Defining the epitope of a blood–brain barrier crossing single domain antibody specific for the type 1 insulin-like growth factor receptor

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Ligand-activated signaling through the type 1 insulin-like growth factor receptor (IGF1R) is implicated in many physiological processes ranging from normal human growth to cancer proliferation and metastasis. IGF1R has also emerged as a target for receptor-mediated transcytosis, a transport phenomenon that can be exploited to shuttle biotherapeutics across the blood–brain barrier (BBB). We employed differential hydrogen–deuterium exchange mass spectrometry (HDX-MS) and nuclear magnetic resonance (NMR) to characterize the interactions of the IGF1R ectodomain with a recently discovered BBB-crossing single-domain antibody (sdAb), VHH-IR5, in comparison with IGF-1 binding. HDX-MS confirmed that IGF-1 induced global conformational shifts in the L1/FnIII-1/-2 domains and α-CT helix of IGF1R. In contrast, the VHH-IR5 sdAb-mediated changes in conformational dynamics were limited to the α-CT helix and its immediate vicinity (L1 domain). High-resolution NMR spectroscopy titration data and linear peptide scanning demonstrated that VHH-IR5 has high-affinity binding interactions with a peptide sequence around the C-terminal region of the α-CT helix. Taken together, these results define a core linear epitope for VHH-IR5 within the α-CT helix, overlapping the IGF-1 binding site, and suggest a potential role for the α-CT helix in sdAb-mediated transcytosis.

Abbreviations

HDX  Hydrogen–deuterium exchange
HDX-MS  Hydrogen–deuterium exchange mass spectrometry
NMR  Nuclear magnetic resonance
HSQC  Heteronuclear single quantum coherence
HMQC  Heteronuclear multiple quantum coherence
SOFAST-HMQC  Band-selective optimized flip-angle short transient HMQC
HNCA, HN(CO)CA, CBCA(CO)NH, HN(CA)CO, and HNCO  A set of heteronuclear 3D NMR experiments for backbone resonance assignments of 15N- and 13C-labelled proteins
NUS  Non-uniform (sparse) sampling
SPR  Surface plasmon resonance
RU  Resonance unit
PDB  Protein data bank
SD  Standard deviation
CDR  Complementarity-determining region

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Human insulin like-growth factor receptor (IGF1R) is a member of a small family of transmembrane receptor tyrosine kinases (RTKs) that also includes the insulin receptor (IR) and a third member, the orphan insulin receptor-related receptor (IRR). Aside from IR, IGF1R is a secondary cell-signaling RTK involved in the regulation of glucose uptake and energy metabolism. IGF1R itself has important functions in fetal and prenatal growth and in cancer proliferation, by promoting the immortalization of transformed cells through binding to two insulin-like growth factors, IGF-1 and IGF-2. Implication of IGF1R in cancer has prompted the development of anti-cancer strategies using neutralizing antibodies and small-molecule inhibitors of the RTK. Both IR and, more recently, IGF1R have emerged as therapeutic targets for brain-specific neurological disorders, especially for drug-delivery across the blood–brain barrier (BBB).

Many diseases of the central nervous system (CNS) have limited treatment options because of the BBB, whose relative impermeability severely impedes the physical delivery of blood-borne drugs to the CNS. The BBB represents a particular hurdle to CNS delivery of antibody-based therapeutics and efficient crossing of large macromolecules must engage endocytotic pathways through a process called receptor-mediated transcytosis (RMT). To date, transferrin receptor (TfR), low-density lipoprotein (LDL), and insulin receptor (IR) have been explored as RMT targets, with IR-targeted therapies in advanced clinical trials. However, targeting these common receptors has substantial development risks related to species-dependent expression in associated with interfering with their normal physiological functions. Our recent work has focused on IGF1R, based on its elevated expression in brain endothelial cells relative to peripheral tissue, and on its established role in the delivery of one of its endogenous ligands, IGF-1, across the BBB.

IGF1R is a heavily glycosylated transmembrane protein with a complex dimeric (αβ) architecture and with the two-chain (α and β) ectodomains held together by tertiary structural interactions along with inter-chain and intra-chain disulfide bonds. The modular architecture of IGF1R includes a leucine-rich region (L1), a cysteine-rich region (CR), the second leucine-rich region (L2) and the N-terminal part of a type-III fibronectin-like domain FnIII-2 in the α-chain and the C-terminal part of FnIII-2 leading to two other fibronectin-like domains FnIII-1/3 in the β-chain of the ectodomain. The intact FnIII-2 domain in the IGF1R precursor also contains a long insertion domain (ID) connecting the α- and β-chains generated by proteolytic cleavage at a site within ID. Extensive structural studies of the IGF1R protein family have revealed a novel mechanism of receptor auto-inhibition and ligand-triggered receptor activation. In such a mechanism, IGF-1 appears to engage only one L1-CR-L2 segment of the dimeric IGF1R along with one of the all-important α-CT helices exposed at the extreme C-termini of the α-chains (hence α-CTs) and to release an auto-inhibiting structure of the ectodomain leading to spontaneous receptor auto-phosphorylation. A significant structural feature of the IGF1R/IGF-1 interaction is the malleability of the receptor architecture and the disparate roles played by the two α-CT helices in IGF1R, one as part of the binding site for IGF-1 and the second remaining apparently free with partial contacts with the L1’ and the FnIII-2 regions of the dimeric receptor. The differing environment of the two α-CT helices elicited by IGF-1 binding to IGF1R has been identified as the structural origin of negative cooperativity observed for the IGF-1/IGF1R ligand-receptor system.

We recently discovered a unique class of single-domain antibodies (sdAbs) specific for the IGF1R ectodomain, among which the single-domain antibody IGF1R-5 (referred here to as VHH-IR5), does not
appear to interfere with normal kinase functions of IGF1R and its interactions with the endogenous ligand\(^{21}\). As such, VHH-IR5 not only has the potential to act as a BBB-crossing shuttle, but may help decipher the fundamental mechanisms of cell signaling through IGF1R and receptor transcytosis. In this work, we focus on elucidating the mode of action of VHH-IR5, with a specific goal of defining its binding epitope on IGF1R in relation to the natural IGF1R ligand IGF-1. Results from differential hydrogen–deuterium exchange mass spectrometry (HDX-MS) and nuclear magnetic resonance (NMR) spectroscopy, and binding studies with IGF1R fragments outline an apparently shared binding site between the sdAb VHH-IR5 and the natural ligand IGF-1 convergent on the α-CT helix motif, an epitope on IGF1R not targeted, to the best of our knowledge, by other antibodies reported to date\(^{28–31}\).

Results

Section I: comparison of IGF-1 and VHH-IR5 binding and functional activation of IGF1R. Previous work has suggested that VHH-IR5, an IGF1R-binding sdAb (of the llama origin\(^{27}\)) with BBB crossing activity, does not interfere with the interaction between IGF1R and its native ligand, IGF-1\(^{21,27}\). Figure 1a shows that saturation of surface-immobilized IGF1R ectodomain (eIGF1R) with VHH-IR5 does not preclude the subsequent binding of IGF-1. Reversing the order of pre-loading produced the same result (Fig. 1b). Qualitatively, pre-loading of either VHH-IR5 (Fig. 1a) or IGF-1 (Fig. 1b) also does not alter the shape of SPR sensograms of the other ligand (Fig. S2a,b). Thus, occupancy of eIGF1R by VHH-IR5 (or by IGF-1) has little effect on IGF-1 (or VHH-IR5) binding, outlining apparently independent and non-competitive binding sites on eIGF1R for these two ligands. Quantitatively, VHH-IR5 has a binding affinity of \(K_D = 0.6\ \text{nM}\) for eIGF1R along with a slow rate of dissociation, \(k_{off} = 3.4 \times 10^{-4}\ \text{s}^{-1}\), as seen by the characteristic plateau for the dissociation phase of the VHH-IR5 sdAb-eIGF1R complex (Fig. S2a). We measured an affinity of \(K_D = 8.2\ \text{nM}\), and a faster dissociation rate constant (\(8.2 \times 10^{-3}\ \text{s}^{-1}\)) for IGF-1 binding to the eIGF1R ectodomain (Fig. S2b), in agreement with previous reported values (\(K_D \approx 10–20\ \text{nM}\))\(^{28}\). VHH-IR5, therefore, has a higher affinity for eIGF1R as compared to IGF-1. In addition, VHH-IR5 is also specific for IGF1R as it does not bind the insulin receptor (Fig. S2c) in contrast to IGF-1 which interact with both (Fig. S2d).

Next, we demonstrated in a cell-based assay that VHH-IR5 binding does not induce significant activation of IGF1R, in sharp contrast to the effects of IGF-1 (Fig. 1c). Very importantly, increasing concentrations of VHH-IR5 had little further effects on the activating level of IGF-1, even when present up to >500-fold molar excess (Fig. 1d), demonstrating essentially mutual independence of these two ligands. This finding is in agreement with previous studies that showed VHH-IR5 did not induce phosphorylation of the downstream kinase AKT\(^{21}\), which is the expected outcome of IGF1R activation by the endogenous ligand IGF-1\(^{21,32}\). Taken together, both biophysical binding (SPR) and cell-based assays indicate that VHH-IR5 and IGF-1 likely have independent binding sites on IGF1R and that VHH-IR5 binding is unlikely to alter the ligand-sensing capabilities of IGF1R on the cell-surface.

Section II: comparison of IGF-1 and VHH-IR5 binding to IGF1R using HDX-MS. We next assessed the global structural consequences of IGF-1 and VHH-IR5 binding to the soluble eIGF1R using differential bottom-up HDX-MS profiles\(^{33}\). This method is extremely sensitive to conformational changes in proteins and has been employed for mapping antibody-epitope interactions\(^{34,35}\). The HDX-MS experiments were performed using the fully-glycosylated form of eIGF1R to preserve native eIGF1R conformations, similar to a strategy utilized for the insulin receptor (IR)\(^{46}\). Overall, 47% sequence coverage was achieved across 134 peptides (Table 1 and Table S1). Despite the resulting gaps in overall sequence coverage (Fig. S4), HDX-MS data still afforded key insights into the eIGF1R-binding characteristics of both protein ligands (Fig. 2). The HDX responses to the native ligand, IGF-1 (Fig. 2a) are similar to what has been reported previously, validating our experimental approach, for example, conformational stabilization was observed when binding were observed at the contact site between the L domain and IGF-1 (residues I9-L16 and L57-L63 specifically), and at the dimer interface between the two L domains (L2/L2’, Y417-L424) and the two FnIII-1 domains (FnIII-1/FnIII-1’, R488-F493) (Fig. 2a, blue regions). There was increased deuteration or conformational destabilization for the L2 residues C323-L331, which may be related to the hinging motion of L1-CR away from the L2 domain observed in the uncomplexed, IGF-1-bound and VHH-IR5-bound states of eIGF1R. Inspection of the unlabelled isotopic profile (Fig. S5b) shows that the observed bimodality is not a result of overlapped profiles from unrelated peptides. However, further comparison of the exchange profiles between the three states was not possible due to inherent variability of the deuteration in this region (Fig. S5a).

Interestingly, many of the dramatic changes in deuteration elicited by IGF-1 were almost entirely absent in response to VHH-IR5 binding (Fig. 2b), suggesting that VHH-IR5 does not induce the same global structural changes in IGF1R. However, both IGF-1 and VHH-IR5 appear to engage the α-CT structural motif located at the C-terminal end of the eIGF1R α-chain, although key differences were observed in the deuteration pattern of the α-CT helix (Fig. 3a) and its immediate surroundings. First, the HDX-MS profile for the IGF-1/eIGF1R interaction (Fig. 3a) closely tracks engagement of the α-CT helix and the associated conformational malleability\(^{38}\). Specifically, an increase in deuteration was observed with residues Y688-E693, which suggests that this sequence segment is conformationally more dynamic relative to apo-eIGF1R, despite elongation of the α-CT helix in the IGF-1 bound state\(^{39}\). This is followed by progressively stronger reduction in deuteration as peptide coverage shifts towards the C-terminus of the α-CT motif, implying the strongest structural stabilization for residues N694–F701.
Section III: epitope mapping of VHH-IR5 using IGF1R insertion domain (ID) fragments containing the α-CT helix.

In order to further define the role of the α-CT helix motif in ligand engagement, we employed NMR to examine the binding behaviour of VHH-IR5 with IGF1R fragments derived from the insertion domain (Table 2). Two fragments, IGF1R674-742 and IGF1R689-742, in the form, respectively, of Ubi-ID and Ubi-s-ID fusion proteins (Table 2) retain the covalent junction between the extreme C-terminus of the α-chain and the N-terminus of the β-chain in IGF1R that is present in pro-IGF1R. Interestingly, both Ubi-ID and Ubi-s-ID, the latter lacking the N-terminal part of the long α-CT helix, have specific binding with VHH-IR5, as shown by localized perturbations in the 1H-15N SOFAST-HMQC map of 15N-labelled Ubi-ID and Ubi-s-ID by

Table 1. Summary of HDX-MS experimental parameters3. *All HDX-MS experiments were carried out with recombinant and SEC-purified eIGF1R (Fig. S3).

| Data set | Control | IGF-1 | VHH-IR5 |
|----------|---------|-------|---------|
| HDX reaction details | PBS, pH = 7.0, 25 °C | | |
| HDX time course (min) | 1, 3, 10, 60 | | |
| # of Peptides found across all samples | 133 | | |
| Sequence coverage | 47% | | |
| Average peptide length/redundancy | 11.7/1.7 | | |
| Replicates (biological or technical) | 3 (technical) | | |
| Repeatability (average standard deviation, %) | 1.6 1.5 1.4 | | |
| Significant differences in HDX | Two-state student T-Test performed at each time point (> 2 SD, p value 0.05) | | |
Figure 2. A global survey of HDX-MS profiles of eGF1R in response to IGF-1 and VHH-IR5 binding. (a) IGF-1 response profile projected on a structural model of the 1:1 eGF1R/IGF-1 complex (PDB: 6PYH). The (αβ)′ monomer, except for α-CT′, is rendered as a transparent surface, while the (αβ) monomer and α-CT′ are shown as a ribbon cartoon. The schematic in the inset shows the domain organization of IGF1R where the (αβ) monomer is colored in green, and (αβ)′ in black. (b) VHH-IR5 response profile projected on the same structural model as in (a) (PDB: 6PYH), except that the structure is rotated counterclockwise by 90°. Significant structural destabilizations are shown in red, stabilizations in blue, lack of significant changes in grey, and missing sequence coverage in black. Residues are colored based on differences in deuterium at a single time point (± 2 SD, p = 0.02). In both (a) and (b), IGF-1 molecules are shown as magenta spheres. Of note, residues of the rhesus eGF1R used for HDX-MS data collection were mapped by BLAST onto the mouse eGF1R before rendering the HDX-MS results onto the 3D structure of mouse eGF1R (PDB: 6PYH).

Figure 3. Detailed analysis of the HDX-MS profiles in the IGF1R α-CT region. (a) HDX profile of the α-CT′ bound to IGF-1 (magenta) (based on PDB: 6PYH). Residues 684-706 of mouse IGF1R α-CT′ (residues 683-705 in rhesus/human eGF1R) are shown as a ribbon cartoon. Significant structure destabilization is shown in red and stabilization in blue, while lack of significant changes in grey and missing coverage in black. (b–e) HDX-MS kinetics of peptides covering the α-CT helix. Data collected in quadruplicate, and error bars represent 1× SD. Deuteration was normalized to theoretical maximum uptake of 45%. Free eGF1R is shown as black circles, eGF1R/IGF-1 as blue diamonds, and eGF1R/VHH-IR5 as red triangles.
unlabelled VHH-IR5 (Fig. S6). Ubiquitin moieties present at the N-terminus of Ubi-ID and Ubi-s-ID served as an internal control for the interpretation of NMR signal perturbations. The ubiquitin protein is not expected to bind VHH-IR5 and indeed the ubiquitin moiety in the fusion protein showed minimal HMQC signal perturbations (Figs. 4 and S6).

The observed NMR HMQC perturbations for the longer ID fragment were localized in the N-terminal region, i.e., residues T675-K683, A686 and E687 (Fig. 4); in other words, further upstream to the α-CT helix characterized by HDX-MS (Fig. 3). Surprisingly, the HSQC map (as compared to SOFAST-HMQC,vida infra) of Ubi-ID appears to lack assignable signals for the entire C-terminal region of the α-CT helix, i.e., residues R689-V702 and E706-E709, including residues E706-R709 within an extended segment after the α-CT helix (Fig. S7). These signals are missing at both near-neutral pH 6.8 and pH 5.5 (Fig. S7), despite that this latter lower pH condition normally favors optimal NMR signal intensities. The more sensitive proton-15N SOFAST-HMQC experiment revealed a few additional crosspeaks of very low intensities (Fig. S8a,b), suggesting that the largely invisible region of Ubi-ID (Figs. 4 and S7) must be a consequence of severe line broadening of the NH and/or 15N NMR signals of the IGF1R689-709 segment. As well, the SOFAST-HMQC spectrum of Ubi-ID became sharper at an elevated temperature (308 vs. 298 K, Fig. S8c), especially in the spectral region where new SOFAST-HMQC signals (i.e., the observable α-CT residues) are located (Fig. S6). These same SOFAST-HMQC signals also experience more significant changes in NH/15N chemical shifts upon temperature elevation (Fig. S8c). Taken together, these NMR lineshape behaviors strongly implicate the α-CT residues as being involved in conformational dynamics such as frame shifts of the α-CT helix (Fig. 3) or other yet-to-be-clarified structural transitions.

Intriguingly, as many as 10 new SOFAST-HMQC peaks appeared from within the IGF1R689-742 fragment of the shorter Ubi-s-ID construct, which can be traced to the sequence segment without significant HMQC signals in Ubi-ID (Fig. S6). All these reappeared SOFAST-HMQC signals exhibited specific perturbations in the presence of VHH-IR5. The binding interactions were then examined in complementary experiments using 15N-labelled VHH-IR5 titrated with unlabelled Ubi-ID and Ubi-s-ID. Figure 5a and b show that 15N-labelled VHH-IR5 responds specifically to binding of Ubi-ID and Ubi-s-ID, in a similar fashion as the 15N-labelled Ubi-ID and Ubi-s-ID responding to unlabelled VHH-IR5 (Fig. S6). Despite lacking the N-terminal residues K674-Y688 of the long α-CT helix, binding of the Ubi-s-ID construct perturbs essentially the same HMQC signals of VHH-IR5 as binding to Ubi-ID (Fig. 5a,b). Very importantly, the same set of perturbed residues in VHH-IR5 include some of CDR2, e.g. T51 and I52 and many in the framework region, e.g. V13, L21, A25, E47, F48, Q83, M84, A93 and V121 (Fig. 5a,b), based on the NMR assignments of free VHH-IR5 (Fig. S9).

To further explore the interactions of VHH-IR5 with the α-CT helix, a short peptide derived from the ID region of IGF1R similar to the fragment VFENFLHNSIFVPR (α-CT697-706) was produced (Pep5, Table 2). Previously, this portion of the α-CT helix was utilized for reconstituting a hybrid IR-IGF1R micro-receptor. More importantly, Pep5 contains all of the amino acids in the α-CT region of IGF1R that were seen to have modified HDX-MS patterns upon VHH-IR5 and/or IGF-1 binding. The synthetic Pep5 (unlabelled) was used here to titrate into 15N-labelled VHH-IR5 for selective observations of responses in 15N-labelled VHH-IR5. Instead of broadening and the resulting disappearance of many HSQC signals of free VHH-IR5, Pep5 binding to VHH-IR5 led to the appearance of new HSQC signals (i.e., those from the Pep5-sdAb complex) accompanying the disappearance of the HSQC signals of the free VHH-IR5 (Fig. 5c). Such widespread differences between free and ligand-bound proteins are characteristic of slow off-rates and indicative of high-affinity binding interactions in

### Table 2. Polypeptide and protein constructs used in NMR and related binding studies.

| Polypeptide/Protein Sequence | Sample Name |
|-----------------------------|-------------|
| (Ubi)sgsQggKTEAEQK AEREEAYRK VFENFLHNSI FVPRPERKRR DVMQVANTTM | Ubi-ID* |
| SSSRNTTAA DTYNITDPEE Lehhhhhh | |
| (Ubi)sgsQggRK VFENFLHNSI FVPRPERKRR DVMQVANTTM | Ubi-s-ID |
| SSSRNTTAA DTYNITDPEE Lehhhhhh |
| MQVKLEESGG GLVQAGGSGLR LSCAASGRTI DNYAMAWSQ APGKDREFVA | Pep5 |
| TIDWGGDGAR YANGSQRRT ISRQNAKSTM YLQHNSLEEPE DZAVYSVAMCA |
| RQSrvNLDVA FYDYWGQSSQ VTYsgsQgklisedImhhhhhh |
| RK VFENFLHNSI FVPR |
| MQIFVKTLTG-KTITLEVEPS-DTIENVKAKIQDKEGIPPPDQ-QRLIFAGKQL-EDGRTLSDYN-IQKESTLHLV-LRLRGG | VHH-IR5 |
| lnhhhhhh |

* (Ubi) denotes the ubiquitin moiety, which has an amino-acid sequence of MQIFVKTLTG-KTITLEVEPS-DTIENVKAKIQDKEGIPPPDQ-QRLIFAGKQL-EDGRTLSDYN-IQKESTLHLV-LRLRGG. Ubi-ID is, therefore, a 157-residue fusion protein and Ubi-s-ID, a 142-residue fusion protein. For Ubi-ID, Ubi-s-ID and Pep5, upper-case letters designate the amino-acid sequence of the uncleaved single-chain human IGF1R precursor1. In lower-case letters are the linker sequence, sgsgsg (for Ubi-ID and Ubi-s-ID), and the c-Myc tag sgseqklised (in VHH-IR5) and a His6 tag added for protein expression and purification. All expressed proteins contain an artificial Met residue at the N-terminus added by the E. Coli host during protein expression.
A smaller peptide fragment of the IGF1R ID such as Pep5 (Table 2), or the related α-CT 691-706 peptide, very likely has a high binding affinity to VHH-IR5, suggesting that it may represent the linear epitope of this sdAb. To confirm our results with Pep5, the epitope of VHH-IR5 was further examined using a library of 15-residue peptides that covers the same 674-742 region of IGF1R contained in Ubi-ID (Table 2). Interestingly, the first peptide KTEAEKQAEEKAHY showed no binding at all to VHH-IR5 (Fig. 6 and Table S2) despite containing all the residues of Ubi-ID with large HSQC signal perturbations (Fig. 4). Binding interactions only become observable for peptides KVFENFLHNSIFVPR, VFENFLHNSIFVPRP, and FENFLHNSIFVPRPE covering IGF1R residues K690-E706, which mirror the strong NMR signal perturbation for residue R704 (Fig. 4). There was essentially no difference between Ubi-ID and Ubi-s-ID for their binding to 15N-labelled VHH-IR5 (Fig. 5) since both carry the same sequence K690-E706 of IGF1R (Table 2). NMR signal perturbations in the N-terminal region of the α-CT helix therefore reflects conformational changes induced by VHH-IR5 binding to the epitope segment as represented by Pep5, which, for yet unknown mechanisms, has severely broadened NMR signals as part of the longer ID fragments in Ubi-ID (Fig. S8). This latter experimental NMR observations may be related to HDX-MS results (Fig. 3), outlining significant structural stabilizations of this important region of IGF1R in response to VHH-IR5 binding.

Discussion
In the search for sdAbs targeting IGF1R, VHH-IR5 has demonstrated the characteristics required for drug delivery across the BBB: efficient BBB transmigration, no detectable impact on the functional activation of IGF1R and no interference with IGF1R/IGF-1 interaction (Fig. 1). The uniqueness of this BBB-crossing sdAb is further highlighted by the nature of the binding epitope of VHH-IR5, revealed here using an array of orthogonal and complementary methodologies. In summary, HDX-MS and NMR results converged on an epitope sequence consisting of residues FENFLHNSIFVPR located near the extreme C-terminus of the IGF1R α-chain (the α-CT). This region of the IGF1R ectodomain demonstrated significant decreases in deuteration in HDX-MS profiles (Fig. 3), equating a structural/conformational stabilization upon binding either driven by allostery or as a direct consequence of binding contacts or both. Of particular interest is the finding that IGF1R fragments containing FENFLHNSIFVPR all retain binding to VHH-IR5 as shown by use of NMR spectroscopy (Figs. 4 and 5). The nature of the VHH-IR5 epitope is further highlighted by two shorter fragments of IGF1R having increased binding affinities for VHH-IR5: RKVFENFLHNSIFVPR (Pep5, Table 2 and Fig. 5) and FENFLHNSIFVPRPE (Fig. 6). These peptide segments terminate, 6 and 4 residues, respectively, before the α-chain C-terminus of IGF1R.
Figure 5. Comparative responses of VHH-IR5 to IGF1R ID fragments. HSQC spectra of $^{15}$N-labelled VHH-IR5 identify effects of interactions with three unlabelled IGF1R ID fragments. In black are the HSQC spectra of free VHH-IR5 collected immediately before additions of Ubi-ID, Ubi-s-ID and Pep5 while superimposed in red are the HSQC spectra of the sdAb complexes at ~1:1 molar ratio. (a) spectral comparisons showing the effects of Ubi-ID binding. (b) spectra showing the effects of Ubi-s-ID binding. Note that Ubi-ID (a) and Ubi-s-ID (b) binding induce almost the same perturbations, i.e. disappearances of many HSQC signals of $^{15}$N-labelled VHH-IR5, especially those of residues T51 and I52 at the beginning of the CDR2 loop and many other residues in the framework region. All perturbed residues were labelled using the resonance assignments of $^{15}$N-labelled VHH-IR5 (Fig. S5). (c) Widespread spectral displacements of $^{15}$N-labelled VHH-IR5 were produced by binding of the peptide fragment Pep5.
IGF1R (Table 2), which is exposed by enzymatic cleavage during IGF1R maturation (Fig. S6)\(^1\). Taken together, the sdAb VHH-IR5 has a prototypical linear epitope\(^4\), which is available on the surface of the IGF1R structure, allowing binding of VHH-IR5 to native IGF1R on the cell surface.

At first glance, the identified VHH-IR5 epitope directly overlaps with part of the IGF-1 binding site, residues F692-N694, L696-N698, and I700-F701 identified previously\(^3\),\(^7\),\(^8\),\(^4\) and seen here again using HDX-MS (Fig. 3). VHH-IR5 binding therefore necessitates availability of this sequence motif in both apo-IGF1R and in the IGF-1/IGF1R complex since these two ligands do not appear to interfere with each other for IGF1R binding (Fig. 1). As a disulfide-linked homodimer, IGF1R contains two potential binding sites and IGF-1 has been shown to engage only one of these sites on full-length IGF1R at biologically relevant concentrations, as a consequence of negative cooperativity\(^2\),\(^5\),\(^6\). Upon binding to IGF1R, IGF-1, along with the L1 domain of one monomer, engages the α-CT' helix of the other monomer, and induces a dramatic structural rearrangement along with the elongation of the helical structure of the free α-CT toward the N-terminal region (residues 670-681)\(^2\). For non-competitive binding to be possible, the second α-CT site must remain competent for binding VHH-IR5 and vice versa for IGF-1. Alternatively, both binding sites can be equally available for independent ligation of IGF-1 and VHH-IR5 as shown by SPR evidence of simultaneous IGF-1 and VHH-IR5 interaction with eIGF1R (Fig. 1a). Interestingly, despite dramatic structural rearrangement, the unligated α-CT of the 1:1 IGF-1/IGF1R complex indeed maintains a surface-exposed configuration\(^2\), leaving this second α-CT available for VHH-IR5 complexation as we observed here for the IGF1R ectodomain (Fig. 1).

The precise mechanism for how VHH-IR5 interacts with the rest of IGF1R without activating downstream signalling events remains largely unknown. However, there are clearly identifiable differences between structural changes of the IGF1R ectodomain in response to IGF-1 and VHH-IR5 binding. The HDX-MS responses in FnIII-1, -2, and L2 of eIGF1R to IGF-1 binding closely mirror the recently-determined symmetric conformation of the IGF1R ectodomain in complex with two IGF-1 molecules\(^3\),\(^7\),\(^8\). Such responses are notably absent upon binding to VHH-IR5. There is, instead, a decrease in deuteration at the L1 domain of the eIGF1R-sdAb complex (Fig. 2b), which suggests structural relaxation of this L1 region subsequent to VHH-IR5 complexation with the α-CT. The same L1 region interacts with the α-CT helix in an apo-form of eIGF1R\(^3\). In the case of VHH-IR5, its main binding effect is in stabilizing the entire helix and limiting its conformational sampling. Taken together, while the IGF-1 binding elicits more global conformational rearrangements in eIGF1R, VHH-IR5 appears to exhibit a localized binding response through a predominantly linear epitope at the α-CT. VHH-IR5 binding may also be negatively cooperative in the sense that it can only bind a single α-CT site on IGF1R. Since VHH-IR5 alone does not activate IGF1R, VHH-IR5 must induce conformational changes in IGF1R that are different from the large structural shifts induced by IGF-1, such that associations of the IGF1R transmembrane domains and the subsequent receptor auto-phosphorylation cannot take place. Such a distinct mode of receptor engagement by VHH-IR5 may underline the structural basis of the transcytosis capacity of this unique sdAb without a dramatic impact on the normal functions of IGF1R. Future structural investigations are needed, particularly with more complete HDX-MS sequence coverages, for a detailed understanding of the novel sdAb-IGF1R interaction.

**Limitations.** This work is focused on assigning the epitope of an IGF1R-specific sdAb VHH-IR5 and comparing the molecular interactions with those observed for the natural ligand IGF-1. Consequently, the ectodomain of IGF1R was selected for the binding studies and for the comparison. There are therefore limitations in the data that prevent detailed assessments of the global architecture of the sdAb-IGF1R complex. Indeed, our HDX-MS data accurately reflect structural shifts in the bound α-CT motif reported by Li et al. for a full-length IGF1R\(^2\). Our HDX-MS findings suggest a limited degree of asymmetry upon IGF-1 binding to the IGF1R ectodomain, which is not in conflict with asymmetric and simultaneous binding of VHH-IR5 and IGF1 in the...
full length receptor. It is also plausible that the low deuterium content may mask the underlying asymmetry in the HDX profiles. We are unable to explore this question in greater depth within the scope of this current study.

**Methods**

**Surface plasmon resonance (SPR).** The binding of IGF-1 and VHH-IR5 to immobilized IGF1R ectodomain (eIGF1R) was determined using BIACORE 3000 (GE Healthcare). Approximately 3000 Resonance Units (RU) of recombining human eIGF1R (R&D Systems, Cat# 391-GR-050) were immobilized on a sensor chip CM5 after the quality of the eIGF1R was confirmed by size-exclusion chromatography (SEC, see Fig. S1a). Immobilization was carried out at 10 μg/mL in 10 mM acetate at pH 4.0 using the amine coupling kit (GE Healthcare). The remaining reactive sites were blocked with 1 M ethanolamine at pH 8.5. An ethanolamine blocked surface was used as a reference. Binding studies were carried out at 25 °C in 10 mM HEPES, pH 7.4 containing 150 mM NaCl, 3 mM EDTA and 0.005% surfactant P20 (Polyoxyethylensorbitan, GE Healthcare). Various concentrations of SEC-purified IGF-1 and VHH-IR5 (Fig. S1b,c) were flowed over the immobilized eIGF1R and reference surfaces at 20 μL/min. The concentrations employed for VHH-IR5 were 0.75, 1, 2.5, 5, 5.75 and 10 nM, while for IGF-1 binding, they were 10, 25, 50, 100 and 250 nM. The SPR co-injections (satisfaction of the surface with injection of analyte 1, followed by injection of a mixture of analyte 1 + analyte 2) utilized VHH-IR5 at 10 nM and IGF-1 at 250 nM. Surfaces were regenerated, after binding of VHH-IR5 with 10 mM glycine at pH 2.0 with a contact time of 24 s. No regeneration was used for IGF-1 binding experiments. Data were analysed with BLAevaluation 4.1 (GE Healthcare). The data was fit to a 1:1 binding model without further attempts at quantifying the binding stoichiometry (Text S1).

**Cell-based IGF1R functional assay.** The receptor functional assay was carried out using the PathHunter eXpress IGF1R kit (DiscoverX), which uses engineered HEK293 cells expressing a (Pro-Link or PK-) tagged IGF-1 Receptor and an Enzyme Acceptor (EA-) tagged SH2 domain. Upon receptor activation, EA-SH2 binds to the phosphorylated PK-IGF1R and reconstitutes an active β-galactosidase enzyme which hydrolyzes a substrate to generate a chemiluminescent readout. Cells were thawed and plated in a 384-well white wall clear bottom plate (Greiner, North Carolina, USA) at 20,000 cells/well in the cell plating 17 reagent media. After a 24-h incubation at 37 °C in an atmosphere with 5% CO2, cells were treated with IGF-1 (R&D, Minneapolis, USA) (with concentrations varying from 0.03—2000 nM), VHH-IR5 (0.03—6000 nM) or 3 nM IGF-1 + VHH-IR5 (0.03–6000 uM) for 180 min at RT. The chemiluminescent substrate was added and cells were further incubated at RT for 60 min. The resulting luminescence was measured using the CLARIOstar plate reader (BMG LABTECH, Ortenberg, Germany).

**Expression and purification of the IGF1R ectodomain.** Codon-optimized cDNAs (Genscript) for the rhesus IGF1R ectodomain (eIGF1R, residues 1-902) with a C-terminal His-tag was subcloned into pTTTM109 expression vector. A pool of CHO cells (CHOBRi55E1-JN, proprietary to The National Research Council of Canada) was initiated and the CHOBRi TM55E1 pool was selected with methionine sulfoximine for approximately two weeks. A fed batch production was then performed in optiFlasks. The eIGF1R protein was purified by immobilized metal ion affinity chromatography followed by size-exclusion chromatography in 1× PBS (HiPrep 26/60Superdex 200) (Fig. S3a). Eluting fractions were monitored by SDS-PAGE (Fig. S3b), pooled, concentrated to 10.17 mg/mL and stored at − 80 °C.

**Peptides and expression of isotope-labelled proteins for NMR Spectroscopy.** Fusion proteins of IGF1R ID fragments, Ubi-ID and Ubi-s-ID, were constructed by adding the sequence of human ubiquitin (Ubi) to the N-terminus of the two IGF1R fragments (Table 2). The DNA sequences were codon-optimized for E. coli, synthesized and inserted into the pD441-SR expression vector (Atum). The plasmids were electroporated into E. coli BL21ompT lon hsdSB,rB-,mB- (K-12(λS)) and selected with methionine sulfoximine for approximately 20 min. The clarified lysate was mixed with 3 mL Ni–NTA resin re-suspended in ~ 150 mL lysis buffer (50 mM sodium phosphate, 300 mM NaCl, pH 8), sonicated on ice for 2 min, and clarified at 10,000 rpm at 4 °C for 30 min. The clarified lysate was mixed with 3 mL Ni-NTA resin by gently vortexing, washed with 100 mL lysis buffer + 20 mM imidazole, and eluted with lysis buffer + 250 mM imidazole. Eluted protein was dialyzed into 50 mM sodium phosphate, 50 mM NaCl at either pH 6.8 or pH 5.5. The peptide Pep5 (Table 2) was synthesized chemically using the solid-phase method (CanPeptide Inc., Pointe-Claire, Quebec, Canada) and purified by use of HPLC. Identities of the purified peptide Pep5 and Ubi-fusion proteins were confirmed by mass spectrometry: Pep5 — theoretical MW of 3033.2; Ubi-ID—theoretical MW of 3482.2; Ubi-s-ID—theoretical MW of 3506.2.

**Bottom up HDX-MS.** HDX-MS was performed by first mixing 20 μM eIGF1R with 25 μM ligand (IGF-1 or VHH-IR5) in PBS at a 1:1 volume ratio. Labelling was initiated by adding 3 uL of 90% D2O (0.1× PBS, pH 6.9)
to 3 μL of an IGF1R-complex at 19 °C. The labelling reaction was quenched by the addition of a 54 μL ice-cold solution of 250 mM TCEP, 2 M Guanidine, 100 mM glycine–HCl at pH 2.0. Quenched and deuterium-labelled samples were immediately injected into a 20 μL sample loop (at 20 pmol eIGF1R/25 pmol ligand). Samples were non-specifically digested online with a Poroszyme Immobilized Pepsin Cartridge (21 × 30 mm, Thermo Scientific) at room temperature for 1.5 min at 75 μL/min, and desalted at 400 μL/min with a C18 PepMap 100 trap at 1 °C (1 × 5 mm, 5 μm, Thermo Scientific) in mobile phase A (0.23% Formic Acid in water, pH 2.55). Peptides were eluted from a BioBasic-18 (50 × 0.32 mm, 5 μm, Thermo Scientific) column with a separation gradient spanning 10–35% mobile phase B (0.23% Formic Acid in Acetonitrile) over 20 min. Samples were injected into Q-TOF Premier mass spectrometer (Waters) in quadruplicate technical replicates, and deuteration assignment and statistical analysis of individual timepoints (2 × SD, p = 0.02) were achieved using MS Studio45. Unlabelled eIGF1R was injected and analyzed by data-dependent acquisition, and peptides were identified with Mascot. Further experimental details can be found in Table 1. Sequence alignment of the rhesus eIGF1R used for HDX-MS on to the mouse eIGF1R was performed with BLAST46 prior to displaying the HDX-MS data on the 3D structure (PDB: 6PYH) of mouse eIGF1R (Figs. 2 and 3). The sequences of rhesus and human eIGF1R are highly homologous and completely in frame with each other.

**NMR spectroscopy.** All NMR experiments were carried out at a sample temperature of 298 K on Bruker Avance-III 600 MHz and 800 MHz NMR spectrometers equipped with a 5 mm PATXI sample probe and Z-field gradient accessories. Assignments of the 1H-15N correlations (i.e. HSQC or HMQC) spectra for Ubi-ID (Table 2) were achieved using 3D HNCA, HN(CO)CA, CBCA(CO)NH, HNCACB, HN(CA)CO, and HNCO spectra47 collected at 600 MHz. The NMR sample was prepared from a 0.5 mM uniformly 15N-/13C-labeled Ubi-ID dissolved in a buffer that was 50 mM in sodium phosphate and 50 mM in NaCl at pH 5.5 followed by the addition of 10% D2O. The SOFAST-HMQC48 pulse sequence with 0.3 s recycle delay was used to record 2D 1H-15N correlation spectra of Ubi-ID and Ubi-s-ID (Table 2) titrated with varying concentrations of the unlabelled sdAb VHH-IR5.

NMR experiments for VHH-IR5 (Table 2) resonance assignments were carried out at 800 MHz with a 0.3 mM uniformly 15N-/13C-labeled VHH-IR5 dissolved in the NMR buffer (see above) at pH 6.8 followed by the addition of 10% D2O. Multi-dimensional heteronuclear NMR data, i.e. HNCA, HN(CO)CA, HNCACB, CBCA(CO)NH, HN(CA)CO, and HNCO were collected using pulse sequences from the Bruker pulse sequence library. HN(CA)CO and HNCO experiments were carried out with a non-uniform (sparse) sampling (NUS) schedule generated using the jittered rejection sampling algorithm48. These NUS schedules were individually optimized for the 13C and 15N dimensions, separately, each using a 50% of sparsity with denser sampling at short evolution times, resulting in, for example, a total of 32 sampled data points from a uniform (Nyquist) grid of 64. Water suppression was achieved using gradient selection for all 3D NMR experiments. The 3D HN(CA)CO and HNCO spectra were processed from time-domain data matrices after filling the NUS data along with extrapolation in each of the indirect dimensions using an in-house implementation of the Convex-Accelerated Maximum Entropy Algorithm (CAMERA)49. The filled 3D NUS-NMR data matrices were processed using the same processing parameters as for complete data matrices, but leaving out linear prediction, by use of NMRPipe50. All other 3D NMR spectra were processed using NMR Pipe with linear prediction for the two evolution dimensions. Analysis of 3D NMR spectra and resonance assignments were achieved using nmrView51.

**Fragment scanning of the IGF1R Insertion Domain.** A total of 55 overlapping peptides of 15 residues in length were derived from the 674–742 sequence segment of human IGF1R using the PepScan procedure6,12. Binding activities of the surface-immobilized peptides were quantified through ELISA using an anti-His antibody specific to a His-tag present in VHH-IR5 (Table 2). Concentration of VHH-IR5 was 10 μg/mL in all loadings.

Data availability
All data generated or analyzed during this study are included in this published article.

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Author contributions
J.S., J.J.H. and J.K. conceived and designed HDX-MS experiments. J.S. performed HDX-MS experiments, interpreted the data and wrote the manuscript. F.N. designed IGF1R fragment constructs, interpreted NMR data and wrote the manuscript. M.A. and L.M. designed specialized plasmids for expressing intact ubiquitin fusion proteins in E. Coli. P.W. produced isotope-labelled proteins, PX. collected NMR data and both analyzed the NMR spectra. K.K. and G.H. designed SPR experiments and H.v.F. collected and analyzed the data. D.S. designed receptor activation experiments and E.B. performed cell-based assays and analyzed the data. K.K. and D.S. designed IGF1R peptide (PepScan) experiments. J.S. and F.N. contributed equally to manuscript drafting and all authors reviewed and edited the manuscript.

Competing interests
The authors declare no competing interests.

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