., 2013).

**Recombinant binders in synthetic biology.** Synthetic biology not only helps in the generation of recombinant
binders but also utilizes them to control gene transcription, protein turnover, or reroute signaling cascades (Lienert et al., 2014). Central claims of the still young field of synthetic biology involve the rational redesign of genetic building blocks to engineer novel cellular functions for a broad range of purposes and applications. In consequence, the conception of antibody derivatives and synthetic binders as cellular affinity modules that can be combined with any other gene adds to the molecular toolbox for synthetic biology. As a result of the widespread use of GFP fusion proteins in all areas of cell biology, the GFP-binding nanobody has often been used for initial proof-of-concept experiments that, however, demonstrate the potential of recombinant binders for future applications in synthetic biology. Thus, a nanobody was combined with the Drosophila melanogaster ubiquitin pathway and used for specific degradation of GFP fusion proteins in vivo (Caussinus et al., 2012). For this purpose, the WD40 domain of an F-box protein that naturally mediates the interaction with the protein to be degraded was replaced with GBP, allowing targeted proteasomal knockdown of GFP fusion proteins (Fig. 4A). In principle, this approach can be adapted with any binder-mediated specificity to subtly control target protein abundance and thus specifically modulate cellular functions in vivo.

Likewise, the F3H assay that was described in the biosensors section for monitoring PPIs in living cells is based on a synthetic binding device and demonstrates the concept of intracellular immunorecruitment. It can be used to target practically any biochemical activity at a defined site of action. This was shown with an engineered PloI kinase, in which the natural targeting sequence had been replaced by GBP, thereby enriching enzymatic activity at intracellular sites of GFP localization (Grallert et al., 2013). Moreover, this GBP-anchoring system can be combined with proteins recognizing particular DNA sequences within the genome to manipulate the localization of chromosomal segments and chromatin domains in living cells. With this approach, topological effects can be studied as rearrangement of chromosomal segments and chromatin domains in living cells.

Figure 4. **Recombinant binders as modular entities in conceptual, cell-based assay design.** (A) A targeted protein degradation via an engineered proteasomal degradation device. The natural WD40 domain that mediates the interaction with a substrate to be degraded via the Skp1-Cul1-F-box protein (SCF) ubiquitin (Ub) ligase complex is substituted with a recombinant binder, allowing targeted ablation of a protein of interest (POI; Caussinus et al., 2012). (B) A scaffold-induced system to manipulate gene activity. Two different binders, recognizing distinct epitopes of a scaffold protein of interest are fused to a DNA-binding protein (DBP) and a transcriptional activator (TA) enabling protein of interest-dependent gene activation (Tang et al., 2013). Some of the pioneering applications described in this review were initially demonstrated with only few recombinant binders. This was in part caused by the limited availability of well-characterized binders against a broad range of antigens. The extremely laborious procedures of library screening to generate specific and reliable binders still remain the major bottlenecks and represent a particular challenge to most cell biology laboratories. Thus, recent efforts in developing new binder formats and automatized screening, together with the establishment of publicly available binder resources (Taussig et al., 2007) to ideally generate binding proteins of defined specificity for specific targets. These studies illustrate the plethora of novel experimental designs enabled by intracellular binders. In general, the small size and intracellular stability of these recombinant binders allow the exchange of natural binding modules and thereby redirect enzymatic activities and redesign cellular structures and signaling pathways.

Conclusions and perspectives

Some of the pioneering applications described in this review are just a click away. Although conventional antibodies are still dominating in classic antibody realms, such as proteomics, Western blot, and immunofluorescence techniques, recombinant formats enable completely novel types of applications in structural biology, living cells, and synthetic biology. The possibility to genetically combine their target affinity with biological executor functions will be of interest not only for cell biology but also for future therapies. Here, the concomitant development of cellular delivery strategies will be crucial to enable direct uptake of functional binders in living cells. In addition, with gene therapy finally realizing its decades-old promises, recombinant binders will increasingly become important as shown for innovations such as chimeric antigen receptors to fight cancers (Kochenderfer et al., 2010).
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