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Will the chemical probes please stand up?†

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In 2005, the NIH Molecular Libraries Program (MLP) undertook the identification of tool compounds to expand biological insights, now termed small-molecule chemical probes. This inspired other organisations to initiate similar efforts from 2010 onwards. As a central focus of the Probes & Drugs portal (P&D), we have standardised, integrated and compared sets of declared probe compounds harvested from 12 different sources. This turned out to be challenging and revealed unexpected anomalies. Results in this work address key questions including; a) individual and total structure counts, b) overlaps between sources, c) comparisons with selected PubChem sources and d) investigating the probe coverage of druggable targets. In addition, we developed new high-level scoring schemes to filter collections down to probes of higher quality. This generated 548 high-quality chemical probes (HQCP) covering 447 distinct protein targets. This HQCP collection has been added to the P&D portal and will be regularly updated as established sources expand and new ones release data.

Introduction

In 2005, the NIH Molecular Libraries Program (MLP) undertook the first large-scale identification of tool compounds to expand biological insights, now termed small-molecule chemical probes.^{1,2} Their systematic generation against a range of molecular targets was a key driver for the establishment of the PubChem database in order to collate structures and data from the initial ten funded screening centres.³ The concomitant screening compound collection was established as the Molecular Libraries Small Molecule Repository (MLSMR) for which PubChem had hosted 255 000 compounds by the end of 2005 and since expanded to 406 000 by 2015 (but updates have ceased). Although 25 of the 64 early compounds were judged to be of equivocal quality by a crowdsourcing assessment in 2009 (ref. 4) the program progressed to 375 probes (see data section below) before ending in 2014. As conceived from the outset, the availability of these compounds and, crucially, their associated characterisation data, have facilitated the exploration of new targets, pathways and therapeutic hypotheses.⁵ Notwithstanding these success stories, the MLP undertaking has been subject to criticisms that remain relevant to

contemporary efforts. These include considerations of the MLSMR fitness-for-purpose as a library accrued in an academic context (compared to arguably better-resourced pharmaceutical company screening collections), persistent probe quality issues and remaining confusion on exactly how many probe compounds the program generated.^{6–8}

A less tangible but equally important success of the MLP is that, from approximately 2010 onwards, it inspired other organisations to also initiate probe discovery with open dissemination. The three most recent announcements (but not yet surfacing data) are EU-OPENSOURCE⁹ with probe development as one of their main objectives, the EUBOPEN¹⁰ consortium aiming to synthesize at least 100 new chemical probes and the Target 2035 initiative to accrue probes for all human targets by 2035.¹¹ As of June 2021, we were able to collect probe data from the sources listed below.

- MLP probes (NIH screening initiative)
- Structural Genomics Consortium (SGC, 3D structure-based)¹²
- Nathanael Gray Laboratory (cancer research focused)¹³
- Chemical Probes Portal (literature curation, expert opinion)¹⁴
- Pharmaceutical companies (offering in-house compounds)¹⁵
- Probe Miner (data filtration for putative probes)¹⁶
- Probes & Drugs (comprehensive collation of probe data)¹⁷

Detailed descriptions of these sources and their individual approaches to probe development are available from their websites and related publications. These are expanded in a recent review¹⁸ as well as articles in this special issue.

This work was conceived to answer the following questions that can not be answered *via* the individual sources:

- How many declared chemical probe structures are there?

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- What is the distribution of their physicochemical properties?
- What are the differences between experimental and calculated probes?
 - What is their representation in PubChem sources?
 - What are their intersections with each other?
 - What are their individual targets?
 - What is their combined human proteome coverage?

Addressing these questions is specifically enabled by the Probes & Drugs portal (P&D, <https://probes-drugs.org>).¹⁷ P&D was designed as a hub for the integration of high-quality bioactive compound sets enabling their analysis and comparison. As the name indicates, the main focus is on probes and drugs but includes additional relevant sets extracted from, or supplied by, recently published specialist databases (e.g. BiasDB¹⁹), vendor sets, ESI† from papers (e.g. kinase inhibitors) or harvested from publication out-links (e.g. the British Journal of Pharmacology “Concise Guide” series^{20–26}). Other high-quality curated bioactive chemistry sources, including ChEMBL,²⁷ BindingDB,²⁸ Guide To Pharmacology (GtoPdb),²⁹ DrugCentral³⁰ and DrugBank³¹ are utilized for compound biological annotation (the latter 3 are also P&D compound sets). The P&D compound database is currently composed of 69 sources including 12 probe-related, 7 drug compilations, 36 academic (non-commercial) sets and 14 precompiled sets from bioactive compound vendors (suggestions for expansion are welcome).

P&D has additional advantages for this study. An important one is that chemical structures from all sources are standardised after importation. This means that our internal comparisons are as rigorous as we can make them (notwithstanding cheminformatic nuances that preclude this from being perfect). We have used the *standardiser* python package³² for salt-stripping, charge neutralization, standardizing common functional groups, preserving stereochemistry and identification of the main active pharmaceutical ingredients (API) in mixtures. Structural uniqueness is based on the InChIKey, a hashed version of the International Chemical Identifier, InChI.³³ Within the text of this article, we have used InChIKeys to designate mentioned compounds. These can be searched not only against P&D and all major databases but also in Google.³⁴ On our website, users can choose to browse and compare the structures in three different forms: 1) standardized; 2) original (*i.e.* as imported from the source) and 3) non-isomeric (*i.e.* core connectivity without stereochemistry). Importantly, we made the considered decision not to submit our entire content to PubChem for two main reasons. The first is to reduce non-obvious circularity which can confound database users.^{35,36} The second is that this enables informative Boolean query combinations to be made between P&D sets and essentially any PubChem source or filtration selects.

As of version 02.2021 (March 2021), P&D contains 77 130 compounds with 4466 labelled as chemical probes by their sources used for this study. However, these should not be

considered equivalent in a canonical sense because they have been generated by divergent approaches. By analysing the structures and associated metadata, we have tried to establish a high-quality subset based on the probe origin and by the application of scoring schemes. In addition, where the data allow plausible assignments, we have compiled target coverage.

Experimental vs. calculated

We chose to internally partition P&D probes into two main categories; experimental and calculated.

Experimental denotes compounds from published probe characterisation experiments. These papers include profiling data for target modulation potency, selectivity, and possible secondary targets. Also important to note is that the experimental data largely originate from a single laboratory and are thus likely to have more consistent and reproducible data (*e.g.* where intra-laboratory assay variation is controlled *via* sufficient replicates and internal standards). Examples of such experimental probe sets include; bromodomains chemical toolbox,³⁷ Chemical Probes Portal,¹⁴ Gray Laboratory Probes,¹³ Nature Chemical Biology Probes, Open Science Probes,¹⁵ opnMe Portal,³⁸ Protein methyltransferases chemical toolbox,³⁹ and SGC Probes.¹²

We use the term calculated here to denote *in silico* evaluation using the combination of public data with a custom scoring function (we intentionally avoided the term predicted in this context because machine learning methods were not used).^{16,40} The evaluation of probes at scale involves comparing public data at different stringencies according to availability. The seven key criteria are: 1) <100 nM target potency *in vitro*, 2) <1 μM for cell-based assays with evidence of direct target engagement, 3) target selectivity >100-fold, 4) absence of structural alerts indicating chemical liabilities 5) identification of an inactive analogue as control with significantly lower potency or inactive against the primary target,⁴¹ 6) an orthogonal probe with a different chemotype against the same primary target, and 7) SAR data to increase confidence in specific target modulation. The simplest approach to assigning a compound as a probe is to use these criteria (or a subset thereof). However, this can be nuanced by weightings (*e.g.* for potency and selectivity) as well as more complex scoring (*e.g.* functions favouring compounds with a wider range of target profiling data). In contrast to experimental probes, the data for the evaluation of the calculated probes from different sources can be less consistent and may not be acquired with the objective of developing and validating a probe *per se*. The calculated sources included here are from *Probe Miner*¹⁶ and the *tool compound set*.⁴⁰

Source descriptions and counts

Those compared in this study are listed in Table 1 with brief descriptions below.



Table 1 Sources with their compound numbers and probe type. “E” refers to experimental and “C” to calculated probe type. The targets column counts distinct probe-target annotations. The total number of compounds/targets represents a distinct number of compounds/targets for all sets combined

	Set	Probe type	Set class	Compounds	Targets
1	Bromodomains toolbox	E	High-quality	25	26
2	Chemical Probes.org	E	High-quality	362	322
3	Gray Laboratory	E	High-quality	53	56
4	MLP	E	Legacy	375	156
5	Nature Chemical Biology	E	Legacy	58	51
6	Open Science Probes	E	High-quality	83	95
7	opnMe Portal	E	High-quality	55	57
8	Probe Miner	C	Calculated	3187	326
9	Methyltransferases toolbox	E	High-quality	19	20
10	SGC Probes	E	High-quality	81	97
11	Tool compound set	C	Calculated	515	392
12	Historical compounds	E	Historical/obsolete	239	—
	Total		(Not historical)	4466	819

MLP and Nature Chemical Biology Probes

The former has been outlined in the introduction. The latter was extracted from articles published in Nature Chemical Biology (although since 2018 this dedicated section of the Journal is no longer available) As legacy collections, probes from these two sources may lack the stringent characterisation of maintained collections. This is reflected in some cases by a) the lack of controls or orthogonal probes, b) unclear potency and selectivity criteria or c) no target annotation.

SGC Probes, Open Science Probes, opnMe Portal, and Gray Laboratory Probes

These organisations apply the currently accepted probe quality criteria and are maintained sets in that P&D has picked up at least some new compounds since their initial release (even if at a variable frequency).

Bromodomains and protein methyltransferases chemical toolboxes

These compounds were extracted from publications focused on the study of bromodomains and methyltransferases. Except for four bromodomain probes developed elsewhere, these also belong to the *SGC set*.

Tool compound set and Probe Miner

These are calculated selections, mainly from ChEMBL, selected *via* probe-likeness criteria. While the tool compound set is a one-off extraction from the publication, Probe Miner is regularly updated. The tool compound set independently includes more than 100 compounds available from the Chemical Probes Portal at the time of its publication.

Chemical Probes Portal (CP portal)

This provides expert usage recommendations based on publication evaluations from a Scientific Advisory Board (SAB, of which one of us, CS, is a member). While content had languished below 200 compounds for some time, this has recently expanded to 362. Importantly, this portal also

lists historical probes (these are also captured as a P&D set, see next section).

Historical compounds

The use of these obsolete compounds is no longer recommended by the CP portal because new data indicates promiscuity or displacement by better tool compounds.⁴² These structures include the well-known and notorious medicinal chemistry time-wasters of staurosporine (HKSZLNNOFSGOKW-FYTWVXJKSA-N), quercetin (REFJWTPEDVJJIY-UHFFFAOYSA-N), resveratrol (LUKBXSAWLPMMSZ-OWOJBTEDSA-N), and curcumin (VFLDPWHFBUODDF-FCXRPKNKRS-N).

Set comparisons

The sets are compared by exact matches in Table 2. The resulting matrix is unique in that no individual source has published a comparable analysis. However, some results were unexpected. The first surprise was the low intersection between MLP and the calculated sets. We attribute this to the 80 compounds from the MLP set without bioactivity data on P&D. In addition, 203 have a primary target potency above the 100 nM threshold. The unexpectedly high overlap between the Chemical Probes Portal and tool compound sets is a consequence of the (already mentioned) inclusion of the former in the latter but without data-supported evaluation. Also surprising is the low overlap between the two calculated sets, even though the data sources and selection criteria were conceptually similar. However, the main goal of the tool compound set was to select effective agonists or antagonists with a high stringency for target selection and cell potency.

Another surprising observation was that, while overlap with historical compounds is reassuringly low, Table 2 indicates there are still nine of these undesirables in Probe Miner, four in the MLP, and two in the high-quality SGC Probes. The first of these, bromosporine⁴³ (UYBRROMMFMPJAN-UHFFFAOYSA-N), was designed to be a pan-bromodomain inhibitor and could usefully be family-selective. The second, GSK-J1 (ref. 44) (AVZCPICCWKMZDT-UHFFFAOYSA-N), an inhibitor of the KDM protein family is



Table 2 A matrix showing the intersections between 12 sources. This was computed using the InChIKey exact match for the standardised structures from the P&D portal. The diagonal figures in white represent the source counts in Table 1

	Set	1	2	3	4	5	6	7	8	9	10	11	12
1	Bromodomains toolbox	25	16	0	0	0	0	1	0	0	21	10	0
2	Chemical Probes.org	16	362	17	2	13	24	10	25	13	43	114	0
3	Gray Laboratory	0	17	53	0	2	0	0	1	0	0	7	0
4	MLP	0	2	0	375	3	0	0	4	0	0	4	4
5	Nature Chemical Biology	0	13	2	3	58	1	0	0	1	4	9	1
6	Open Science Probes	0	24	0	0	1	83	12	2	0	5	0	0
7	opnMe Portal	1	10	0	0	0	12	55	1	0	3	2	0
8	Probe Miner	0	25	1	4	0	2	1	3187	1	2	32	9
9	Methyltransferases toolbox	0	13	0	0	1	0	0	1	19	19	11	0
10	SGC Probes	21	43	0	0	4	5	3	2	19	81	26	2
11	Tool Compound Set	10	114	7	4	9	0	2	32	11	26	515	1
12	Historical Compounds	0	0	0	4	1	0	0	9	0	2	1	239

not cell-permeable. The SGC Probes resource has noted this and consequently now recommends a pro-drug of GSK-J1, GSK-J4 (WBKCKEHGXNWYMO-UHFFFAOYSA-N) for cell-based assays.

Dataset compilation

Merging individual sets resulted in 4466 structurally distinct probe compounds (*i.e.* as unique InChIKeys). This includes 940 (21%) experimental plus 3670 (82.2%) calculated probes including 3178 from Probe Miner. The overlap of 143 (3.2%) compounds is mainly due to the inclusion of the Chemical Probes Portal in the tool compound set. The full 4466 set includes 275 labelled as drugs (reported to be in clinical phases) with 103 labelled as approved by FDA, EMA and other agencies. The full set also includes 29 PROTACs (Proteolysis Targeting Chimeras) from PROTAC-DB⁴⁵ and Chemical Probes Portal, 60 covalent binders from CovalentInDB,⁴⁶ and 21 biased GPCR ligands from BiasDB.¹⁹

Our analysis also established that 132 compounds were flagged with one or more structural alerts from either a) PAINS filters,^{47,48} b) aggregators,⁴⁹ c) cellular assay nuisance compounds⁵⁰ or d) historical compounds. Of the 60 stereoisomers, 54 originate from the Probe Miner set. Compared to small-molecule approved drugs extracted from ChEMBL (as a set in P&D) probes are generally larger and more complex (Fig. 1).

This correlates with higher target selectivity that, in turn, is reflected in the number of associated targets (Fig. 2). However, these values could be biased by approved drugs accumulating more cross-screening data and hence a wider range of secondary targets.

Target mapping

The majority of probes have primary targets specified in their sources. In most cases these are supported by quantitative



Fig. 1 Physico-chemical properties distribution (top: molecular weight, middle: calculated logP, bottom: TPSA) for experimental probes (blue), calculated probes (orange) and small-molecule approved drugs set from ChEMBL (green) (x: property range, y: compounds percentage). The properties were calculated by RDKit.





Fig. 2 The number of associated targets for probes (blue) and the small-molecule approved drugs set from ChEMBL (green) (x: axis number of targets, y: compounds percentage). Only compounds with at least one associated target were included encompassing 4257 probes and 2005 approved drugs.

in vitro binding data (e.g. K_i , IC_{50} or K_d). Some may also have secondary targets with a data-supported potency below that against the primary target (we have avoided using the term “off-target” since there are few cases where secondary targets have been mechanistically assigned as a side-effect or toxicity liabilities). The 132 probes without primary target annotation were, in most cases, directed against viruses, bacteria, cell lines or pathways. They also predominantly belonged to the MLP and Nature Chemical Biology legacy sets. For the remainder, we collated 819 single and multi-component protein targets with 549 for both experimental and calculated with an overlap of 279. In total these constituted 807 distinct single protein identifiers (*i.e.* UniProt IDs⁵¹), 544 for experimental and 535 for calculated probes with 272 in-common. The human Swiss-Prot target count was 796.

In practice, the number of protein targets is below 819, since multi-component targets may be variably annotated against either a protein subunit, the complex target, or both. For example, probes directed against the BCR-ABL1 fusion protein may be annotated with the fusion protein (of which

there are several length forms in TrEMBL but not Swiss-Prot) or ABL1 (P00519) or, in the case of ~20 probes from the Probe Miner set, BCR (P11274). Other complicating examples are the cyclin-dependent kinases (CDKs), where the probes may be annotated by sources with one of the human CDKs, a CDK in complex with a specific cyclin, or both.

The highest probe target numbers (5 experimental and 225 calculated) have been assigned against mTOR (MTOR, P42345). Next in rank are histone deacetylase 1 (HDAC1, Q13547) with 2/161 experimental/calculated, epidermal growth factor receptor (EGFR, P00533) with 7/103 and estrogen receptor (ESR1, P03372) with 1/92. The larger number for calculated probes reflects their more frequent origin from panel screening papers and consequent higher average compounds: target ratio of 9.8 for the Probe Miner set compared to ~1.0 for experimental probes. For these, the highest assigned target numbers are BRD4 (O60885) and the BRD3/BRD2 (Q15059/P25440) subfamily pair with 18 and 15 probes, respectively. This is a consequence of the declared SGC Probes focus on epigenetic regulators. The family distribution of all annotated targets is shown in Fig. 3.

The differences in Fig. 3 reflect inherent biases. For example, predicted probes have almost double the number of GPCRs⁵² but, compared to predicted probes, proportionally fewer kinases and epigenetic regulators. This is likely to be due to the challenges of optimising single-target selective ligands within these target families.^{37,53} However, probes with intra-family selectivity can also be experimentally useful but run the risk of being rejected by quality scoring weighted towards single-target selectivity.

Target intersections in UniProt

Having assigned target IDs to both probe sets we compared these with informative cross-references in UniProt.⁵¹ We selected 4213 protein IDs (as human Swiss-Prot entries) based on the union (OR operator) of the four high-quality

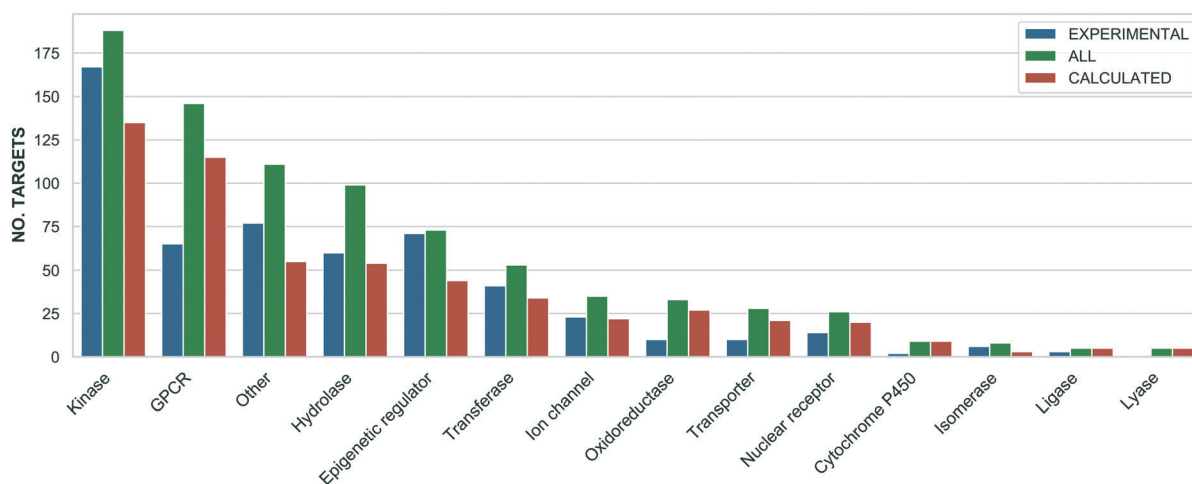


Fig. 3 A bar chart showing the target families distribution separately for all (green), experimental (blue) and calculated (orange) probes (x: target family, y: number of targets). The assignments are based on the ChEMBL and Guide to Pharmacology target classification.



curated chemistry-to-target databases (already mentioned) of ChEMBL, BindingDB, GtoPdb and DrugCentral. These IDs represent liganded targets for 21% of the UniProt proteome of 20 395 (although this drops to 19 205 for HUGO Gene Nomenclature Committee annotation). The comparative protein sets we also selected were from the four target development levels (TDLs) of the Pharos resource for Illuminating the Druggable Genome (IDG).⁵⁴ This facilitates exploration of both the characterised and the understudied (or “dark”) regions of the human proteome with a view to expanding functional insights and finding new drug targets. We initially selected the combination of the Tchem⁵⁴ (1593 proteins) known to bind small molecules (other than approved drugs) with target-class specific potency thresholds plus the Tclin⁵⁴ (659 proteins) as targets of approved drugs. The union of these two is 2221 proteins. The intersections of these four lists are shown in Fig. 4.

The two notable features are:

1. Probes have activity against 53 proteins not in Tclin or Tchem.
2. The Swiss-Prot liganded proteome includes 743 probe targets.

Analysis with other TDL sets established that of the 659 approved targets 265 were also covered by probes. However, there were no intersections between probes and the 6368 Tdark proteins. This implies that the current probes may have already expanded Tchem but there is no overlap with dark targets.

Source errors

Despite curatorial diligence, low levels of annotation errors, including the transitive inheritance of author mistakes extracted from papers, inevitably creep into the bioactivity databases we have used as sources.⁵⁵ During manual cross-checking we identified the following error types affecting approximately 40 probe entries:

- Substitution of a biochemical for a cell-based assay as well as *vice versa* cases (the most common problem).

- Concentration unit errors for secondary target activity (*i.e.* the compound was thus not selective).
- Erroneous target annotations that falsely indicate potent secondary target activity.
- Potency values assigned to only a subunit in a multi-component target (*i.e.* thus technically without supporting bioactivity data).
- Some sources had incorrectly assigned human TrEMBL partial sequence entries as targets rather than the human Swiss-Prot IDs (although only three cases were found).

As an operating principle P&D fixes any unequivocal errors we spot. At the same time, we notify the originating sources about these errors that could otherwise persist and proliferate between databases. However, we have found the speed with which these are fixed at source has been variable (although this is clearly dependent on build cycle times and release versions).

Probe scoring schemes

We optimised four different scoring schemes to support users for probe triage and selection. As explained, the Probe Miner (PMIS) and P&D probe-likeness scores (PDPS) are data-supported. The other two scores are expert opinion-based and thus more subjective. These are abstracted from the Chemical Probes Portal rating for use in cells (CPOC) and in organisms (CPOO). These represent a summed rating of chemical properties, primary targets, secondary targets and in most cases, expert judgments. The conceptual difference with data-supported scores is that these are calculated for compound-target pairs. Thus, a single compound can have multiple scores against each of its assigned targets. Users may thus select the most suitable of these pairings but the probes can also be used for intra- or inter-family selectivity. For comparability, all scores are normalized to between 0 and 100%.

The PMIS, ranging from 0 to 1, combines partial scores for 1) potency, 2) selectivity 3) activity in cells, 4) SAR data 5) availability of an inactive analogue and 6) a PAINS score. Nevertheless, probe suitability is not prescribed by the score value but by so-called minimum quality criteria. These include 100 nM potency in biochemical assays, 10-fold target selectivity, and 10 μ M potency in cell-based assays (but not necessarily evidence of intra-cellular primary target engagement). The Probe Miner set of 3187 compounds meeting these criteria thus have a PMIS between 0.38 and 0.85. A more detailed description is given in the Probe Miner publication.¹⁶

The PDPS, scaled from 0 to 1, incorporates partial scores in common with PMIS but adds in orthogonal probes. The comparison of both scoring schemes is shown in Table 3. Unlike the Probe Miner selectivity score, P&D also highlights target sub-family selectivity beyond just single proteins. The probe-likeness of a compound is closely related to the PDPS value. Each compound with a score above 0.7 is labelled as P&D-approved based on the available data. The score is capped to not exceed 0.7 unless it passes all three core criteria (*i.e.*, *in vitro* potency, cell potency and selectivity).



Fig. 4 Venn diagram of P&D targets against selected human UniProt cross-references and two Pharos TDLs.



Table 3 Comparison of PMIS and PDPS. For parameters with a defined range, the score is 0% for values below the minimum and 100% for values greater than the maximum. Within this range, there is a linear relationship between the value and the score

Parameter	PMIS value range (weight) <i>note</i>	PDPS value range (weight) <i>note</i>
Potency (biochemical)	5–10 [–log(<i>M</i>)] (4)	6.5–7 [–log(<i>M</i>)] (2)
Selectivity	Complex selectivity score normalized per target (8)	10–30-Fold (2)
Potency (cell-based)	5 [–log(<i>M</i>)] (2) <i>without the evidence of primary target engagement</i>	5.5–6 [–log(<i>M</i>)] (2) <i>with the evidence of primary target engagement</i>
Inactive analogue	Binary (1)	Binary (1)
Orthogonal probe	—	Binary (1)
SAR	Binary (1)	—
Structural alert	Binary (1) <i>PAINS</i>	Binary (1 and –3 for historical compounds) <i>PAINS, aggregators + other nuisance compounds in cellular assays, historical compounds</i>
Probe-likeness determination	Independent of the score value, compounds labelled as possible suitable probes if they meet minimum quality criteria (100 nM potency, 10 μM cell potency, 10-fold selectivity)	Compounds labelled as P&D approved for PDPS >70%
Probe-like compounds count	3187	1109

Compounds labelled as historical are down-weighted by subtracting 0.3 and thus cannot be labelled as P&D-approved. Currently, there are 1109 probes labelled as P&D approved. More details on this are given on the P&D FAQ page.⁵⁶ The CPOC and CPOO scores (from 0 to 4 stars) are based on Scientific Advisory Board (SAB) reviews. These may be accompanied by comments and usage recommendations. However, there is currently a review backlog in that out of 362 compounds, 274 have been rated for use in cells and 225 for use in model organisms.

In Table 4, we review three examples of compounds with assigned probe scores. The first is a selective RIPK1 inhibitor, *GSK2982772* (ref. 57) (LYPAFUINURXJSG-AWEZNQCLSA-N), highly scored by all three probe sources. The second is BET family bromodomain inhibitor from SGC Probes, *JQ-1* (ref. 58) (DNVXATUJJDPFDM-KRWDZBQOSA-N), scored highly by P&D and Chemical Probes Portal, but as a family-selective probe with lower PMIS. The latter is the only P&D approved

probe with low ratings from the Chemical Probes Portal (not scored by Probe Miner), *AGI-5198* (ref. 59) (FNYGWXSATBUBER-UHFFFAOYSA-N), was proposed as a prototypical IDH-1 R132H inhibitor. However, this was surpassed in potency and characterization details by the more recent *GSK864* (ref. 60) (DUCNNEYLFQFSW-PMERELPUSA-N) which also has an inactive control for proof of target involvement. However, as a second, distinct chemotype, this probe could be used for corroborative phenotypic assays.

Score comparisons

To extend our systematic comparison of scoring, we introduced quality thresholds. PDPS was set at 70% as used for P&D approved probes. For CPOC and CPOO, we raised this to 75% (equivalent to 3 out of 4 stars in the original rating system). For the PMIS, there is no clear threshold and

Table 4 Three selected chemical probes with assigned probe scores from P&D, Probe Miner and Chemical Probes Portal

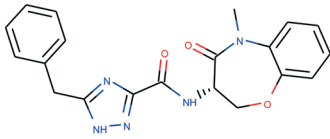
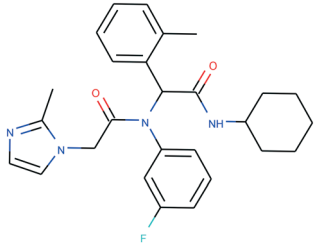
			
Name	GSK2982772	JQ-1	AGI-5198
PDPS	100%	100%	86%
PMIS	70% (<i>in Probe Miner set</i>)	48% (<i>not in Probe Miner set</i>)	—
CPOC	100%	100%	50%
CPOO	83%	75%	42%



Table 5 Matrix showing the intersections between six different probe scores and probe types. This was computed using the InChIKey exact match for the standardised structures from the P&D portal

	Set	1	2	3	4	5	6
1	CPOC [$\geq 75\%$]	206	106	55	10	203	86
2	CPOO [$\geq 75\%$]	106	108	28	6	107	43
3	PDPS [$\geq 70\%$]	55	28	1109	265	150	1013
4	PROBE MINER AND PMIS [$\geq 60\%$]	10	6	265	1282	20	1282
5	Experimental probe	203	107	150	20	940	144
6	Calculated probe	86	43	1013	1282	144	3670

since the highest value is 85%, a setting of 75% would leave only 54 compounds from more than 3000. We thus chose to set the Probe Miner threshold at 60%, thus leaving 1282 compounds, a similar number to P&D approved probes.

The comparison between different scoring schemes (Table 5) highlights differences between judgment-based and calculated scores. One of the reasons for these differences is data incompleteness, for example where affinity data is only for the presumed primary target thereby precluding selectivity assessment.

Even the experimental probes included 195 compounds without bioactivity data. In addition, we found 142 compounds annotated against single targets without selectivity data. The union of these represents 36% of the experimental probes that cannot thus be properly scored. Another reason for differences arises between stringent criteria-based evaluation and expert judgment.

We also found differences between calculated scores for the 265 compounds-in-common between the 1109 (P&D) and 1282 (Probe Miner) sets. This could be attributed to the differences in the scoring methodology but also the associated data (*i.e.* not all experimental probes have PMIS). While Probe Miner currently employs bioactivity data from ChEMBL and BindingDB, P&D uses more recent versions and complements these with smaller data sources such as GtoPdb. This is reflected in a high P&D score for 150 experimental probes (with 55 highly-rated by CPOC) while Probe Miner detects 20 (including 10 based on the CPOC).

High-quality chemical probes set

As an outcome of this study, we have compiled a high-quality chemical probes subset (HQCP). We have used the PDPS for the addition of P&D approved experimental probes plus those P&D approved calculated probes that are in at least one established tool compound set. We have thus partitioned four compound sets from P&D:

1. *Concise Guide to Pharmacology 2019/20* is a set extracted from a biennial series of publications providing concise overviews of the key properties of ~1800 human drug targets with an emphasis on selective pharmacology.^{20–26}

2. *Kinase chemogenomics set* is a collection of narrow-spectrum small molecule kinase inhibitors assembled by the SGC-UNC to study the biology of dark kinases. This is the most diverse and highly annotated public collection of kinase inhibitors.⁶¹

3. *Kinase inhibitors* were extracted from a series of Molecular Cell papers by Wang and Gray summarising recently-reported kinase inhibitors.^{62,63}

4. *Novartis Chemogenomic Library – NIBR MoA Box* was compiled *via* data mining and institutional crowdsourcing. It is regularly updated and used widely both within Novartis and by their external collaborators.⁶⁴

We used the quality criteria in Table 6 to select 548 probes for HQCP (451 experimental, 208 calculated with 114 in common). The intersections are shown in Table 7 including the EU-OPENSOURCE Bioactive Compound Library and Drug Repurposing Hub⁶⁵ set. As non-commercial bioactive libraries, these are included in P&D as relevant for probe research.

The HQCP set contains 42 approved drugs with 102 clinical candidates, 27 PROTACs, 15 covalent binders and 10 compounds tagged with a structural alert (four for aggregation and six for PAINS). The overlap between HQCP and the calculated sets is largest for the P&D approved probes with 244 out of 1109 compounds, but these were also partly used for the HQCP selection. From the complete Probe Miner set (*i.e.* without the PMIS threshold applied), there are 66 compounds from nearly 3200 meeting the Probe Miner minimum quality criteria. On the other hand, there are 153 compounds from 515 in the tool compound set, mainly from the inclusion of the Chemical Probes Portal compounds in the tool compound set.

The intersections between the three bioactive screening libraries (Novartis Chemogenetic Library, EU-OPENSOURCE

Table 6 The criteria used for the selection of HQCP. The count column contains the number of compounds matched by the criterion. The total number represents the union of all criteria

Criterion	Count
1 Belong to one of the high-quality probe sets (except Chemical Probes Portal)	256
2 CPOC or CPOO score at least 75% (<i>i.e.</i> three out of four stars in the original Chemical Probes Portal rating system)	208
3 P&D approved experimental probes	150
4 P&D approved probes belonging to one of the non-commercial high-quality sets (<i>Concise Guide To Pharmacology, Kinase Chemogenomic Set, Kinase Inhibitors, and Novartis Chemogenetic Library</i>)	177
5 Not labelled as a historical compound	–2
Total	548



	Set	1	2	3	4	5	6	7	8	9	10
1	HQCP	548	451	208	244	66	153	103	210	185	186
2	Experimental probes	451	940	144	150	34	118	81	217	235	221
3	Calculated probes	208	144	3670	1013	3187	515	138	210	222	279
4	P&D approved	244	150	1013	1109	702	331	83	127	129	156
5	Probe Miner	66	34	3187	702	3187	32	67	93	99	148
6	Tool Compound Set	153	118	515	331	32	515	77	131	132	144
7	Concise Guide to Pharmacology	103	81	138	83	67	77	2536	637	622	1065
8	Novartis Chemogenetic Library	210	217	210	127	93	131	637	4185	743	1274
9	EU-OPENSOURCE Library	185	235	222	129	99	132	622	743	2464	1401
10	Drug Repurposing Hub	186	221	279	156	148	144	1065	1274	1401	6764

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