# Large scale prospective evaluation of co-folding across 557 Mac1-ligand complexes and three virtual screens

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# Abstract

Accurate prediction of ligand-bound protein complexes and ranking them by affinity are central problems in drug discovery. While deep learning co-folding methods can help address these challenges, their evaluation has been hampered by the difficulties in assessing independence from training data and insufficiently large test sets. Here we test the ability of co-folding methods to predict the structures of 557 ligands bound to the SARS-CoV-2 NSP3 macrodomain (Mac1) that were determined after the training cut-off dates. AlphaFold3 (AF3), Boltz-2, and Chai-1 each reproduced >50% of the Mac1 ligand poses to better than 2 Å RMSD of experiment. Despite the potential for co-folding to describe protein conformational changes that stabilize ligand binding, we did not find that common conformational rearrangements, including peptide flip and a large loop opening, were recapitulated by the co-folding prediction. For AF3 and Chai-1, ligand pose prediction confidence weakly, but significantly, tracked experimental potency, while DOCK3.7 energies were only weakly correlated. Boltz-2 affinity predictions showed the strongest correlation with measured potency and, after calibration, achieved lower mean absolute error than a baseline predictor. We next assessed whether co-folding scores could rescore docking hit-lists to distinguish true ligands from non-binders among hundreds of molecules prospectively experimentally tested against AmpC β-lactamase, the dopamine D4 and the $\sigma_2$ receptors. AF3 ligand pose confidence values did not separate true ligands from high-scoring false-positives as effectively as docking scores or Boltz-2 affinity predictions did. Taken together, the modest, but independent correlations of docking score and co-folding confidence or affinity suggests that integrating physics-based and deep-learning and approaches may help with hit prioritization and subsequent optimization in structure-based ligand discovery.

# Introduction

Deep Learning methods such as AlphaFold2 (Jumper et al. 2021) and RosettaFold (Baek et al. 2021) revolutionized protein structure prediction. Key to their wide adoption has been their extensive testing, first in well-controlled prospective tests like CASP (Kryshtafovych et al. 2021; Ashizawa et al. 2025) and subsequently in many studies where the methods found new applications and new experimental validation (Scardino et al. 2023; Goverde et al. 2023). Notably, these methods can provide protein structural models for small molecule ligand discovery by docking/virtual screening, leveraging physics-inspired functions to score the resulting complexes (Seok and Tiwary 2025; Brocidiacono et al. 2025; Huang et al. 2006; Michino et al. 2025; Xie et al. 2025). Retrospective analyses of AlphaFold2 models suggested that the accuracy of the predicted ligand binding poses was lower than experimental structures (Karelina et al. 2023; Holcomb et al. 2023). However, prospective use of AlphaFold2 models and experimental structures (determined after the training cutoff) to identify new ligands revealed that both approaches yielded similar quality and quantity of ligands (Lyu et al. 2024; Díaz-Holguín et al. 2024). These results indicate that AF2 models may sample alternative low energy conformations that are compatible with distinct ligands from the experimentally resolved conformations, reconciling the poor retrospective recall with the prospective successes.

More recently, the advent of co-folding methods like AlphaFold3 (AF3) (Abramson et al. 2024), Chai (Boitreaud et al. 2024), and Boltz (Wohlwend et al. 2025) that leverage diffusion-based approaches for multi-component (e.g. protein and small molecule) complex predictions, have promised to extend the impact of deep learning methods from protein structure prediction to small molecule ligand-bound protein complexes (He et al. 2025). The advantages co-folding may have over traditional docking include its abilities to explore conformational variability of the protein on-the-fly and to predict ligand affinity, reportedly as well as or better than a gold-standard physics-based method, free energy perturbation (FEP) (Passaro et al. 2025). However, multiple studies have highlighted how co-folding success can track with structural similarity to training structures, consistent with substantial structural memorization (Xu et al. 2025; Škrinjar et al. 2025). An innovative approach introduced adversarial mutations to fill binding sites to show that co-folding models will still produce high confidence poses even when interactions are physically impossible (Masters et al. 2025). These results highlight pervasive memorization and hallucination in current co-folding methods and suggest that alternative computational approaches are needed to quantify the physical basis of the protein-ligand interactions.

Still, co-folding has potential for synergizing with many aspects of structure-based design and more traditional computational chemistry approaches. Two attractive applications are the prediction of ligand-bound complexes, around which structure-based ligand design is articulated (Erlanson et al. 2025); (Backus et al. 2016), and the re-scoring of large library docking hit-lists. Co-folding is orders of magnitude slower than most docking programs, which can address libraries of hundreds-of-millions to billions of compounds on CPU clusters. The method nevertheless is fast enough to re-rank virtual screening “hit-lists” and may be better at categorizing certain types of ligands (e.g. those that stabilize alternative conformations) than the simpler physics-based methods. Moreover, the errors and biases from co-folding affinity estimation methods (Passaro et al. 2025) may be independent from those in docking, suggesting that each method may have different strengths in distinguishing likely from unlikely ligands. However, partly because of the lack of widely adopted community prospective tests, akin to CASP (Gathiaka et al. 2016) and partly because of the challenges of developing, testing, and potentially optimizing new chemical matter for ligand activity studies (Ackloo et al. 2022), co-folding methods have not yet been subject to the rigorous, large-scale testing that helped establish the utility, domain of applicability, and ultimately optimization of the early protein structure-prediction methods.

It therefore seemed interesting to test co-folding predictions of the co-complexed structures and affinities of ligands in a drug-discovery series from outside of their training sets, and to test their ability to

distinguish likely from unlikely ligands from large library virtual screens (Cabeza de Vaca et al. 2025). Here, we evaluated the performance of three widely-used co-folding methods, AF3, Chai-1, and Boltz-2 against two benchmarks that speak to domain utility and are largely outside the benchmarks against which they trained. First, we tested the ability of the co-folding methods to recapitulate the crystallographic structures of 557 different ligands bound to the SARS-CoV-2 Macrodomain (Mac1), an emerging antiviral target (Suryawanshi et al. 2025). This dataset represents the scale and depth typical of the early phase of industrial structure-based drug design campaigns. Second**,** we investigated the ability of the co-folding methods to distinguish true- from false-positives in hit lists from recent large library, prospective docking against three well-behaved targets: AmpC β-lactamase, the σ2 receptor, and the dopamine D4 receptor (Lyu et al. 2019; Alon et al. 2021; Liu et al. 2025). In the campaigns against these targets, between 500 and 1500 molecules were prospectively tested across a range of docking ranks identifying ligands with previously unexplored chemotypes as well as finding many docking false positives. Contemporaneous work has highlighted the importance of using AF3 in the same experimental datasets from ultra-large library docking campaigns (Menon et al. 2025). Our analyses reveal the strengths and limitations of different co-folding methods versus traditional docking and point to potential synergies that can improve probe and lead discovery pipelines.

# Results

## A set of 557 previously unreported Mac1 ligands complexes

As part of a drug discovery effort (Suryawanshi et al. 2025; Jaishankar et al. 2025; Correy et al. 2025; Flowers et al. 2025; Gahbauer et al. 2023; Schuller et al. 2021), we tested thousands of novel small molecules as inhibitors of Mac1, a new target for SARS-CoV2 and other viruses. This led to potent inhibitors that are active *in vivo* as anti-virals (Suryawanshi et al. 2025; Jaishankar et al. 2025). As part of this work, we determined x-ray crystal structures for ~1000 small molecules bound to Mac1, typically at close to 1 Å resolution, but only a limited number of these were disclosed in the papers describing the more advanced candidates. Moreover, only up to 326 of our molecules, largely consisting of initial fragments and early hit-to-lead material, were available in public databases prior to the training cutoffs for the co-folding methods (e.g. 2021-09-30 for AF3, 2023-06-30 for Boltz-2). Apart from their role in driving inhibitor optimization for Mac1, we thought that these structures might be a resource for evaluating computational approaches against a set that is representative of the scale and depth often seen in industrial structure-based drug design campaigns.

From our internal dataset, we identified 557 ligands where we determined X-ray structures that have not been disclosed previously (**Fig. 1**a, **STable 1**). These molecules all bind in the active site where they compete with the substrate, ADP-ribosylated (ADP-r) peptides. There are several stereotyped interactions across this set, which have been described in previous fragment and lead optimization efforts (Gahbauer et al. 2023; Schuller et al. 2021; Correy et al. 2025), including: interactions with backbone NH groups of Phe156 and Asp157 (the oxyanion site), aromatic packing against the side chain of Phe156 (mimicking the adenosine ring of ADP-r), and hydrogen bonds to Asp22 and Ile23 (the upper active site) (**Fig. 1**a). Because these ligands were synthesized during an active campaign and many did not meet evolving thresholds for single point % inhibition, only 202 have fully measured dose response $IC_{50}$ estimations by a HTRF-based peptide displacement assay. For these compounds, the measured potencies range from sub-micromolar to >500 uM (**Fig. 1**b). This distribution is skewed towards weaker affinities as more potent molecules were progressed and disclosed in papers describing *in vivo*-active lead molecules (Suryawanshi et al. 2025; Jaishankar et al. 2025).

The diversity of the 557 ligands is important for understanding the performance of the co-folding methods. By topological similarity, using ECFP4-based pairwise Tanimoto coefficients (Tc), the ligands were diverse, with pairwise Tc clustering between 0.2 to 0.4 (**SFig.1**a, **STable 2**). For context, a scaffold-hop for the ECFP4-based fingerprint is thought to occur at Tc values below 0.35 (Cramer et al. 2004; Muchmore et al.

2008), while random similarity occurs around Tc values of 0.25 (Hert et al. 2008). As the Tc metric has well-known liabilities that can mask what a chemist would see as similarity, we also calculated pairwise Maximum Common Substructures (MCS) and found a distribution that centers around 44.8% (**SFig.1**b, **STable 2**). Clustering based on pairwise MCS > 35% revealed only four distinct cluster heads and a limited number of outliers (**Fig. 1**c, **STable 2**, **SFig.2**). To quantify the potential for model performance to be driven by memorization, for each ligand in our set, we retrieved the most similar ligand that was deposited prior to the training cutoff date by both MCS and Tc (**Fig. 1**d). Here, the Tcs have a trimodal distribution with a small number having Tc > 0.4 (indicating relatively high similarity for this fingerprint), and two larger distributions centered around 0.35 (indicating scaffold hops) and 0.2 (indicating low similarity to any ligand). The MCS similarities spanned a broad range centered around ~60%, consistent with the majority of the dataset representing a small number of conserved scaffolds bearing diverse peripheral substitutions.

## Co-folding can accurately reproduce poses of ligands dissimilar to those trained

Next we assessed how well co-folding methods could predict the structures of the 557 complexes. Three co-folding methods (AlphaFold3, Chai-1, and Boltz-2) were given the SMILES strings of each ligand and the sequence of the enzyme as inputs. In parallel, we performed classic docking with DOCK3.7 using the SMILES of the ligand and the three dimensional structure of the enzyme bound to an early inhibitor (PDB: 5SQW, (Gahbauer et al. 2023) that is roughly contemporaneous with the ligands in this datasets. To assess whether ligands were correctly predicted in the right area of the active site (Nittinger et al. 2025), we used a threshold of ligand center of mass < 2.5 Å after alignment of protein structures: 519 out of 550 AlphaFold3 poses (94%), 509 out of 551 Chai-1 poses (92%) and 511 out of 548 Boltz-2 poses (93%). With DOCK3.7, only 83% (450 out of 544 poses) passed this cutoff. In addition, roughly 1.5% of 557 co-folded or docked molecules did not pass the RDkit-based sanitization pipeline necessary for RMSD comparisons. Therefore, the total number of poses compared for prospective prediction differed slightly among all co-folding and docking methods (**STable 3**).

Across the co-folding methods, AF3 and Chai-1 performed qualitatively similar with a roughly trimodal distribution centered around ~1 Å (the mode), 2 Å and 4 Å RMSD values (**Fig. 2**a). Using the field standard, 2 Å cutoff for correct predictions, AF3 and Chai-1 predicted ~70% of the ligands correctly. Boltz-2 performed less well, with more structures shifting to the 2 Å distribution, and an overall predictive accuracy of 52%. This result was surprising because the later training cut-off date for Boltz-2 meant that it could potentially benefit from additional Mac1 ligand-bound structures. The distribution for docking was more unimodal centered around an RMSD of 2Å, yielding a predictive accuracy of 41%. At least for this set, the co-folding models outperformed DOCK3.7 in accurately predicting the poses of bound ligands.

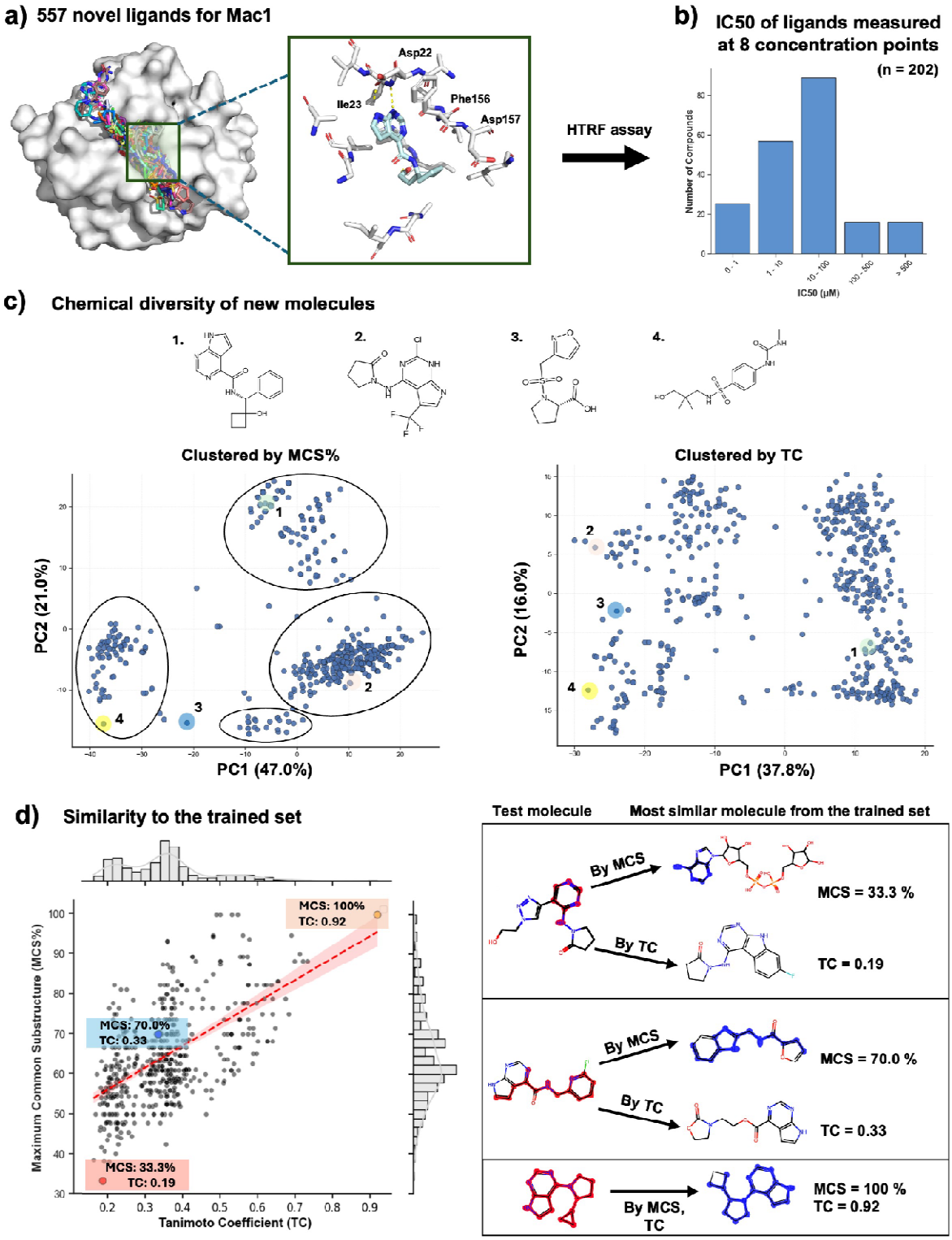



**Figure 1**: **Structure diversity of 557 previously unreported ligands co-folded with the SARS-CoV-2 Mac1 (NSP3 macrodomain 1).** **a)** Global superposition of all 557 ligands within the Mac1 binding pocket highlighting residues involved in key interactions. **b)** The affinity distribution of 202 molecules determined by HTRF (drug-response curves at 8 concentration points). **c)** Principal Component Analysis on pairwise MCS% and Tc to illustrate the chemical diversity of new molecules in the space. **d)** The highest MCS% vs. ECFP-4 Tc values in the training set for each Mac1 compound. At right are examples of the low similarity, scaffold hop, and high similarity distributions, respectively.

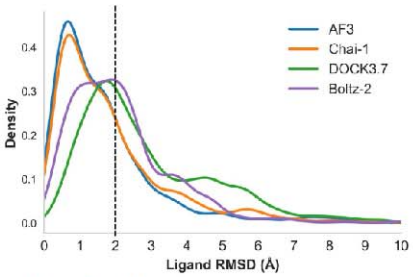
Next we investigated the potential for co-folding methods to identify alternative conformations of the protein that are stabilized by ligand binding. This application is a potentially large differentiator relative to virtual

screening, which generally uses a fixed structure. Although protein alignment gave generally low global C-alpha RMSDs (range 0.1 Å – 0.4 Å; **STable 3**), rare local and high magnitude conformational changes observed experimentally were not captured by co-folding methods. Previously published structures indicated two characteristic states with significant conformational changes: a twisted state, characterized by a peptide backbone flip around Phe156, and an open state, characterized by a ~8 Å loop rearrangement including Gly130 and Ala129 (**Fig. 2**b, **SFig.3**, **STable 4**). The open state has also been observed in the human MacroD1 enzyme (PDB 2X47, (Chen et al. 2011) and PDB 6LH4, (Yang et al. 2020)), suggesting it might be a common feature across macrodomains. We observed that only 4/19 twisted states and 0/20 open states were correctly predicted by AF3. Interestingly, AF3 predicted 8 false positive twisted states, where the ligand was experimentally observed in the normal ground state. This result suggests that there is only a limited ability to predict conformational adaptations, which admittedly may be, at least in part, stabilized by crystal contacts. We next asked how often the predictions recapitulated hydrogen bonding interactions in the Mac1 active site (**SFig.3**a). Among the dominant interactions, we found a low false positive rate for hydrogen bonds with D22 and I23 at the top of the active site, and mispredictions in both directions for hydrogen bonds with F156 and D157 at the oxyanion site (**SFig.3**b). Thus, despite failing to predict large conformational changes and aspects of the detailed hydrogen bonding interactions, AF3 still correctly placed these ligands with low RMSDs relative to the crystal structure pose (Errington et al. 2025).

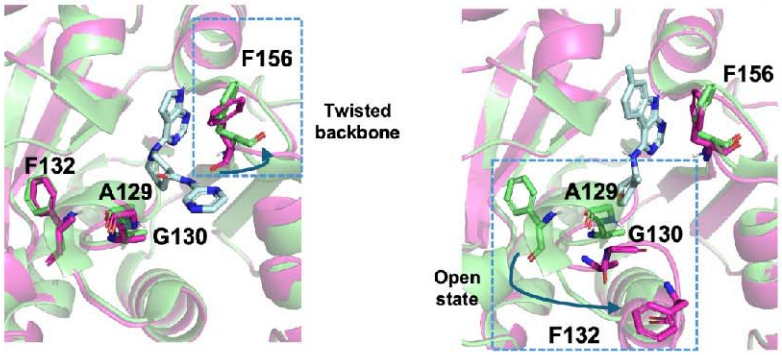
Finally, we asked how much of co-folding success might be explained by memorization of Mac1 ligands deposited prior to the training date (**Fig. 1**d). We measured the correlation between AF3 ligand RMSD, and either the Tc (ECFP4) or %MCS of the most similar ligand within the training set, finding no meaningful signal (**Fig. 2**c). Examples illustrated have low similarity in both metrics, and mac-x5015, mac-x3538 are not from the commonly shared purine scaffold. Nonetheless, even for these cases, AF3 co-folded poses are recovered to within 1 Å accuracy. The weak correlations between RMSD and both similarity metrics illustrate the ability to predict binding modes for ligands dissimilar from the training set.

a)

| Prediction method | # passed RDkit-based sanitization | # COM distance < 2.5 Å | Success rate (%) RMSD < 2 Å |
|---|---|---|---|
| AlphaFold3 | 550 | 519 | 72.2 (402/557) |
| Chai-1 | 551 | 509 | 66.6 (371/557) |
| Boltz-2 | 548 | 511 | 52.1 (290/557) |
| DOCK3.7 | 544 | 450 | 40.9 (228/557) |




b)




| | F156 (Twisted backbone) | A129, G130 (Open state) |
|---|---|---|
| Alternate conformations (total) | 19 | 20 |
| Success rates with AF3 (% correctly predicted) | 4/19 (21%) | 0/20 (0%) |
| False positives among normal conformations | 8/538 (1.5%) | 0/537 (0%) |

c)

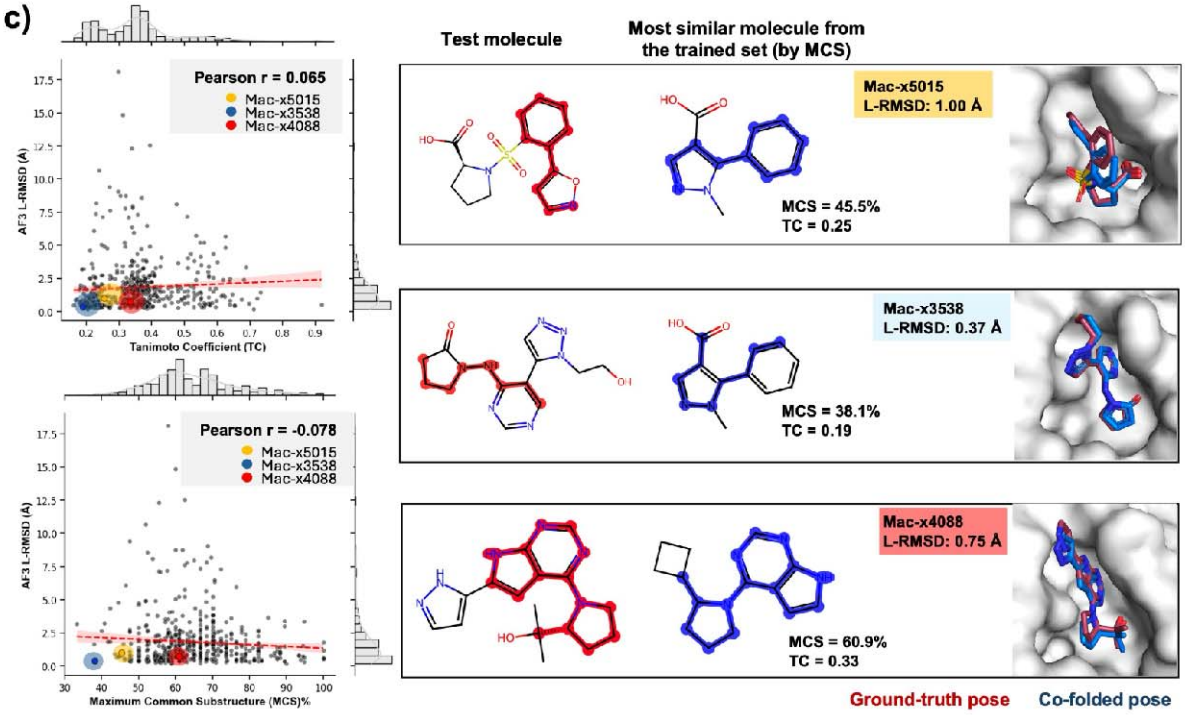



**Figure 2: Co-folding outperforms DOCK3.7 in predicting the ligand pose for > 500 ligands against Mac1 binding site.** a) Histogram distributions of ligand heavy-atom RMSD (L-RMSD) between predicted and crystallographic poses obtained with AF3, Chai-1, Boltz-2 and DOCK3.7, with success rate defined as RMSD < 2 Å, b) Demonstration of alternate conformations of Mac1 binding pocket (twisted backbone and open state) and ability of AF3 to capture these conformational changes (residues in green are from PDB ID: 5SQW, and residues in pink have altered conformations), c) AF3 L-RMSD pose recovery is compared with TC ECFP4 and MCS% and test molecules with poor similarity to the closest molecule in the training set (selected based on MCS%).

## Model confidence and energy scores reflect pose reproduction and affinities

A central question for any co-folding or docking method is whether its internal confidence or energy scores track with experimental pose accuracy and affinity. We used the structure confidence scores reported by AF3, the per atom Predicted Local Distance Difference Test (pLDDT) scores for ligand atoms, and for Chai-1 the interface predictive template modeling (ipTM) score. While these metrics are intended to estimate the model confidence in accuracy, they are not explicitly designed to predict affinity. In contrast, the Boltz-2 affinity module predicts $IC_{50}$ ($pIC_{50}$) which should correlate with affinity, whilst the DOCK3.7 score, though in kcal/mol, is not expected to correlate strongly with experimental affinity (Shoichet 2004; Irwin and Shoichet 2016; Tirado-Rives and Jorgensen 2006). Boltz-2 affinity score and DOCK energies do not explicitly estimate the accuracy (RMSD) of the prediction.

We first generated Receiver Operating Characteristic (ROC) curves to measure the retrieval of True Positives (good score, RMSD < 2Å) versus False Positives (good score, RMSD > 2Å) (**Fig. 3**a). We observed the best performance for AF3 L-pLDDT (AUC=0.76) and Chai-1 ipTM (AUC=0.73). This indicates that the internal model confidence has predictive power and, indeed, we observed that these scores are correlated with RMSD (**SFig.4**). Perhaps not surprisingly, the performance of the DOCK3.7 energy (AUC=0.67) and Boltz-2 $pIC_{50}$ (AUC=0.55) scores, which are designed to predict affinity, was poorer than those designed to predict accuracy. The DOCK3.7 scores correlated with Mac1 ligand RMSD with a Pearson *R* of 0.35 ($p < 0.001$), which is in line with expectations from historical analyses of docking studies (Wierbowski et al. 2020). However, Boltz-2 affinity scores did not correlate with the predicted pose (Pearson $r = -0.03$, $p = 0.49$), which likely results from the fact that its internal structure prediction confidence module is separated from the affinity module (Passaro et al. 2025). Indeed, using the Boltz-2 structure prediction confidence metric (iPTM), yields an AUC of 0.73, comparable to AF3 and Chai-1. Since confidence scores correlated with RMSD, we also investigated how each method was able to predict poses and scores depending on similarity of molecules to the trained set (**SFig.5**). As before (**Fig. 2**c), the performance of co-folding and docking scores did not show a strong dependence on novelty relative to the training set. This result suggests that co-folding can accurately reproduce poses of ligands dissimilar to those on which it trained, and that the model internal confidence scores reflect how well poses are reproduced.

Representative examples of correct and incorrect pose predictions across co-folding and docking methods are illustrated with mac-x3927 and mac-x4091, respectively (**Fig. 3**b,c). Both compounds exhibited comparable experimental IC□□ values (20 µM for mac-x4091 and 42 µM for mac-x3927) and similarity to molecules in the training set (**Fig. 3**b). The molecule consistently predicted correctly (mac-x3927) was a small, rigid fragment comprising a purine scaffold rigidly linked to a fluorobenzene moiety, favoring a well-defined binding mode. In contrast, the compound mispredicted by all methods (mac-x4091) was larger and more conformationally flexible, with an alkyne linker prone to flipping, potentially altering the orientation of the fluorobenzyl ether substituent. Although this group packs against a crystal contact, the high predictive confidence/affinity scores across co-folding methods (AF3 L-pLDDT = 86.4, Chai-1 ipTM = 0.91) are likely explained by the adenosine mimic purine group where predictions recapitulate key interactions observed in the experimental structure.

To understand if prediction errors between methods are correlated, we directly compared the agreement between AF3 to Chai-1, Boltz-2 or DOCK3.7 predictions with the agreement between AF3 and experimental ground truth (**SFig.6**). We observed that while there are a small number of structures where AF3 predicts much better than other methods (high AF3-other prediction RMSD, low AF3-experiment RMSD), most large errors are either missed in a similar way (low AF3-other prediction RMSD, but high AF3-experiment RMSD) or are both extreme mispredictions (high AF3-other prediction RMSD and AF3-experiment RMSD). Consistent with this result, in comparing RMSD relative to experimental structures for all methods (**SFig.7**), we observed that Chai-1 and AF3 exhibited the strongest agreement ($r = 0.72$), whereas poses from DOCK and Boltz-2 diverged more ($r = 0.45$ for AF3 vs. DOCK, and $r = 0.52$ for AF3 vs. Boltz-2). Collectively, these results

suggest that the co-folding methods (AF3, Chai-1) may be complementary to more physics-based approaches like DOCK3.7 as the errors are less correlated than observed for the different cofolding methods.

Second, we asked how well these scores correlate with experimentally measured potency (**Fig. 4**a). The correlation was best for the Boltz-2 $pIC_{50}$ (r = 0.6), significantly outperforming DOCK when tested with Friedman and Conover tests with Bonferroni corrections (p = 0.0057). Prior to any linear correction, the mean absolute error (MAE) in affinity prediction was ~1.6 kcals (1.19 pIC units), with most affinities predicted as too potent. Empirically correcting with a linear fit reduced this error to ~0.7 kcals (0.54 pIC units). To put these results in context, simply predicting the mean pIC50 from the dataset for every compound would yield a baseline MAE of 0.93 kcal because of the relatively limited dynamic range of the dataset. The DOCK3.7 energy correlated more weakly with potency (r = -0.225), consistent with previous studies that have observed only weak correlations with affinity from virtual screening energies (Kinnings et al. 2011; Jones et al. 2021). The AF3 co-folding confidence metrics correlated slightly better than docking with potency (r = 0.314, **Fig. 4**a), but the post-hoc Conover tests with Bonferroni corrections showed the difference was not statistically meaningful. Despite this, a correlation between pose accuracy and potency was more relevant with AF3 (r = -0.297, **Fig. 4**b), as more accurate poses tend to be more confidently predicted (**SFig.4**). In contrast, there was no correlation between docking accuracy and potency (**Fig. 4**b). Boltz $pIC_{50}$ had the highest correlation between pose accuracy and affinity (r = -0.387). This result is interesting because in Boltz-2 affinity prediction and confidence are decoupled architecturally and have little correlation with each other across the Mac1 dataset (**SFig.4**).

Collectively, these results suggest that the categorization abilities of the co-folding confidence metrics, and their correlation with affinity, could mark an important advance. Still, the Mac1 set, as large as it is, represents only a single system with ligands that share a limited set of pharmacophores. We wanted to broaden testing to more targets and more diverse ligands, such as those encountered in virtual screening hit lists with their focus on new chemotype discovery.

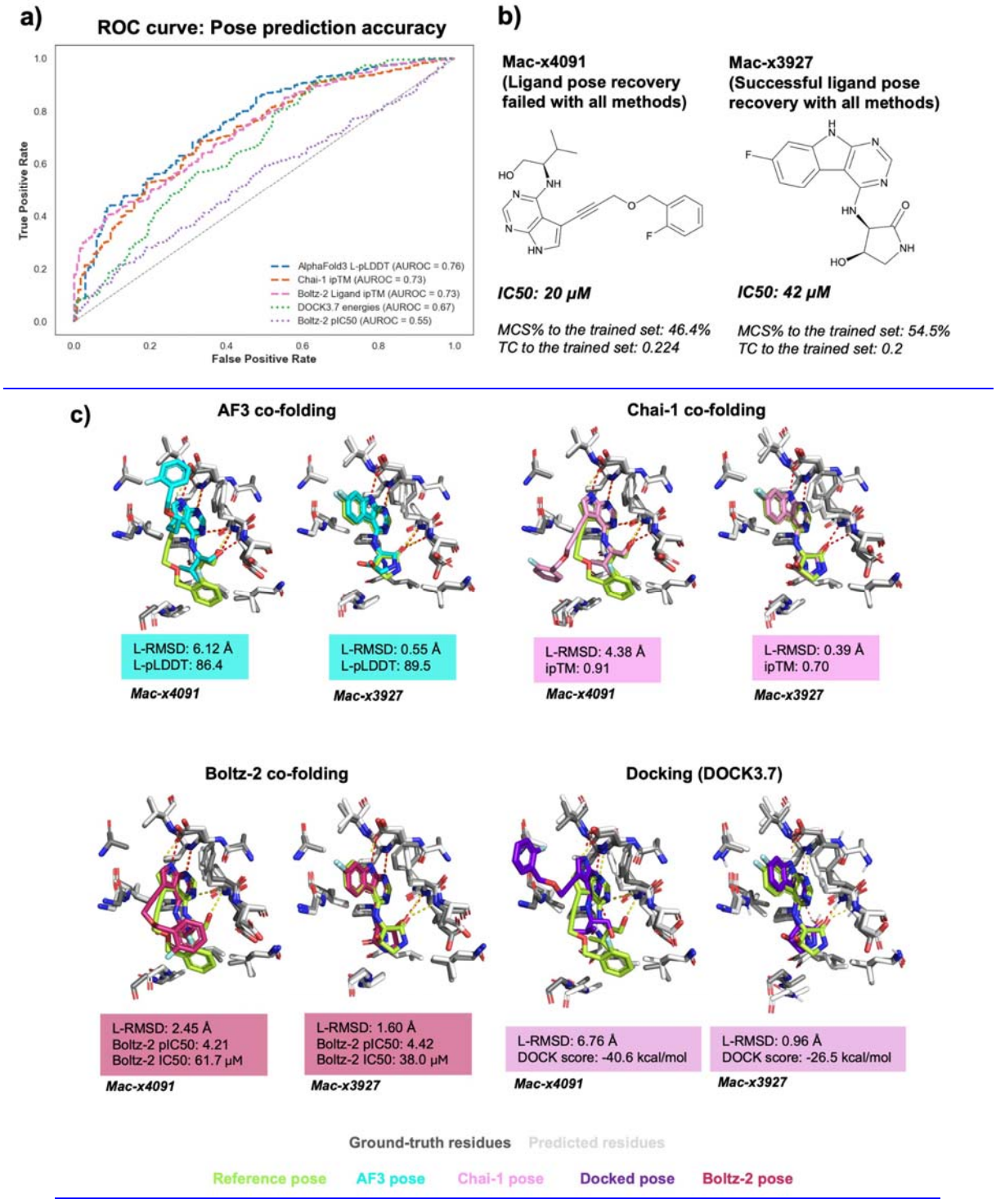



**Figure 3: Complementarity between co-folding confidence metrics and ligand-pose accuracy.** a) Receiver Operating Characteristic (ROC) curves for affinity/energy and confidence scores in AlphaFold3, Chai-1 and Boltz-2, with the success rate being defined as L-RMSD < 2 Å. b) Chemical structure of a compound that was challenging to model with all docking, co-folding methods (mac-x4091), and one that was well-modelled with all methods (mac-x3927), c) Overlay of co-folded (AlphaFold3, Chai-1, Boltz-2) and docked ligand poses on ground-truth, for hits with similar $IC_{50}$, but different structures. Hydrogen bonds with co-folded, docked ligands are shown in red, and those with crystal ligands are shown in yellow.

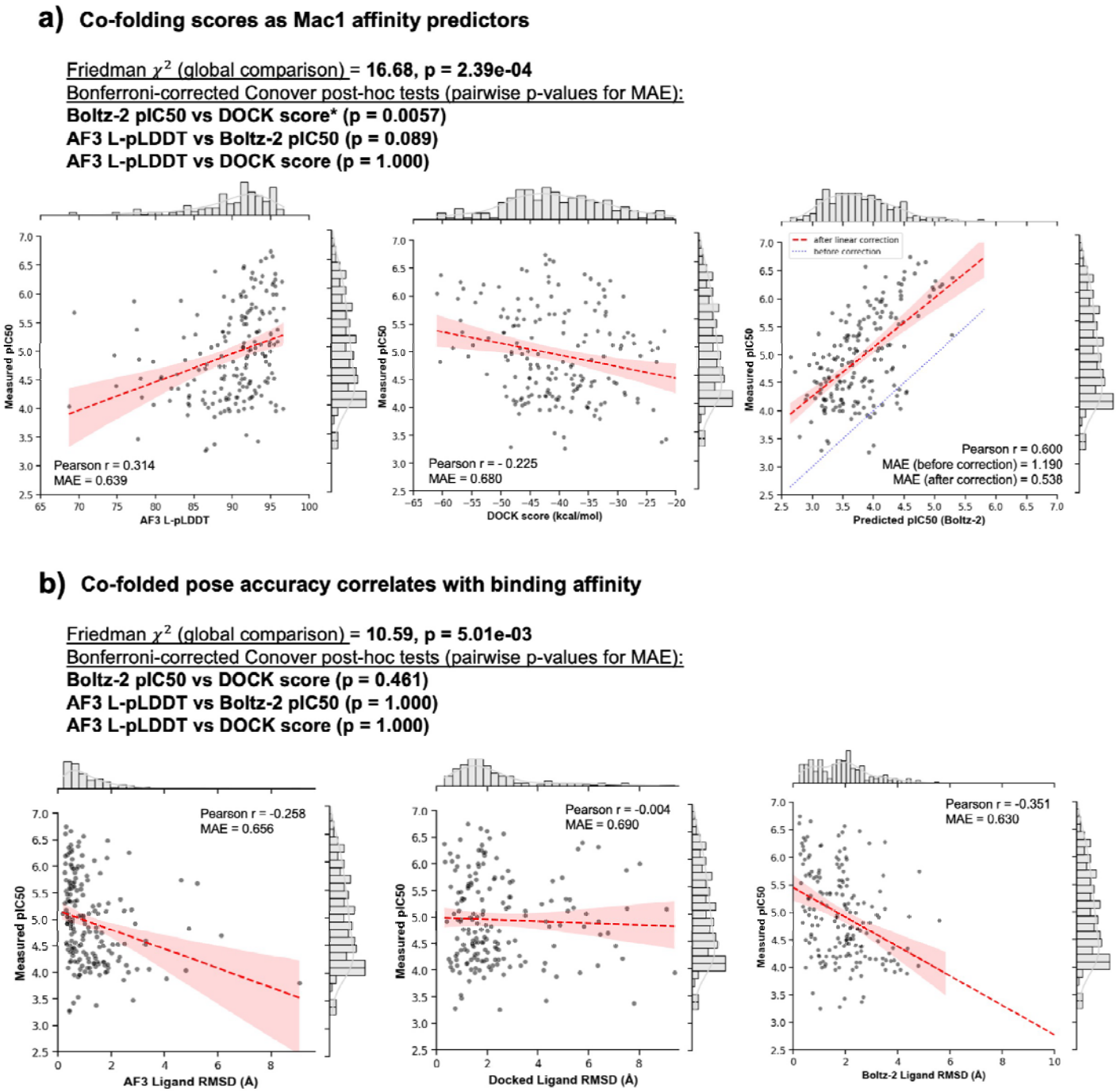



**Figure 4: Co-folding scores and pose accuracy correlate with Mac1 binding affinity for 202 newly synthesized Mac1 ligands.** a) AF3 ligand pLDDT (L-pLDDT) and Boltz-2 predicted pIC50 correlate with experimental pIC50 more strongly than DOCK energy scores, b) Lower co-folded ligand RMSDs associate with higher affinity, while docked pose RMSDs do not. Pearson's Correlation Coefficients and Mean Absolute Errors (MAE) are shown. The baseline MAE was 0.68 pIC (or 0.93 kcals), measured against a fixed average pIC50 (pIC50 = 4.95). Friedman tests with Conover post-hoc comparisons and Bonferroni corrections are used to compare MAE between methods.

## Co-folding struggles to reliably re-rank Docking Hit-lists

Encouraged by the ability of co-folding to reproduce ligand structures and affinities, we were interested to see if the methods could also improve hit identification from large library docking screens. In these screens, hundreds of millions to billions of molecules are docked against a target and scored for complementarity to a binding site. High ranking molecules are then synthesized and tested for binding. While large library docking has successfully discovered potent novel chemotypes in recent studies (Lyu et al. 2019; Gorgulla et al. 2020; Stein et al. 2020; Sadybekov et al. 2022; Fink et al. 2022; Singh et al. 2023; Liu et al. 2025), the method is plagued by false positives (molecules that rank well but are inactive). We reasoned that if co-folding can predict the structures and activities of true ligands confidently, perhaps it could distinguish docking true binders from false positives.

To test this idea, we analyzed experimentally tested compounds from three large and heavily tested docking campaigns: 506 experimentally-tested compounds from a 490-million molecule screen against the $\sigma_2$ receptor (Alon et al. 2021), 1,293 tested compounds from a 1.7-billion molecule screen against AmpC β-lactamase (Liu et al. 2025), and 541 tested compounds from a 138-million molecule screen against dopamine D4 receptors (Lyu et al. 2019) with AF3 (the best performing model from the Mac1 set), DOCK scoring function, and the Boltz-2 $pIC_{50}$ score. These test sets differ from the Mac1 benchmark in three ways. First, they are a mix of molecules that would turn out to be true binders (247 in AmpC, 201 in $\sigma_2$ and 205 in Dopamine D4) and high-ranking docking false positives as judged by measured potencies (worse than 400 μM in AmpC, worse than 2.0 μM against the $\sigma_2$ receptor, and worse than 7.8 μM against the dopamine D4 receptor). For these molecules, the challenge is not only predicting the bound structures of true ligands, but distinguishing true ligands from decoy molecules that by many biophysical criteria fit the binding site well. Second, the docking molecules are more diverse than the Mac1 set (**SFig.2**), which largely falls into four substructure clusters representing conserved pharmacophores. The docking sets were selected, in the original studies, for their topological diversity, and their dissimilarity to previously known ligands. For each target, the ligands group into more than 10-times the number of clusters than the Mac1 dataset, consistent with a greater diversity. Indeed, the average pairwise ECFP4 Tc values within the three sets vary from 0.12 to 0.15, all well-below scaffold-hop cut-offs for this fingerprint (**SFig.2**, **STable 2**). Finally, the sets were all selected for novelty and there was little similarity between the ligands in the hit-lists and known ligands, which ranged between 0.11 and 0.19 by Tc, and between 33 to 42 by MCS%. These ranges were lower than our prior Mac1 dataset (where Tc is distributed around 0.36, and MCS% is distributed around 60.1%), and the lack of similarity with the training set for target systems with hit-lists could be explained by having relatively fewer ligand structures to compare to in the PDB (**SFig.8**, **STable6**). These conditions, while representing new and difficult tests for co-folding, are the common case for structure-based ligand discovery campaigns. For validation, there are more measured $IC_{50}$ values, but they are admittedly not as deeply supported by structural information as the Mac1 set.

We observed the strongest discrimination between active and inactive molecules in the $\sigma_2$ dataset, which contains 201 confirmed actives (apparent $K_i$ values between 2.5 nM and 1.5 μM) and 305 inactive false positives (**Fig. 5**a). DOCK3.7 scores well-separated binders from non-binders (median –55.0 vs –42.5 kcal $mol^{-1}$; AUC = 78.8), though the correlation between docking score and $pK_i$ for true ligands was relatively weak (r = –0.211). AF3 L-pLDDT showed a poorer ability to discriminate between true ligands and false positives (median 75.9 vs 74.1, AUC = 56.1). Intriguingly, despite the worse discriminatory ability, the AF3 correlation with binders $pK_i$ was not much worse than docking's (r = 0.183). Against the σ set, Boltz-2 $pIC_{50}$ performed better still, with high discriminatory power (median 6.70 vs 5.38, AUC = 83.8) and the strongest correlation of affinity predictions against true positives (r = 0.486). Among active molecules, Boltz-2 $pIC_{50}$ had a MAE of only 0.7 kcal and a linear calibration reduced that error to 0.54 kcal (**SFig.9**). This absolute estimation of $IC_{50}$ was

superior to its performance by ~0.5 kcal on the Mac1 dataset; however, it is restricted only to true positives and the performance could be much worse if the false positives were accounted for in the comparison. Since inactive predictions were in the same $pIC_{50}$ range as active molecules, as an optimistic limit of the performance that includes inactive molecules, we assigned an $IC_{50}$ of 4 µM, which is 2X our threshold for declaring a non–binder, to all false positives and calculated the MAE. With this assumption, the MAE is 0.92 kcal (0.56 kcal after linear calibration). A less generous assumption would be to randomly assign an $IC_{50}$ between the $pK_i$ threshold (2 µM) and $pK_i$ = 1 (100 mM). Under this assumption, the MAE is 1.66 kcal (1.55 kcal after linear calibration) (**SFig.10**), which is in line with previously published estimates of the accuracy of this method (Passaro et al. 2025) and only a slight improvement over the MAE (1.84 kcal) produced by predicting the dataset average affinity for each datapoint.

For AmpC (**Fig. 5**b), a total of 1,293 high-ranking molecules were experimentally tested for inhibition (Liu et al. 2025). Among these were 247 molecules that turned out to be true inhibitors. We observed that docking (AUC = 76.5, median -85.6 vs -82.8 kcal/mol) was much stronger than Boltz-2 (AUC = 68.1, median 5.16 vs 4.87) or AF3 (AUC = 60.5, median 76.6 vs 72.0) in discriminating actives from inactives. The correlation of metrics to $pK_i$ among actives was relatively weak for DOCK3.7, AF3 and Boltz-2 in this case. The Boltz-2 $pIC_{50}$ values for AmpC were particularly high in absolute error 1.81 kcal; however, linear correction reduced this error to only 0.42 kcal. Including inactive molecules degraded performance to 2.33 kcal (0.32 kcal with linear correction) under the optimistic and 3.29 kcal (1.04 kcal with linear correction) under the random $IC_{50}$ assumptions (**SFig.10**). Similarly, this represents a small reduction relative to the MAE (1.07 kcal) produced by predicting the dataset average affinity for each datapoint

The dopamine receptor also proved to be more challenging than $\sigma_2$, particularly for AF3. With D4 with 205 confirmed actives (apparent $K_i$ values between 3.1 nM and 7.8 µM) and 336 inactive false positives with, we observed that actives were distinguished from docked false positives by docking (-63.9 vs -52.0 kcal/mol) and Boltz-2 (6.15 vs 5.72) scores, both with AUC ~ 71. In contrast, AF3 L-pLDDT values showed little separation (65.4 vs 66.2) and an AUC = 46.4, which is slightly worse than random (**Fig. 5**c). Among active molecules, the docking correlation was slightly stronger for D4 than it was for $\sigma_2$ (r = -0.247), but performance degraded for AF3 (r = -0.109, notably a positive correlation is expected) and Boltz-2 $pIC_{50}$ (r = 0.284). Despite this reduced linear correlation for the Boltz-2 $pIC_{50}$, the MAE was 0.89 kcal (0.55 kcal after correction). The performance degraded to 1.15 kcal (0.57 kcal after linear correction) under the optimistic assumption of 2X $IC_{50}$ for all false positives and to 1.97 kcal (1.4 kcal after linear correction) under the randomly assigned assumption (**SFig.10**). This result is also a small improvement over the MAE (1.46 kcal) produced by predicting the dataset average affinity for each datapoint

An alternative way of looking at how well the scores can identify active ligands is to perform a hit enrichment analysis using normalized ranking scores (pProp, the negative logarithm of the percentage rank (Liu et al. 2025): a molecule that ranks among the top 0.1% of the docked library would have a pProp of 3, one that ranked among the top 0.01% of the docked library would have a pProp of 4, and so forth). pProp assigned here is based on the ranking within the docked list, for direct comparison between scoring methods and largely confirmed the trends above (**SFig.11**). Both DOCK3.7 and Boltz-2 clearly enriched actives across most targets, with hit rates plateauing at 40 - 50% for high-ranking compounds. AF3 L-pLDDT showed meaningful and comparable enrichment only for AmpC, while providing no enrichment signal for σ□ and D4. Overall, the success of the co-folding methods at predicting structures and correlating with affinities among true Mac1 inhibitors was far less apparent in their ability to distinguish true- from false-positives among docking hit-lists, and to correlate with affinities among these highly diverse molecules. Indeed, the methods showed little advantage over docking, a method that is about four orders of magnitude faster. An important caveat is that the hit-lists were composed of molecules prioritized by docking in the first place, giving it an advantage on these particular sets.

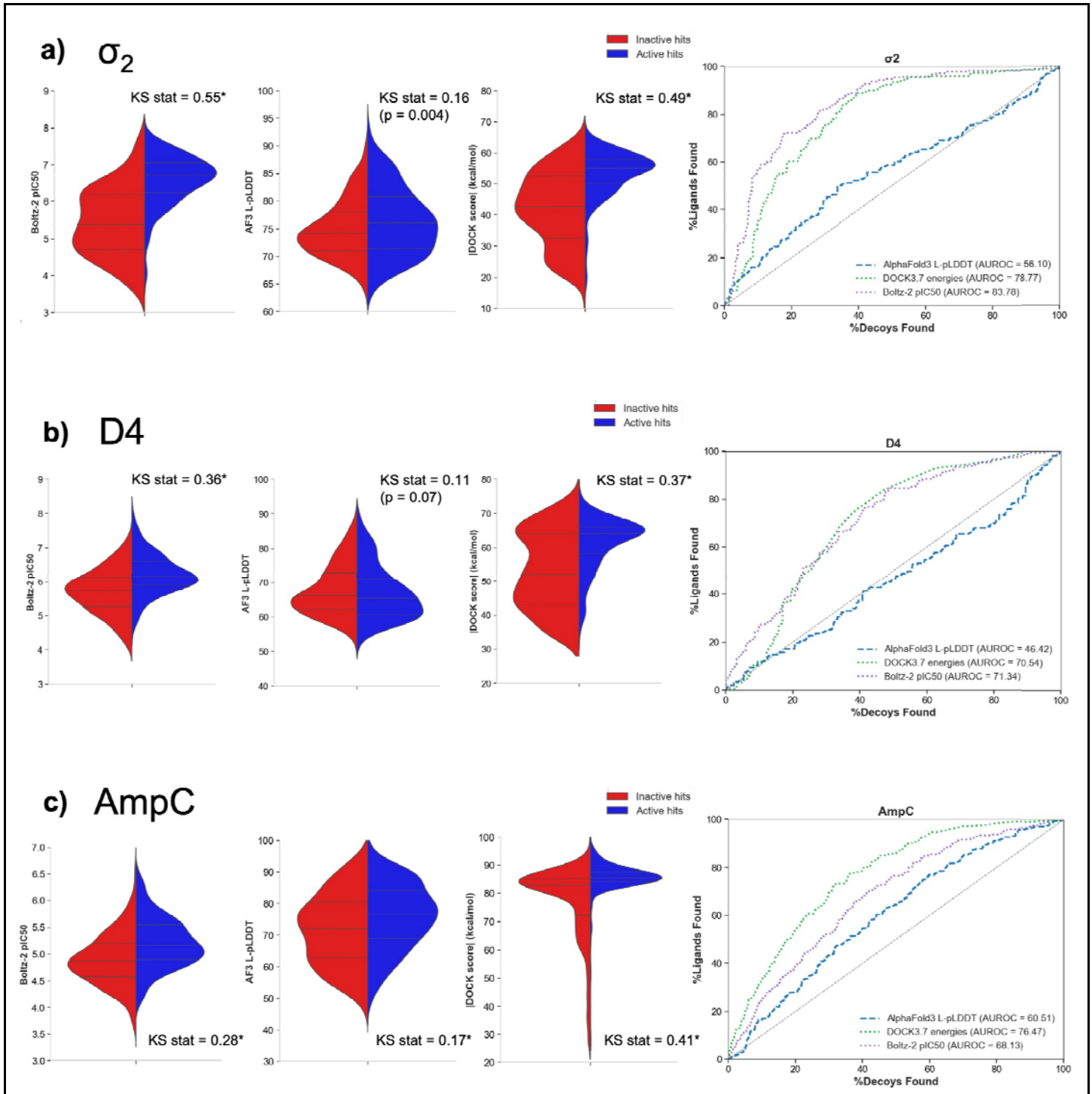



**Figure 5** **Classification of known active hits from docked false positives from an initially-screened docked molecules.** The leftmost panel shows affinity scores from Boltz-2 (Boltz-2 $pIC_{50}$), the middle panel for AF3 L-pLDDT and the right panel for absolute value of DOCK energies (kcal/mol) from DOCK3.7. Docked hit lists from a) σ□, b) Dopamine D4, c) AmpC are grouped into known hits and non-hits, and score distributions are shown in violin plots. Komolgorov-Smirnov test and its KS-statistic (KS stat) is used to indicate meaningful separation of two distribution curves. Asterisks indicate the p-value of KS statistic < 0.001. ROC curves show an ability to distinguish known actives from the docked list with different scoring methods.

# Discussion

A striking observation from this study was the ability of the co-folding methods to predict ligand structures across 557 previously unpublished Mac1 complexes, and for the co-folding confidence metrics to correlate with ligand affinities. By both criteria all co-folding methods were meaningfully better than docking, at least as represented by DOCK3.7. Affinity prediction by Boltz-2 also performed well with mean absolute error that was ~0.2kcal better than a baseline expectation of simply using the mean $IC_{50}$ value. This small improvement was a trend that also held up in the large docking sets as well. Conversely, when confronted with high-ranking molecules from large-library docking, a diverse mix of true- and false-positives with thousands of novel molecules prospectively tested against three unrelated targets, co-folding was not meaningfully better than docking at ligand categorization or at affinity prediction, indeed often worse. This despite the four log-orders greater speed of the docking and its well-known approximations and errors, long a topic of regret in the community (Jorgensen and Tirado-Rives 2005; Irwin and Shoichet 2016). These apparently contradictory results may be reconciled by the different tasks of predicting geometries for molecules that were all true ligands versus that of categorizing binders and non-binders within a highly diverse docking hit-list intentionally dissimilar to known ligands.

It may be that success and relative failure in these different tasks points also to a complementarity between physics-based docking and co-folding in the related, but different tasks of ligand discovery and optimization. The weak correlation between the methods (**SFig.12**) further suggests complementarity between deep learning and physics based approaches. Docking and related methods are designed for categorization among diverse, novel sets of candidate molecules, drawn from the vast new make-on-demand space (Gentile et al. 2020; Radaeva et al. 2023; Sadybekov et al. 2022; Sadybekov and Katritch 2023; Gorgulla et al. 2021; Lyu et al. 2019, 2023; Liu et al. 2024, 2025), whereas the co-folding methods, relying on learned patterns, may be relatively less effective. Where they have complementary strengths is in predicting the accurate geometries of ligands, once known, and their ranking within a series, both of which are crucial for affinity maturation once a series is discovered and with both of which docking has historically struggled. It is in these two linked but different aspects of ligand discovery and ligand optimization where the true complementarity between the methods may lie, at least for now.

Several caveats merit airing. The Mac1 benchmark, while diverse by ECPP4 Tc, is nevertheless dominated by four groups when clustered by MCS and benefits from a large training set of fragment molecules bound to the protein in the training set (Schuller et al. 2021). This may inflate apparent pose recovery rates, although Boltz-2 predicted poses were poorly recapitulated even with its much later training cutoff dates. Unexpectedly, even when the co-folding methods got important conformations of the enzyme binding site wrong, the ligands were still often posed correctly, speaking to a non-physical aspect of the ML methods that others have also described (Masters et al. 2025; Škrinjar et al. 2025). Even though the prospective docking hit lists included thousands of diverse molecules, they still represent only three targets, and so offer a limited view of molecular recognition. Finally, comparing co-folding to docking based on hit-lists themselves selected by docking is arguably unfair to co-folding. Counter-balancing this is the inclusion, in each of the three hit lists, of molecules that had mediocre and poor docking scores intentionally selected to test the correlation between docking score and hit-rate. Here too, the correlation between co-folding score and likelihood to bind, what we sometimes call a “dock-response-curve” was no better than docking’s, often worse (**SFig.11**).

These caveats should not obscure the central lessons from this study. Against a benchmark of 557 topologically diverse Mac1 ligand-bound structures, determined to true atomic resolution and largely dissimilar to structures already in the PDB, each of the co-folding methods predicted ligand poses and interactions with high-fidelity to the experimental structure. For Boltz-2, there was a strong and highly significant correlation between predicted affinity and experimental $IC_{50}$ (compared to a modest correlation for AF3 L-pLDDT and experimental $IC_{50}$). In both cases, co-folding performance was much better than that of DOCK3.7, although the

correlation was only significantly different for Boltz-2 $pIC_{50}$ versus experimental $pIC_{50}$. Conversely, when the more homogeneous benchmark of true ligands was replaced by a diverse set of docked molecules, composed of what would turn out to be hundreds of true ligands and thousands of false positives, the co-folding methods did no better than docking and often did worse. These observations are supported by a fascinating study on some of the same ligand sets as investigated here, using AlphaFold3, reaching similar conclusions (Menon et al. 2025). The relatively strong performance by docking is surprising given its well-known weaknesses and its much reduced computational cost (and therefore greater speed). What emerges are approaches that may be complementary in their roles in ligand discovery (docking) and structure-based ligand optimization (co-folding), areas that will merit future research and application.

# Methods

## Mac1 X-ray crystallography

Ligands were soaked into SARS-CoV-2 Mac1 crystals ($P4_3$ crystal form (Schuller et al. 2021)), and data collected and reduced, as described previously (Gahbauer et al. 2023; Correy et al. 2025). We modeled ligands into PanDDA event maps (Pearce et al. 2017) using ligand coordinates and restraints generated with phenix.elbow (Moriarty et al. 2009).

## Ligand Similarity, Chemical Diversity and Memorization Analyses

To test whether pose-prediction success reflected latent memorization, we quantified chemical similarity between test and training ligands using two complementary metrics. First, extended-connectivity fingerprints (ECFP4) were computed for all molecules, and Tanimoto coefficients (TC) were used to identify, for each test ligand, the most similar compound present in the training corpus (Chai-1 entries deposited before 1 December 2021 and AlphaFold3 entries before 30 September 2021). The training date cutoff for Boltz-2 is 1 June 2023 (**STable 6).**

As another measure of structural similarity, the maximum common substructure percentage (MCS%) was calculated using the rdFMCS algorithm in RDKit, reporting the proportion of heavy atoms shared between two ligands. To evaluate chemical diversity within benchmark datasets, pairwise TC or MCS% values were computed across all ligands. Best-First Clustering (BFC) based on these similarity metrics was used to estimate the number of unique chemotypes represented in each dataset. Principal component analysis (PCA) was used to visualize the resulting chemical space, and to identify an appropriate threshold for MCS-based scaffold grouping. Within the Mac1 benchmark set, applying an MCS cutoff of 35% yielded four distinct scaffolds, in agreement with PCA-derived cluster separation. The same clustering threshold was subsequently applied to define scaffold families for the AmpC, σ□ and D4 benchmark datasets.

## Molecular Docking of Mac1 compounds

The modelled orthosteric site of the Mac1 model was available from the PanDDA analysis group deposition in the Protein Data Bank (PDB), with PDB ID: 5SQW (Gahbauer et al. 2023). We used this structure because the inhibitor (Z5014193706) was the most potent molecule with a structure determined around the same time as the ligands in this dataset were tested. This structure was used to generate matching spheres, which are later used by DOCK3.7 to pre-sample ligand conformations into binding sites by superimposing ligand atoms on spheres in the site. 45 matching spheres were used, and the docked ligands were scored by summing receptor-ligand electrostatic and van der Waals energies and corrected for context-dependent ligand desolvation. Receptor structures were protonated using Reduce. Partial charges from the united-atom AMBER force field were used for all receptor atoms. Potential energy grids for the different energy terms of the scoring function were precalculated using AMBER for the van der Waals term and the Poisson-Boltzmann method QNIFFT for electrostatics. Context-dependent ligand desolvation was calculated from the Generalized Born method. Ligands were protonated at pH7.4 (ChemAxon), and each protomer was rendered into 3D using Corina, followed by conformational sampling using Omega. For the docking campaign of 557 novel compounds, each molecule was sampled in about 2,719 orientations and, on average, 105 conformations by. Overall, over 290 million complexes were sampled and scored.

## Co-folding

All three co-folding methods (AlphaFold3, Chai-1 and Boltz-2) were available from following sources:

- AlphaFold3: https://github.com/google-deepmind/alphafold3
- Chai-1: https://github.com/chaidiscovery/chai-lab.git
- Boltz-2: https://github.com/jwohlwend/boltz

Ligand SMILES and amino acid sequences for the target protein were used to perform co-folding in an array of NVIDIA A40 GPUs. Multiple Sequence Alignment (MSA) was performed with the Jackhammer module. Co-folding tools will generate predicted protein-ligand complex poses (mmCIF files), and different confidence metrics for these predictions. Chai-1 and AlphaFold3 generated global confidence scores (pTM, ipTM), whilst AlphaFold3 had local atom-specific metrics, such as, pLDDT, PAE and contact probabilities in large vectors. To put an emphasis on prediction of ligands, we computed a ligand-centric pLDDT (L-pLDDT) by taking an arithmetic mean of pLDDT on ligand atoms, and a ligand-centric PAE (L-PAE) by taking an average of all PAE elements that contain ligand atoms. Minimum PAE (mPAE) was obtained by scanning all PAE elements that involve ligand atoms, and finding the smallest error. Boltz-2 generated all the confidence metrics in Chai-1 and AlphaFold3, but its affinity module generated an output of the binding likelihood (probability metric) and the predicted binding affinity (Boltz-2 $pIC_{50}$) of compounds. Scripts to prepare input .json, .fasta files, post-process output files, are available in https://github.com/jongbin99/Cofolding.

## Benchmarking Co-folding on Experimentally Validated Compounds

With novel compounds discovered by fragment-based screening, 557 Mac1 complexes were resolved by the crystallography work. The same sequences were used to perform co-folding with AlphaFold3, Chai-1 and Boltz-2. For further sets of analyses with docked hit lists on targets from the Large-Scale Docking (source: lsd.docking.org)(Hall et al. 2025), only AlphaFold3 and Boltz-2 were used. For each system, we redefined the threshold for true hits and non-binders as described herein:

- AmpC β-lactamase: Among 1.7 billion docked molecules, the top scoring 1% were selected (n = 1,293) (Liu et al. 2025). We selected ones with apparent $K_i$ < 400 μM (n = 247) as true hits, and the rest were regarded as non-binders.
- $\sigma_2$ : True hits among these docked molecules were previously defined as displacing greater than 50% of $^3$H ditolylguanidine ([$^3$H]DTG) to $\sigma_2$ (Alon et al. 2021), but in our work, the cutoff is adjusted to 25% displacement, giving a less stringent threshold to include more molecules into the space of true hits. Single concentration point measurement (at 1 μM) is converted to an apparent $K_i$ value using Cheng-Prusoff equation ($K_i = \frac{IC_{50}}{1+\frac{[S]}{K_m}}$), with pre-calculated values of $\frac{[S]}{K_m}$ based on radioligand displacement (Alon et al. 2021). This yields a non-binding cutoff of an apparent $K_i$ < 2.0 $\mu M$.
- Dopamine D4: For the dopamine receptor D4, 138 million molecules were docked and then prioritized synthesizable high-ranking docked molecules (n = 549). True hits from these molecules were defined as displacing more than 50% of $^3$H-N-methylspiperone (Lyu et al. 2019), but similarly to $\sigma_2$, we adjusted the cutoff to 25% displacement. Apparent $K_i$ values are calculated similarly to $\sigma_2$ case, but at 10 μM. This yields a non-binding cutoff of an apparent $K_i$ < 7.8 μM.

## Ligand Pose Evaluation via RMSD against crystal structures

Predicted ligand binding modes were benchmarked against experimentally determined reference poses from ultra-high-resolution (~1.0 Å) crystal structures. The Mac1 protein receptor was first superimposed by least-

squares fitting of all Cα atoms using PyMol. The heavy-atom root-mean-square deviation (RMSD) between the aligned co-folded ligand and its crystallographic counterpart was then calculated. Before RMSD evaluation, both predicted and reference ligands were confirmed using a uniform RDKit-based sanitization pipeline (Buttenschoen et al. 2024). SMILES strings were converted to RDKit Mol objects, chemically validated (valence, aromaticity), and assigned bond orders from a canonical SMILES-derived template. We then check that ground-truth (experimental) and co-folded ligand coordinates for all molecules to have valid chemistry as described with the SMILES-derived template. A substructure match then enabled a one-to-one correspondence of heavy atoms between the ground-truth and co-folded poses for comparison. The crystal ligand and co-folded ligand had the center-of-mass (COM) calculated separately, and measured the distance between each COM. If the distance exceeds 2.5 Å, we would expect the RMSD to be large, indicating that ligand poses are not modelled accurately in an orthosteric pocket (Nittinger et al. 2025). For all the predicted ligand poses, rdMolAlign.CalcRMS from Rdkit.Chem is used for calculating the ligand RMSD between co-folding and the ground-truth crystal pose.

## Hit-rate curves for different scoring methods

To get hit-rate curves for co-folding and docking scores across three targets of docked hit-lists (AmpC, □2, D4), molecules in the list are ordered by increasing score (for DOCK) and decreasing score (for L-pLDDT and Boltz-2 $pIC_{50}$). A rolling window of 100 was used for AmpC and □2, and 50 for D4 (Liu et al. 2025), calculating the hit rate as a fraction of molecules with experimental binding affinity that match our definition for true hits described above (refer to Benchmarking Co-Folding on Experimentally Validated Compounds in Method section). The docked hit-list is a mixture of known actives and non-binders that include false positives based on docking scores. For direct comparison of hit classification performance between DOCK energies, AF3 L-pLDDT and Boltz-2 $pIC_{50}$, the pProp (the negative base 10 logarithm of the fractional rank) was assigned within the list, instead of the billion-scale full docking list. For each window, we used a 95% Wilson confidence interval with normal approximation for the hit rate.

## Statistical Analysis

Pearson's correlation coefficients were used throughout the paper to quantify linear associations between two variables in the study. We have also reported a mean absolute error of data points with respect to the regression line for Boltz-2 affinity $pIC_{50}$ against the measured $pIC_{50}$. The mean absolute errors are also calculated for the perfect fit (Pearson r = 1) and for the fixed measured $pIC_{50}$ (Pearson r = 0), to give an indication of how accurate the Boltz-2 scores are with respect to actual binding energies (in kcal), in terms of correlations and absolute errors. Friedman and post-hoc Conover tests were used to compare how each of AF3, Boltz-2 and DOCK3.7 score predicts for actual binding affinity, and Holm-Bonferroni correction for pairwise comparison p-values (Ash et al. 2025). The distribution of scores for known actives and docked false positives were compared using the Kolmogorov-Smirnov (KS) non-parametric statistical test, as shown in violin plots, and the KS statistic value indicates how distant the two distributions are, and p-values show the extent of statistically detectable differences. All analyses were performed using Python packages.

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# Competing Interests

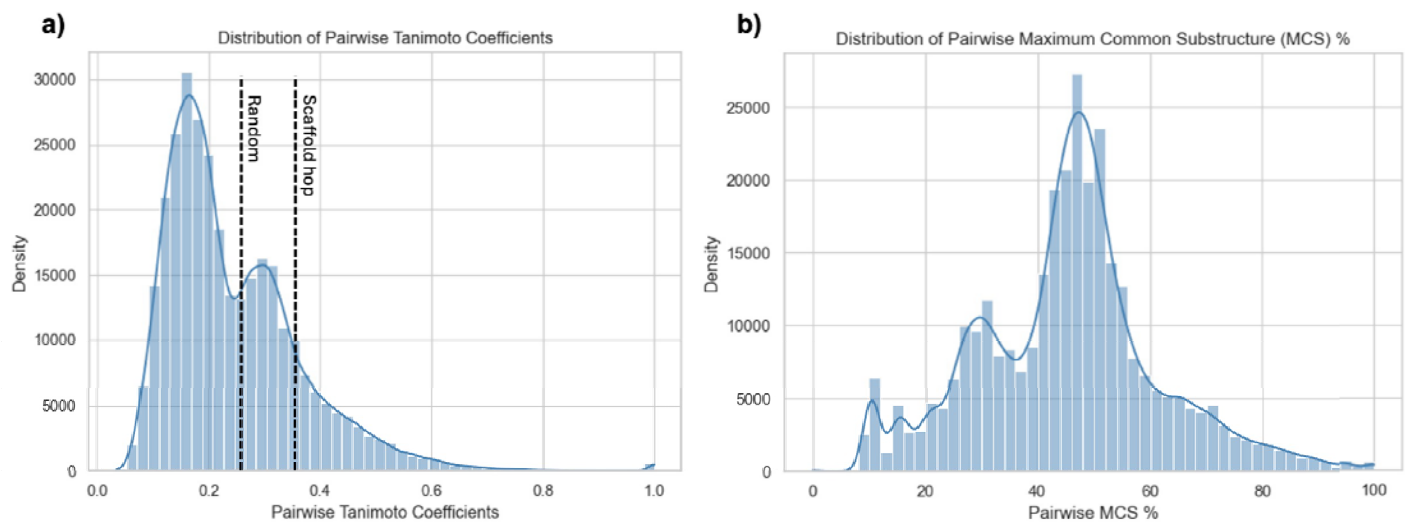
N.J.K.: The Krogan laboratory has received research support from Vir Biotechnology, F. Hoffmann-La Roche, and Rezo Therapeutics. N.J.K. has financially compensated consulting agreements with Maze Therapeutics and Interline Therapeutics. He is on the Board of Directors and is President of Rezo Therapeutics and is a shareholder in Tenaya Therapeutics, Maze Therapeutics, Rezo Therapeutics, GEn1E Lifesciences, and Interline Therapeutics.
A.R.R.: A.R.R. is a co-founder and holds equity in BridgeBio Oncology Therapeutics, Tatara Therapeutics, and Elgia Therapeutics.
A.A.: A.A. is a co-founder of Tango Therapeutics, Azkarra Therapeutics, and Kytarro; serves on boards and advisory boards as listed; receives research support from SPARC; and holds patents on the use of PARP inhibitors held jointly with AstraZeneca.
B.K.S.: B.K.S. is co-founder of BlueDolphin LLC, Epiodyne Inc, and Deep Apple Therapeutics, Inc., and serves on the SRB of Genentech and the SAB of Schrodinger LLC and Vilya Therapeutics.
J.S.F.: J.S.F. holds equity in Relay Therapeutics, Impossible Foods, Arda Therapeutics, Profluent Bio, Interdict Bio (co-founder), Vilya Therapeutics, and Edison Scientific Inc and is a paid consultant for Relay Therapeutics, Profluent Bio, Vilya Therapeutics, and Monimoi Therapeutics.
All other authors declare no competing interests.

# Author contributions

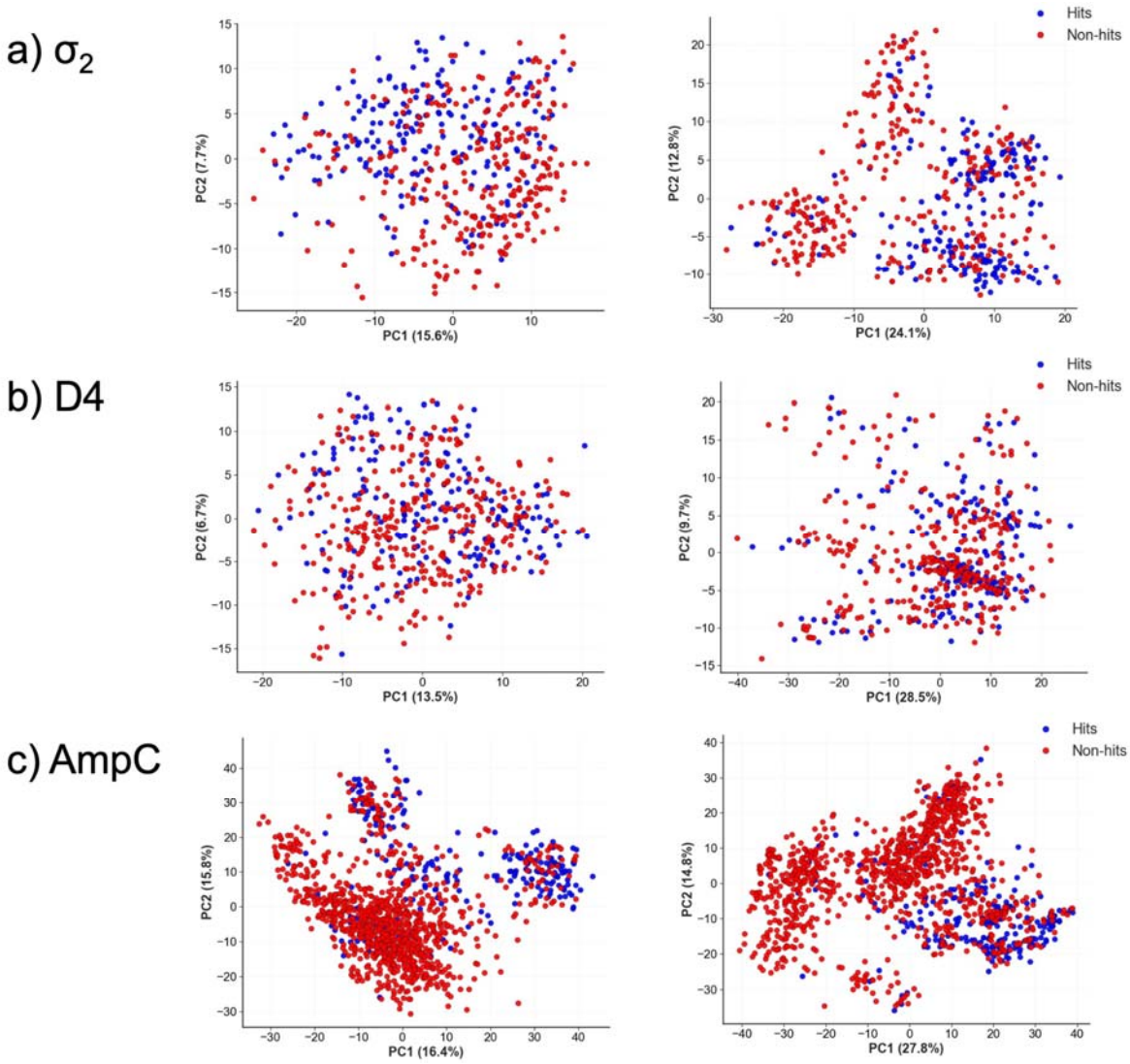
Cofolding: J.K., B.W.H., R.R.
X-ray crystallography: G.J.C.
HTRF assay: Y.U.D., M.G.V.S., M.E.D.
Compound design: G.J.C., M.M.R., T.T., R.L.G., P.J., R.J.N., E.R.H., A.R.R.
Compound synthesis: T.T., R.L.G., P.J.
Docking: M.M.R.
Data analysis: J.K., B.W.H., O.M.
Supervision and project management: S.G.
Supervision and manuscript editing: N.J.K., A.A.
Writing and editing: J.K., G.J.C., M.M.R., A.R.R., A.A., B.K.S., J.S.F.

# Supplementary Information

- **Supplementary Table1** : X-ray data statistics will be released after a small delay to preserve the potential for blind predictions by any new methods.
- **Supplementary Table2** : Pairwise similarity between each dataset (Mac1, AmpC, σ2, D4) based on ECFP-4 TC and MCS%.
- **Supplementary Table3** : 557 new Mac1 ligands with all co-folded, docked poses and scores.
- **Supplementary Table4** : Ligand interactions (hydrogen bonds only) on ground-truth and AF3 predicted models influenced by residue flexibility
- **Supplementary Table5** : Data collection of all docked hits on AmpC, σ□, D4.
- **Supplementary Table6** : PDB accessions by cutoff date



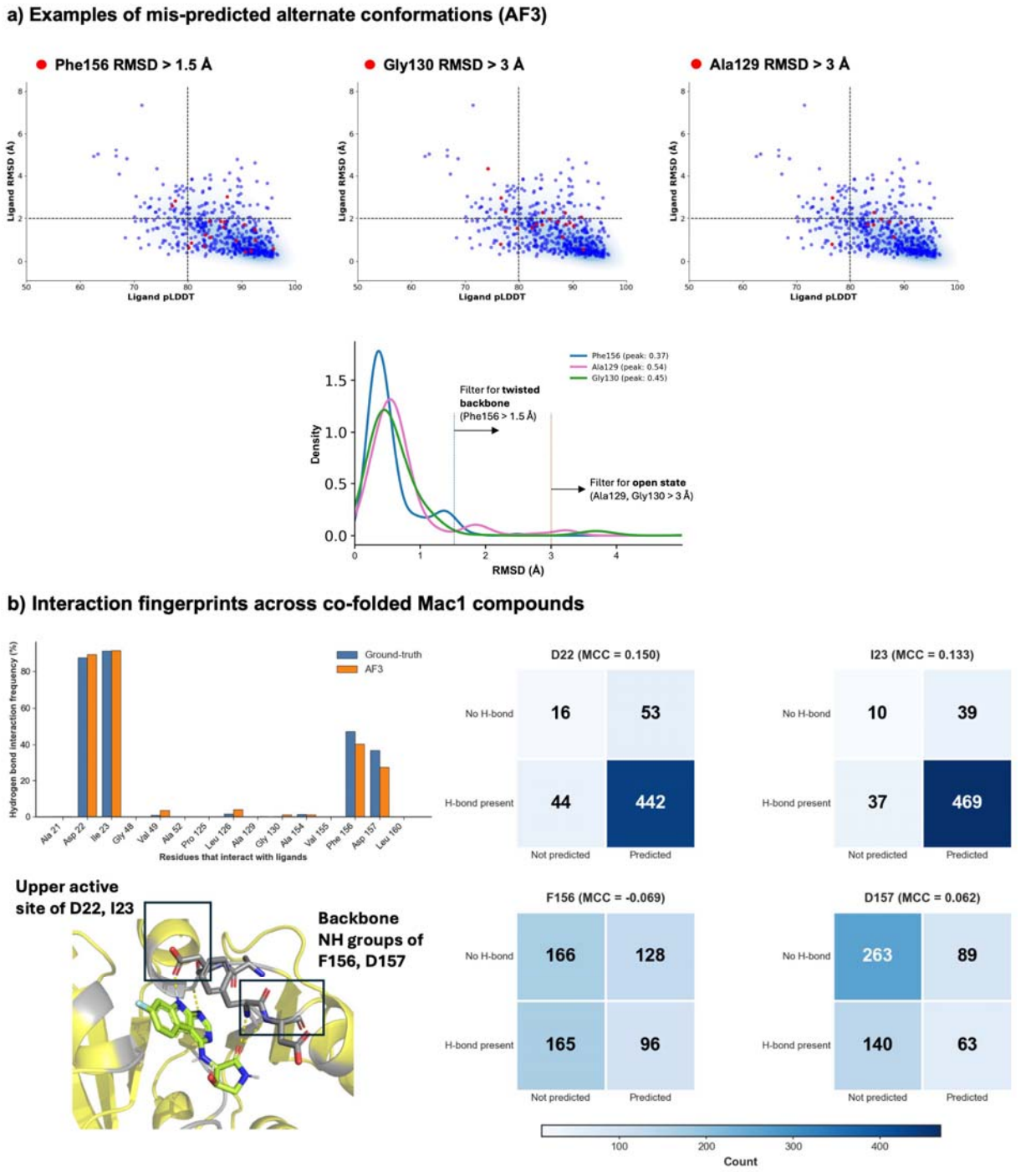

**Supplementary Figure1**: **<u>Distribution of pairwise similarity for Mac1 ligands</u>** using a) ECFP4 Tanimoto Coefficients (TC), with lines indicating values for a “scaffold hop” or random distributions. b) Maximum Common Substructure (MCS) similarity values.




| Target proteins | # known ligands | # total | # cluster heads by TC > 0.35 | # cluster heads by MCS > 35% |
|---|---|---|---|---|
| **$\sigma_2$** | 201 | 506 | 348 | 46 |
| **D4** | 205 | 541 | 410 | 47 |
| **AmpC** | 247 | 1,293 | 500 | 62 |
| **Mac1*** | 557 | 557 | 30 | 4 |

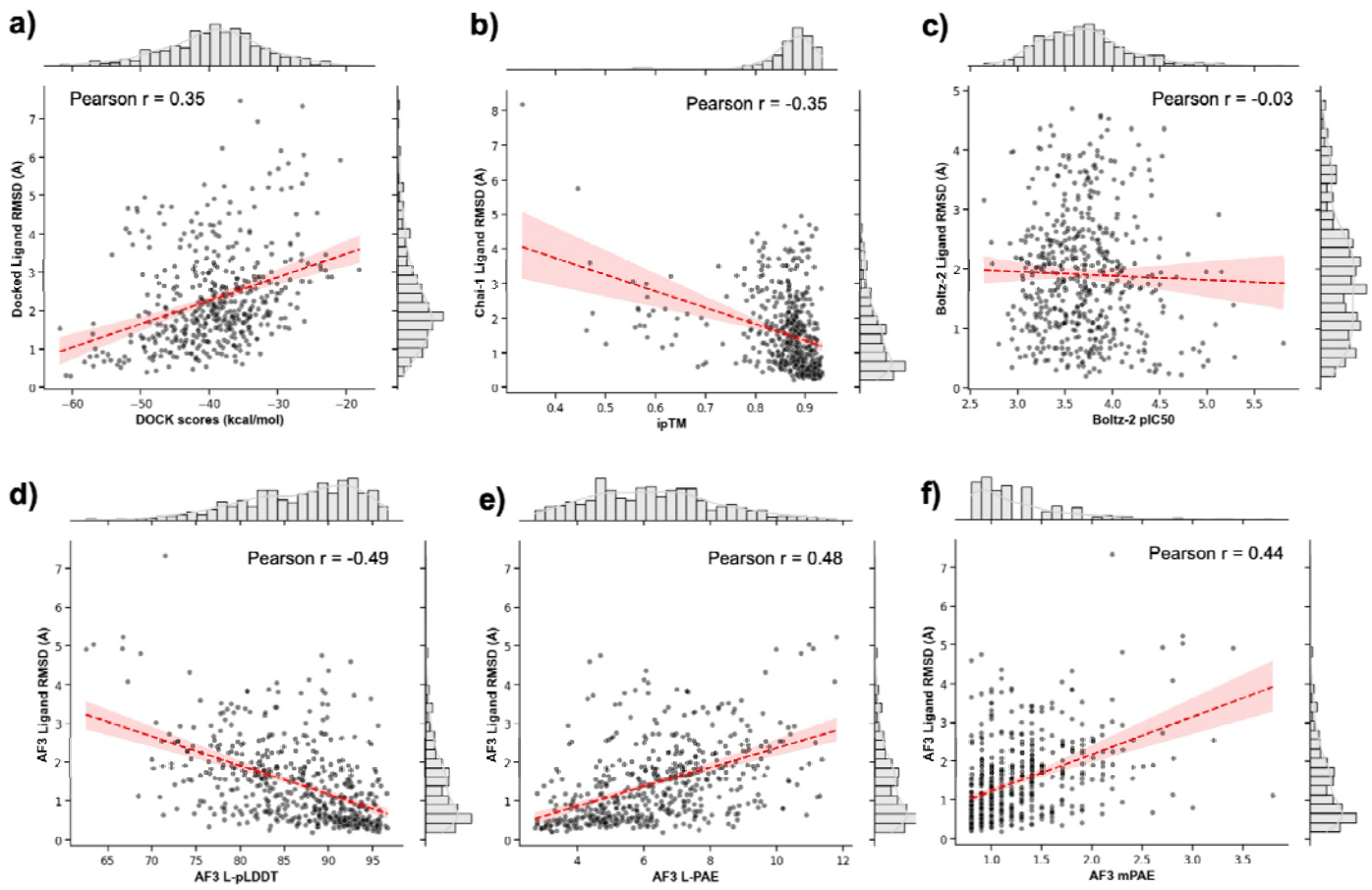
**Supplementary Figure2: Benchmarked datasets are clustered based on TC and MCS% to predict the number of scaffold series.** Principal Component Analysis (PCA) plots are made from making pairwise comparisons of molecules in the docked hit lists of a) $\sigma_2$, b) D4, c) AmpC for similarity using two different metrics: TC (first column) and MCS% (second column). Red points indicate docked false positives in the hit list, and blue points indicate known actives. Mac1 clustering plots were shown in Fig.1 and an asterisk indicates we only have known ligands for the Mac1 set. The number of scaffolds based on cluster heads are

determined by either TC > 0.35, or MCS > 35%. The table gives the summary of the clustering analysis of the datasets used.

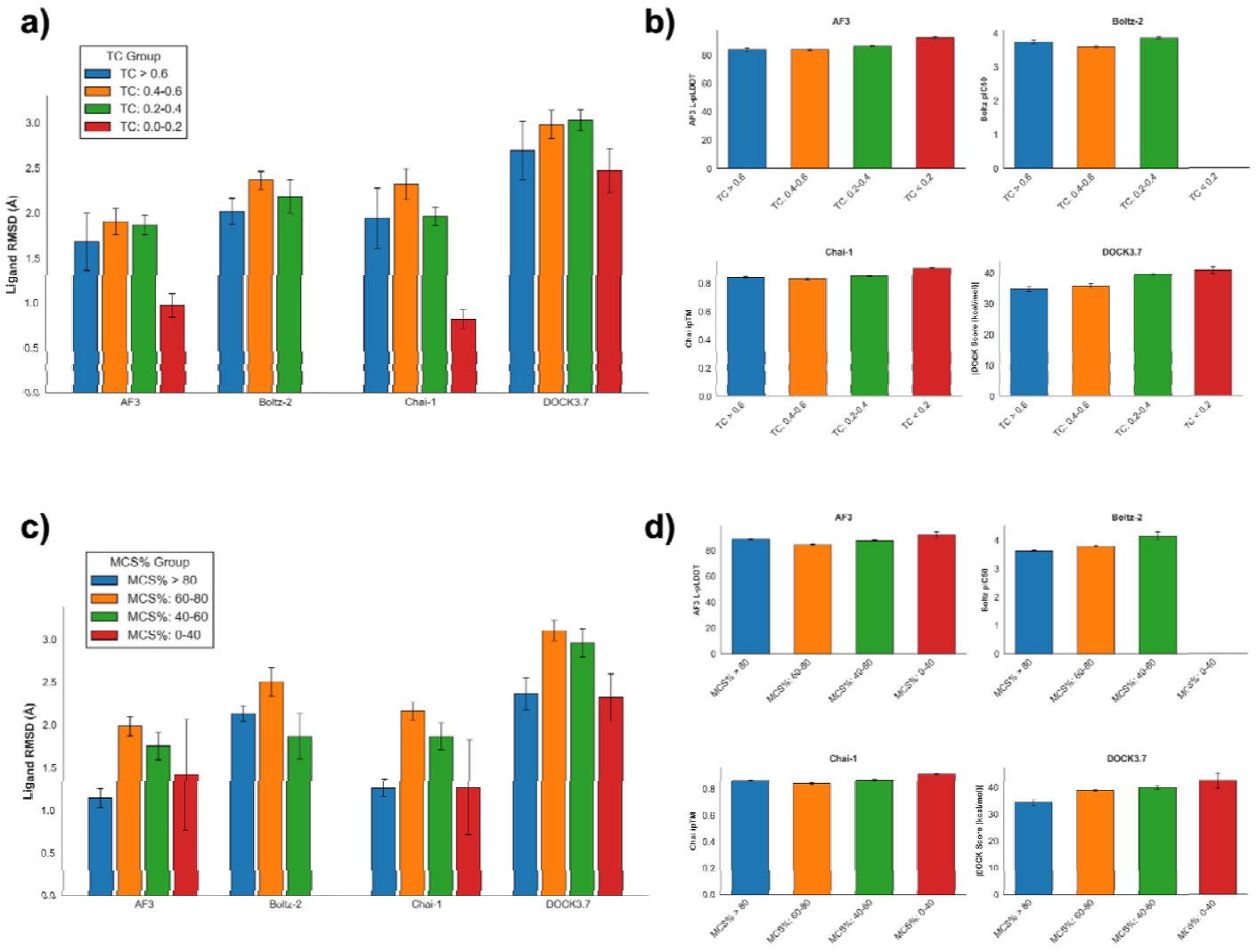



**Supplementary Figure3: <u>Detailed predictions of alternate residue conformations and interactions in co-folded poses.</u>** a) Mis-prediction of alternate conformations of the binding pocket, including twisted backbone (Phe156 RMSD > 1.5 Å) and open structure (Gly130 RMSD, Ala129 RMSD > 3 Å) did not produce false positive or false negative pose prediction. The density distribution plot shows RMSD between residues in PDB ID: 5SQW, and 557 complexes obtained from crystallography, a filter that was used to define alternate conformations. b) Hydrogen bonds between ligands and residues within 5 □ were counted for

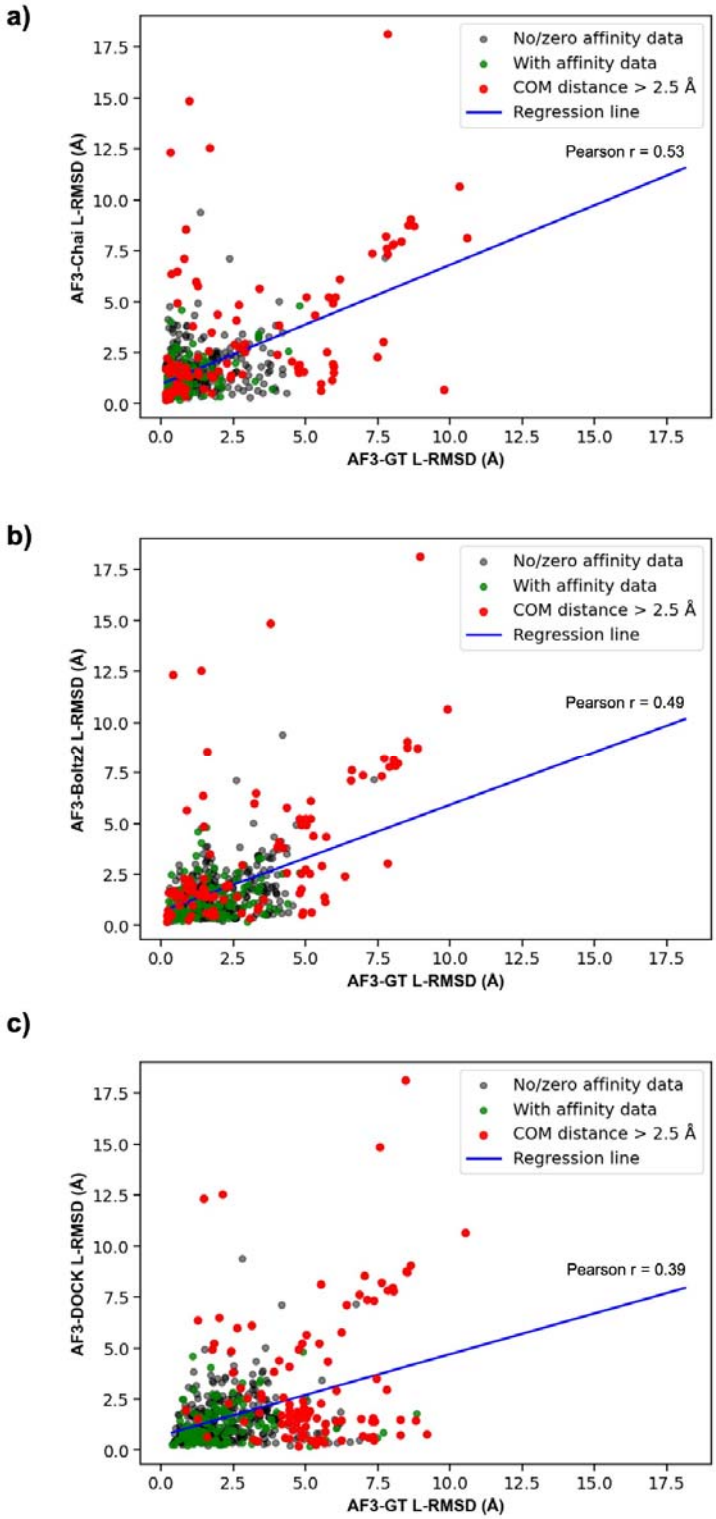
crystal, and for AF3 co-folded structures. Interactions shown with 4 hotspot residues (D22, I23, F156, D157) were compared and confusion matrices show absolute counts of True positives, True negatives, False positives and False negatives. Matthews Correlation Coefficients are calculated for each residue.



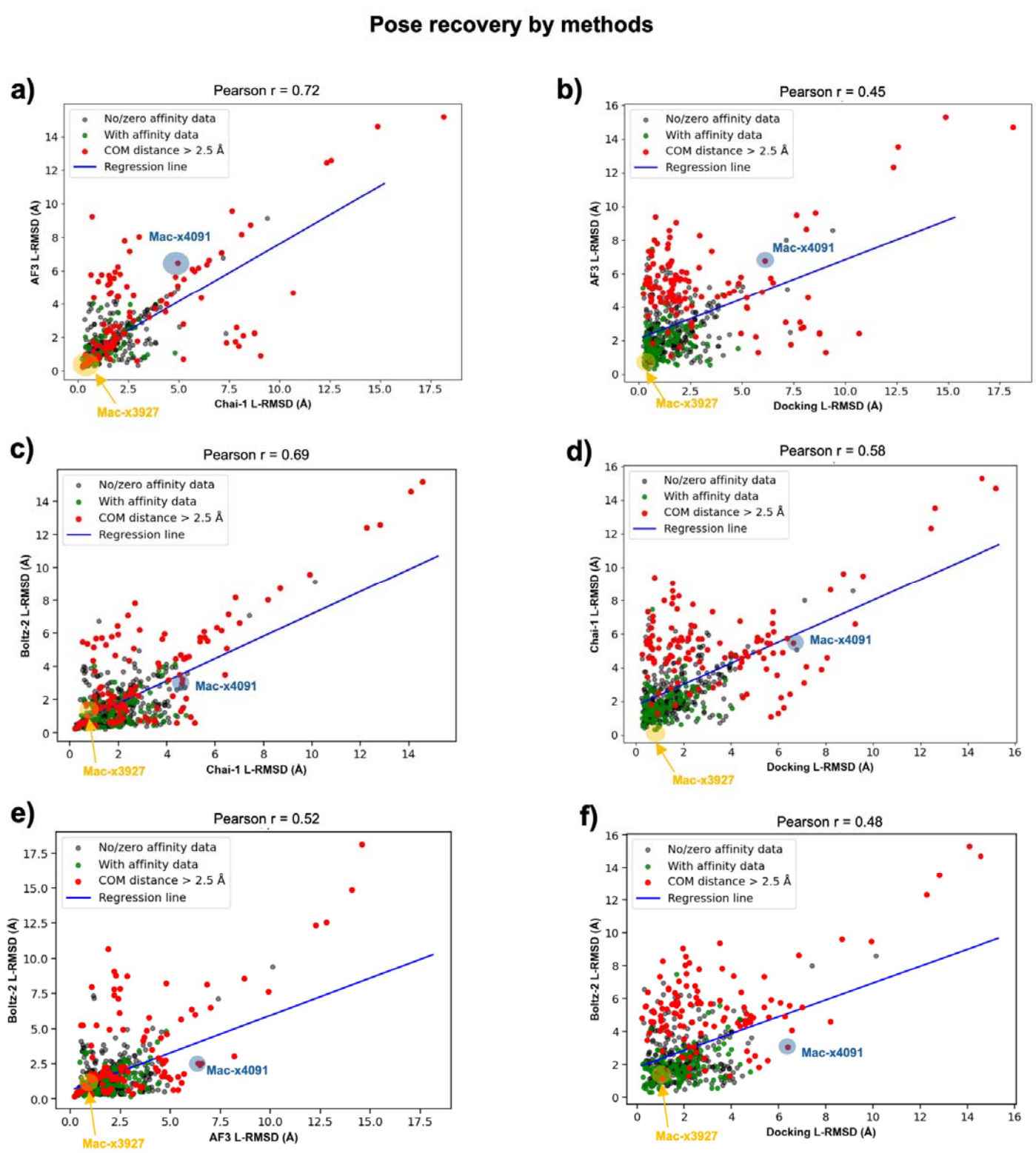

**Supplementary Figure4: <u>Correlations between Mac1 pose recovery with co-folding/docking scoring metrics.</u>** a) Docked pose RMSD is compared with DOCK scores (in kcal/mol), b) Chai-1 RMSD is compared with interface predictive TM-Score (ipTM), and c) Boltz-2 pose RMSD is compared with predicted Boltz-2 $pIC_{50}$ affinity score. AF3 RMSD is compared with three possible scoring metrics from AF3: d) Ligand-specific pLDDT (L-pLDDT), e) Ligand-specific PAE (L-PAE), f) minimum PAE (mPAE).



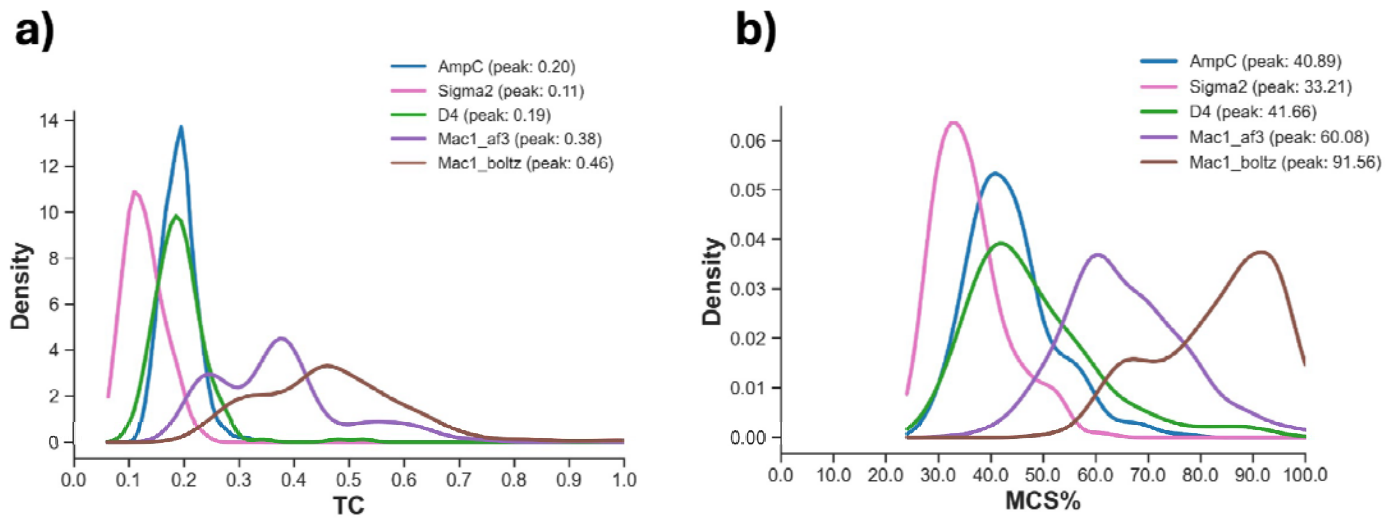

**Supplementary Figure5: <u>Pose accuracy and co-folding, docking scores are compared for different similarity bins.</u>** a) Ligand RMSD between co-folded poses and docked poses are compared to the ground-truth for different Tanimoto Coefficient (TC) bins ( < 0.2, 0.2-0.4, 0.4-0.6, > 0.6), b) Scores for each method (AF3 L-pLDDT, Chai-1 ipTM, Boltz-2 $pIC_{50}$, DOCK3.7 energies) are compared for different TC bins. c) Ligand RMSD for different Maximum Common substructure (MCS%) bins ( < 40, 40-60, 60-80, > 80), d) Scores for each method are compared for different MCS% bins. TC and MCS% of new molecules to the trained set were calculated for each model, and for DOCK3.7 (the only non-co-folding method), the similarity was calculated against the AF3 trained set.



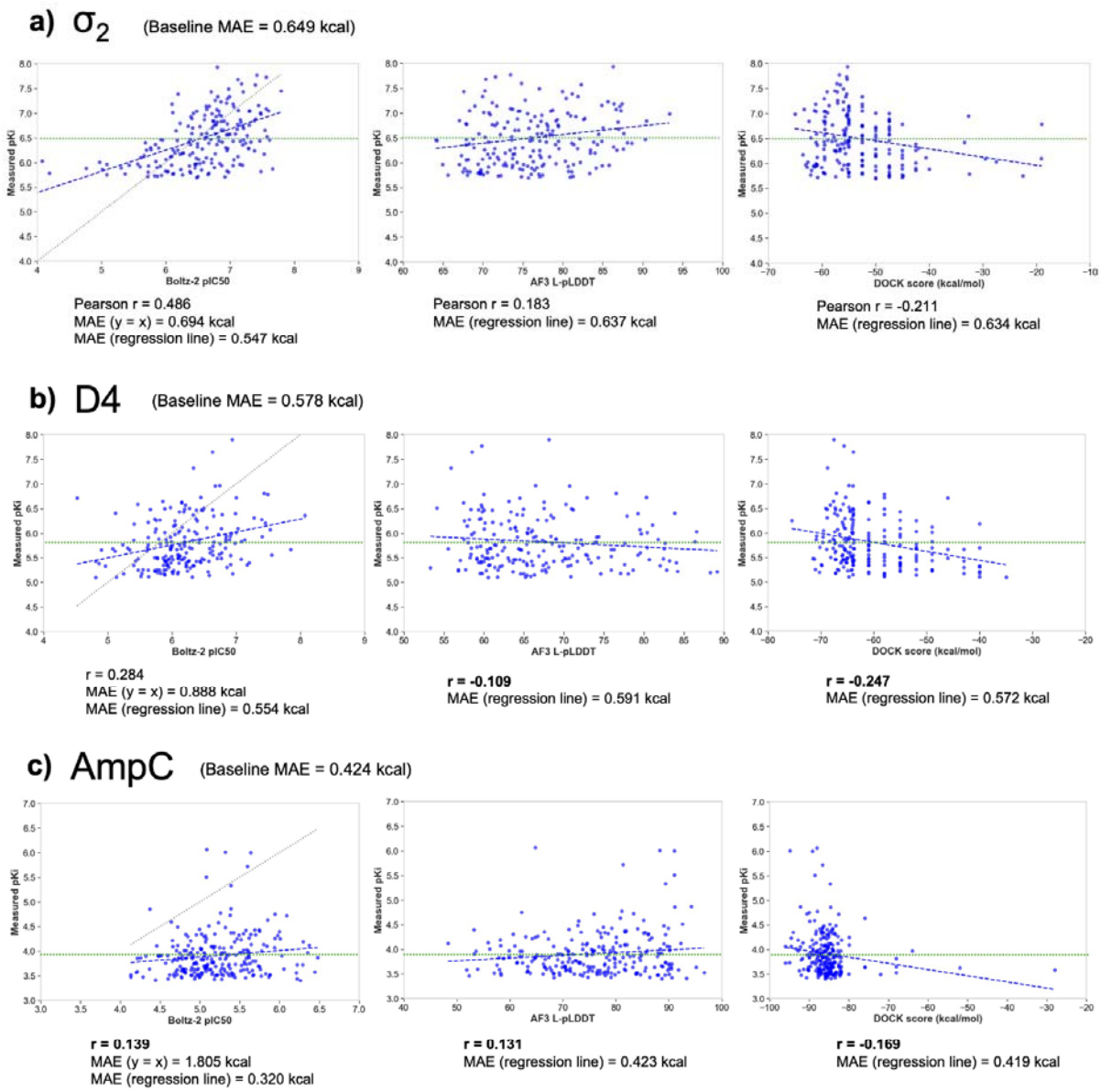

**Supplementary Figure6**: **<u>Correlation between AF3 pose-prediction error and errors from other co-folding and docking methods for 557 Mac1 compounds.</u>** a) AF3-Ground truth L-RMSD vs AF3-Chai L-RMSD, b) AF3-Ground truth L-RMSD vs AF3-Boltz 2 L-RMSD, c) AF3-Ground truth L-RMSD vs AF3-DOCK L-RMSD. Ligands with affinity data (n= 202) are marked in green, those with high COM distance are marked in red, and molecules with no affinity data are marked as grey. Pearson correlation coefficients and regression lines are shown.




**Supplementary Figure7: <u>Pose recovery by methods of co-folding and docking against the ground-truth pose.</u>** a) AF3 vs. Chai-1, b) AF3 vs. DOCK3.7, c) Boltz-2 vs. Chai-1, d) Chai-1 vs. DOCK3.7, e) Boltz-2 vs. AF3, f) Boltz-2 vs. DOCK3.7. Molecules indicated (Mac-x4091, Mac-x3927) are exemplary ligands highlighted in **Fig. 3**, and L-RMSD values quoted here are all compared against the ground-truth pose.




**Supplementary Figure8: <u>Actives from the benchmarked datasets are compared with resolved structures used to train models like AF3 and Chai-1.</u>** Actives from AmpC (n = 247), D4 (n = 205), $\sigma_2$ (n = 201), Mac1 (n = 557) were compared with corresponding trained sets, to give an indication how similar the actives were to those trained by co-folding models, based on a) Tanimoto coefficients, and b) Maximum Common Substructures (%). AF3 and Chai-1 have similar cutoff dates (indicated as Mac1_af3), but Boltz-2 has been trained with more recent PDB and is labelled as Mac1_boltz. Exact PDB IDs used for each model system is shown in Supplementary Table 6.

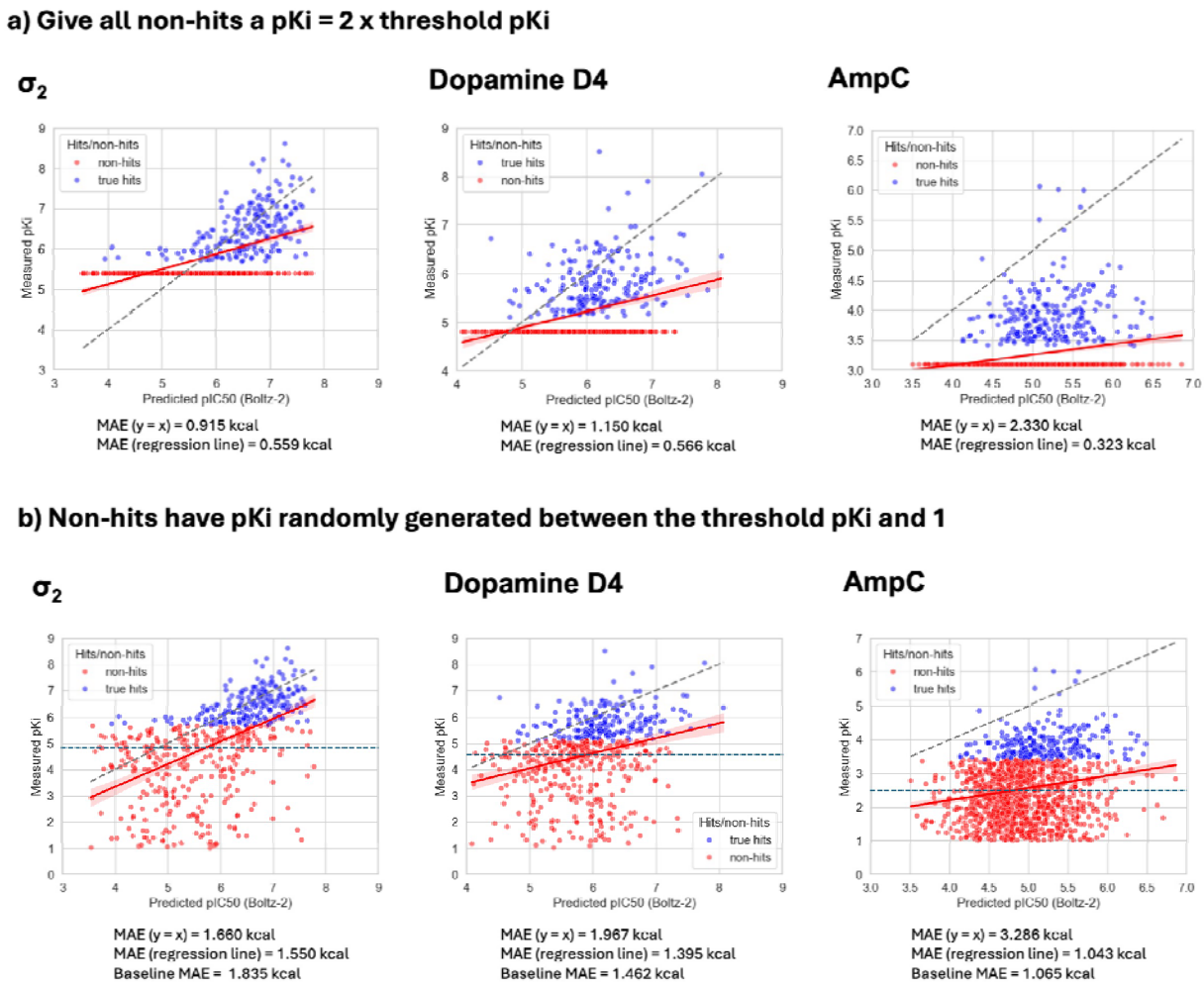



| Bonferroni-corrected p-values | Sigma2 | AmpC | D4 |
|---|---|---|---|
| MAE_AF3 vs MAE_Boltz-2 | 0.0722 | 1.00 | 1.00 |
| MAE_AF3 vs MAE_DOCK | 1.00 | 1.00 | 1.00 |
| MAE_Boltz-2 vs MAE_DOCK | 0.0674 | 1,00 | 1.00 |

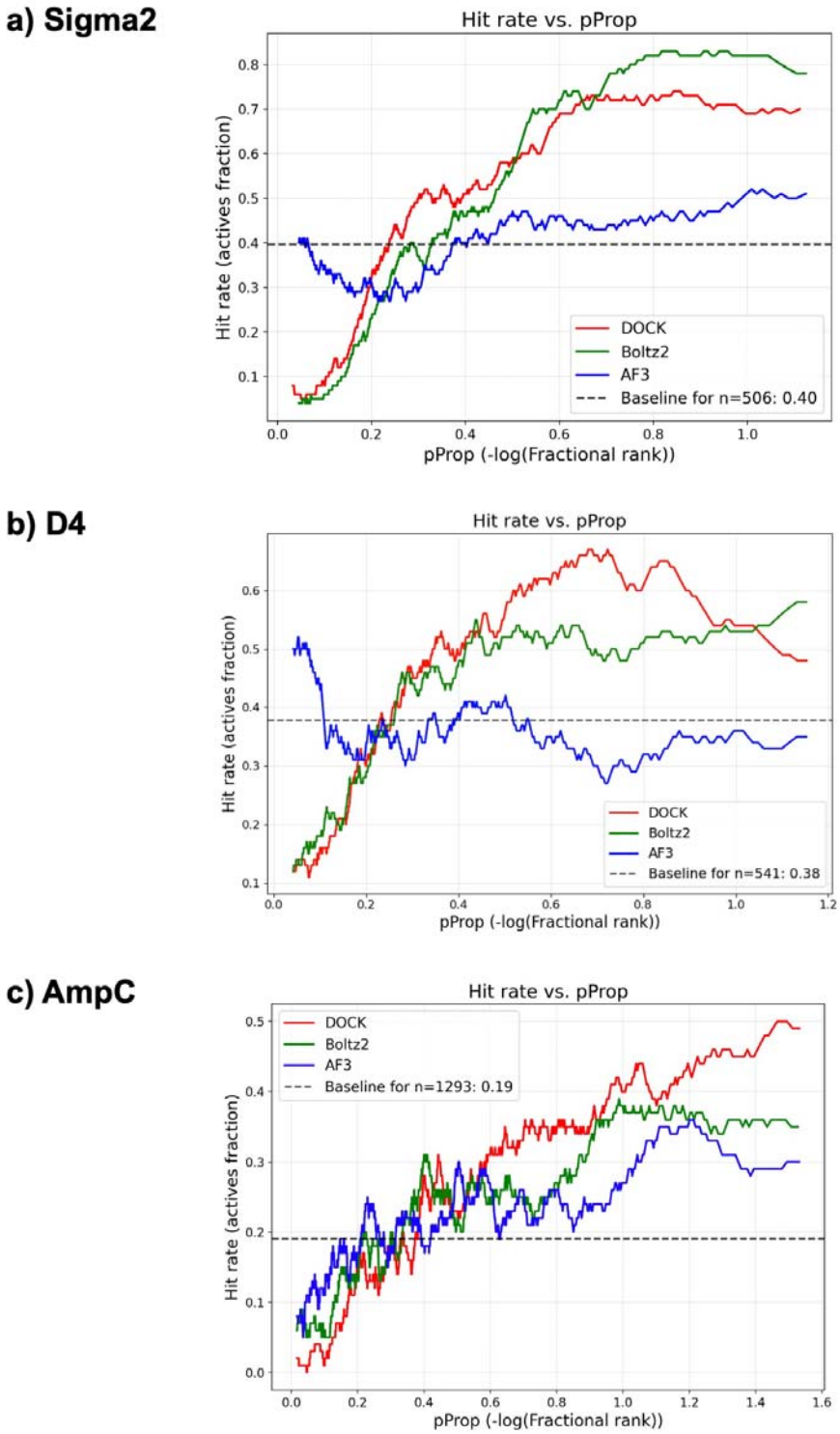
| | | | |
|---|---|---|---|
| **Friedman $\chi^2$** | **18.79 (p = 8.31e-05)** | **0.073 (p = 0.964)** | **0.48 (p = 0.786)** |

**Supplementary Figure9**: **Discriminative power of co-folding and docking scores by correlation against experimentally measured *$pK_i$* across different targets:** a) σ□ (n = 201 actives), b) D4 (n = 205 actives), c) AmpC (n = 247 actives). The leftmost panel is for Boltz-2 $pIC_{50}$ affinities, the middle panel for AF3 L-pLDDT and the rightmost panel for DOCK scores. All these scores are compared against measured *$pK_i$* values. Mean Absolute Errors and Pearson correlation coefficients are calculated. Blue lines show the corrected regression line, black dotted line before the correction, and green horizontal line shows the baseline at an average measured pKi value. Friedman test with Conover post-hoc comparisons and

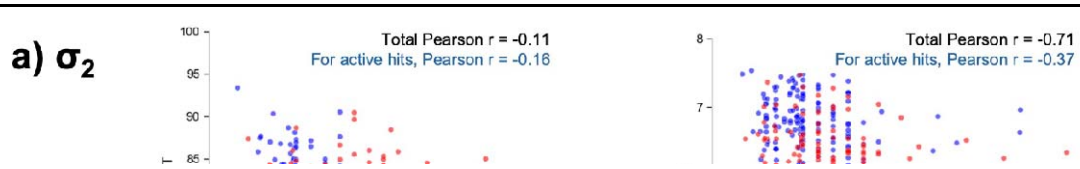
Bonferroni corrections compare MAE between methods for each target system.




**Supplementary Figure10: <u>Different methods of treating randomly assigned $pK_i$ values for docked false positives in σ□, D4, AmpC docked lists</u>.** Mean absolute errors (before linear correction is shown in grey dotted line, after linear correction is the red regression line and blue line indicates the baseline when we predict all values at measured $pK_i$) between the measured $pK_i$ and Boltz-2 $pIC_{50}$ affinity scores when a) all non-hits are assigned a $pK_i$ = 2 x $pK_{i,threshold}$; b) non-hits are randomly assigned a $pK_i$ value between the $pK_{i,threshold}$ and 1. Baseline MAE is only quoted for b), since the error could be misinterpreted when all non-binders' $pK_i$ values are fixed.




**Supplementary Figure11: Hit rate curves for co-folding and docking scores for the three experimental benchmark datasets.** Hit rate curves are plotted over a rolling window (window = 100 for AmpC, σ□ and 50 for D4) after ranking with pProp from the docked hit lists, from three different targets: a) σ□ (201 actives, 305 non-binders), b) Dopamine D4 (205 actives, 336 non-binders), c) AmpC β-lactamase (247 actives, 1,046 non-binders).



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