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A Census of Human Soluble Protein Complexes


SUMMARY

Cellular processes often depend on stable physical associations between proteins. Despite recent progress, knowledge of the composition of human protein complexes remains limited. To close this gap, we applied an integrative global proteomic profiling approach, based on chromatographic separation of cultured human cell extracts into more than one thousand biochemical fractions that were subsequently analyzed by quantitative tandem mass spectrometry, to systematically identify a network of 13,993 high-confidence physical interactions among 3,006 stably associated soluble human proteins. Most of the 622 putative protein complexes we report are linked to core biological processes and encompass both candidate disease genes and unannotated proteins to inform on mechanism. Strikingly, whereas larger multiprotein assemblies tend to be more extensively annotated and evolutionarily conserved, human protein complexes with five or fewer subunits are far more likely to be functionally unannotated or restricted to vertebrates, suggesting more recent functional innovations.

INTRODUCTION

Protein complexes are stable macromolecular assemblies that perform many of the diverse biochemical activities essential to cell homeostasis, growth, and proliferation. Comprehensive characterization of the composition of multiprotein complexes in the subcellular compartments of model organisms like yeast, fly, worm, and bacteria have provided critical mechanistic insights into the global modular organization of conserved biological systems (Hartwell et al., 1999), accelerated functional annotation of uncharacterized proteins via guilt by association (Hu et al., 2009; Oliver, 2000), and facilitated understanding of both evolutionarily conserved and disease-related pathways (Vidal et al., 2011). How the ~20,000 or so proteins encoded by the human genome are partitioned into heteromeric “protein machines” remains an important but elusive research question, however, as less than one-fifth of all predicted human open reading frames are currently annotated as encoding subunits of protein complexes in public curation databases (Ruepp et al., 2010).

Loss-of-function mutations in genes encoding the subunits of protein complexes typically give rise to similar phenotypes or, through genetic interaction, amplify the phenotypic effects of other alleles in functionally linked sets of genes. Identifying the membership of protein complexes, therefore, addresses a crucial layer in the hierarchical functional organization of biological systems that links the core biochemistry of a functioning cell to the general physiology of an organism and is fundamental to deciphering the relationship between genotype and phenotype. Although bioinformatics analyses have been used to predict evolutionarily conserved human protein-protein interactions (PPIs) on a large scale (Ramani et al., 2008; Rhodes et al., 2005), most of these associations remain to be verified experimentally.

Affinity purification (AP) of tagged exogenous proteins coupled with tandem mass spectrometry (MS) is an effective method for isolating and characterizing the composition of stably associated human proteins in experiments ranging from dozens to hundreds of different “baits” (Behrends et al., 2010; Bouwmeester et al., 2004; Ewing et al., 2007; Hutchins et al., 2010; Jeronimo et al., 2007; Mak et al., 2010; Sardiu et al., 2008; Sowa et al., 2009). Likewise, immunoprecipitation can be used...
to systematically isolate endogenous human protein complexes from human cell lines (Malovannaya et al., 2011). Nevertheless, the limited availability of high-quality antibodies or sequence-verified complementary DNA (cDNA) clones suitable for targeted protein complex enrichment precludes scale-up required for unbiased assessment of the molecular association networks underlying human cells. Hence, despite considerable successes in the comprehensive identification of protein complexes in model organisms (Butland et al., 2005; Gavin et al., 2002, 2006; Guruharsha et al., 2011; Ho et al., 2002; Hu et al., 2009; Krogan et al., 2006; Kühner et al., 2009), clone-based protein purification techniques remain challenging for proteome-scale studies of physical interaction networks in mammalian cells. Conversely, although traditionally used to isolate discrete complexes with specific assayable biochemical properties (e.g., enzymatic activity), classical biochemical fractionation procedures have been used to resolve biological mixtures as a means of ascertaining the collective composition of human protein complexes present in certain subcellular compartments (Ramani et al., 2008; Wessels et al., 2009).

Here, we have combined extensive, scaled-up biochemical fractionation with in-depth, quantitative mass spectrometric profiling and stringent computational filtering to resolve and identify endogenous, soluble, stably associated human protein complexes present in cytoplasmic and nuclear extracts generated from cultured cells. Although the resulting reconstructed high-quality physical interaction network shows strong overlap with existing curated and experimentally derived sets of annotated protein complexes, it contains many predicted subunits and previously unreported complexes with specific functional, evolutionary, and disease-related biological attributes. To our knowledge, this resource represents the largest experimentally derived catalog to date of human protein complexes from cell culture, measured using a single standardized assay, and a reliable first draft reference of the basic physical wiring diagram of a human cell.

RESULTS

High-Throughput Complex Fractionation and Detection by Tandem Mass Spectrometry

To isolate human protein complexes in a sensitive and unbiased manner, we subjected cytoplasmic and nuclear soluble protein extracts isolated from human HeLa S3 and HEK293 cells grown as suspension and adherent cultures, respectively, to extensive complementary biochemical fractionations. These two widely studied laboratory cell lines have been used as models of human cell biology for many decades (Graham et al., 1977; Masters, 2002), providing a rich biological context for interpreting the resulting proteomic data. Stably interacting proteins that cofractionated together were identified subsequently by nanoflow liquid chromatography-tandem mass spectrometry (LC-MS/MS). We optimized our entire experimental pipeline, illustrated schematically in Figure 1A, by using a multi-pronged strategy to minimize two major confounding issues: limited dynamic range (i.e., preferential detection of high-abundance components) and “chance” coelution (i.e., cofractionation of functionally unrelated proteins).

To address the former concern, we performed extremely deep biochemical fractionations by employing multiple orthogonal separation techniques to better resolve distinct protein complexes. As a primary separation technique, we employed nondeaturing high-performance multibed ion exchange chromatography (IEX-HPLC) by using four different empirically optimized analytical column combinations (see Experimental Procedures) and shallow salt gradients unlikely to perturb nonionic protein associations (Havugimana et al., 2007). In parallel, we applied complementary sucrose gradient centrifugation and isoelectric focusing technologies to capture salt-sensitive protein assemblies. In total, we collected 1,163 different fractions in a total of eight nuclear and five cytosolic extract fractionation experiments (for details see Table S1 available online), which were each subjected to label-free shotgun sequencing (duplicate LC-MS/MS analyses) using highly sensitive ion trap-based mass spectrometers (see Experimental Procedures).

We identified 5,584 distinct human proteins (Figure 1C; estimated theoretical false discovery rate of 1% at both the protein and peptide levels based on a statistical model [Kislinger et al., 2003]; see Experimental Procedures for details). Despite the underrepresentation of membrane proteins in the starting cell extracts, this coverage encompasses about half of the experimentally verified human proteome (Figure S1B) (Nagaraj et al., 2011). This included 989 proteins detected exclusively in nuclear fractions (of which 376 were annotated transcription or chromatin-related factors) and 1,006 with links to human disease (e.g., annotated in a public database like OMIM). Only 1,632 (29%) of the identified proteins had biochemical annotations as subunits of previously reported protein complexes (corresponding to 64% of all existing human protein entries) in the CORUM curation database (Figure S1C; Ruepp et al., 2010). Due to the extensive fractionation, we observed minimal bias in terms of protein abundance beyond that reported for previously annotated complexes or the experimentally defined human proteome (Figure 1D).

Next, to minimize the possibility of chance coelution, rather than simply identifying the proteins present in each fraction, we quantified variation in protein abundance based on the observed patterns of spectral counts recorded across all of the collected fractions to determine the extent to which pairs of proteins coeluted. As shown in Figure 1B, these experimental profiles were highly reproducible (i.e., average Spearman rank correlation coefficients >80% between replicate experiments; Figure S2), even using alternate methods of mass spectrometric quantification (i.e., extracted MS1 peak intensities were largely consistent with spectral counting; Figure S2D). To objectively evaluate the biochemical data, we calculated a stringent summary statistic, termed the coapex score, for each pair of proteins identified by LC-MS/MS by determining the number of fractionation experiments in which the proteins showed maximum (modal) abundance in the same exact peak fraction.

To assess the effectiveness of our cofractionation approach, we performed an initial validation by examining the coelution profiles and coapex scores obtained for a reference set of 20 well-known human protein complexes reported in CORUM. As illustrated by the representative HeLa nuclear extract IEX-HPLC profiles shown in Figure 1B, the subunits of these complexes...
complexes typically coeluted in the same biochemical fractions. Of the 155 components detected by mass spectrometry, most (85%; 499/585) of the detected subunit pairs of the reference complexes had high coapex similarity scores (i.e., coeluted together in at least two or more experiments), validating the overall efficacy of the fractionation procedures we used to isolate native protein complexes and the general correctness of the protein identification and quantification pipeline.

Reconstruction of a High-Confidence Cocomplex Interaction Network

Despite the consistency in coelution of annotated complex members, certain functionally distinct complexes occasionally exhibited overlapping chromatographic elution profiles (e.g., splicing factor 3b and Coatomer complexes; Figure 2A), presenting a potential source of spurious interactions. Although this artifact was minimized to a certain degree by performing multiple independent fractionation experiments, we used an integrative computational approach to further improve deconvolution (Figure 2B). Because physically interacting cocomplexed proteins often perform related biological functions (Alberts, 1998) and are often evolutionarily coconserved (Hartwell et al., 1999), we devised a machine learning procedure (Figure 2B; see Experimental Procedures for details) to score and select higher-confidence physical interactions based on both the experimentally measured coelution profiles and the existence of additional

Figure 1. Integrative Cofractionation Strategy Used to Identify Human Soluble Protein Complexes

(A) Cell extracts were extensively fractionated using different biochemical techniques (IEX, ion exchange chromatography; IEF, isoelectric focusing; SGF, sucrose density gradient centrifugation). Coeluting proteins were identified by mass spectrometry, and a coelution network was generated by calculating profile similarity (see Extended Experimental Procedures).

(B) Cofractionation (IEX-HPLC) profiles of annotated subunits of 20 representative human protein complexes from HeLa nuclear extract. Shading indicates normalized spectral counts (SPC). Peak apex and adjacent peaks are shown.

(C) Hierarchical clustering of 5,584 proteins identified by LC-MS/MS.

(D) Protein abundance levels corresponding to components of our identified coeluting proteins (red line), reconstructed complexes (blue), or annotated CORUM complexes (black) estimated from the reported HeLa proteome (Nagaraj et al., 2011). See also Figure S1 and Table S1.
supporting functional association evidence inferred from correlated evolutionary rates (Tillier and Charlebois, 2009) and functional genomics data sets compiled for H. sapiens, S. cerevisiae, D. melanogaster, and C. elegans (see Table S6 for details).

First, for each of the 13 fractionation experiments, we calculated correlation measures between all possible pairs of proteins to capture their tendency to coelute. In addition to the coapex summary statistic, to account for mass spectrometry sampling error, we devised a weighted cross-correlation function to account for slight variation in the protein profiles measured in each experiment. To account for low spectral values, we also generated a Poisson noise model before calculating Pearson correlation scores, deeming the coelution profiles of protein pairs measured with low spectral counts as less predictive of genuine physical interactions (Figure S5). Only protein pairs...
with a correlation score of at least 0.5 by at least one of these measures in one or more experiments were considered for further analysis, reducing the total number of pairs from over 15 million initially to the roughly 800,000 pairs with reasonable biochemical evidence.

To improve the assignment of interaction probabilities, we also exploited the predictive power of correlated protein evolutionary rates (Tillier and Charlebois, 2009), messenger RNA (mRNA) coexpression, and domain co-occurrence and, via orthology, fly protein-protein interactions (based on binary yeast two-hybrid assay studies) and extensive physical and functional associations reported previously for yeast and worm (see Experimental Procedures) (Lee et al., 2011). The discriminatory power of the procedure was further improved by penalizing those interactions that lacked independent supporting evidence—and that were thus more likely to correspond to cases of “chance” colocation—by integrating evidence from these functional association data (Figure 2B). A feature selection algorithm was used to select the most informative data sets (Table S2) in addition to the biochemical correlation scores, and the resulting features were used to estimate the probability of interaction to protein pairs using a cross-validated random forest classifier.

For training, we used the CORUM curated set of human protein complexes as our base reference, filtered for those complexes annotated as having been observed by biochemical methods. As many CORUM complexes are highly overlapping due to redundancy in existing annotations, we combined complexes sharing subunits (Simpson coefficient >0.5 between complexes). We used half of the resulting 324 nonredundant reference complexes (Table S3) as the training set for cocomplex probability prediction, defining gold standard positive interactions as pairs of proteins in the same complex and inferring gold standard negatives between proteins in different complexes. (The other half of the reference complexes was withheld for subsequent use as an independent training set for cluster optimization, as described below.)

Although the biochemical data were a prerequisite for scoring, the performance curves shown in Figure 2C indicate that the inclusion of the additional functional genomic information substantially increased recall at the same level of precision compared to classifiers based on the profiling data alone. Moreover, the integration of this additional supporting functional evidence removed the bulk of spurious, intercomplex interactions (Figure 2D). Another advantage of our bioinformatic pipeline is that the results of the feature selection algorithm (Table S2) can be explored to examine the impact of each data set. For example, we find generally that sets of smaller biochemical fractionations using different separation techniques, although individually yielding a higher PPI false discovery rate, collectively provided more information on protein complex composition than deeper fractionations using a single separation method.

As an alternate measure of reliability, we compared our scored human protein interactions to a recently reported network of Drosophila cocomplex protein interactions (Guruharsha et al., 2011), which had not been used for building the classifier. Strikingly, despite using vastly different experimental methods and scoring schemes, we observed a remarkably good overall correlation (Spearman r = 0.40; n = 11,675 orthologs mapped using Inparanoid). Even after removing interactions supported by alternate Drosophila data, high-scoring fly pairs matched high-scoring pairs in our analysis and were strongly enriched for reference-positive cocomplex members (Figure 2E).

Finally, in order to remove any remaining false positive interactions, we further denoised our cocomplex data set by pruning loosely connected interactions using a computational diffusion procedure calibrated by protein colocalization semantic similarity scores (Pesquita et al., 2009; Yang et al., 2012) to enforce local network topologies more consistent with annotated complexes from the withheld portion of the reference Corum complexes (see Experimental Procedures). Benchmark precision and recall versus the holdout set of known reference complexes (Figure 2F) were significantly higher than those reported for a smaller, recently published set of affinity-purified human protein complexes (Hutchins et al., 2010), validating the reliability of our scoring procedure.

Applying a PPI score threshold of 0.75, which corresponds to an estimated false discovery rate of 21.5% (i.e., well below the ~40% reported for AP/MS-based analyses of protein complexes in model organisms [Gavin et al., 2006; Krogan et al., 2006; Kühner et al., 2009]), we thus derived a high-confidence set of 13,993 cocomplex interactions among 3,006 unique human proteins (Table S2), most of which (8,691 PPI) have not been reported before (i.e., are not publicly annotated). It is worth reiterating that all of these physical interactions were directly supported by the experimental biochemical cofractionation data; the addition of functional data and denoising served only to flag candidates lacking either functional support or topological support within the network (Table S2). The interaction probability scores may be underestimated, however, because the reference “gold standards” used for learning are imperfect (Jansen and Gerstein, 2004).

**Construction and Validation of Protein Complexes from the Probabilistic Interaction Network**

In order to define complex membership, we partitioned the high-confidence probabilistic physical interaction network by using the cluster growth algorithm ClusterONE (Nepusz et al., 2012), which outperformed other clustering methods on the denoised PPI network (Table S5). In total, the clustering predicts 622 discrete putative complexes encompassing 2,634 distinct proteins (Table S3). Complex membership size distribution approximated an inverse power law with a median of four subunits (Figure S4A). The majority (62%; 385/622) of the complexes have not been annotated (i.e., only 237 are currently curated in a public database like CORUM; Figures 3A and 3C). Although the fraction of curated components varies, we also recapitulated 258 previously reported complexes (Figure 3C), including several well-known membrane-associated complexes, such as the coat protein I and II (COPI/II) vesicle transport complexes that shuttle cargo between the Golgi and endoplasmic reticulum. Strikingly, most (67%; 335) of the 500 smaller putative complexes with five or fewer components, including the bulk (74%; 83) of the 112 predicted heterodimers, have never been curated before (Figure 3C).

Both independent experimental validation based on more traditional immunoprecipitation or coaffinity purification methods
and orthology mapping support at least 21 of these putative complexes (i.e., not in any reference database) (Table S3; see Supplemental Information for details). For example, Guruharsha et al. (2011) recently reported 299 cocomplex interactions based on pull-down experiments of 43 affinity-tagged human proteins present in 41 of our complexes, of which 143 interactions map precisely to our predicted complexes, representing a 47.8% validation rate (which may be an underestimate, as Guruharsha et al. [2011] do not report human interactions that fall outside the fly interologs examined in their study). Likewise, the results of Malovannaya et al. (2011), who used large-scale immunoprecipitation to isolate native human protein complexes, show excellent agreement to 123 of our complexes (i.e., Benjamini-corrected hypergeometric p ≤ 0.05), including 42 (34%) of our complexes that are not curated in CORUM (Figure 3B and Table S3). Figure 3D summarizes the highly significant overlap of our inferred complexes with these fully independent data sets, with enrichments ranging from 4- to 477-fold more than chance, thus broadly and systematically validating our network of derived human protein complexes.

By design, insoluble membrane-associated (hydrophobic) protein complexes were largely missed in this study, and the proteins assigned to complexes had a higher average transcript abundance (Figures S2A and S2B). Moreover, in an effort to control the false positive rate, our conservative clustering algorithm, ClusterONE, underweighted small clusters of size 2 or 3 for lack of sufficient association evidence, likely contributing to the prominence of complexes with four subunits in Figure 3A. But we did not observe any significant bias toward negative (pI < 7) or positive (pI > 7) charge as compared to complexes curated in CORUM (Figure S4B).

Figure 4 shows the broad functional diversity of the predicted complexes (a navigable map is available online for close visualization of individual clusters and their supporting cocomplex interactions). Consistent with biological expectation (Hartwell et al., 1999; Lage et al., 2007; Oliver, 2006; Vidal et al., 2011),
Figure 4. Global Map of High-Confidence Human Protein Complexes

(A) Schematic of the global network of inferred human soluble protein complexes (colored by membership) with representative examples and supporting PPI highlighted.

(B) Putative complexes with two or more components with human disorder associations annotated in UniProt (UniProt Consortium, 2011), Online Inheritance of Man (OMIM) (Hamosh et al., 2005), or the Genetic Association Database (GAD) (Becker et al., 2004). Inset table shows highly significant interaction overlap.
the subunits of the complexes were significantly enriched for related biological functions, transcriptional regulatory motifs, and pathological processes (Figure 4B, inset table). Compared to the entire set of identified proteins, the clustered proteins also showed enrichment for posttranslation modifications linked to cellular regulation, like acetylation (Benjamini-corrected \( p \leq 10^{-41} \)) and phosphorylation (\( p \leq 10^{-5} \)). Many of the complexes are linked to core cellular processes, such as mRNA splicing (\( p \leq 10^{-15} \)) or transcription (\( p \leq 10^{-5} \)), that either are essential in human (\( p \leq 10^{-136} \)) or that have RNA interference (RNAi)-induced phenotype in cell culture (e.g., cell division arrest, \( p \leq 10^{-31} \)) or are associated, via orthology, with similar mouse, yeast, or worm mutant phenotypes (Figure 4B, inset table; see Table S4 for details).

(i.e., shared annotated edges) with phenotypic data sets that reveals that protein subunits of the same predicted human complex tend to exhibit similar disease and genetic associations in human populations (see Extended Experimental Procedures), RNAi phenotypes in cell culture (Neumann et al., 2010), mutational and RNAi phenotypes in other species (via orthology), and shared transcriptional regulatory motifs (Xie et al., 2005). See also Figure S4C and Table S4.

Clinical and Biological Implications of the Reconstructed Human Protein Complexes

Consistent with this strong tendency for proteins in the same complex to be affiliated with similar mutational and RNAi phenotypes, subunits of the predicted human protein complexes were much more likely than chance (\( p \leq 10^{-46} \)) to have links to a documented clinical pathology (Figure 4B; see Table S4 for details), with disease-associated proteins distributed broadly among the complexes (Figures 4B and S4C). Closer examination of the interaction subnetworks—comprising known human disease genes with genes that currently lack annotation or that have not previously been associated with any human disorders (Figure 4B)—highlights the utility of the map.

One such example is shown in Figure 5A, illustrating the case of the human developmental disorder Cornelia de Lange syndrome (CdLS). Mutations in three subunits of the cohesin complex (SMC1A, SMC3, and NIPBL) have been linked to CdLS (Pie´ et al., 2010), implicating an additional component (RAD21) as a candidate CdLS locus, and are consistent with at least one unmapped CdLS locus residing on chromosome 8 (DeScipio et al., 2005). The link to RAD21 provides a likely...
explanation for the occasional overlap of Langer-Giedion syndrome (LGS) clinical presentation with CdLS, as all LGS patients are at least partially defective for RAD21 (see e.g., McBrien et al., 2008; Wuyts et al., 2002). Similarly, RAD18, a homolog of SMC3 and SMC1A, may play a role in CdLS that is consistent with unmapped CdLS deletions within chromosome 3p25 (DeScipio et al., 2005). Reports coinciding with the preparation of this manuscript confirm that RAD21 mutations do indeed lead to a CdLS-like syndrome (Deardorff et al., 2012), supporting the use of the complex map to prioritize promising candidate genes for human diseases.

Similarly, participation in the same complex suggests shared functions; the map can thus be used to predict new biochemical functions for proteins and other types of functions. We experimentally validated one such case for a ribosome-associated subcomplex containing BOP1, RRS1, GNL3, EBP2, FTSJ3, and MK1671P, and we first confirmed the interactions by affinity tagging/purification and mass spectrometry (Figure 5B). BOP1, EBP2, and the yeast ortholog of RRS1 are known to participate in maturation of the large 60S ribosomal subunit, suggesting that the other factors likewise engage in ribosome assembly, which is consistent with the nucleolar localizations of GNL3, FTSJ3, and MK1671P. Supporting a role in ribosome biogenesis, short interfering RNA knockdowns of FTSJ3, MKI671P, and, to a lesser extent, GNL3, perturbed 60S formation in cell culture, decreasing the ratio of free 60S to 40S subunits (Figure 5C). Taken together, these data support roles in ribosome biogenesis for these proteins and confirm the utility of the map for identifying biological functions.

**Conservation of Human Protein Complexes**

Estimates based on sequence similarity across orthologs indicate that the components of the complexes we detect are generally more ancient and have higher conservation on average than most human proteins (Figure 6A; see Table S3 for details). Using orthology relationships derived from well-established sources and calculating evolutionary rates and ages for all human proteins as a base distribution for gauging the emergence of complexes (see Extended Experimental Procedures), we found that many complexes appear to be quite ancient and slowly evolving (Figure 6B). Strikingly, however, most (60%; 376/622) human complexes likely arose with vertebrates, i.e., orthologs not present in invertebrates or fungi (Table S3). Hence, our analyses suggest a major shift/expansion in the ancestral protein interaction network coincident with the emergence of vertebrates.

Given the availability of experimentally derived networks of fly and yeast protein complexes, we could directly examine the evolutionary conservation of protein complexes across animals by comparing our network of human complexes with the extensive maps of 556 fly protein complexes recently reported for *D. melanogaster* (Guruharsha et al., 2011) and 720 yeast protein complexes documented for *S. cerevisiae* (Babu et al., 2012). Roughly one quarter (24%; 149/622) of the predicted human protein complexes showed statistically significant overlaps with complexes reported for these models (Figure 6B, inset; see Table S3 for details), with half of the subunits having clear orthologs (Figure 6C); the remaining components presumably represent genuine differences or incomplete orthology annotations.

The functional significance of unannotated ancestral human complexes supported by conservation in yeast or fly (Table S3 and Figure 6) warrants further investigations. At least one such complex, a multisubunit transfer RNA (tRNA)-splicing ligase (Popow et al., 2011), was characterized recently. The interaction between DDX1 and C14orf166 was detected at high confidence both in our data set (probability score 0.889) and in the Guruharsha et al. (2011) fly cocomplex data, and the other respective associated complex subunits likewise show significant overlap (Benjamini-corrected p value 1.1 × 10^-1). Additional examples of complex conservation are similarly supported by independent experimental evidence, e.g., such as the matching tissue specificities of the putatively interacting proteins endoplasmic glucosidase 2β (Figure 6D), which form an uncharacterized complex conserved in both the fly and human maps.

Functional enrichment analysis of ancient complexes in comparison to vertebrate-specific ones also reveals intriguing biological trends. For example, we expected ancient, core cellular functions to be depleted among vertebrate-specific complexes. Consistent with this expectation, we find proteins associated with the ribosome (p < 10^-67, 113 proteins) and RNA polymerase II (p < 10^-27, 45 proteins) to be highly enriched only among conserved complexes. However, we also observe several notable variations from this hypothesis. For example, compared to the genomic background, mitochondrial proteins are more highly enriched among proteins assigned to vertebrate complexes than among those assigned to conserved complexes; 159 vertebrate proteins have a mitochondrial Gene Ontology Biological Process (GO BP) annotation (p < 10^-31) versus only 81 proteins assigned to conserved complexes (p < 10^-5). Similarly, proteins annotated as being part of the splicing apparatus are enriched in both conserved (p < 10^-33, 63 proteins) and vertebrate complexes (p < 10^-11, 43 proteins), which is consistent with an ancient function gaining additional complexity in vertebrates (e.g., increased alternative splicing). Our study therefore offers a unique perspective into the functional conservation and diversification of protein complexes across animals.

**Protein Abundance, Ubiquity, and Complex Subunit Stoichiometries**

Consistent with the documented origins of the HeLa and HEK293 cells analyzed in this study, the complexes we identified were significantly enriched for epithelial markers (p < 10^-183, UniProt tissue annotations). Explicit comparison of results across the two cell lines used in this study provided little evidence for tissue-specific or cell-type-specific complexes (see Supplemental Information). Most proteins were detected in both cell line fractionations, which is consistent with the similar protein and mRNA expression patterns observed in these cell lines (Figure S1), whereas the few proteins detected uniquely in one cell line or the other did not preferentially assort into tissue-specific complexes (Figure S2). The vast majority of complex components are universally expressed in 11 cancer cell lines (Geiger et al., 2012) (Figure S3A) and show high and largely invariant expression in an mRNA sequencing (mRNA-seq) study of 16 normal human tissues (EBI accession number...
Indeed, complex subunits are considered near ubiquitous (p < 10^{-11}; protein information resource [PIR] tissue specificity annotations) and are expressed in the top quartiles of 1,045 of 7,067 neoplastic and normal tissue CGAP EST libraries (1% false discovery rate [FDR]), including normal kidney (p ≤ 10^{-10}), muscle (p ≤ 10^{-20}), liver (p ≤ 10^{-12}), brain (p ≤ 10^{-20}), vascular (p ≤ 10^{-10}), bone (p ≤ 10^{-15}), and embryonic tissue (p ≤ 10^{-31}). Consistent with this, genes encoding complex subunits also tend to share common upstream transcriptional regulatory motifs (p ≤ 10^{-8}; Figure 4B, inset table). Proteins mapped to complexes showed no major bias in abundance over the complete set of human proteins identified by mass spectrometry (Figure 1D).

The pervasiveness of ubiquitously expressed protein complexes argues strongly for broad relevance to basic human cell biology. Although often coexpressed, the subunit stoichiometries of human protein complexes in vivo are largely unknown and have never been systematically measured globally. Because...
all reconstructed complexes are supported by the same set of extensive experimental mass spectrometry data, we could estimate subunit stoichiometries based on the ratios of recorded spectral counts after correcting appropriately for protein size and composition (see Extended Experimental Procedures). Although only approximate ratios were inferred and peaked at ~1:1 (Figure 7A), such as between known ribosomal subunits (Figures 7B and 7C), the results highlight intriguing deviations in subunit abundance (Table S2). An example drawn from the proteasome is illustrative: whereas the median stoichiometry of core α and β enzymatic subunits is close to the expected 1:1 ratio, the median of stoichiometries of core to non-ATPase regulatory subunits deviated significantly at ~4:1 (Mann-Whitney p ≤ 10^-16; Figures 7D and 7E). Hence, these data suggest a rich source of information about the physical organization of human proteins.

**DISCUSSION**

The biochemically based interaction data obtained in this integrative proteomic study have enabled the identification of both 364 previously unannotated protein complexes (i.e., predicted complexes with no statistically significant match to complexes in public databases) encompassing 1,278 human proteins, many of which are linked to human disease, and unexpected components and interactions for well-studied, widely conserved nuclear and cytoplasmic protein machineries, such as ribosome biogenesis, with clear biological implications. Most of the high-confidence protein interactions provided in this resource have not been previously reported in public interaction databases and hence motivate mechanistic investigations of specific biological systems.

Prior to this work, experimental knowledge regarding soluble protein complex membership in human cells has generally been ad hoc or focused on specific subcellular systems. Our relatively unbiased integrative approach, wherein biochemical evidence (cofractionation) of soluble native macromolecules was combined with genomic inferences (imputed functional associations), provides an inclusive snapshot of human protein

**Figure 7. Protein Complex Stoichiometries**

(A) Overall distribution of derived intracomplex component stoichiometries. (B and C) Estimated subunit stoichiometries within and between proteins of the large and small ribosome subunits agree on average with the expected 1:1 ratio. Boxes summarize first quartile, median, and third quartiles, whiskers represent ±1.5 IQR, and circles represent outliers. (D and E) Estimated protein subunit stoichiometries within and between proteasomal proteins. Intrusionsubunit stoichiometries within the core, ATPase, or nonATPase regulatory subunits agree well with the expected 1:1 ratio, but stoichiometries observed between these complexes deviate significantly from 1:1 (ATPase:non-ATPase, Mann-Whitney p ≤ 10^-3; core:ATPase, p ≤ 10^-16; core:non-ATPase, p ≤ 10^-10). See also Table S2.
complexes present under a standardized cellular context, thus serving as a reference against which future process- or cell-type-specific or dynamic interaction data sets can be compared. Information gleaned from orthology proved to be an important resource in separating true pairwise interactions from putative false positives and, in turn, could reasonably be expected to bias our results toward conserved complexes. In fact, although we do find conserved complexes as expected, we also find a majority that are not conserved (in fly and yeast) and that seemingly have arisen with vertebrates (i.e., Figure 6B). The slower rate of evolution of the subunits we report for our protein complexes is also a feature of other human PPI networks, such as in CORUM, and thus, our predictions of broad complex conservation, albeit incomplete, are not just artifacts of our methodology.

The fact that we detected little evidence of tissue specificity for most of the derived human protein complexes and few cell-type-specific components likely reflects undersampling by our mass spectrometry procedures, which is a common limitation of LC-MS/MS. At the level of predicted PPI (which are derived from multiple biochemical fractions), differences in the proteomic profiles generated for the two cell lines lie within the variance observed between biological replicates of the same cell line (Figures S1 and S2). Yet it is clear that differential interactomes and the contextual rewiring of PPI networks are major determinants of cell behavior and phenotypes. The complexes we report undoubtedly undergo differential rewiring in response to environmental, physiological, developmental, or disease states. With further refinements to our experimental procedures, our interaction mapping strategy has the potential to interrogate changes in interaction space in a systematic manner in the future.

To enable exploitation of these data by the scientific community, we have generated a dedicated web database of human protein complexes (http://human.med.utoronto.ca) that contains all the data generated in this study in an easily navigated format. These include all of the supporting information for each of the pairwise protein interactions obtained through integration of our cofractionation data with public genomic evidence, a list of the 5,584 proteins detected in each of the 1,163 biochemical fractions collected, and the subunit composition of the 622 putative protein complexes obtained through clustering of our generated high-confidence interaction network. This “first pass” draft of the soluble, stably associated human protein “complexome” provides a glimpse into the global physical molecular organization of human cells, which is likely to be perturbed in pathological states.

**EXPERIMENTAL PROCEDURES**

**Cell Culture and Extract Preparation**

HeLa S3 (ATCC#: CCL-2.2) and HEK293 (ATCC#: CRL-1573) soluble nuclear and cytoplasmic protein extracts were prepared by conventional methods (see Extended Experimental Procedures). Prior to fractionation, lysates were treated with 100 units/ml Benzonase (Novagen Inc.) to remove nucleic acids and clarified by centrifugation to remove debris.

**Biochemical Fractionation and Proteomic Analysis**

We performed weak anion-exchange and mixed-bed ion exchange, both with and without a heparin precolumn to enrich for nucleic-acid-binding proteins. In total, 1,095 chromatography fractions were collected (see Extended Experimental Procedures). Isoelectric focusing was carried out on a MicroRotofor Liquid-Phase IEF cell (Bio-Rad) according to the manufacturer’s protocol, with 40 fractions collected across a pH range. Sucrose density gradient centrifugation was performed as previously described (Ramani et al., 2008), with 28 fractions collected.

Proteins were acid precipitated and trypsin digested, and the peptide mixtures were fractionated and sequenced by using nanoflow liquid chromatography-electrospray tandem mass spectrometry. Spectra were collected on an LTQ linear ion trap (ThermoFisher Scientific) (majority) or LTQ Orbitrap Velos hybrid mass spectrometer and searched against a UniProt human target-decoy sequence database by using SEQUEST (Eng et al., 2008) (see Extended Experimental Procedures). The LC-MS/MS identifications were filtered to a 1.0% protein and peptide theoretical FDR.

**Bioinformatics Analyses**

Protein cofractionation networks were scored by correlation analysis (Pearson correlation, weighted cross-correlation, coapex) based on the protein spectral counts recorded across each set of fractions (see Extended Experimental Procedures). Weighted networks were likewise constructed based on functional evidence reported in HumanNet (Lee et al., 2011), omitting human protein interaction data to minimize circularity that might bias our association predictions. A coevolution network (Tillier and Charlebois, 2009) based on correlated evolutionary rates was built to account for additional associations not covered in HumanNet.

For the machine-learning classifier, we used the fast random forest implementation in Weka (see Extended Experimental Procedures) to integrate all generated networks. Cross-validated decision trees were learned and benchmarked by using independent training and test sets of CORUM reference complexes (Ruepp et al., 2010). We denoised the network by using a diffusion procedure to delete interactions lacking network topology support and by calibrating the diffused interaction scores with Gene Ontology (Cellular Component) normalized semantic similarity scores (see Extended Experimental Procedures).

Clusters were defined by using the ClusterONE algorithm with parameter settings chosen to yield the highest maximum matching ratio (Nepusz et al., 2012) between the predicted complexes and set of cluster-training complexes (see Extended Experimental Procedures).

Stoichiometries calculation is shown in Extended Experimental Procedures.

**ACCESSION NUMBERS**

The interaction data have been deposited into BioGRID and are also publicly accessible via a dedicated web portal (http://human.med.utoronto.ca/).

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Extended Experimental Procedures, five figures, and six tables and can be found with this article online at http://dx.doi.org/10.1016/j.cell.2012.08.011.

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