Perspectives on Codebook: sequence specificity of uncharacterized human transcription factors

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SUMMARY

We describe an effort ("Codebook") to determine the sequence specificity of 332 putative and largely uncharacterized human transcription factors (TFs), as well as 61 control TFs. Nearly 5,000 independent experiments across multiple *in vitro* and *in vivo* assays produced motifs for just over half of the putative TFs analyzed (177, or 53%), of which most are unique to a single TF. The data highlight the extensive contribution of transposable elements to TF evolution, both in *cis* and *trans*, and identify tens of thousands of conserved, base-level binding sites in the human genome. The use of multiple assays provides an unprecedented opportunity to benchmark and analyze TF sequence specificity, function, and evolution, as further explored in accompanying manuscripts. 1,421 human TFs are now associated with a DNA binding motif. Extrapolation from the Codebook benchmarking, however, suggests that many of the currently known binding motifs for well-studied TFs may inaccurately describe the TF's true sequence preferences.

KEYWORDS: Transcription factor, TF, ChIP-seq, HT-SELEX, GHT-SELEX, SELEX, SMiLE-seq, Motif, DNA-binding specificity, PWM, PBM, Codebook

Introduction and motivations

The human genome encodes >1,600 putative transcription factors (TFs), defined as proteins that bind specific DNA sequence motifs and regulate gene expression[1]. These DNA binding motifs are most commonly modelled as a Position Weight Matrix (PWM) that describes the relative preference of the TF for each nucleotide base pair in the binding site[2,3], and can be visualized as a sequence logo[4]. Several hundred putative human TFs still lack DNA binding motifs[1], and even for well-characterized TFs, it remains controversial whether the reported motif model is accurate[5,6], and to what degree the TF's sequence specificity contributes to binding site selection in living cells[7,8]. These uncertainties are due in part to the fact that different methods for measuring TF binding, and for deriving PWMs from these data, can have different inherent limitations and biases[2]. Such shortcomings represent fundamental hurdles for the analysis of gene regulation, as well as a myriad of related tasks in genome analysis, including the interpretation of conserved genomic elements and sequence variants, or genetic engineering such as synthetic enhancer design.

To address these issues, we analyzed a large majority of the as-yet uncharacterized human TFs[1], as well as several dozen previously studied control TFs[9,10], using a panel of assays that provide different perspectives on DNA sequence specificity. This unprecedented effort generated what we believe is the largest uniform data structure of its kind. We refer to this international collaborative project as the "Codebook/GRECO-BIT Collaboration": the reagent set and laboratory experiments were initiated as the "Codebook Project", alluding to the fact that TFs decode individual "words" in the genome, and the existing Gene REgulation COnsortium Benchmarking IniTiative, GRECO-BIT, was then engaged for much of the data analysis and benchmarking.

In this paper, we present an overview of the data collection and its analysis, the resulting data, several major outcomes and findings of the Codebook study, and examples of prevalent phenomena and applications. We also introduce web resources that can be used to access the primary and processed data, including the PWMs. Accompanying manuscripts provide greater depth regarding biological findings, new assays, and intriguing TF families, as well as methods for identifying binding patterns (i.e. PWM derivation) and PWM benchmarking (**Table S1**).

Codebook reagents, assays, and data structure

Figure 1 provides a schematic of the Codebook project. We chose 332 putative TFs (i.e., "Codebook TFs") (**Table S2**) for study by starting with a previously described list of 427 hand-curated "likely" human TFs that lacked known motifs or any large-scale DNA binding data[1]. We removed 95 C2H2 zinc finger (C2H2-zf) proteins for which we were already aware of unpublished data (mainly from our prior collaboration with ENCODE[11]). As of June 2024, most of these putative TFs still lack motifs, outside of the Codebook study: of the 332, only 107 have PWMs on Factorbook[12] and/or HOCOMOCO[13]. Many of these motifs appear to be simple repeats, or common cofactor motifs (such as CTCF, REST, and CRE sites) (examples in **Figure S1**), but among the 107, 59 have at least one PWM that appears plausible for representing specificity of the TF (see below).

136 Among the 332 Codebook TFs, 180 contain C2H2-zf domains, while another 103
137 contain another type of well-known DNA-binding domain (DBD). Forty-nine did not have
138 an established DBD at the outset of the study; these were mainly identified as
139 sequence-specific in studies of individual proteins or regulatory sites[1]. We
140 simultaneously analyzed 61 control TFs, encompassing 29 well-characterized TFs
141 representing diverse human DBD classes[9], and an additional 32 C2H2-zf proteins for
142 which published ChIP-seq data were available and had led to a binding motif[10]. For
143 these controls, we incorporated the published SMiLE-seq and ChIP-seq data, rather
144 than repeating the experiments.

145 To study the 332 Codebook proteins, we manually designed 716 protein-coding inserts,
146 corresponding to full-length coding regions of the dominant isoform, and one or more
147 DBDs (or subsets of C2H2-zf domain arrays), if there was a known DBD (**Table S3**).
148 We employed up to three different expression vectors for each insert, as required for the
149 different assays in **Figure 1**, resulting in a total of 1352 new distinct constructs (**Table**
150 **S4**). One of the assays, GHT-SELEX (Genomic high-throughput SELEX), is a new
151 variant of HT-SELEX which is performed with fragmented genomic DNA. As described
152 in the accompanying manuscript[14], GHT-SELEX yields peaks, analogous and often in
153 agreement with ChIP-seq. GHT-SELEX thus provides a new perspective that bridges *in*
154 *vitro* and *in vivo* DNA binding. HT-SELEX and GHT-SELEX were performed with
155 multiple protein sources (mammalian cell extracts, and two different systems for *in vitro*
156 transcription/translation) whereas SMiLE-seq and PBMs were performed with only one
157 protein source. Multiple replicates were performed in many cases, for all assays.

158 The full Codebook data structure is composed of a total of 4,873 technically successful
159 experiments (i.e. they produced data that could be analyzed by at least some
160 subsequent processes) (**Table S5**), The Codebook data structure, experimental
161 information, and PWMs (see below) are accessible at multiple sources (see **Data**
162 **Availability**). Each experiment corresponds to one of the Codebook constructs (or one
163 of the control constructs), analyzed using one of the assays, with one of the protein
164 sources. Not every protein or every insert was analyzed in every assay, by design. For
165 example, the ChIP-seq data only utilize full-length proteins, while Protein Binding
166 Microarray data include only DBD constructs. Long human C2H2-zf domain arrays
167 typically fail in PBMs, and such experiments were omitted. We note that, in general,
168 experiments that are technically successful may not yield motifs that are specific to the
169 TF assessed and supported by other data types (see below). For example, ChIP-seq
170 can detect both indirect and non-sequence-specific DNA binding, as we explored
171 separately[15]. We also emphasize that the *in vitro* assays described here were
172 conducted with unmethylated DNA. We explored the sensitivity of a subset (79) of the
173 Codebook TFs to DNA methylation in an accompanying study, however, which
174 introduces the methylation-sensitive SMiLE-seq variant (meSMS)[16]. DNA binding
175 interactions of 17 of the 79 were impacted by methylation, encompassing inhibition (10)
176 and increased binding or alternative binding sites (7); these data were not incorporated
177 in the analyses described herein.

178 **Motifs are obtained from most C2H2-zf proteins, and half of those containing**
179 **other DBD classes, but only a few proteins with previously unknown DNA binding**
180 **domains**

181 We next derived and examined motifs as PWMs for all the experiments in a semi-
182 automated expert curation format, to identify "approved" experiments (i.e. experiments
183 that contained clear enrichment of credible binding motifs (see **Methods**)). This effort is
184 described in detail in a separate manuscript that describes motif benchmarking, data
185 sets, and success measures[17], and also introduces a web resource that makes all of the
186 motifs available for browsing and download. Briefly, our primary approach was to ask
187 whether similar motifs were obtained for the same protein from different assays and
188 whether the PWMs scored highly by a panel of criteria, including predictive capacity in
189 other data types (depicted schematically in **Figure 1**, *bottom left*), adapting a previously
190 described motif benchmarking framework[18]. To increase our ability to derive motifs that
191 would score highly across data sets, we employed ten motif discovery tools, ranging
192 from the widely used MEME suite[19] to approaches based on machine learning or
193 biophysical modeling, such as ExplaiNN[20] and ProBound[21], thus producing hundreds of
194 motifs per TF. In total, 177 Codebook TFs were associated with "approved" datasets
195 (**Figure 1,** *bottom right*), and a total of 1,072 experiments associated with these 177
196 TFs were approved (**Tables S2** and **S5**). 59/61 controls were also approved, suggesting
197 a low per-TF false-negative rate.

198 The 177 Codebook TFs for which there are approved experiments are dominated by the
199 C2H2-zf domain class, for which 67% (121/180) had approved experiments. These
200 proteins typically contain an array of C2H2-zf domains that bind DNA in tandem[22].
201 Some C2H2-zf domains can bind RNA, protein, or other ligands[23-25]. The Codebook
202 outcome indicates that most C2H2-zf proteins are indeed DNA-binding, although it does
203 not rule out their other activities. Experiments for roughly half (50/103, or 49%) of
204 Codebook TFs in other established DBD classes were also successful. Lack of
205 approved experiments for a putative TF could represent false negatives, which could
206 arise from lack of an obligate binding partner, a requirement for epigenetically modified
207 DNA, lack of requisite post-translational modification in our experiments, or limitations of
208 the methods. Alternatively, they could represent true negatives which are not
209 unexpected; some *bona fide* DBD classes are known to have subtypes that lack
210 sequence specificity (e.g. HMG[26]). Among the Codebook proteins lacking a well-
211 established DBD, only 6/49 (12%) yielded approved experiments (and thus motifs)
212 (discussed in more detail below), suggesting that many of them may indeed lack
213 sequence specificity.

214 We emphasize that our approval process was intentionally conservative, and many
215 experiments were not approved despite being informative in some way (e.g. ChIP-seq
216 yielding reproducible peaks, but no motif, which could indicate indirect association
217 through other TFs or chromatin binding; these are explored in an accompanying
218 manuscript[15]). We also note that our success criteria assume that the sequence
219 preferences of TFs can be represented by PWMs. It is conceivable that uncharacterized
220 TFs could instead recognize interspersed sequence patterns or other features of the
221 DNA sequence that are not readily captured by PWM models or short k-mers.

Diversity and complexity among Codebook TF motifs

To gain an overview of the Codebook TF motifs, and to generate a representative PWM set, we next used expert curation to select a single PWM that is (i) high-performing among all "approved" experiments[17] (see **Methods**), (ii) representative of other high-performing PWMs for the same TF, (iii) consistent with expectation for the class of TF (e.g. the C2H2-zf "recognition code"[27]), and (iv) high information content (IC) (i.e. with a "tall" sequence logo), provided it does not compromise PWM performance. The PWM selected in this process is typically not the highest scoring by criterion (i) alone, as our extensive process typically generated dozens of high-performing PWMs from which to choose, for approved experiments[17]. **Table S6** shows sequence logos for these curated PWMs and their properties; the PWM IDs are given in **Table S2**, and all PWMs can be downloaded (see **Data Availability**). Notably, no data type or motif derivation method stood out as highly preferred by the curators, who were blinded to the source (i.e. data type and derivation method for the PWMs).

Figure 2 shows an overview of similarity[28] among the curated PWMs. Small clusters along the diagonal mostly correspond to the handful of paralogs analyzed (e.g. TIGD4 and 5, SP140 and SP140L, DACH1 and 2, CAMTA1 and 2, and ZXDA, B, and C). In the middle of **Figure 2** is a set of eight TFs that mainly bind CG dinucleotides, leading to similarity in DNA-binding, and in the lower right is a group of five AT-hook proteins that have similar preferences to A/T containing sequences. Most of the Codebook TF PWMs are unlike each other, however, and display a low similarity to any other known PWM[17] (examples are shown in **Figure 2**). This result is partly explained by the large number of C2H2-zf proteins, which are known to differ in their DNA-contacting "specificity residues"[29]. Regardless, a large majority of the Codebook TF motifs are apparently new, and all previous analyses in human regulatory genomics would have been unaware of the ~150 visibly distinct, curated motifs described here.

For dozens of TFs, the curated PWM had a degenerate appearance, i.e. there are few or no positions at which a specific base is absolutely required. Indeed, for fifty-two of them, no individual base at any position achieved a bit score of $\geq$1.4 in the curated PWM (equivalent to roughly >10% of aligned binding sites having a variant base at that position) (**Figure S2A**). Systematically increasing the information content (IC) (i.e., "unflattening" the sequence logo, and increasing the specificity) of the low-IC curated PWMs almost universally reduced performance (**Figure S2B,C**), indicating that the degeneracy is required for accuracy. We also found that, overall, IC is not predictive of motif performance in the benchmarking effort[17]. It is counterintuitive that degeneracy (i.e. lower inherent specificity) would lead to better predictive capacity, but we note that similar findings by others support the validity of the result[30-32].

We propose several explanations for this observation. First, lower IC tends to make affinity distributions across all possible k-mers less digital (i.e. it removes all-or-nothing dependence on specific base positions), which could facilitate the gradual evolution of *cis*-regulatory sequences. Second, homomeric binding (possibly via "avidity"[33]), which a body of evidence suggests is a widespread mechanism[14,34], should reduce reliance on optimal specificity to a single binding site, and strong binding sites may evolve more

265 readily if weak binding sites tend to occur more frequently (and are selected). Third,
266 motif degeneracy may be a consequence of forcing a single PWM to represent the
267 specificity of TFs that, in reality, recognize multiple related motifs. For example, the
268 dependency of binding energy on both enthalpy and entropy can lead to two distinct
269 sequence optima[35]; in another example, different spacings of bZIP half-sites cannot be
270 represented by a single PWM[36]. Consistent with this last possibility, the accompanying
271 manuscript[17] finds that combining multiple PWMs (by Random Forests) typically
272 produces models that are more accurate across platforms, relative to any single PWM.

273 The C2H2-zf proteins present a special case in which a single TF might be anticipated
274 to require multiple PWMs, because long C2H2-zf domain arrays could utilize different
275 segments of the array to bind to either overlapping or distinct sites[37]. Until now,
276 however, examples were sparse and anecdotal. In an accompanying manuscript[14], we
277 present evidence that C2H2-zf proteins often bind multiple sequence motifs that
278 correspond to different subsets of the extended motif predicted by the recognition code
279 (i.e. protein-sequence-based computational prediction of C2H2-zf-domain specificities),
280 consistent with varying usage of the C2H2-zf domains at different genomic binding sites
281 being commonplace.

282 **Underappreciated DNA-binding domains**

283 The six Codebook proteins that were lacking canonical DBDs, yet yielded "approved"
284 experiments and thus motifs (CGGBP1, NACC2, TCF20, PURB, DACH1, and DACH2),
285 appear to represent cases of DBDs that were poorly described at the outset of the
286 study. We and others have recently described CGGBP1 as the founding member of an
287 extensive family of eukaryotic TFs derived from the DBDs of transposons[38,39]. NACC2
288 contains a BEN domain, which over the last decade has been clearly established as a
289 sequence-specific DBD[40,41]. TCF20 contains a potential AT-hook[42] (below the
290 conventional Pfam scoring threshold), and yielded an AT-hook-like motif. PURB is
291 composed largely of three copies of the PUR (Purine-rich-element binding) domain; it
292 yielded a motif on four different PBM assays (resembling ACCnAC/GTnGGT), which is
293 unlike its previously established binding site (CTTCCCTGGAAG)[43]. The sequence
294 specificity of this protein thus remains enigmatic.

295 DACH1 and DACH2 are paralogs that yielded very similar motifs (**Figure 3A**). They
296 contain a SKI/SNO/DAC domain, shared with their *Drosophila* counterpart Dachshund,
297 from which their name is derived. A Forkhead-like motif (different from the one we
298 obtained) was previously described for DACH1[44], but to our knowledge, no other
299 homolog has been reported as being sequence-specific. The SKI/SNO/DAC domain
300 includes a helix-turn-helix (HTH), a feature found in many DBDs. Alphafold3[45] predicts
301 that the HTH inserts into the major groove precisely at the PWM-predicted binding site
302 within an extended DNA sequence (**Figure 3A**). Interpro[46] lists over 7,000 proteins
303 containing SKI/SNO/DAC domains, entirely in metazoans, with specific expansions in
304 several fish lineages, particularly barbels and salmonids[47] (**Figure 3A**). SKI/SNO/DAC
305 therefore may represent an expansive class of poorly-characterized DBDs.

306 In addition to these six examples, the sequence specificity of SLC2A4RG and ZNF395 –
307 both C2H2-zf proteins – appears to reside in their C-clamp. The domain is also present
308 in TCF7L and LEF proteins, where it is known to bind DNA alongside their HMG
309 domains[48]. Alphafold3[45] predicts that the single C2H2-zf domains in SLC2A4RG and
310 ZNF395 are not the main determinants of DNA-binding (although they may contact the
311 major groove), but instead that a region corresponding to the C-clamp model on the
312 SMART database of protein domains[49] binds the major groove precisely at the PWM-
313 predicted binding site within an extended DNA sequence (**Figure 3B**). There is one
314 additional human TF matching the C-clamp model, ZNF704, with a published PWM that
315 is virtually identical to that of SLC2A4RG and ZNF395 (CCGGCCGG)[50] (**Figure 3B**).
316 Like the SKI/SNO/DAC domain, the C-clamp is found broadly across animals[46], and
317 may therefore also represent a large class of unexplored DBDs.

318 **Widespread contribution of transposons to the human TF repertoire**

319 Sixteen of the Codebook TFs (and two controls) that yielded approved experiments
320 possess a DBD that has been co-opted from a DNA transposon: CGGBP1[39], five
321 proteins containing BED-zf domains[51], six with the related CENBP or Brinker domains[52],
322 two with transposon-derived Myb/SANT domains[53], one with a MADF domain, and
323 FLYWCH1[54]. The PWMs obtained for CENPB/Brinker TFs are often long (**Figure 3C**). A
324 striking example is JRK, a TF that is derived from an ancient domesticated Tigger
325 element DBD[55], and is found broadly in mammals[47]. All DNA transposons, including
326 Tigger, have been extinct in the human lineage for over 40 million years[56]. Remarkably,
327 genomic binding of JRK is enriched for binding to a subset of Tigger elements, and the
328 consensus sequence for these same elements has a PWM-predicted binding site for
329 JRK in the terminal repeats of these elements (**Figure 3C**), consistent with its presumed
330 ancestral role in transposition. We speculate that JRK may represent a case of co-
331 option in which the same DNA transposon simultaneously introduced both a multitude of
332 *cis*-regulatory elements, and the TF that binds them.

333 The Codebook data also underscore that many TFs bind preferentially and intrinsically
334 to specific repeat classes. These interactions are explored in greater detail in the
335 accompanying manuscripts[14,15]. Binding to endogenous retroelements is known to be a
336 common property of the KRAB-domain-containing C2H2-zf (KZNF) subfamily *in vivo*[27],
337 but until now it has not been clear that the recruitment is defined almost entirely by the
338 sequence specificity of the KZNFs alone. The combination of assays run here,
339 particularly GHT-SELEX, extends earlier observations by pinpointing the exact binding
340 sites, and demonstrating that these proteins typically have high specificity for these
341 elements, because they bind preferentially to precisely the same elements *in vitro*.
342 Binding preferentially to retroelements is not limited to KZNFs, but includes other C2H2-
343 zf proteins and other classes of TFs. For example, binding sites for TIGD3, a
344 transposon-derived TF which is closely related to JRK, are enriched for binding to L1s,
345 SINEs, and DNA transposons[15].

Codebook PWMs predict TF binding in independent data and across cell types

The Codebook project was conducted over a period of nearly six years, and during this time, several large-scale studies aimed at systematic ChIP-seq analysis of human TFs (e.g. ENCODE) were published[11,57,58]. Combined, the ENCODE data portal[59] and GTRD[60], a compilation database, contain ChIP-seq and ChIP-exo peak data for 214 of the Codebook proteins, including 105 that were among the 166 with either "approved" Codebook ChIP-seq experiments (**Table S7**), or with ChIP-seq replicates that yielded reproducible peak sets[15]. We grouped both types of ChIP-seq data in our study and compared them to the external data. We first asked whether Codebook peak sets overlapped with these external peak sets for the same TF. Among the major ENCODE cell lines, the highest overlap values (Jaccard index) were found with experiments utilizing the same cell type (HEK293 cells) (**Figure S3A,B**). Slightly lower Jaccard values were obtained for experiments performed in HepG2 and other cell types, which would be expected given the altered chromatin profiles in different cell types, but over one-third were still clearly nonrandom (Jaccard > 0.1) (**Figure S3C**). Overlap scores with published K562 data, which dominate the external ChIP data due to a single large ChIP-exo study[58], were much lower, overall (**Figure S3D**). We conclude from these analyses that the Codebook ChIP-seq data provide mainly new information.

We next asked how effectively the Codebook PWMs predict binding of TFs to peak sets in the published datasets. Consistent with the fact that the Codebook and external peaks often overlap, the Codebook PWMs had a median AUROC of 0.71 on the external HEK293 data, and were nearly as effective in predicting peak sets in other cell types (**Figure S3E**), illustrating that the Codebook PWMs are predictive across studies and cell types. We also asked how the predictive capacity of the Codebook PWMs compared to PWMs that appear in the latest versions of Factorbook[12], JASPAR[61], and HOCOMOCO[13] (**Table S8**). We identified 19 TFs with at least one successful Codebook ChIP-seq experiment and Codebook PWM, at least one external ChIP-seq experiment, and at least one PWM from an external database. In most cases, both the Codebook and external PWMs scored well on both Codebook and external peak sets (**Figure S3F,G**), supporting the validity of both PWMs and both peak sets. For seven proteins, low scores were obtained in at least some tests, however. For four of them, the independent Codebook *in vitro* data support the Codebook PWM; for two of the others, the external PWM scores poorly on Codebook peaks, while the Codebook PWM scores well on Codebook and external peak sets (**Figure S3H**). We conclude that the Codebook PWMs are generally more reliable than those published previously, likely because they are aided by confirmation of PWM performance across multiple data types that were not available in previous studies

Codebook TF binding sites suggest functions for tens of thousands of conserved elements

Together, the Codebook assays and PWMs can be used to pinpoint genomic loci that are bound directly by each TF *in vivo* (i.e., in ChIP-seq), by identifying those that are also bound *in vitro* (i.e., GHT-SELEX), and that contain a PWM hit, thus allowing base-level resolution. We refer to these as "triple overlap" (TOP) sites, which are taken as the

overlap of the three sets (ChIP-seq, GHT-SELEX, and PWM hits) after applying optimized score thresholds for each (see **Methods** for details). This process produced a median of 455 TOP sites for 101 Codebook proteins, and a median of 3,014 TOP sites for 36 control TFs.

To gauge functionality of the TOP sites, we examined whether the pattern of per-nucleotide conservation[13] at each site is consistent with the TF's sequence preference driving local sequence constraint (see **Methods** for details). **Figure 4A** shows several examples illustrating that this approach readily detects apparent conservation of PWM hits, for both control and Codebook TFs. In total, 85/101 Codebook TFs (as well as 33/36 controls) displayed conservation of at least one TOP site (FDR < 0.1), and in total we identified 121,785 such conserved TOP sites ("CTOP" sites) (83,621 for Codebook TFs and 38,164 for controls), encompassing 1,577,298 bases. These results, summarized in **Figure S4** and in greater detail in an accompanying manuscript[15], provide strong support for the functional importance of Codebook TF binding sites in the genome.

Many of the CTOP sites were either overlapping or adjacent to CTOP sites for the same or other TFs. We grouped them into 50,375 clusters, based on proximity (allowing a maximum of 100 bases, to capture binding to different segments of what may be the same regulatory element). Codebook TFs with the largest number of CTOP sites were typically associated with CpG islands, which represented 37.5% of all the clusters (**Figure 4B**). The majority of protein-coding promoter CpG islands (58.7%, 7,892/13,427) contained CTOP sites, with an average of 4.3 CTOP sites per CpG island. Moreover, 59/101 (58%) of all Codebook TFs had at least one CTOP site within a CpG island. An example CTOP that overlaps a CpG island is shown in **Figure 4C**.

The extent of specific, conserved, and intrinsic occupancy of CpG islands by many TFs of diverse classes is, to our knowledge, unexpected. The abundance of CG dinucleotides in CpG islands has been attributed primarily to their lack of methylation in the germline, rather than primary sequence constraint[62]. There is one class of TFs (the CXXC proteins) that is known to specifically recognize unmethylated CG dinucleotides and to modulate chromatin at promoters[62], and we do observe this property for the CXXC proteins KDM2A, CXXC4, FBXL19, and TET3. Intriguingly, however, many of the Codebook TFs with CTOP sites in CpG islands recognize elaborate C/G rich motifs, rather than CG dinucleotides (**Figure 4C**).

CTOP clusters were also found in non-CpG island protein-coding promoters (**Figure 4B**) (855/6,606 such promoters, defined as -1000 to +500 relative to TSS). These clusters are not dominated by any specific TFs**,** although some TFs are more prevalent than others (e.g. CTOPs for the controls ELF3 and CTCF, and Codebook TF ZBTB41, are each found in ~10% of all non-CpG promoters) (**Figure 4D**). **Figure 4E** shows an example of one such non-CpG promoter cluster, occurring early in the first intron of the TSPAN31 gene, which exhibits apparent conserved spacing and orientation of multiple Codebook TF binding sites. In contrast, CTOP clusters outside of promoters and CpG islands often contain just one or two CTOP sites (**Figure 4B)**. One example is a very strongly conserved intergenic ZNF689 binding site found in an L1ME1 transposon; this

site is just over 100 bp from a predicted enhancer containing a CTCF binding site (**Figure 4F**).

A total of 42,200 distinct CTOP clusters (out of 50,375) overlapped catalogued conserved elements (UCSC PhastCons track), thus indicating a likely biochemical function for these elements. For the remaining 8,175, detection of functional elements from base-level scores is now augmented by the TF binding information. Relatively few CTOP clusters overlapped with known enhancers, however: only 4,768 are found in the extensive GeneHancer annotation set[63], and 2,819 overlap with HEK293 enhancers (defined by ChromHMM[15]). This low overlap could be attributed to the relatively rapid evolution of enhancers[64], or to lack of complete knowledge of enhancer identities. We also note that, even for well-studied TFs, most TOP sites were classified by our methods as not conserved, and that roughly half of the Codebook TFs had few or no conserved TOP sites (particularly the aforementioned retroelement-binding KZNFs) (**Figure S4**). Lack of conservation does not demonstrate that a sequence is not a functional binding site, however, as turnover in functional genomic binding sites of TFs is common[65]. This result is nonetheless consistent with the notion that many TF binding sites are coincidental, redundant, or serve(d) a purpose other than host genome regulation. In the accompanying manuscript[15], we explore potential functions for proteins that frequently bind non-conserved sites in genomic “dark matter”.

Relationships between Codebook TFs, SNVs and chromatin

Because the CTOP sites are evolutionarily constrained, we reasoned that they might also be less frequently associated with human sequence variation, and indeed, 92.6% of CTOPs lack SNPs and other common short variants, while only 82.1% of unconserved TOPs are variant-free. Both are depleted of common SNPs, however, when examined separately (Fisher’s exact test $p \sim 2.4 \times 10^{-307}$ and odds ratio = 0.657, $p \sim 0$ and ratio = 0.872, respectively). The CTOP SNPs also have a lower impact on PWM scores: on average, the relative PWM score for SNP-containing CTOP sequences declines by 0.027, while PWM scores for unconserved TOPs decline by 0.057 (median declines of 0.011 and 0.0285, respectively). CTOPs are furthermore depleted of common short indels (Fisher’s exact test, $p \sim 1 \times 10^{-150}$, ratio = 0.77), while unconserved TOPs (which often overlap with simple repeats) are enriched ($p < 1 \times 10^{-150}$, ratio = 3.318), relative to genomic background. The depletion of common SNPs is consistent with ongoing purifying selection of CTOPs within recent human populations, and the association of SNPs with specific TFs should provide a ready means for directed study of the functionality of the encompassed SNPs.

We reasoned that the GHT-SELEX and ChIP-seq experiments would also allow direct assessment of allele-specific binding (ASB) of TFs, by quantifying allelic imbalance of read counts at SNVs. We note that the data were not initially intended for this purpose, and caveats included relatively low read counts, linked SNVs, and the fact that HEK293 has an abnormal karyotype and was derived from a single individual. Nonetheless, there was sufficient coverage in the sequencing data to make 925,003 variant calls overlapping with dbSNP common SNPs (889,820 variant calls from 362 ChIP-seq experiments and 35,183 from 374 GHT-SELEX multi-cycle experiments), at 122,364

475 unique genomic locations (**Figure 5A**, **Figure S5A**, **Table S9**). 10,009 of these genomic
476 locations were associated with 12,056 ASBs of 160 Codebook TFs and 46 positive
477 controls in ChIP-seq (10,571 ASBs) or GHT-SELEX (1,485 ASBs) samples, i.e. there
478 was a significant imbalance in the sequencing reads for the two alleles overlapping the
479 respective SNPs. Among these ASBs, 3,569 also overlapped a PWM hit for the TF, and
480 for 2,367 of them, the read count imbalance was concordant with the change in PWM
481 scores, i.e. the allele with the higher read count also has a higher PWM score (**Figure**
482 **S5A,B, Table S9**). (ASBs that do not overlap a PWM hit may be linked to a "causative"
483 SNV, which may act indirectly). ASBs for control TFs were strongly enriched with
484 previously-known ASBs of those TFs (ADASTRA database, odds ratio of 5.7, $p < 10^{-15}$,
485 Fisher's exact test)[66], and nearly three-quarters of ASBs coincided with eQTLs (GTEx
486 database, odds ratio of 1.2, $p < 10^{-15}$, Fisher's exact test)[67] (**Figure S5C**), supporting the
487 reliability of the detected ASBs as well as the validity of detected PWM hits.

488 Compared to whole-length peaks, TOP regions had an increased density of variant calls
489 (~258 sufficiently covered variants per Mb in TOPs, versus 52 per Mb for peaks), and a
490 larger fraction of ASB calls in SNVs (30%, compared to 9% for full peaks), presumably
491 due to detection bias from higher ChIP-Seq or GHT-SELEX coverage at the TOPs.
492 Nonetheless, variants in TOPs had a significantly higher predicted effect on protein
493 binding (i.e. PWM score change) for both controls and Codebook TFs ($p < 2.22 \times 10^{-5}$
494 and $p < 2.98 \times 10^{-12}$, Mann-Whitney U test), relative to full peaks or non-ASB SNPs
495 overlapping PWM hits (**Figure 5B**). Thus, the ASBs in TOPs are more likely to induce
496 an effect than those elsewhere within peaks, presumably because they represent direct
497 TF binding.

498 Among the mechanisms connecting TF binding to biological function are TF-mediated
499 chromatin state changes. Hence, in heterozygotes, variant-dependent TF binding may
500 co-occur with allele-specific chromatin accessibility variants (ASVs) (**Figure 5A**), which
501 are SNVs with imbalanced read counts in ATAC-seq and/or DNase-seq experiments.
502 To ask whether the Codebook TFs may be involved in control of ASVs, we utilized the
503 UDACHA database, which contains ASVs from 577 ATAC-seq and 321 DNase-seq
504 datasets from individual cell types[68] (**Table S9, Figure S5D**). Using a multi-tiered
505 procedure (see **Methods**), we identified cases in which (1) ASVs in a specific cell type
506 overlap significantly with PWM hits for a TF in the Codebook motif collection, (2) the
507 change in the PWM score is concordant with the read imbalance in the ASVs, (i.e.
508 stronger predicted binding is associated with more accessible chromatin), and (3) the
509 concordance is significant across cell types detected in step (1). This procedure
510 identified 53 TFs whose PWM hits were found often at, and concordant with, ASVs
511 (**Figure S5E**). Twenty of these TFs were positive controls including well-known pioneers
512 or activators (such as SOX2, GABPA, or JUN/FOS-family TFs), while 33 were
513 previously unexplored Codebook TFs, including ZNF70, GRHL3, MYPOP, SP140(L),
514 and DMTF1. An example ASV for ZNF70, in a region upstream of the PTMS gene that
515 is annotated with multiple ENCODE enhancer elements is shown in **Figure 5C**.

516 For 34 of these 53 TFs, there was at least one ASV-overlapping TOP site (the non-TOP
517 sites may represent sites that are not bound in HEK293). To assess whether ASVs in
518 PWM hits have a greater effect at TOP sites than in other regions, we first removed

519 cases in which the TF does not appear to impact chromatin directly, by grouping the
520 TFs into ASV-concordant (i.e. having overall concordance between ASVs and PWM hits
521 in ChIP-seq or GHT-SELEX peaks; 18 TFs), and others (16 TFs). We separated the
522 ASV-concordant group into Codebook and control TFs. For each of the groups, we then
523 calculated the concordant-to-discordant ratio for loci that corresponded to PWM hits that
524 are non-ASV for that TF, ASV, ASV in TF's peaks, and ASV in TOPs, and observed an
525 overall monotonic increase in concordance (**Figure 5D**). Thus, the highest-confidence
526 Codebook TF binding sites for these TFs are those most likely to impact the chromatin
527 state. Moreover, the fraction of ASVs within PWM hits also increased monotonously as
528 the ASV confidence increased, and the ASVs preferably occur at binding site positions
529 that are most important for the PWM score (**Figure 5E, Figure S5F**), further supporting
530 relevance of the TF sequence preferences.

531 Overall, the Codebook motifs provide a valuable resource for SNV interpretation,
532 including identification of mechanisms that underpin variation in chromatin and
533 transcription.

534 **Lessons from Codebook: prospects for a complete human TF motif collection**

535 Codebook yielded several clear outcomes, and guidance for future efforts. The high
536 success rate is particularly striking. We obtained motifs for 177 previously
537 uncharacterized human TFs, a number larger than the entire TF repertoire for many
538 eukaryotes[69]. The selected PWMs for most of these TFs are unique, and unlike any
539 previous TF motif. Most are from C2H2-zf proteins, and most C2H2-zf proteins analyzed
540 were successful. Thus, a majority of putative and uncharacterized human TFs are *bona*
541 *fide* TFs, and not annotation errors. We envision that the data produced will be broadly
542 and immediately useful for a variety of applications. Motifs (especially as PWMs) are a
543 standard component of the computational genomics toolkit, due to their utility in a range
544 of tasks ranging from identification of key regulatory factors to building and interpreting
545 models of gene expression[70-73]. For example, differential binding of TFs to noncoding
546 SNVs (Single Nucleotide Variants) is thought to be a major mechanism by which these
547 variants contribute to phenotypic differences[74], and the Codebook data therefore
548 provide vital new information for the analysis of *cis*-regulatory variation.

549 A key technical demonstration of the Codebook project is that the simultaneous
550 application of multiple experimental strategies and multiple motif-derivation and motif-
551 scoring strategies was highly beneficial. No single experiment type or data analysis
552 approach dominated all others, or was universally successful, although specific assays
553 were more or less advantageous for different classes of proteins (as evident in **Figure**
554 **1**). For example, PBMs were uniquely successful with AT-hook proteins, while ChIP-seq
555 and SELEX variants were most successful for C2H2-zf proteins. We caution that there
556 are confounding variables limiting what conclusions can be drawn regarding the
557 strengths and weaknesses of experimental platforms. The protein production and
558 purification method can differentially impact success of specific DBD classes, even
559 when the same assay is used, and the different assays we employed were tied to
560 different affinity tags and expression systems. Data pre-processing (i.e. read filtering

561 and background estimation) is an additional variable that we did not systematically
562 explore, but is known to impact all of the assays used here.

563 As noted above, a subset of the Codebook TFs, as well as other poorly characterized
564 TFs, have been analyzed by others since our study began. To evaluate the current
565 scope of known human TF specificities, we surveyed JASPAR, HOCOMOCO, and
566 Factorbook for PWMs for putative TFs that were not included in this study or not found
567 among 177 Codebook successes. These databases reported PWMs for 107 proteins,
568 63 of which we had tested, and 44 were among the 95 putative TFs not included in our
569 experiments. We manually curated these external PWMs, using procedures similar to
570 those we applied to our own data, to assess whether they are likely to represent the
571 *bona fide* specificity of the TF analyzed. Many of them were comprised of simple
572 repeats (which are common artifacts in virtually all assays) or appeared to correspond
573 to indirect binding and/or recruitment by other TFs in ChIP-seq (See **Table S8** for
574 annotations and classification, and **Figure S1** for examples of nonspecific, concordant,
575 and likely correct PWMs in the external datasets).

576 Based on this curation, 33 additional human TFs (i.e. beyond the 177 described here)
577 have at least one plausible motif available in datasets that have been performed since
578 our 2018 TF census[1], leading to a total of 1,421 human TFs now with characterized
579 sequence specificities (**Figure 6** and **Table S10**). Altogether, only 175 proteins with
580 conventional DBDs now lack known sequence specificity. Not all proteins with such
581 domains are necessarily TFs; for example, one systematic trend we observed is that
582 almost all 36 proteins we tested with only a single C2H2-zf domain failed in every assay
583 (**Figure 6**). At the same time, however, new DBD classes continue to appear, such as
584 the aforementioned BEN, CGGBP, Dachshund, and C-clamp. Some TFs may bind only
585 to methylated DNA, and ongoing advances in the prediction of protein and protein-DNA
586 structures[45] have the potential to identify additional candidates for sequence-specific
587 DNA binding. Thus, while completion of the objective to obtain a motif for every human
588 TF now appears much closer, the list of likely human TFs continues to evolve.

589 Many of the Codebook TFs are now among the best characterized human DNA-binding
590 proteins in terms of their sequence specificity. As illustrated in the accompanying
591 papers (**Table S1**), and consistent with previous benchmarking efforts[18,32], validation
592 across platforms can lead to very different conclusions regarding PWM reliability.
593 Moreover, obtaining *in vivo* and *in vitro* binding to the genome facilitates
594 disentanglement of direct and indirect binding, as well as the contribution of the cellular
595 environment. Obtaining *in vitro* binding data to both genomic-sequence and random-
596 sequence DNA can provide insight into the importance of local sequence context. Only
597 a small handful of the 1,000+ previously characterized TFs have such a combination of
598 data types. A much better perspective on human gene regulation and genome function
599 and evolution could presumably be obtained from generation of such data for all human
600 TFs.

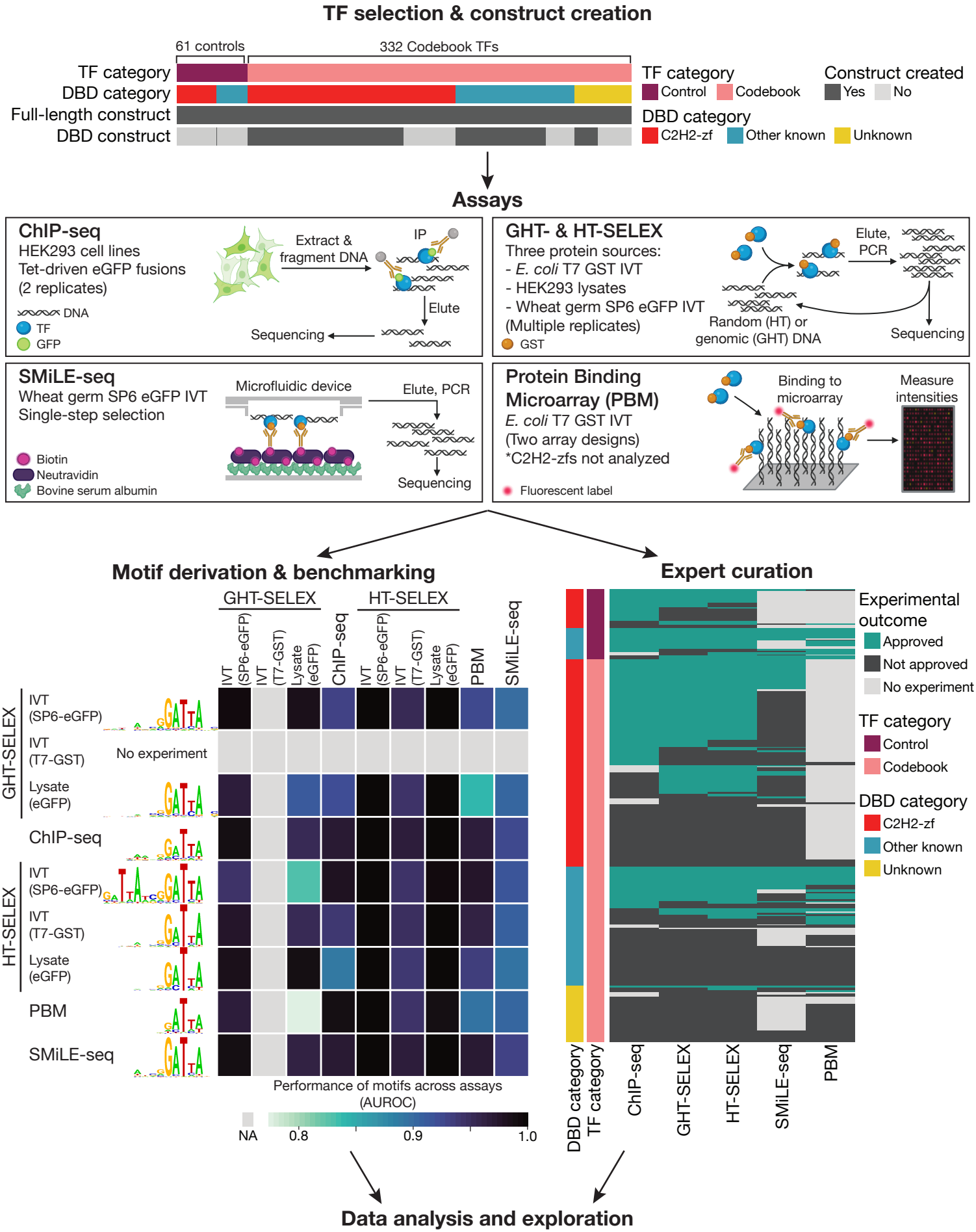
TF selection & construct creation
61 controls
332 Codebook TFs
TF category
DBD category
Full-length construct
DBD construct
TF category
Control
Codebook
Construct created
Yes
No
DBD category
C2H2-zf
Other known
Unknown
Assays
ChIP-seq
HEK293 cell lines
Tet-driven eGFP fusions
(2 replicates)
DNA
TF
GFP
Extract & fragment DNA
IP
Elute
Sequencing
GHT- & HT-SELEX
Three protein sources:
- E. coli T7 GST IVT
- HEK293 lysates
- Wheat germ SP6 eGFP IVT
(Multiple replicates)
GST
Elute, PCR
Random (HT) or genomic (GHT) DNA
Sequencing
SMiLE-seq
Wheat germ SP6 eGFP IVT
Single-step selection
Biotin
Neutravidin
Bovine serum albumin
Microfluidic device
Elute, PCR
Sequencing
Protein Binding Microarray (PBM)
E. coli T7 GST IVT
(Two array designs)
*C2H2-zfs not analyzed
Fluorescent label
Binding to microarray
Measure intensities
Motif derivation & benchmarking
GHT-SELEX
HT-SELEX
IVT (SP6-eGFP)
IVT (T7-GST)
Lysate (eGFP)
ChIP-seq
PBM
SMiLE-seq
No experiment
Performance of motifs across assays (AUROC)
NA
0.8
0.9
1.0
Expert curation
Experimental outcome
Approved
Not approved
No experiment
TF category
Control
Codebook
DBD category
C2H2-zf
Other known
Unknown
DBD category
TF category
ChIP-seq
GHT-SELEX
HT-SELEX
SMiLE-seq
PBM
Data analysis and exploration

Figure 1. Codebook project overview. ***Top***, Categories of 393 TFs assayed and their associated constructs. ***Middle***, Graphical summary of assays employed. ***Bottom left***, Example of performance (as AUROC) of the best performing PWM for TPRX1, for each combination of experiment type – one for motif derivation (rows), and one for motif testing (columns). ***Bottom right***, Depiction of the approval process for each individual experiment, including comparison of motifs and/or binding sites between replicates, evaluation of motifs across experiments, and motif similarity between related TFs (see **Experiment evaluation by expert curation**). Heatmap shows approved experiments for all 393 TFs across all experiment types.

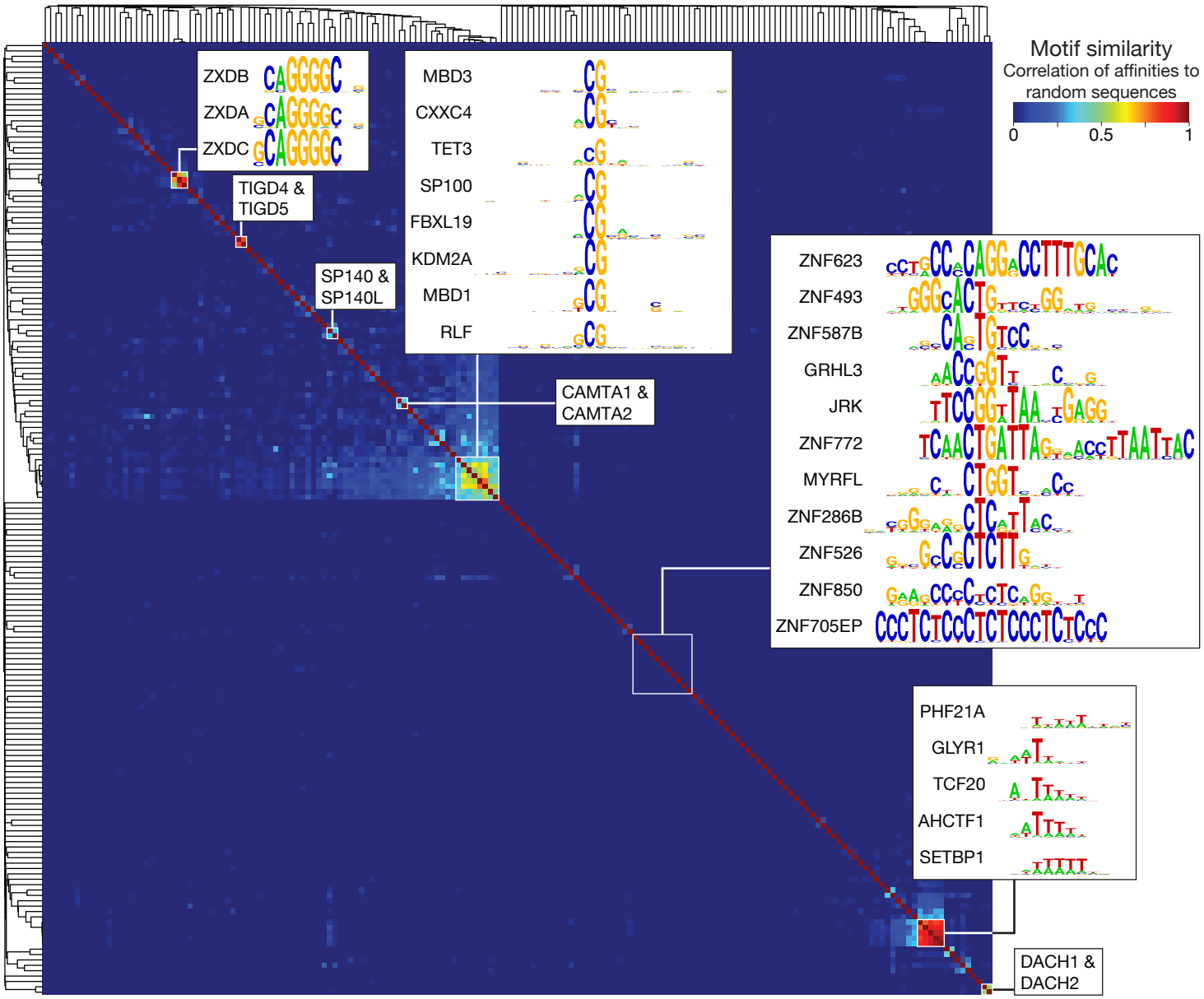
Motif similarity
Correlation of affinities to random sequences
0
0.5
1
ZXDB
ZXDA
ZXDC
TIGD4 & TIGD5
SP140 & SP140L
MBD3
CXXC4
TET3
SP100
FBXL19
KDM2A
MBD1
RLF
CAMTA1 & CAMTA2
ZNF623
ZNF493
ZNF587B
GRHL3
JRK
ZNF772
MYRFL
ZNF286B
ZNF526
ZNF850
ZNF705EP
PHF21A
GLYR1
TCF20
AHCTF1
SETBP1
DACH1 & DACH2

Figure 2. Similarity of Codebook TF motifs. Symmetric heatmap displaying the similarity between expert-curated PWMs for each pair of Codebook TFs, clustered by Pearson correlation with average linkage. The PWM similarity metric is the correlation between pairwise affinities to 200,000 random sequences of length 50, as calculated by MoSBAT[28]. Pullouts and labels illustrate specific points in the main text.

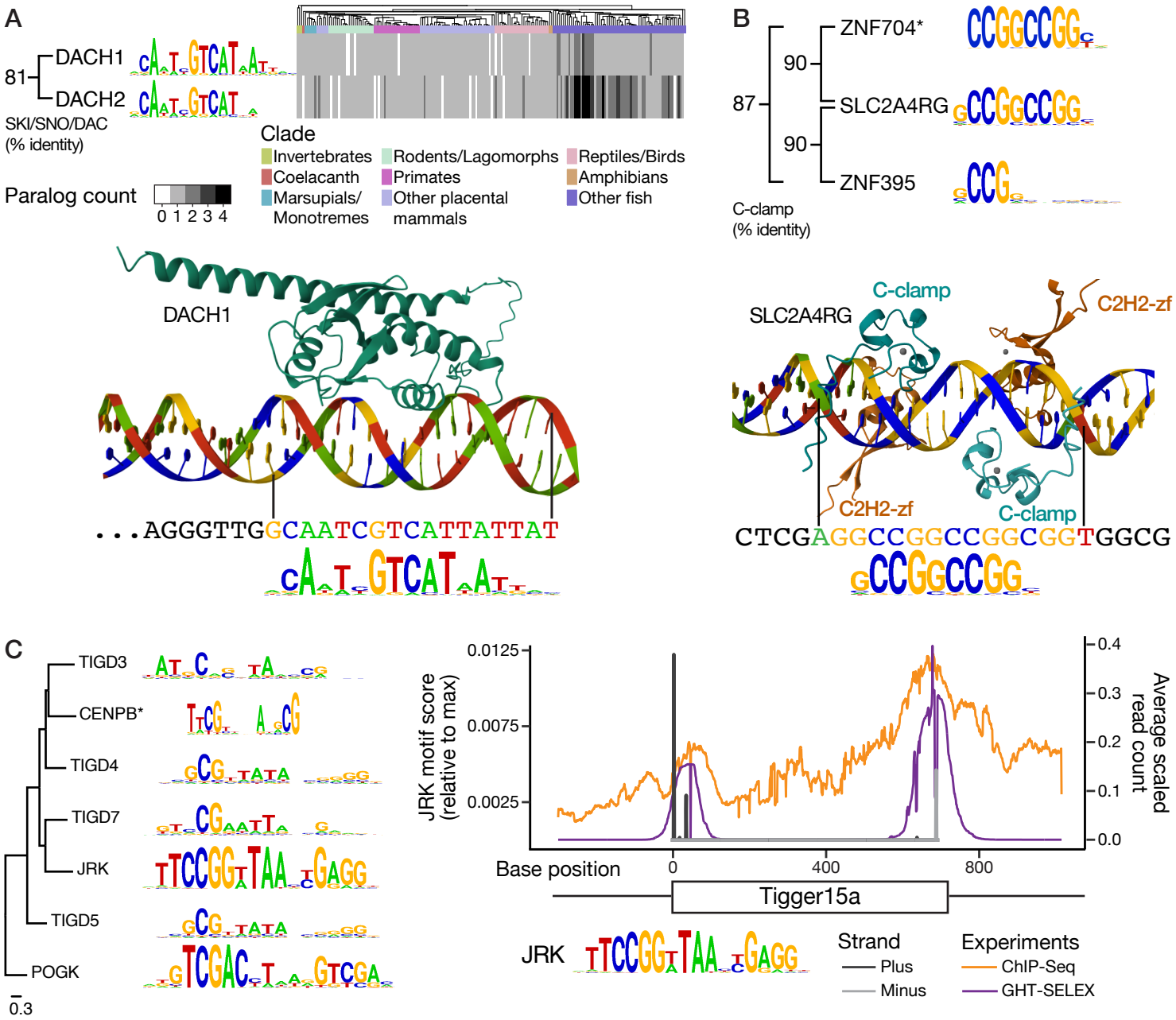
A
81
DACH1
DACH2
SKI/SNO/DAC
(% identity)
Paralog count
0 1 2 3 4
Clade
Invertebrates
Coelacanth
Marsupials/ Monotremes
Rodents/Lagomorphs
Primates
Other placental mammals
Reptiles/Birds
Amphibians
Other fish
B
87
90
90
ZNF704*
SLC2A4RG
ZNF395
C-clamp
(% identity)
DACH1
...AGGGTTGGCAATCGTCATTATTAT
SLC2A4RG
C-clamp
C2H2-zf
C2H2-zf
C-clamp
CTCGAGGCCGGCCGGCGGTGGCG
C
TIGD3
CENPB*
TIGD4
TIGD7
JRK
TIGD5
POGK
0.3
JRK motif score
(relative to max)
0.0125
0.0075
0.0025
Average scaled
read count
0.4
0.3
0.2
0.1
0.0
Base position
0
400
800
Tigger15a
JRK
Strand
Plus
Minus
Experiments
ChIP-Seq
GHT-SELEX

Figure 3. Neglected DNA-binding domains. Overview of new motifs for previously understudied TF families. **A,** *Top,* Number of DACH1 and DACH2 orthologs (union of one-to-one and one-to-many) across Ensembl v111 vertebrates and selected invertebrates. Species order reflects the Ensembl species tree. *Bottom,* AlphaFold3-predicted structure of the DACH1 SKI/SNO/DAC region (residues 130 – 390) bound to an HT-SELEX ligand sequence with a high-scoring PWM hit. **B,** *Top,* Sequence logos and sequence relationships of human C-Clamp domains (*ZNF704 motif from [50]). *Bottom*, AlphaFold3-predicted structure of two full-length SLC2A4RG proteins bound to a CTOP sequence with flanking sequences (chr17:48,048,369-48,048,401), and four Zn^{2+} ions (grey). The remainder of the proteins (beyond the C-clamp and C2H2-zf domains) are hidden, for visual simplicity. **C.** *Left,* Sequence logos of human TFs that are derived from the domestication of *Tigger* and *Pogo* DNA transposon DBDs elements and have known DNA binding motifs. Tree is a maximum-likelihood phylogram from FastTree[92], using DBD sequence alignment with MAFFT L-INS-I[93], rooted on POGK, which is derived from an older family of Tigger-like elements[94,95]. Sequence logos are Codebook-derived, except for CENPB[96]. *Right,* average per-base read count over Tigger15a TOPs in the human genome, for JRK ChIP-seq (orange) and GHT-SELEX (purple), with sequences aligned to the Tigger15a consensus sequence. JRK PWM scores at each base of the Tigger15a consensus sequence are shown in black (plus strand) and grey (minus strand).

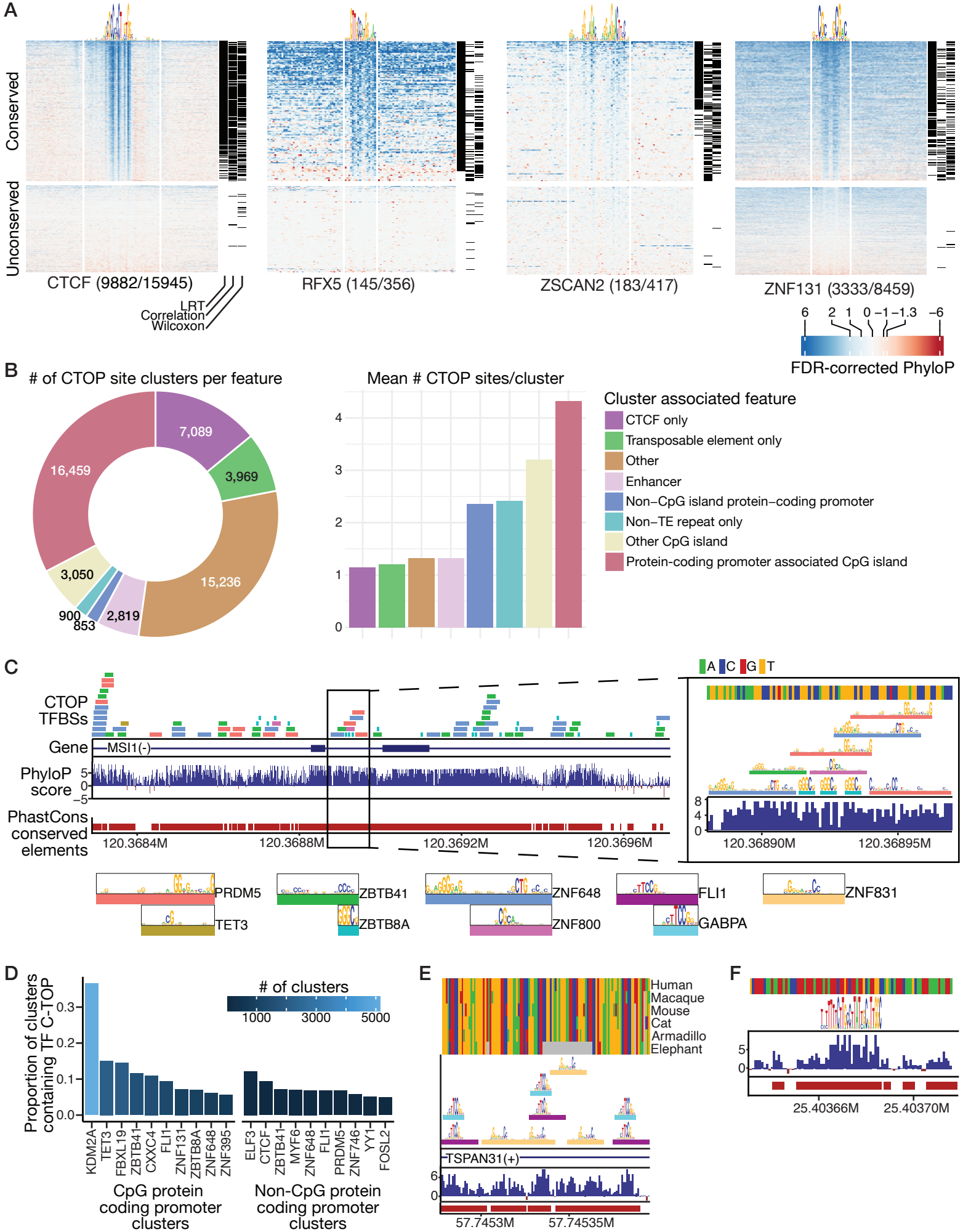
A
Conserved
Unconserved
CTCF (9882/15945)
RFX5 (145/356)
ZSCAN2 (183/417)
ZNF131 (3333/8459)
LRT
Correlation
Wilcoxon
6 2 1 0 −1 −1.3 −6
FDR-corrected PhyloP
B
# of CTOP site clusters per feature
7,089
3,969
15,236
2,819
853
900
3,050
16,459
Mean # CTOP sites/cluster
Cluster associated feature
CTCF only
Transposable element only
Other
Enhancer
Non−CpG island protein−coding promoter
Non−TE repeat only
Other CpG island
Protein-coding promoter associated CpG island
C
CTOP TFBSs
Gene
MSI1(-)
PhyloP score
PhastCons conserved elements
120.3684M
120.3688M
120.3692M
120.3696M
A C G T
120.36890M
120.36895M
PRDM5
TET3
ZBTB41
ZBTB8A
ZNF648
ZNF800
FLI1
GABPA
ZNF831
D
Proportion of clusters containing TF C-TOP
# of clusters
1000 3000 5000
KDM2A TET3 FBXL19 ZBTB41 CXXC4 FLI1 ZNF131 ZBTB8A ZNF648 ZNF395
CpG protein coding promoter clusters
ELF3 CTCF ZBTB41 MYF6 ZNF648 FLI1 PRDM5 ZNF746 YY1 FOSL2
Non-CpG protein coding promoter clusters
E
Human
Macaque
Mouse
Cat
Armadillo
Elephant
TSPAN31(+)
57.7453M
57.74535M
F
25.40366M
25.40370M

Figure 4. Conservation of Codebook TF binding sites and association with genomic features. A, Heatmaps of phyloP scores over the PWM hit and 50 bp flanking for TOP sites for four TFs (two controls and two Codebook TFs). Statistical test results (see main text and **Methods**) are indicated at right. **B,** *Left,* Donut plot displays the proportion and number of clusters of conserved TOP (CTOP) sites that overlap the genomic features indicated. *Middle,* Bar plot displays the mean # of individual CTOPs contained within clusters that overlap the examined genomic regions. **C.** A 1,420-base, CpG-island-overlapping CTOP cluster (chr12:120368293-120369713). Zoonomia 241-mammal phyloP scores and Multiz 471 Mammal alignment PhastCons Conserved Elements are shown. **D,** Bar plot of the frequency of TFs with CTOPs that occur most frequently in CTOP clusters that overlap CpG and non-CpG protein coding promoters, respectively. **E,** CTOP cluster overlapping the non-CpG promoter at chr12:57,745,278-57,745,396. **F,** CTOP site for the KRAB-C2H2-zf protein ZNF689, overlapping an L1ME4a located at chr16:25,403,631-25,403,717.

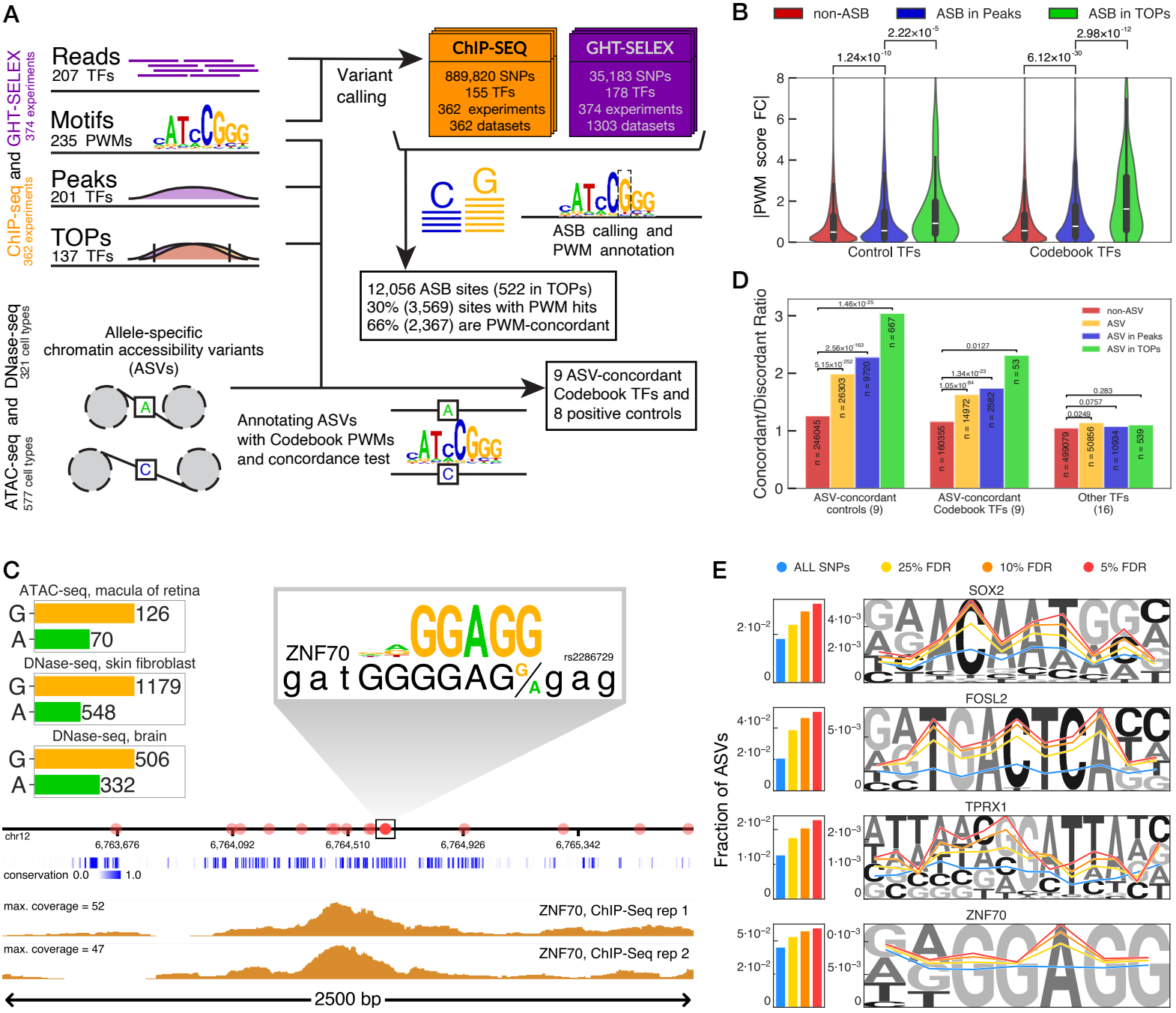
A
ChIP-seq and GHT-SELEX
374 experiments
Reads
207 TFs
Motifs
235 PWMs
Peaks
201 TFs
TOPs
137 TFs
Variant calling
ChIP-SEQ
889,820 SNPs
155 TFs
362 experiments
362 datasets
GHT-SELEX
35,183 SNPs
178 TFs
374 experiments
1303 datasets
ASB calling and PWM annotation
12,056 ASB sites (522 in TOPs)
30% (3,569) sites with PWM hits
66% (2,367) are PWM-concordant
DNase-seq
321 cell types
ATAC-seq
577 cell types
Allele-specific chromatin accessibility variants (ASVs)
Annotating ASVs with Codebook PWMs and concordance test
9 ASV-concordant Codebook TFs and 8 positive controls
B
non-ASB
ASB in Peaks
ASB in TOPs
1.24×10^-10
2.22×10^-5
6.12×10^-30
2.98×10^-12
|PWM score FC|
Control TFs
Codebook TFs
C
ATAC-seq, macula of retina
G 126
A 70
DNase-seq, skin fibroblast
G 1179
A 548
DNase-seq, brain
G 506
A 332
ZNF70
GGAGG
rs2286729
g a t GGGGAG G/A g a g
chr12
6,763,676
6,764,092
6,764,510
6,764,926
6,765,342
conservation 0.0 1.0
max. coverage = 52
ZNF70, ChIP-Seq rep 1
max. coverage = 47
ZNF70, ChIP-Seq rep 2
2500 bp
D
Concordant/Discordant Ratio
non-ASV
ASV
ASV in Peaks
ASV in TOPs
1.46×10^-25
2.56×10^-163
5.15×10^-252
0.0127
1.34×10^-23
1.05×10^-84
0.283
0.0757
0.0249
n = 246045
n = 26303
n = 9720
n = 667
n = 160355
n = 14972
n = 2582
n = 53
n = 499079
n = 50856
n = 10334
n = 539
ASV-concordant controls (9)
ASV-concordant Codebook TFs (9)
Other TFs (16)
E
ALL SNPs
25% FDR
10% FDR
5% FDR
SOX2
FOSL2
TPRX1
ZNF70
Fraction of ASVs

Figure 5. Allele-specific transcription factor binding and chromatin accessibility. **A,** Scheme of the analysis: identification of allele-specific binding sites (ASBs) from Codebook ChIP-Seq and GHT-SELEX data and annotation of allele-specific chromatin accessibility variants (ASVs) with the Codebook motifs. **B,** Distribution of PWM score (log-odds) fold changes between alleles for non-ASB SNPs, ASBs in peaks, and ASBs in TOPs. *Left*, 32 positive control TFs, *Right*, 85 Codebook TFs. P-values: Mann-Whitney U test. **C.** An example ASV for ZNF70, in chr12:6,763,200-6,765,850, around 1kb upstream of the PTMS gene. Onset shows the exact location of the ASV (with A/G alleles) together with the corresponding PWM hit. Allelic read counts for three available ATAC- and DNase-seq samples are shown on the side. **D**. The ratio of concordant-to-discordant PWM hits for <SNP, TF> pairs for non-ASVs (red), all ASVs (yellow), ASVs overlapping with peaks (blue), and ASVs in TOPs (green). P-values: Fisher's exact test. **E**. *Left*, Fraction of ASVs overlapping with PWM hits for four example TFs, using 4 different thresholds on ASV significance: all SNPs (blue), 25% FDR ASVs (yellow), 10% FDR ASVs (orange), and 5% FDR ASVs (red). *Right*, Fraction of ASVs at each location within the genome-wide PWM hits of the representative TFs using four thresholds (same colors as in bar plots).

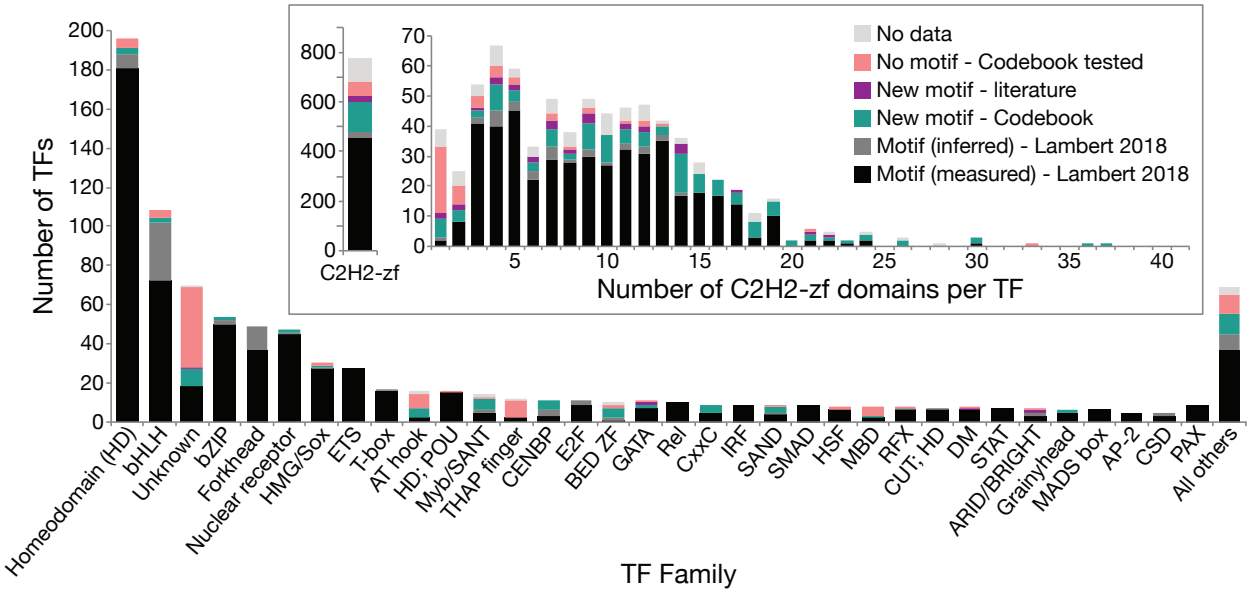
Number of TFs
TF Family
Homeodomain (HD)
bHLH
Unknown
bZIP
Forkhead
Nuclear receptor
HMG/Sox
ETS
T-box
AT hook
HD; POU
Myb/SANT
THAP finger
CENBP
E2F
BED ZF
GATA
Rel
CxxC
IRF
SAND
SMAD
HSF
MBD
RFX
CUT; HD
DM
STAT
ARID/BRIGHT
Grainyhead
MADS box
AP-2
CSD
PAX
All others
C2H2-zf
Number of C2H2-zf domains per TF
No data
No motif - Codebook tested
New motif - literature
New motif - Codebook
Motif (inferred) - Lambert 2018
Motif (measured) - Lambert 2018

Figure 6. Motif coverage of human TFs, by DBD family. TFs are categorized into structural classes based on Lambert et al.[1]. See **Table S10** for underlying information.

601 **METHODS**

602 **Plasmids and inserts.** Sequences and accompanying information are given in **Table**
603 **S3**, and the relationships between constructs, samples, and experiments are compiled
604 in the information provided online at codebook.ccbr.utoronto.ca. Briefly, we selected
605 Codebook TFs (and their DNA-binding domains catalogued) from information
606 accompanying Lambert 2018[1]) and posted at https://humantfs.ccbr.utoronto.ca. Inserts
607 named with an "-FL" suffix correspond to the full-length ORF of a representative isoform
608 of the protein. Those with a "-DBD" suffix contain all of the predicted DBDs in the protein
609 flanked by either 50 amino-acids, or up to the N or C-terminus of the protein. Those with
610 a "-DBD1", "-DBD2" or "-DBD3" suffix contain a subset of the DBDs present in the
611 proteins; these were designed manually, mainly for large C2H2-zf arrays. Inserts were
612 obtained as recoded synthetic ORFs (BioBasic, US) flanked by AscI and SbfI sites, and
613 subcloned into up to three plasmids: (i) pTH13195, a tetracycline-inducible, N-terminal
614 eGFP-tagged expression vector with FLiP-in recombinase sites[10]; (ii) pTH6838, a T7-
615 promoter driven, N-terminal GST-tagged bacterial expression vector[75], and (iii)
616 pTH16500 (pF3A-ResEnz-egfp), an SP6-promoter driven, N-terminal eGFP-tagged
617 bacterial expression vector, modified from pF3A–eGFP[9] to contain the two restriction
618 sites after the eGFP.

619 **Protein production.** Each experiment used a protein expressed from one of the
620 following systems: (a) FLiP-in HEK293 cells (catalog number: R78007), induced with
621 Doxycycline for 24 hours, used for inserts in pTH13195; (b) PURExpress T7
622 recombinant IVT system (NEB Cat.#E6800L), for inserts in pTH6838; or (c) SP6-driven
623 wheat germ extract-based IVT (Promega Cat#L3260), for inserts in pTH16500.

624 **DNA binding assays.** We followed previously-described methods for ChIP-seq[10],
625 PBMs[32], and SMiLE-seq[9]. Detailed descriptions of GHT-SELEX, HT-SELEX, ChIP-seq,
626 and SMiLE-seq data collection and initial analysis are found in the accompanying
627 papers (**Table S1**). For PBMs, we analyzed proteins on two different PBM arrays (HK
628 and ME), with differing probe sequences[76].

629 **Data processing and motif derivation.** The accompanying paper[17] describes motif
630 derivation and evaluation in detail. Briefly, after initial data processing steps, we
631 obtained a set of 'true positive' (likely bound) sequences for each individual experiment.
632 (721 / 4,873) experiments were removed at this step, due to a low number of peaks, or
633 other technical issues, as documented in **Table S5**). We then applied a suite of tools to
634 a training subset of the data from each experiment, and tested the resulting motifs on a
635 test subset of the data from the same experiment, and also on the independent data for
636 the same TF (i.e. the test sets from all other experiments done for the same TF). We
637 employed a binary classification regime for all experiments and all motifs, and scored
638 the motifs by a variety of criteria such as the areas under the receiver operating
639 characteristic (AUROC) or the precision-recall curve (AUPRC).

640 **Experiment evaluation by expert curation.** To gauge the success of individual
641 experiments, we employed an "expert curation" workflow with an initial voting scheme in
642 which a committee of annotators gauged whether individual experiments should be

643 “approved”, i.e. included in subsequent analyses. All experiments were examined by at
644 least three annotators. A subcommittee (AJ, IVK, and TRH) jointly resolved all cases of
645 disagreement among initial annotators (~300 experiments), and then reviewed all
646 approved experiments. Annotators had available an early version of the MEX portal
647 (https://mex.autosome.org) containing results of all PWMs scored against all
648 experiments, and were tasked with gauging whether the experiments yielded PWMs
649 that were similar across experiments, or scored highly across experiments. Annotators
650 also considered whether the motif was consistent with those for other members of their
651 protein family (e.g. BHLHA9 yielded an E-box-like motif, CAnCTG), and/or similar
652 between closely related paralogs (e.g. ZXDA, ZXDB, and ZXDC all yielded similar
653 motifs). We also considered whether (and how many) “peaks” were obtained from ChIP-
654 seq or GHT-SELEX, and whether these peaks were common to independent
655 experiments (e.g. both ChIP-seq and GHT-SELEX). Annotators were further given a
656 measure of similarity between Codebook PWMs and any PWMs in the public domain,
657 as well as enrichment of known or suspected common contaminant motifs in any
658 experiment.

659 **Post-evaluation peak processing.** After identification of “approved” experiments, we
660 re-derived peaks sets for ChIP-seq and GHT-SELEX experiments in order to obtain a
661 single peak set for each TF, as described in the accompanying papers[14,15]. Briefly, for
662 ChIP-seq we repeated the peak calling using MACS2 and experiment-specific
663 background sets, using a procedure previously described[10], then merged the peak sets
664 for replicates of the same TF with BEDTools merge[77] (see accompanying manuscript[15]:
665 “ChIP peak replicate analysis and merging”). We derived GHT-SELEX peaks using a
666 novel method that calculates enrichment of reads in each cycle, and treats different
667 experiments as independent statistical samples in order to obtain a single enrichment
668 coefficient per peak[14].

669 **Expert motif curation.** For this study, to identify a single representative PWM for each
670 TF, we first compiled a set of highest-scoring candidate PWMs for each TF (as
671 summarized above and elsewhere[17], then ran additional tests with them, utilizing the
672 reprocessed peak data, and manually evaluated the outputs. We first took the union of
673 three sets of 20 PWMs for each TF: the 20 PWMs with the highest AUROC (as
674 calculated elsewhere[17]) on (i) any approved ChIP-seq experiment for the given TF, (ii)
675 any approved GHT-SELEX experiment for the given TF, and (iii) any approved HT-
676 SELEX experiment for the given TF. These PWMs were selected regardless of the data
677 set from which they were derived. We then reassessed these PWMs against ChIP-seq
678 and GHT-SELEX data with two parallel methodologies. First, we recalculated AUROC
679 for each of the candidate top PWMs on the merged, thresholded sets of ChIP-seq
680 peaks ($P < 10^{-10}$)[15] using AffiMX[28] to score each peak. We generated negative sets
681 using BEDTools shuffle[77] with the *-noOverlapping* option to create sets of random
682 genomic regions with the same number of peaks, and the same peak width distribution
683 as the corresponding ChIP peak sets. We used the same technique to calculate
684 AUROC values for GHT-SELEX, with thresholded peak sets (using a “Kneedle”[78]
685 specificity value of 30 in the sorted enrichment values[15]). In parallel, we calculated the
686 Jaccard index to measure the overlap between PWM hits (identified by MOODS[79] with -
687 p 0.001) vs. the ChIP-seq peaks, and GHT-SELEX peaks, as two separate

688 measures. The overlap in each case was maximized by applying different thresholds on
689 the peak sets and choosing the cutoff at which the Jaccard index was the highest[14]. We
690 then applied expert curation (by a committee consisting of AJ, TRH, AF, KUL, RR, MA,
691 and IY) to choose a single representative PWM with high performance on all compiled
692 scores that, all else equal, also reflects reasonable expectation from the DBD class
693 (including recognition-code predicted motifs, see accompanying manuscript[14]) and has
694 high information content.

695 **Motif degeneracy analysis.** We adjusted the information content (IC) of PWMs on a
696 per-base-pair basis, with all locations boosted equally, by incrementally scaling weights
697 (e.g. probabilities in the PWM) until the PWM reached an adjusted to an average IC of 1
698 bit per base pair. The script, "logo_rescale.pl", is available at
699 https://gitlab.sib.swiss/EPD/pwmscan.

700 **Comparison to external peak sets and PWMs.** We downloaded comparison peak
701 sets from GTRD[60] and ENCODE (4.12.2023)[59], for all Codebook TFs. We then divided
702 this date into four categories corresponding to cell type: HEK293/HEK293T, HepG2,
703 K562, and other cells. Then, for each combination of TF and cell type category, we
704 selected a single peak set. We preferentially selected the peak sets from GTRD,
705 because it contains systematically derived peak sets; we also note that GTRD contains
706 the majority of ENCODE consortium experiments, together with many non-ENCODE
707 experiments. When multiple experiments were available for a TF in a cell type category,
708 we selected the experiment with higher counts. If multiple computational methods had
709 been used to derive peak sets for the selected experiment, we chose the peak set using
710 a preferential order MACS, GEM, SISSRS, PICS and PEAKZILLA. See **Table S7** for
711 identifiers and metadata of the reference datasets.

712 For PWM scoring, the external peak sets were used as downloaded, with the exception
713 of peak sets that were generated with the GEM peak caller, which have a peak width of
714 1, and were therefore expanded 250 bases in both directions. For Codebook data, we
715 used the merged and thresholded Codebook ChIP peak sets as in "Expert motif
716 curation". We generated negative peak sets for each ChIP-seq peak set using
717 BEDTools shuffle[77] with the *-noOverlapping* option to create sets of random genomic
718 regions with the same number of peaks and the same peak width distribution as the
719 corresponding ChIP peak sets. We downloaded PWMs for all Codebook TFs from
720 JASPAR[80] (2024 version), HOCOMOCO[13] (Version 12) and Factorbook[12] (downloaded
721 15.12.2023). We scanned Codebook and external peak sets (and corresponding
722 negative sets) with the expert curated Codebook motifs as PWMs using AffiMX[28], and
723 calculated AUROC values. Additionally, for the 19 Codebook TFs with a successful
724 Codebook ChIP-seq experiment, a Codebook PWM, an external ChIP-seq experiment,
725 and an external PWM, we compared the performance of PWMs across the different
726 peak sets as follows. We first selected a single external PWM for each of the 19 TFs by
727 scanning each PWM for a given TF on each external peak set for the same TF and
728 identifying the PWM that produced the highest AUROC. We then used these highest
729 scoring PWMs to scan the corresponding Codebook data and calculate AUROC values.

730 **TOP (Triple Overlap) and CTOP (Conserved Triple Overlap) peak set analyses.** To
731 obtain TOP sites, we first identified thresholds for ChIP-seq peaks, GHT-SELEX peaks,
732 and PWM score "peaks" that maximize the three-way Jaccard metric (overlap/union) of
733 the three sets, with the thresholds calculated for each TF independently. We converted
734 PWM hits (derived from MOODS[79] using a p-value cut-off of 0.001) into peaks by
735 merging neighboring matches with a distance less than 200bp and re-scoring them
736 using the sum-of-affinities for clusters. We then identified TOPs were as peaks
737 exceeding these thresholds in all three sets, and overlap in all three sets. To obtain
738 CTOP sites, we then extracted PhyloP scores for each base at each TOP site (and 100
739 flanking bases) from the Zoonomia consortium[81], removed sites overlapping the
740 ENCODE Blacklist[82] or protein coding sequences (due to the skew in phyloP scores
741 caused by codons), and applied three different statistical tests for significance of phyloP
742 scores over the PWM hit: two that test correlation between the IC and the phyloP value
743 at each base position of the PWM (using either Pearson correlation or linear
744 regression), and one that tests for higher phyloP scores over the PWM hit (Wilcoxon
745 test). Greater detail on these specific operations is given in the accompanying
746 manuscripts[14,15].

747 **Intersection of TOPs/CTOPs and genomic features.** We first clustered all CTOPs
748 using BEDTools merge[77], with a max distance of 100 bp, then intersected with the
749 following genomic feature sets: basic canonical protein coding promoters from
750 GENCODE version 44[83], defined as 1000 bp upstream and 500 bp downstream of the
751 canonical TSS; the "Unmasked CpG Island" track, PhastCons Conserved Elements
752 from the Multiz 470 Mammalian alignment, and RepeatMasker track from UCSC[84];
753 ChromHMM HEK293 enhancers[15]. We classified promoters as CpG island or non-CpG
754 island based on the GENCODE basic TSS being within +/- 50 bp of a CpG island from
755 the unmasked track. We classified the CTOP clusters as associated with a single type
756 of genomic feature in the following order of priority: CpG island associated with a protein
757 coding promoter; other CpG islands; a non-CpG island-associated protein-coding
758 promoter; an enhancer; containing a CTCF binding site but not overlapping a CpG
759 island, promoter or enhancer; overlapping a transposable element and none of the
760 previous categories; overlapping a non-TE repeat and none of the prior categories; and
761 "Other" for CTOP clusters not intersecting any examined features.

762 **SNV analyses.** *TOPs and CTOPs*. For analysis of common variants, we intersected
763 TOPs with the common short variants from dbSNP version 53, defined as a minor allele
764 frequency of >= 1% in the 1000 Genomes project[85]. We determined genomic overlap
765 enrichment between CTOPs/unconserved TOPs and dbSNP variants using the Fisher's
766 Exact Test implemented in BEDTools[77].

767 **Variant calling for allele-specific binding analysis.** We performed variant calling on
768 our GHT-SELEX and ChIP-seq datasets by mapping raw ChIP-Seq and pre-trimmed
769 GHT-SELEX reads[17] for 207 TFs to the hg38 human genome assembly using *bwa-mem*
770 (v.0.7.1) with default settings (workflow is shown in **Figure S5A**). Next, we used
771 *filter_reads.py* from *stampipes* (https://github.com/StamLab/stampipes/tree/encode-
772 release/, accessed Sept 2022) to filter out reads with >2 mismatches and mapping
773 quality <10. Then, we used a previously-described approach[86] for SNV calling and read

774 counting: (1) *samtools reheader* (v.1.16.1) was used to set the identical sample SM field
775 in all alignment files; (2) SNP calling was performed using *bcftools mpileup* (v.1.10.2)
776 with *--redo-BAQ --adjust-MQ 50 --gap-frac 0.05 --max-depth 10000* and *bcftools call*
777 with *--keep-alts --multiallelic-caller*; (3) the resulting SNPs were split into biallelic records
778 using *bcftools norm* with *--check-ref x -m* - followed by filtering with *bcftools filter -i*
779 *"QUAL>=10 & FORMAT/GQ>=20 & FORMAT/DP>=10" --SnpGap 3 --IndelGap 10* and
780 *bcftools view -m2 -M2 -v snps* leaving only biallelic SNPs covered by 10 or more reads;
781 (4) SNPs were annotated using *bcftools annotate* with *--columns ID,CAF,TOPMED* and
782 dbSNP (v.151)[87] (5) heterozygous variants located on the reference chromosomes with
783 GQ ≥20, depth ≥10, and allelic counts ≥5 on each allele were filtered with *awk* (v.5.0.1),
784 (6) *WASP* (v.0.3.4)[88] was used with *bwa mem* and *filter_reads.py* to account for
785 reference mapping bias, (7) *count_tags_pileup_new.py* was used to obtain allelic read
786 counts with *pysam* (v.0.20.0), (8) *recode_vcf.py* was used to convert the resulting BED
787 files to VCF. In total, we made 925,003 candidate variant calls supported by five reads
788 for both alleles and listed in the dbSNP common subset[87].

789 **ASB calling and analysis.** ASB calling was performed independently for GHT-SELEX
790 and ChIP-seq data. To account for aneuploidy and copy-number variation, the profiles
791 of relative background allelic dosage were reconstructed with BABACHI (v.2.0.26) using
792 default settings ([89], Abstract O3). The allelic imbalance was estimated with MIXALIME
793 (v.2.14.7)[68] starting with *mixalime create*. Next, we fitted a marginalized compound
794 negative binomial model (MCNB) using *mixalime fit* specifying MCNB and setting --
795 *window-size* to 1000 and 10000 for GHT-SELEX and ChIP-Seq, respectively, taking into
796 account lower coverage and SNP counts in GHT-SELEX. Finally, we used *mixalime test*
797 followed by TF-wise *mixalime combine* to obtain the TF-specific ASB calls (**Figure**
798 **S5A**).

799 We then identified ASBs that overlap a PWM hit (P-value < 0.001) for the associated
800 TF. For those ASBs, we calculated the PWM score for both alleles and estimated the P-
801 value of those scores against a uniform background distribution for each allele using
802 PERFECTOS-APE[90]. The fold-change between allele P-values (P1/P2) was then
803 calculated with the P-value of the more abundant allele as the numerator. ASBs with a
804 log2(fold-change) >=1 were labelled "strongly concordant", i.e., the allele we observed
805 to be bound more often is consistent with the PWM score (**Figure S5B**).

806 To assess the enrichment of Codebook ASBs within GTEx eQTLs[67] and ADASTRA
807 ASBs[66] we combined the ASB P-values from ChIP-Seq and GHT-SELEX data across all
808 TFs and datasets (*logitp* method[91]) to generate a single P-value for each TF (**Figure**
809 **S5C**).

810 **Analysis of allele-specific chromatin accessibility.** In this analysis, we relied on 321
811 and 577 cell type-specific chromatin accessibility datasets derived from DNase- and
812 ATAC-Seq experiments, respectively, and available in the UDACHA database (Release
813 IceKing 1.0.3)[68]. We identified 4,048 instances in which ASVs in a specific cell type
814 overlap significantly with PWM hits (P<0.0005) for a TF in the Codebook motif collection
815 (236 PWMs) (Right-tailed Fisher's exact test P < 0.05, and requiring 10 or more
816 overlapping PWM hits) (**Figure S5D**). Then, for each ASV in each combination of TF

817 and cell type passing the PWM enrichment filter, we asked whether the change in the
818 PWM score is concordant with the read imbalance in the ASVs, e.g. whether a higher
819 PWM score at a given locus corresponds to a higher read count, and we assigned a P-
820 value for each combination of TF and cell type, using a right-tailed Fisher's exact test,
821 including only sites with at least two-fold change in PWM-predicted affinity. Finally, to
822 obtain a single significance estimate per TF, we combined these P-values for each TF
823 across the different cell types passing the first stage, i.e. for which the overlap between
824 PWM hits and ASVs is significant (Fisher's method, considering DNase-Seq and ATAC-
825 Seq data separately and FDR-adjusted). TFs passing FDR < 0.05 in the final stage
826 were considered ASV-concordant.

827 To further verify the concordance between ASVs and Codebook motifs, we selected 34
828 (out of 53 TFs) with at least one TOP region overlapping ASVs, and re-evaluated the
829 concordant-to-discordant ratio for ASVs within peaks and TOP regions (see **Results**
830 and **Figure 5C**). For this analysis, for each TF, we picked the most significant ASV at
831 each unique genomic position (SNP) from all available cell types, and performed a right-
832 tailed Fisher's Exact Test (**Table S9**). At this stage, we considered SP140 and SP140L
833 jointly they share short and highly similar DNA-binding motifs.

DATA AVAILABILITY

The sequencing raw data for the HT-SELEX and GHT-SELEX experiments have been deposited into the SRA database under identifiers PRJEB78913 (ChIP-seq), PRJEB76622 (GHT-SELEX), and PRJEB61115 (HT-SELEX). Genomic interval information generated for the GHT-SELEX and ChIP-seq have been deposited into GEO under accessions GSE280248 (ChIP-seq) and GSE278858 (GHT-SELEX). PWMs can be browsed at https://mex.autosome.org and downloaded at https://doi.org/10.5281/ZENODO.8327372. An updated list of human TFs is available at https://humantfs.ccbr.utoronto.ca. Information on constructs, experiments, analyses, processed data, comparison tracks, and browsable pages with information and results for each TF is available at https://codebook.ccbr.utoronto.ca.

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DECLARATION OF COMPETING INTERESTS

O.F. is employed by Roche.

872 **SUPPLEMENTARY TABLES**

873 **Table S1. Accompanying manuscripts.** Table lists the 5 studies performed by the
874 Codebook Consortium, providing basic information for each of the manuscripts,
875 including title and author list.

876 **Table S2. TF list and assay success.** Table lists the Codebook proteins and positive
877 control TFs that were analyzed in the Codebook studies and provides metadata and
878 information on whether they showed sequence-specific DNA binding activities in
879 different types of experiments, together with the ID of the representative PWM selected
880 in this study, if any.

881 **Table S3. List of inserts used in this study.** Table provides the amino acid sequence
882 and type (full-length or DBD) for the 716 inserts used in the Codebook studies.

883 **Table S4. List of plasmids used in this study.** Table lists the plasmid backbone and
884 insert for each of the 1,387 plasmids used in the Codebook studies.

885 **Table S5. List of experiments performed in this study.** Table lists the 4,873
886 experiments performed on Codebook and control TFs, along with 20 additional GFP
887 control experiments. The experiment ID, experiment type, TF assayed, expert curation
888 result, and plasmid ID are listed for each experiment. Each experiment is mapped to its
889 ID in an accompanying manuscript[17], and 9 additional experiments used only in an
890 accompanying manuscript[17] are listed.

891 **Table S6. Representative PWMs.** Table shows logo representations for the PWMs
892 that were selected as the representative for each of the TFs (i.e. the expert-curated
893 motifs) and provides metadata describing the role of the TF in the study, DBD that it
894 belongs to, source of the experimental data and motif derivation approach.

895 **Table S7. External peak datasets**. Table lists external peak location datasets obtained
896 from GTRD database and ENCODE consortium, that were used in the comparisons
897 carried out in this study.

898 **Table S8. External PWM datasets**. Table lists PWM identifiers, manual curation and
899 other metadata for external motifs available from the databases Jaspar, HOCOMOCO
900 and Factorbook.

901 **Table S9. ASE and ASV data.** Allele-specific binding sites detected in Codebook data
902 and motif annotation of allele-specific chromatin accessibility events.

903 **Table S10. Updated census of human transcription factors and their motif**
904 **coverage**. Table is modified from Lambert et al. to display an updated motif coverage of
905 human TFs.

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A

ARID1 Factorbook ENCSR491EBY
YVSYGCCMYCTGSTG

CTCF(CisBP)

KDM5B Factorbook ENCSR000AQA
GCCGCCATCTY

YY1 (CisBP)

DRAP1 Factorbook ENCSR765MKZ
YKSYSATTGGYYSN

NFY (CisBP)

GATAD2A Factorbook ENCSR160QYK
RTKRTGCAAYM

CEPBP (CisBP)

GATAD2A Factorbook ENCSR925BFV
HWRWGYAAACA

Forkhead (FOXO1, CisBP)

MBD1 Factorbook ENCSR396QWK
CGCTGTCCRYGGTGCTGAA

REST (CisBP)

B

GATAD2A Factorbook ENCSR160QYK
WGATAAGV

GATAD2A (Codebook)

ZBED2 Jaspar MA1971.1

ZBED2 (Codebook)

ZNF623 Jaspar UN0210.1

ZNF623 (Codebook)

C

ZNF592 Factorbook ENCSR701AQS
AGYRACTCCATCTTG

No result in codebook,
validated in multiple cells

ZNF577 HOCOMOCO
ZN577.H12CORE.0.P.B

Not tested in Codebook, specific
and consistent for a C2H2-zf

GATAD2B Factorbook ENCSR547LKC
HNNDNWNYCTTATCTVYHNHY

No result in codebook,
consistent with orthologs

Figure S1. Examples of evaluation of external PWMs. A, Cases in which the external PWM matches that of a well-studied TF that is a frequent "contaminant" motif in ChIP-seq[97]. In each example, the top sequence logo represents the external PWM, and the bottom sequence logo represents a highly-similar CisBP PWM. **B,** Cases in which the external PWM (top in each example) is consistent with the Codebook PWM for the same TF (bottom in each example). **C,** External PWM sequence logos that cannot be explained as known contaminants or artifacts, some of which are supported by multiple lines of evidence, and thus appear accurate.

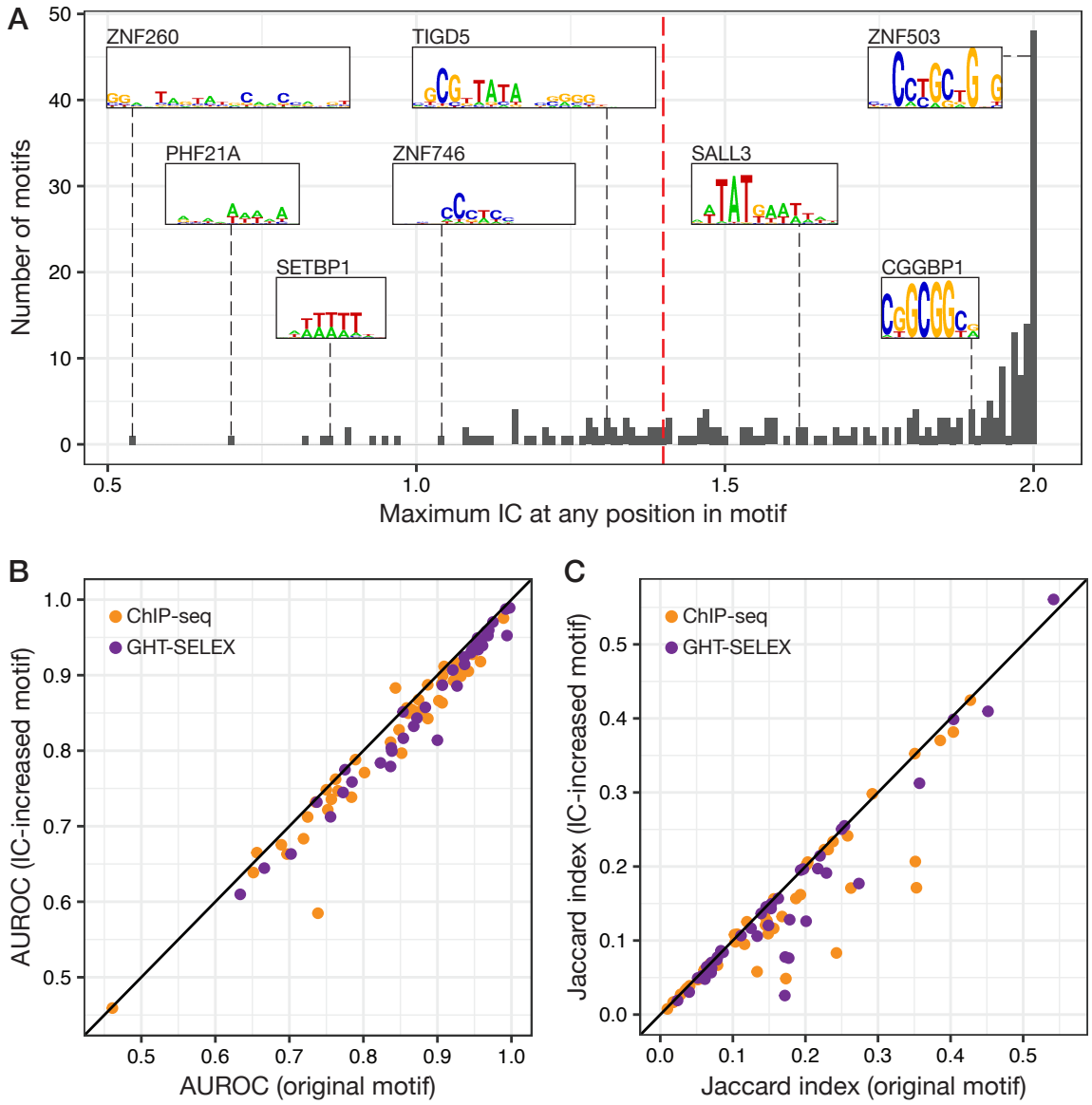
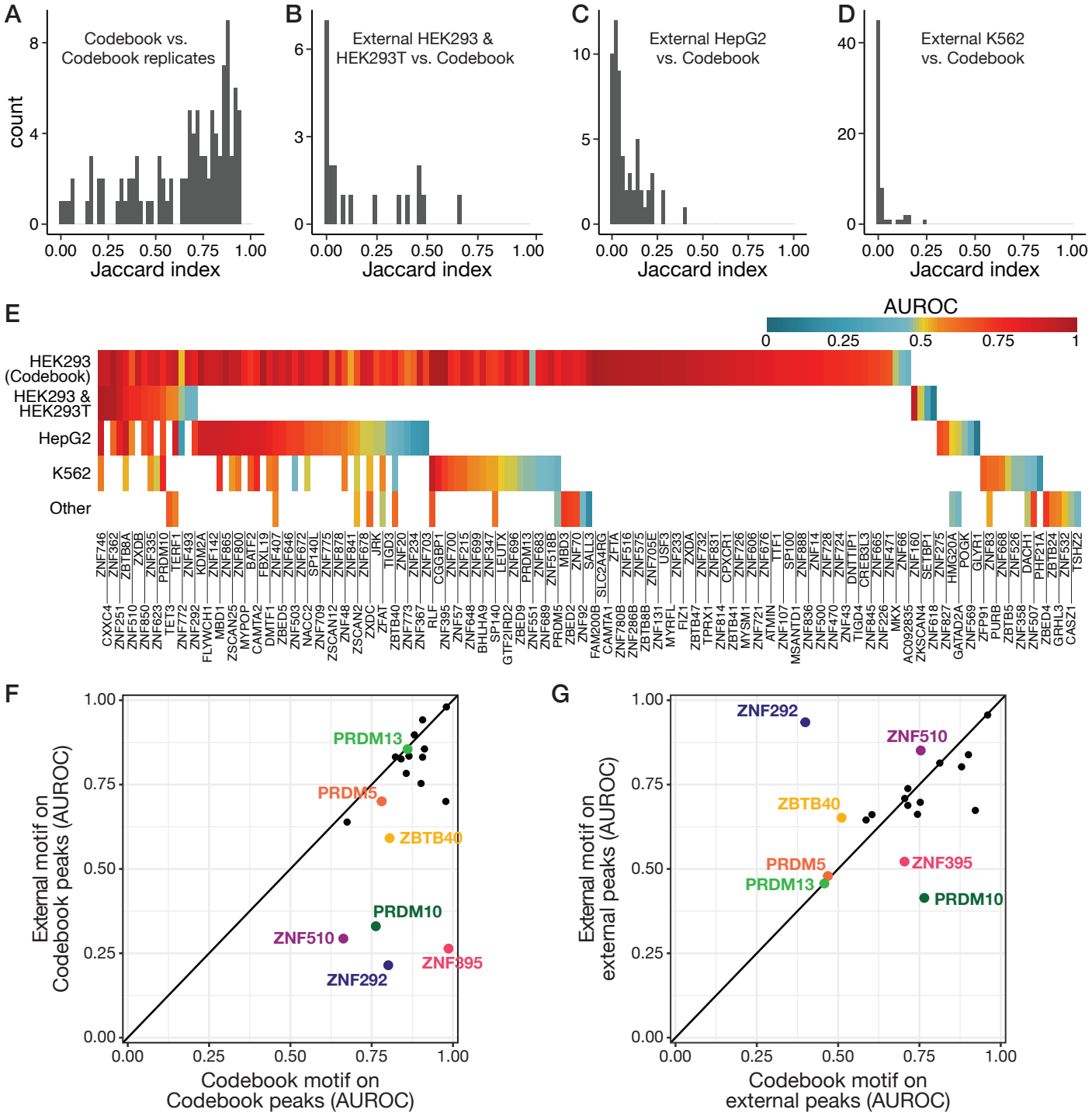

A
ZNF260
TIGD5
ZNF503
PHF21A
ZNF746
SALL3
SETBP1
CGGBP1
Number of motifs
Maximum IC at any position in motif
B
ChIP-seq
GHT-SELEX
AUROC (IC-increased motif)
AUROC (original motif)
C
ChIP-seq
GHT-SELEX
Jaccard index (IC-increased motif)
Jaccard index (original motif)

Figure S2. Motif degeneracy analysis. A, Histogram displays the maximum information content (IC) for any position within the expert-curated PWM for all Codebook and control TFs. Logos are shown for TFs at various maximum positional IC values, for illustration. Red dashed line indicates an IC of 1.4. **B,** and **C,** comparison of original PWMs to IC-increased PWMs for the 52 TF PWMs for which no base position exceeded an IC of 1.4. **B,** AUROC scores for original vs. IC-increased PWMs, discriminating ChIP-seq or GHT-SELEX peaks vs. random genomic background loci. **C,** Maximum Jaccard index for ChIP-seq or GHT-SELEX peak sets; using the approach described for optimized TOPs in **Methods**, for original vs. IC-increased PWMs.


A
Codebook vs. Codebook replicates
count
Jaccard index
B
External HEK293 & HEK293T vs. Codebook
Jaccard index
C
External HepG2 vs. Codebook
Jaccard index
D
External K562 vs. Codebook
Jaccard index
E
AUROC
HEK293 (Codebook)
HEK293 & HEK293T
HepG2
K562
Other
F
PRDM13
PRDM5
ZBTB40
PRDM10
ZNF510
ZNF395
ZNF292
External motif on Codebook peaks (AUROC)
Codebook motif on Codebook peaks (AUROC)
G
ZNF292
ZNF510
ZBTB40
PRDM5
PRDM13
ZNF395
PRDM10
External motif on external peaks (AUROC)
Codebook motif on external peaks (AUROC)

H

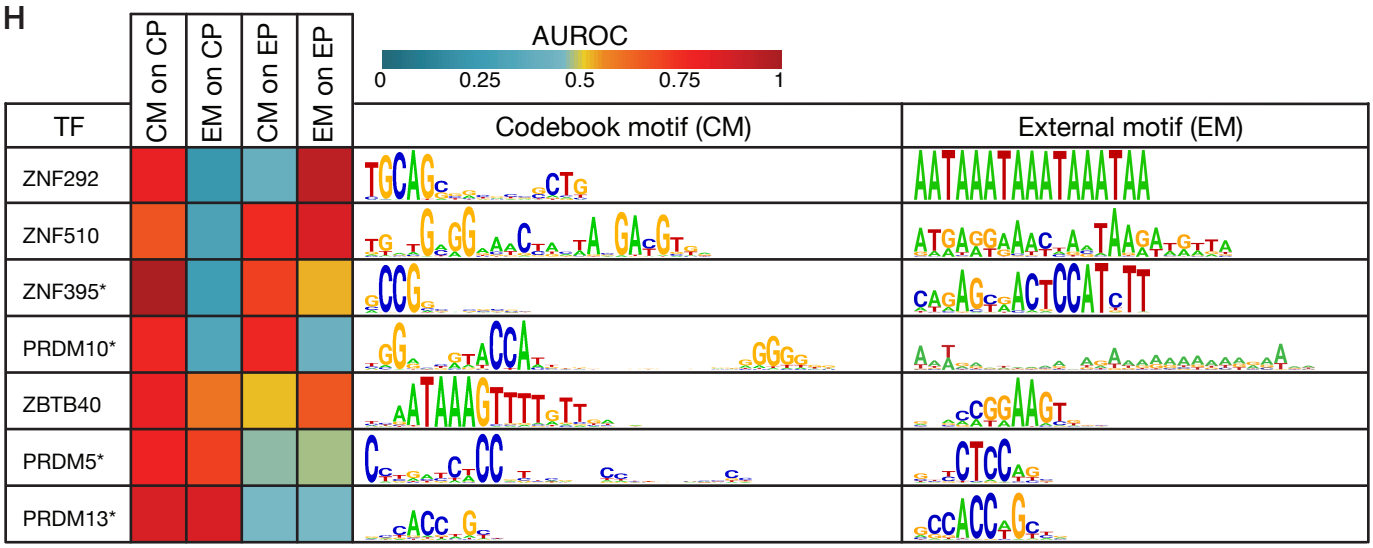

AUROC
0
0.25
0.5
0.75
1
TF
CM on CP
EM on CP
CM on EP
EM on EP
Codebook motif (CM)
External motif (EM)
ZNF292
ZNF510
ZNF395*
PRDM10*
ZBTB40
PRDM5*
PRDM13*

Figure S3. Comparison to external ChIP-seq datasets and PWMs. A-D, Histograms of Jaccard indices measuring the overlap between two ChIP-seq peak sets for the same TF: **A,** Codebook ChIP-seq replicates; **B, C, D**: Codebook ChIP-seq vs. external ChIP-seq performed in HEK293 cells (**B**), HepG2 cells (**C**), or K562 cells (**D**). **E,** AUROC scores for expert curated Codebook PWMs (columns), discriminating ChIP-seq peaks vs. random genomic background loci. Rows show different cell types. **F, G,** comparison of Codebook and external PWMs at the task of discriminating ChIP-seq peak sets from random sequences (as in **E**), for the 19 TFs that have a Codebook peak set (CP), a Codebook motif (CM), an external peak set (EP), and an external motif (EM), for Codebook ChIP-seq data (**F**) and external ChIP-seq data (**G**). The seven TFs with an AUROC of < 0.55 on either axis of either plot are highlighted. **H,** Sequence logos for the seven TFs highlighted in **F** and **G**. All Codebook PWMs shown are supported by ChIP-seq, GHT-SELEX, and HT-SELEX. Asterisk indicates that the Codebook PWM is additionally supported by SMiLE-seq data.

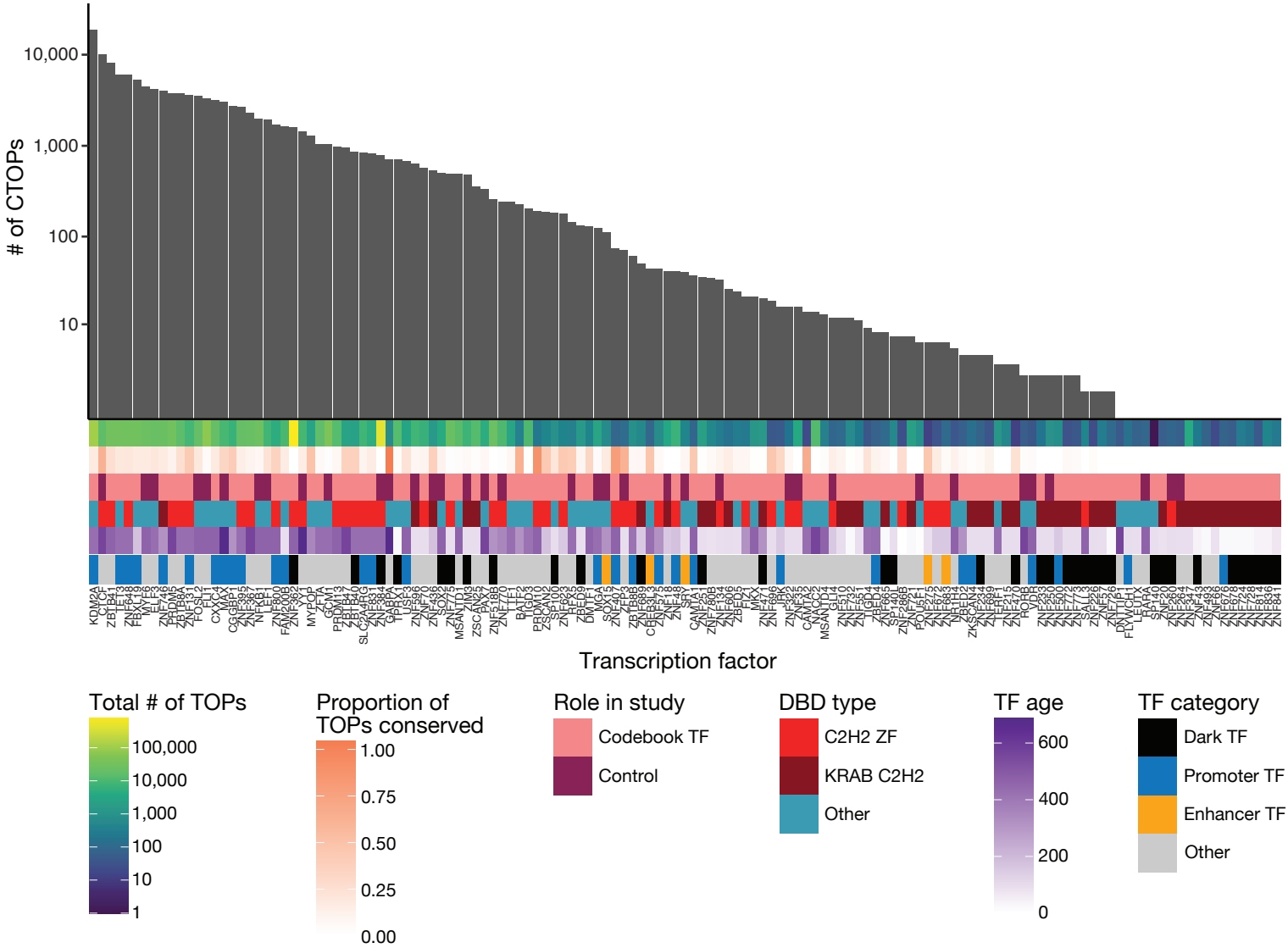

# of CTOPs
10,000
1,000
100
10
Transcription factor
Total # of TOPs
100,000
10,000
1,000
100
10
1
Proportion of TOPs conserved
1.00
0.75
0.50
0.25
0.00
Role in study
Codebook TF
Control
DBD type
C2H2 ZF
KRAB C2H2
Other
TF age
600
400
200
0
TF category
Dark TF
Promoter TF
Enhancer TF
Other

Figure S4. Number of CTOP sites per TF. Bar graph displays the number of individual CTOP sites obtained for each TF. Heatmap and annotations below indicate other properties of each TF and its TOP sites.

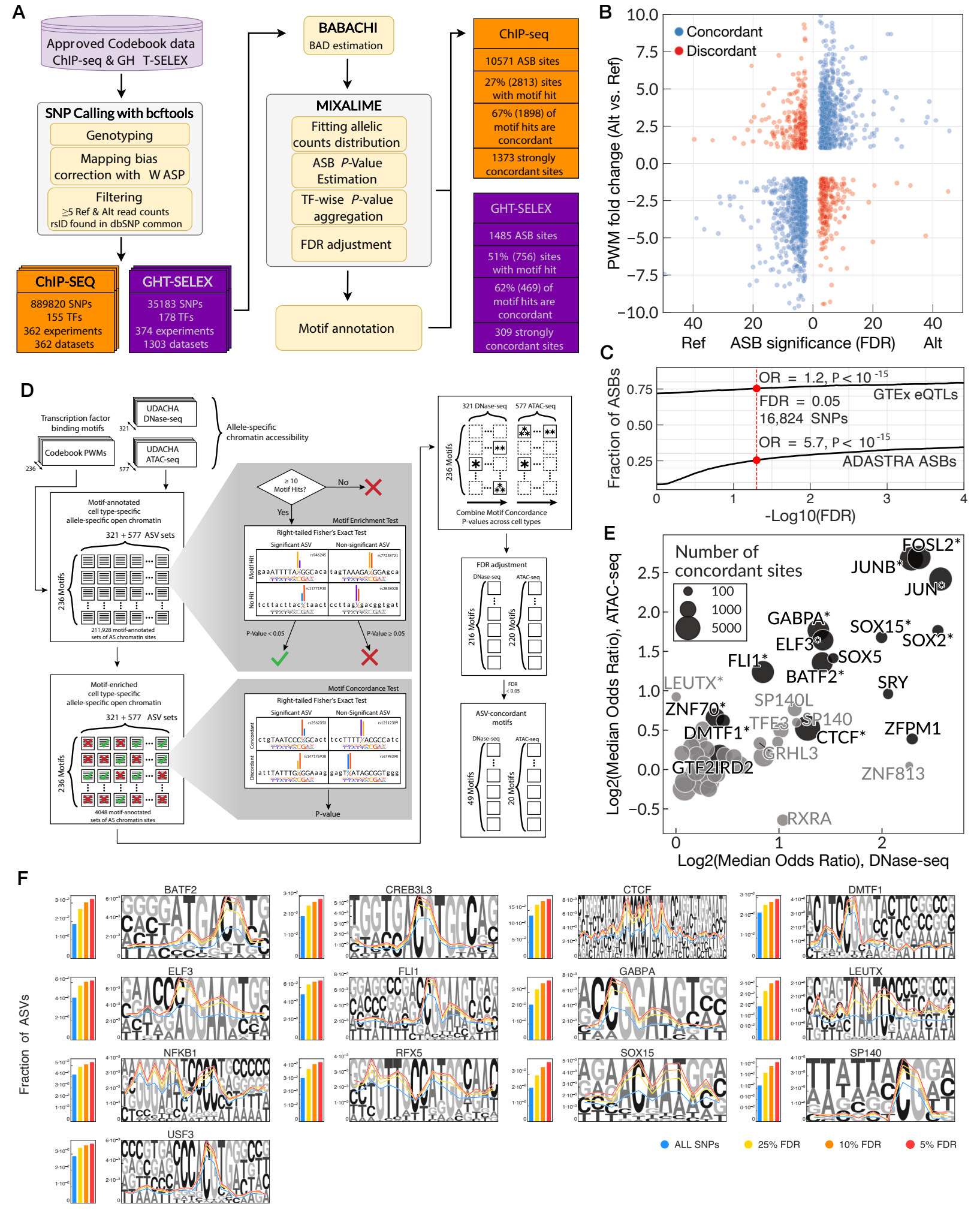

A
Approved Codebook data
ChIP-seq & GH T-SELEX
SNP Calling with bcftools
Genotyping
Mapping bias correction with W ASP
Filtering
≥5 Ref & Alt read counts
rsID found in dbSNP common
ChIP-SEQ
889820 SNPs
155 TFs
362 experiments
362 datasets
GHT-SELEX
35183 SNPs
178 TFs
374 experiments
1303 datasets
BABACHI
BAD estimation
MIXALIME
Fitting allelic counts distribution
ASB P-Value Estimation
TF-wise P-value aggregation
FDR adjustment
Motif annotation
ChIP-seq
10571 ASB sites
27% (2813) sites with motif hit
67% (1898) of motif hits are concordant
1373 strongly concordant sites
GHT-SELEX
1485 ASB sites
51% (756) sites with motif hit
62% (469) of motif hits are concordant
309 strongly concordant sites
B
Concordant
Discordant
PWM fold change (Alt vs. Ref)
Ref
ASB significance (FDR)
Alt
C
Fraction of ASBs
OR = 1.2, P < 10^-15
FDR = 0.05
16,824 SNPs
GTEx eQTLs
OR = 5.7, P < 10^-15
ADASTRA ASBs
-Log10(FDR)
D
Transcription factor binding motifs
Codebook PWMs
UDACHA DNase-seq
UDACHA ATAC-seq
Allele-specific chromatin accessibility
Motif-annotated cell type-specific allele-specific open chromatin
321 + 577 ASV sets
236 Motifs
211,928 motif-annotated sets of AS chromatin sites
≥ 10 Motif Hits?
No
Yes
Motif Enrichment Test
Right-tailed Fisher's Exact Test
Significant ASV
Non-significant ASV
Motif Hit
No Hit
P-Value < 0.05
P-Value ≥ 0.05
Motif-enriched cell type-specific allele-specific open chromatin
4048 motif-annotated sets of AS chromatin sites
Motif Concordance Test
Concordant
Discordant
P-value
321 DNase-seq
577 ATAC-seq
Combine Motif Concordance P-values across cell types
FDR adjustment
DNase-seq
ATAC-seq
216 Motifs
220 Motifs
FDR < 0.05
ASV-concordant motifs
49 Motifs
20 Motifs
E
Number of concordant sites
100
1000
5000
FOSL2*
JUNB*
JUN*
GABPA*
SOX15*
SOX2*
ELF3*
SOX5
FLI1*
BATF2*
SRY
LEUTX*
SP140L
ZNF70*
TFE3
SP140
ZFPM1
DMTF1*
CTCF*
GRHL3
GTF2IRD2
ZNF813
RXRA
Log2(Median Odds Ratio), ATAC-seq
Log2(Median Odds Ratio), DNase-seq
F
BATF2
CREB3L3
CTCF
DMTF1
ELF3
FLI1
GABPA
LEUTX
NFKB1
RFX5
SOX15
SP140
USF3
Fraction of ASVs
ALL SNPs
25% FDR
10% FDR
5% FDR

Figure S5. Identifying allele-specific TF binding in HEK293 cells and analyzing allele-specific chromatin accessibility events using Codebook motifs.
A, Codebook ASB calling workflow: SNP calling with bcftools, mapping bias correction with WASP, background allelic dosage reconstruction with BABACHI, statistical scoring of the allelic imbalance with MIXALIME, and motif annotation with PERFECTOS-APE. **B,** Motif concordance of Codebook ASBs. X-axis: ASB significance (i.e., allelic preference; $\log_{10}$ FDR, *minus side*: preference for Ref, *plus side*: preference for Alt). Y-axis: $\log_2$ PWM score fold-change between Alt vs. Ref. The plot shows only strongly concordant and strongly discordant sites with |log2(Fold Change)| ≥ 1. **C,** Fraction of Codebook ASBs (combined) coinciding with GTEx eQTLs and ADASTRA known ASBs at different FDR thresholds for ASB calling. Fisher's exact test odds ratios (OR) and P-values for ASBs at 5% FDR (covering 16,724 SNPs, dashed line) are labeled on the plot. **D,** Workflow for detection of TFs involved in allele-specific chromatin accessibility. UDACHA DNase-seq and ATAC-seq ASVs across different cell types were annotated with Codebook motifs, followed by motif enrichment and motif concordance analysis, combining the resulting P-values across the cell types, and FDR correction for multiple tested motifs. Central call-outs: details of the motif enrichment and motif concordance test using SP140 motif for illustration. SNPs (rs946245, rs77238721, rs11771930, rs2838028, rs2562353, rs12112389, rs147176938, rs6798390) illustrating the cells of the 2x2 contingency tables are actual UDACHA ASVs with or without motif hits of selected TFs. **E,** Scatterplot of Median Odds Ratios of PWM scores within the ASVs enriched in and concordant with the PWM matches. Motifs significant for both DNase-seq and ATAC-seq (black), or just one assay (gray). The asterisk denotes TFs that exhibit significant enrichment considering peaks-supported PWM hits only. **F,** *Bar plots*: Fraction of ASVs overlapping with PWM hits for 13 TFs, using 4 different thresholds on ASV significance: all SNPs (blue), 25% FDR ASVs (yellow), 10% FDR ASVs (orange), and 5% FDR ASVs (red). *Line plots*: Fraction of ASVs at each location within the genome-wide PWM hits of the representative TFs (P-value<0.001) using four thresholds (the same colors as in bar plots). SNP: single-nucleotide polymorphism, ASB: allele-specific binding, ASV: allele-specific chromatin accessibility variant.