A Distant Supervision Corpus for Extracting Biomedical Relationships Between Chemicals, Diseases and Genes

Dongxu Zhang*,†, Sunil Mohan*,†, Michaela Torkar‡, Andrew McCallum¹

¹ University of Massachusetts, Amherst, Massachusetts, USA
‡ Chan Zuckerberg Initiative, Redwood City, California, USA

Abstract
We introduce ChemDisGene, a new dataset for training and evaluating multi-class multi-label document-level biomedical relation extraction models. Our dataset contains 80k biomedical research abstracts labeled with mentions of chemicals, diseases, and genes, portions of which human experts labeled with 18 types of biomedical relationships between these entities (intended for evaluation), and the remainder of which (intended for training) has been distantly labeled via the CTD database with approximately 78% accuracy. In comparison to similar preexisting datasets, ours is both substantially larger and cleaner; it also includes annotations linking mentions to their entities. We also provide three baseline deep neural network relation extraction models trained and evaluated on our new dataset.

Keywords: Corpus, Information Extraction, Linked Data, Weakly-supervised Learning, Relation Extraction

1. Introduction
Biomedical researchers have used systems of experimentally confirmed interactions between chemicals, diseases, genes/proteins and other entities, for understanding disease mechanisms for diagnosis, e.g. [Lee et al. (2019)], for drug repurposing [Morselli Gysi et al. (2021)], and even for understanding the health hazards associated with spaceflight [Nelson et al. (2021)]. These knowledge graphs (KGs) are often built by integrating manually curated databases like CTD and DrugBank who use domain experts to extract observed interactions from research publications and other sources. While the information in these databases is high in precision, with the growing publication rate their recall is low [Baumgartner et al. (2007)]. To improve coverage, researchers have resorted to automated mining of biomedical interactions from research texts, to supplement their KG [Himmelstein et al. (2017)], or even to build the entire KG, e.g. [Crichton et al. (2020)].

The bioinformatics community recognized that machine learning Relation Extraction (RE) models could help the manual curation task, and the BioCreative workshops introduced the first shared task and manually labeled ‘gold standard’ dataset for training and evaluating models for extracting protein-protein interactions from full text articles in 2006 [Krallinger et al. (2006)]. Several such labeled corpora have followed, primarily focusing on extracting relationships from abstracts. However, labeling of relationships requires domain experts and is slow and expensive. Consequently, most labeled corpora are small, and focus on a small number of entity types and relationships.

In this paper, we introduce ChemDisGene, a new dataset of biomedical research abstracts labeled with pairwise interactions between Chemicals, Diseases and Genes/Gene-products. It contains two sub-corpora:
• A large corpus of ~ 80k abstracts with distant labeling of 14 relation types. This corpus is automatically derived from CTD [Davis et al. (2020)], thus allowing for a larger size more suitable for training deep learning models. However, relationships are distantly labeled because relationships in CTD are associated only with a paper, and not with a specific text passage within the paper.
• A smaller corpus of 523 abstracts, manually annotated with relationships by domain experts. This corpus is aimed primarily for testing models trained on the CTD-derived corpus, and the relationships here are also distantly labeled.

A previous version of the CTD-derived corpus was introduced in [Verga et al., 2018]. ChemDisGene adds a manually annotated component, and includes several improvements to the derivation process:
• More recent updates (2021 February) from CTD.
• Entity linking uses PubTator Central [Wei et al., 2019] with significantly improved models for recognizing Chemicals (+66.3% improvement in F1 score), Diseases (+3.8%) and Genes/Proteins (+8.2%) over the previous PubTator model.
• The previous dataset was randomly split into training, dev and test, while in ChemDisGene these splits are based on paper publication date, to better simulate a real world scenario.
• A cleaner extraction of binary relationships from complex nested relationships captured by CTD.

The rest of this paper describes how the labeled cor-

* Equal contribution
† CTD: Comparative Toxicogenomics Database
‡ https://go.drugbank.com

https://github.com/chanzuckerberg/ChemDisGene
2. Methodology

A note on terminology: we will use relation to refer to the predicate schema $r(T_s, T_o)$, where $r$ is the relation type, and $T_s, T_o$ are the argument entity types: Chemical, Disease or Gene. A relationship is a ground instance $r(e_s, e_o)$ of a relation, with argument entities $e_s \in T_s, e_o \in T_o$.

The ChemDisGene dataset comprises a large corpus automatically derived from CTD, and a smaller curated corpus manually labeled by domain experts.

2.1. Derivation from CTD

Comparative Toxicogenomics Database (CTD) is a public knowledge base containing manually curated interactions between chemicals, genes, diseases and phenotypes (Davis et al., 2020). CTD curators regularly scan new research publications to identify those interactions that are the primary contributions of each paper (Davis et al., 2011). These are then encoded using a hierarchical ontology of ~ 50 Chemical–Gene interaction classes, and two types each for Chemical–Disease and Gene–Disease interactions (phenotypes are not covered in our dataset). Each interaction is expressed using relation types from these interaction classes, along with the argument entities, and recorded with a reference to the paper from which it was extracted (but no reference to any text within the paper). Entities are also identified using public ontologies: MeSH for Chemicals, MeSH and OMIM for Diseases, and NCBI Gene for Genes and Gene-products.

While CTD curators scan full papers to extract these relationships, we limited the text in ChemDisGene to only the title and abstract. Starting with the February 2021 dump of CTD, we obtained abstracts for all referenced articles from PubMed. Each abstract was processed through PubTator Central (PTC) to identify and link mentions of chemical, disease and gene/gene-product entities. We then performed a ‘distant alignment’ of the annotated abstracts with the relationships linked to each paper in CTD: relationships whose entities were not detected in the abstract were discarded. This yielded a dataset of abstracts with linked entity mentions, and distantly linked relationships.

This distant linking of relationships to aligned abstracts is noisy due to the following sources of error: (i) entity recognition models in PTC, whose F1 scores for each entity type are in the range 0.84–0.90 (Wei et al., 2019), (ii) even if the entities of a relationship are correctly identified in the abstract, the corresponding interaction may not have been mentioned in the abstract text, and (iii) an abstract may mention some relationships that are not extracted by CTD. To measure these sources of error, we selected a subset of aligned abstracts for manual curation (see §2.2).

Relations in CTD are organized into a class hierarchy, with some relation classes qualified by a ‘degree’. ChemDisGene includes 10 of these classes, which combined with the degrees defines 18 relation types:

- **Chemical-Disease:**
  - marker/mechanism: A chemical that correlates with a disease.
  - therapeutic: A chemical that has a known or potential therapeutic role in a disease.

- **Chemical-Gene:** Each qualified by a degree.
  - activity: An elemental function of a molecule. Degrees: increases, decreases, or affects when the direction is not indicated.
  - binding: A molecular interaction (affects).
  - expression: Expression of a gene product (increases, decreases, affects).
  - localization: Part of the cell where a molecule resides (affects).
  - metabolic processing: The biochemical alteration of a molecule’s structure, not including changes in expression, stability, folding, localization, splicing, or transport (increases, decreases, affects).
  - transport: The movement of a molecule into or out of a cell (increases, decreases, affects).

- **Gene-Disease:**
  - marker/mechanism: A gene that may be a biomarker of a disease or play a role in the etiology of a disease.
  - therapeutic: A gene that is or may be a therapeutic target in the treatment of a disease.

In some cases, CTD defines a finer granularity of Chemical-Gene interactions. Because their occurrence is rare, they would be harder for a model to recognize, so we abstracted them to the levels described above. The relationships in CTD also include complex and nested biomedical interactions involving multiple entities. For ChemDisGene we only extracted binary relationships. In particular, (a) we omitted CTD’s “cotreatment” relation type because it is non-binary, and (b) we implemented a cleaner extraction of binary relationships from nested interactions (see example in fig. [1]).

The previous CTD-derived dataset in Verga et al. (2018) used the same relation types for Chemical-Disease and Gene-Disease interactions, but a different set of 10 relation types for Chemical-Gene. With three years of new publications, the distribution of relation types in CTD has changed, affecting our selection.

The derivation of relationships from CTD in Verga et al. (2018) did not take into account nesting levels in complex interactions: in the example in fig. [1] the previous dataset would also extract reaction-decreases between the chemical ‘24-hydroxycholesterol’ and the
into 10% of the CTD-derived corpus, which was then split as 'null' documents with no relationships. This forms abstracts that did not align with any CTD relationships.

As a final step, we added some randomly sampled abstractor "gene 'ITGB1', even though the corresponding indica-

expression-increases(24-hydroxycholesterol, ITGB1)
expression-increases(27-hydroxycholesterol, ITGB1)
expression-increases(cholest-5-en-3 beta,7 alpha-diol, ITGB1)

Figure 1: An example showing extraction of binary relationships from a complex nested interaction in CTD.

gene ‘ITGB1’, even though the corresponding indicator “inhibits the reaction” is at a different nesting level. As a final step, we added some randomly sampled abstracts that did not align with any CTD relationships as ‘null’ documents with no relationships. This forms 10% of the CTD-derived corpus, which was then split into train, development (dev) and test sets by publication year (2018 as dev and years 2019, 2020 as test).

2.2. Curation

As described above, the relationship labels in the CTD-derived corpus are noisy. To perform more reliable testing of RE models, we selected some documents for manual annotation: 303 sampled from the test split, and an additional 252 documents from CTD that were also included in the DrugProt corpus (Martin Krallinger and Valencia, 2021), to enable future comparative analyses. These were distributed for annotation by five biologists, each document assigned randomly to three curators.

We developed a web-based annotation tool which displayed for each document, the title and abstract, all the linked chemicals, diseases and genes/gene-products, and their mentions in the text, and all the relationships derived from CTD. Annotating a document involved two tasks: (i) review each relationship derived from CTD, and either reject or approve it, and (ii) add all other established relationships expressed in the document. Relationships mentioned in the abstract without any conclusions were excluded from annotation. In keeping with our goal of a realistic dataset, 44 of these documents had no CTD-derived relationships.

We developed annotation guidelines (published with the dataset) that describe the steps in the curation process and the types of pairwise interactions curated in this dataset, including brief definitions and real-world example statements that do or do not support a specific relation type. These guidelines underwent multiple rounds of revisions through 4 iterations of practice annotations. During the practice phase, all 5 curators were given the same set of documents to curate (15–30 per cycle); annotation disagreements and questions were clarified during multiple workshops, and feed-

A: Don’t record investigated or motivating relationships that remain unknown and hypothetical.

“Gene A is a therapeutic target for treatment of Disease X; it may therefore have a potential role in treatment of Disease Z.”

Record a relationship between Gene A and Disease X; but not between Gene A and Disease Z.

B: Inferring a relationship across sentences.

“We have previously identified a panel of fusion genes in aggressive prostate cancers. In this study, we showed that . . . CCNH-C5orf30 and TRMT11-GRK2 gene fusions were found in breast cancer, colon cancer, . . . ”

Record a ‘Gene-Disease: marker/mecanism’ relationship between C5orf30 and prostate cancers.

Figure 2: Two examples from the curation guidelines. Colors identify disease and gene mentions.
yielding a total of 523 annotated documents.

On analyzing the singletons, we noticed that some of these differed only in degree from an approved relationship in the same document: 45 were abstractions (degree affects) and 7 refinements (degree increases or decreases) of an approved relationship. These singletons were then automatically rejected.

In this annotation task, when one annotator does not identify a particular relationship that was found by another, it could be for one of two reasons: (i) both annotators noticed the same text passage but disagreed on whether it expressed the relationship, or (ii) the first annotator did not notice the passage that the second annotator used to identify the relationship. To resolve this ambiguity, all the singleton relationships were reviewed by an annotator not originally assigned to that document, followed by a second review by the curation manager to ensure consistency. Relationships approved in this phase were added to the curated data.

Table 1: General statistics for the CTD-derived corpus.

|                  | Train | Dev  | Test |
|------------------|-------|------|------|
| Nbr. abstracts   | 76,942| 1,521| 1,939|
| ... with no rels.| 7,244 | 397  | 436  |
| Nbr. rels.       | 167,005| 3,290| 5,116|
| ... unique rels. | 93,801| 3,127| 4,801|
| Total Entity mtns| 1,532,117| 36,114| 49,839|
| Chemicals        | 686,102| 13,986| 19,895|
| Diseases         | 478,397| 8,962 | 11,750|
| Genes            | 367,618| 13,166| 18,194|
| Unique Entities  | 14,991 | 1,894 | 2,345|
| Chemicals        | 7,187  | 759  | 999  |
| Diseases         | 2,413  | 283  | 287  |
| Genes            | 5,391  | 852  | 1,059|

3. ChemDisGene Corpus Statistics

3.1. The CTD-derived Corpus

The median of number of tokens (split by space) in abstracts is 214. And about 99.8% of abstracts have less than 512 tokens. Other statistics for the CTD-derived corpus are shown in table 1 and the distribution of the number of relationships per document in fig. 3. About 80% of the documents have 3 or fewer relationships, followed by a long thin tail. The dev and test splits have a higher proportion of documents with no relationships. There are an average of 2.2 relationships per document, with over 9,000 entity pair occurrences with multiple relation type labels in the same document.

Counts for each relation type are shown in table 2. Unique numbers count unique argument-entity pairs. Four Chemical-Gene relation types (activity-affects, metabolic-processing-affects, transport-affects, and transport-increases) were omitted from the CTD-derived corpus because of their low incidence. However, they are included in the curated corpus for completeness, making the annotation task a little easier.

3.2. The Curated Corpus

The curated corpus contains 523 documents: 271 from CTD-derived’s test split, and an additional 252 documents taken from DrugProt, that are not in the CTD-derived corpus. Twenty seven of these documents had no relationships derived from CTD. Manual annotation rejected 22% of all CTD-derived relationships, leaving 64 documents with no approved CTD-derived relationships. This indicates a fairly high 78% confidence in the automatically derived relationships.

There are a total of 1,279 approved CTD-derived relationships (avg. 2.4/doc), 2,632 approved new relationships (5.0/doc). The distribution of the 18 types of relations in this corpus is shown in table 3.

The 3,911 approved relationships (3,806 unique) in the curated corpus involve 1,875 unique entities: 670 unique Chemicals, 318 Diseases and 887 Genes. Figure 4 shows the distribution of number of approved relationships in each document. As expected, the CTD-derived approved relationships are more skewed to the left than the new added relationships.
Table 2: Nbr. of relationships (instances) for each relation type in the CTD-derived corpus.

| #  | Relation type                                      | Total          | Unique          |
|----|---------------------------------------------------|----------------|-----------------|
|    |                                                   | Train          | Dev             | Test            | Train | Dev   | Test  |
| 1  | Chemical-Disease : marker/mechanism               | 66,155         | 559             | 754             | 27,706 | 486   | 602   |
| 2  | Chemical-Disease : therapeutic                   | 34,775         | 250             | 410             | 16,093 | 245   | 398   |
| 3  | Chemical-Gene : activity - decreases             | 5,555          | 101             | 232             | 4,128  | 97    | 232   |
| 4  | Chemical-Gene : activity - increases             | 6,152          | 127             | 174             | 4,133  | 120   | 157   |
| 5  | Chemical-Gene : binding - affects                | 3,123          | 67              | 77              | 2,024  | 65    | 73    |
| 6  | Chemical-Gene : expression - affects             | 1,247          | 51              | 160             | 1,206  | 51    | 158   |
| 7  | Chemical-Gene : expression - decreases           | 10,204         | 480             | 923             | 8,487  | 467   | 905   |
| 8  | Chemical-Gene : expression - increases           | 19,810         | 919             | 1,570           | 14,685 | 878   | 1,491 |
| 9  | Chemical-Gene : localization - affects           | 1,448          | 50              | 73              | 1,216  | 50    | 70    |
| 10 | Chemical-Gene : metabolic_processing - decreases | 1,653          | 101             | 116             | 1,313  | 100   | 111   |
| 11 | Chemical-Gene : metabolic_processing - increases | 4,640          | 175             | 293             | 3,507  | 172   | 283   |
| 12 | Chemical-Gene : transport - increases            | 1,962          | 92              | 108             | 1,405  | 88    | 96    |
| 13 | Gene-Disease : marker/mechanism                  | 9,388          | 301             | 219             | 7,384  | 292   | 218   |
| 14 | Gene-Disease : therapeutic                       | 893            | 17              | 7               | 514    | 16    | 7     |

Table 3: Frequency distribution of relation types in curated corpus (each column sums to 100%). Empty frequencies indicate some relations are rare in CTD.

| #  | Relation type                                      | Approved, New | Approved, CTD |
|----|---------------------------------------------------|---------------|---------------|
| 1  | Chemical-Disease : marker/mechanism               | 16.6          | 16.4          |
| 2  | Chemical-Disease : therapeutic                   | 10.4          | 12.0          |
| 3  | Chemical-Gene : activity - affects                | 1.2           |               |
| 4  | Chemical-Gene : activity - decreases             | 8.3           | 7.3           |
| 5  | Chemical-Gene : activity - increases             | 8.7           | 7.8           |
| 6  | Chemical-Gene : binding - affects                | 4.3           | 6.7           |
| 7  | Chemical-Gene : expression - affects             | 2.8           | 0.6           |
| 8  | Chemical-Gene : expression - decreases           | 10.4          | 13.1          |
| 9  | Chemical-Gene : expression - increases           | 11.8          | 18.4          |
| 10 | Chemical-Gene : localization - affects           | 0.8           | 1.5           |
| 11 | Chemical-Gene : metabolic_processing - affects   | 0.8           |               |
| 12 | Chemical-Gene : metabolic_processing - decreases | 1.7           | 1.5           |
| 13 | Chemical-Gene : metabolic_processing - increases | 3.0           | 4.0           |
| 14 | Chemical-Gene : transport - affects              | 0.3           |               |
| 15 | Chemical-Gene : transport - decreases            | 0.6           |               |
| 16 | Chemical-Gene : transport - increases            | 1.1           | 0.9           |
| 17 | Gene-Disease : marker/mechanism                  | 14.1          | 9.3           |
| 18 | Gene-Disease : therapeutic                       | 2.9           | 0.5           |

3.3. Inter-Annotator Agreement

Commonly used measures of inter-annotator agreement are defined for tasks where the units being classified or measured are precisely specified. As noted in (Kilicoglu et al., 2011), identifying all relationships expressed in a text does not match this paradigm. This task could be decomposed into the following steps: (i) find relationship indicators in the text, (ii) identify the entity mentions each indicator refers to, and (iii) map the expressed relationship to the appropriate ontological term. Here the space of possible annotations is clearly defined only for step (iii). In step (ii) the space would be clearly specified only if we presented the annotators with every pair of linked mentions. The set of possible relationship indicators in a document, in step (i), is also not presented to the annotators. When a relationship is identified by only one of two curators reviewing the same text, it could be because either the first one did not ‘notice’ the same sentence, or actually saw it and rejected it. This inherent ambiguity causes a problem even for measures that allow varying number of annotations per unit.

Similar to (Kilicoglu et al., 2011), we evaluated each curator’s annotations against a reference, using precision, recall and F1 scores, as more feasible and intuitively understandable metrics for our use case. We used the ‘majority approved’ relationships (§2.2) as the reference dataset. The annotator agreement metrics (ta-
4. The Relationship Extraction Task

4.1. Task definition

The document-level relation extraction (RE) task in ChemDisGene is to identify all relationships \( r(e_s, e_o) \) expressed in a document, comprised of the title and abstract texts, that are the primary contributions of that article. We consider 14 binary relation types (from the CTD-derived corpus) among chemical, disease and gene/gene-product entities. All mentions of these entities in the text are identified and linked to the corresponding ontologies. This is a distant supervision (relationships are associated with documents, but not specific entity mentions) multi-label (a document, and a pair of entities, may have more than one relationship) classification task. For evaluation, we use Micro/Macro F1 scores where per-relation thresholds are tuned on the dev set, and average precision where thresholds are not required.

4.2. Models

In our experiments, we trained and evaluated three baseline methods on ChemDisGene.

BRAN Bi-affine relation attention networks \(^{(Verga et al., 2018)}\) is one of the first papers to tackle document level (distant supervision) relation extraction in the biomedical domain. It uses multiple self-attention + convolutional neural network (NN) layers to encode the text input, then leverages per-relation biaffine transformation to calculate mention level scores of the query \( r(e_s, e_o) \), and a logsumexp layer to capture the most significant signal among mention pairs. In our experiment, we omitted BRAN’s NER joint loss in order to analyze its core RE module.

PubMedBert \(^{(Gu et al., 2021)}\) is a BERT-based pretrained language model \(^{(Devlin et al., 2019)}\) trained from scratch on PubMed abstracts. For relation extraction, we first get each entity’s embeddings by max-pooling over PubMedBert’s encoding of all the entity’s mentions. Then concatenated embeddings of candidate argument entity pairs are processed through a feed-forward NN to predict scores for each relation type.

PubMedBert + BRAN. This model combines the stronger text encoder of PubMedBert with the relation detection layers of BRAN. The model structure is: Input \( \rightarrow \) PubMedBertEncoder \( \rightarrow \) Biaffine \( \rightarrow \) logsumexp \( \rightarrow \) logits.

4.3. Empirical Results

Table 5 shows overall performance of the three baseline models on ChemDisGene. Performance metrics are shown for the test split of the CTD-derived corpus, and separate metrics on the curated corpus for approved relationships derived from CTD, and for all approved relationships, which also includes new relationships added by the curators.

From the table, our best model PubMedBert + BRAN has 43.8 Micro F1 and 50.6 average precision on the ‘all relationships’ curated test set, indicating the difficulty of this task. The pretrained language model adds significant improvement over BRAN. And the biaffine transformation and logsumexp layer are also complementary to the pretrained language model.

Compared with the CTD-derived test set, the performance decreases significantly on the curated test set, indicating the necessity of evaluation on expert-labeled data. We also observe that Macro results are lower than Micro, indicating that performance varies across different relation types. In Table 6, we see that relation types with low frequencies in the training data tend to perform poorly. The particularly bad performance of our model on Chemical-Gene: expression-affects is also caused by distraction from two similar but common Chemical-Gene relation types: expression-increases and expression-decreases.

Performance of baseline models on ‘BRAN’ dataset.

We also trained and tested the baseline models on the CTD-derived dataset from \(^{(Verga et al., 2018)}\), referred to as the ‘BRAN’ dataset (Table 5). As described above (2.1), there are several differences in this dataset that account for the different performance results from those on ChemDisGene (Table 5). Perhaps the most important one is that in the BRAN dataset, abstracts from test and dev splits are randomly selected, whereas in ChemDisGene 271 abstracts are assigned based on publication date. The ChemDisGene test set also includes more documents with no relationships. While this makes ChemDisGene more challenging, it also reflects a more realistic scenario for applying such RE models. The relative order of performance of the three baselines is the same on both datasets.

Comparing the performance on CTD-derived and All relationships in curated corpora.

From Table 5 we can see that while model precision increases when tested on all approved relationships from the curated corpus, compared to the performance on just CTD-derived approved relationships, the recall of all models drops significantly. A main reason is that the training

We trained all three baselines on ChemDisGene training set with hidden dimension 128, and we tuned the hyper-parameters such as learning rate \(1e-5, 1e-4\) and weight decay = \([0, 1e-4]\) over the distant supervision dev set.

| Relationships  | A     | B     | C     | D     | E     |
|----------------|-------|-------|-------|-------|-------|
| All            | 0.85  | 0.84  | 0.83  | 0.88  | 0.88  |
| CTD-derived only | 0.99  | 0.96  | 0.97  | 0.99  | 0.96  |
| New only       | 0.76  | 0.77  | 0.69  | 0.82  | 0.83  |

Table 4: Agreement F1 scores for the 5 annotators (A-E) against the ‘approved’ reference subset.
data only includes CTD-derived relationships, which are selected by CTD to be the ‘primary’ contributions of the paper. While this is mostly determined within the context of other publications, there might be a signal in the wording (an area for further investigation).

Curators were asked to reject CTD-derived relationships when the entities involved were incorrectly linked. This probably accounts for the small difference in models’ performance between the CTD-derived and curated corpora.

5. Related Work

5.1. Distant Supervision Biomedical Corpora

As described above (see §1.2.1). ChemDisGene offers a reworking of the derived corpus introduced in (Verga et al., 2018), focusing on a cleaner derivation from an updated CTD with better entity linking. The number of abstracts also increased by ~20k.

A well known manually labeled biomedical corpus is BC5-CDR (Li et al., 2016), which identifies a single relation type between Chemicals and Diseases, distantly labeled in 1,500 abstracts. BC6-PM (Islamaj Do˘gan et al., 2019) is another manually annotated distant supervision corpus, for Protein-Protein interactions. It has a total of 1,232 abstracts, but only one relation type.

The GDA dataset (Wu et al., 2019) takes a similar approach to CTD-derived, to derive a Gene-Disease associations dataset from the DisGeNET database, using PubTator to link entity mentions. Abstracts are distantly labeled with a single relation type.

5.2. Direct Supervision Biomedical Corpora

DrugProt (Miranda et al., 2021), (Martin Krallinger and Valencia, 2021), is the most recent manually annotated corpus of biomedical research abstracts covering multiple (13) relation types between Chemicals and Genes/Gene-products. ChemDisGene uses a different set of 14 relation types between Chemicals and Genes, derived from CTD. These relations generally describe the observed effect of an interaction. For example, a Localization relation is recorded when the interaction between a Chemical and Gene product affects the part of the cell where the molecule resides. In contrast, the DrugProt relation classes are defined by the specific type of interaction between a Chemical and Gene/Gene-product: they distinguish between ‘Direct’ and ‘Indirect’ regulation (where possible), and the subclasses focus on the direction of the interaction (‘Upregulator’, ‘Downregulator’). The subclasses for direct regulation are highly granular, differentiating between ‘Activator’, ‘Agonist’, ‘Antagonist’, etc.

As an example, the relationship expressed in the text “bisphenol P Chemically showed estrogen receptor-Gene antagonistic activities” would be annotated as Chemical-Gene: antagonist. DrugProt would record it as Chemical-Gene: antagonist.

Our ChemDisGene manually curated corpus is smaller, but also includes relationships between Chemicals and Diseases, and Diseases and Genes. All entity mentions are identified and linked by the models in Pubtator Central, and relationships are distantly labeled, associated with a document but not specific entity mentions. The curated corpus contains ~6 approved relationships per abstract, distinguishing between primary contributions (derived from CTD) and other (‘new’) relationships. Most other manually annotated corpora used in biomedical RE tasks are also directly supervised, and cover fewer relation types, typically between fewer types of entities. As another example, Drug-drug interaction (DDI) (Herrero-Zazo et al., 2013) specifies 4 relation types among drugs, on sentences extracted from 1025 documents.

5.3. Other RE Corpora

In the general domain, there exist several RE benchmarks for sentence level, document level and few-shot scenarios. SemEval 2010 task 8 (Hendrickx et al., 2010) includes ten semantic relation types between nouns over ~11k sentences. The TAC relation extraction dataset (TACRED) (Zhang et al., 2017), includes multi-segmented, linked mentions. TACREL (Ali et al., 2020) and Re-TACRED (Stoica et al., 2021) provides cleaner versions of TACRED. DOCRED (Yao et al., 2019) is a document level relation extraction dataset on the WikiPedia domain, with 5053 manually annotated documents and 100 relation types. FewRel (Han et al., 2018) is a relation extraction benchmark for few-shot scenario, based on WikiPedia. A newer version (Gao et al., 2019) includes Biomedical relations as a domain adaptation task.

5.4. Relation Extraction Models

Traditional RE models have focused on classifying the entity interaction in a sentence. For example, Zeng et al. (2014) encoded sentences and entity pairs with convolutional neural networks and position embeddings. Soares et al. (2019) finetuned Bert with self-supervised signals from entity linking, and applied the model to downstream RE tasks. There is also previous work targeting longer text passages such as cross sentence relation extraction (Quirk and Poon, 2017), or document level distant supervision RE (Verga et al., 2018). Sahu et al., 2019 Christopoulou et al., 2019.

Sahu et al. (2019) and Christopoulou et al. (2019) encode graphs generated from each document for RE. In contrast, BRAN (Verga et al., 2018) uses transformers to encode the text sequence and then evaluates each mention pair of candidate argument entities. All these
Table 5: Performance of baseline models on ChemDisGene CTD-derived ‘dev’, ‘test’ and curated corpora.

| Model                     | P       | R       | F1      | Avg. P | P       | R       | F1      |
|---------------------------|---------|---------|---------|--------|---------|---------|---------|
| CTD-derived corpus: ‘dev’ split / ‘test’ split |         |         |         |        |         |         |         |
| BRAN                      | 32.1 / 31.7 | 46.3 / 44.2 | 37.9 / 36.9 | 28.4 / 27.9 | 25.9 / 23.6 | 32.3 / 30.1 | 28.2 / 26.0 |
| PubmedBert                | 50.3 / 49.6 | 59.3 / 56.1 | 54.5 / 52.6 | 50.3 / 50.1 | 43.6 / 39.0 | 50.3 / 48.4 | 44.9 / 41.7 |
| PubmedBert + BRAN         | 53.9 / 53.9 | 61.0 / 57.3 | 57.3 / 55.6 | 54.0 / 54.3 | 45.0 / 42.7 | 54.1 / 50.4 | 48.7 / 44.4 |
| Curated corpus: CTD-derived relationships only / All relationships |         |         |         |        |         |         |         |
| BRAN                      | 24.4 / 41.8 | 45.8 / 26.6 | 31.8 / 32.5 | 28.1 / 33.5 | 20.3 / 37.2 | 35.7 / 22.5 | 24.5 / 25.8 |
| PubmedBert                | 43.0 / 64.3 | 61.7 / 31.3 | 50.7 / 42.1 | 50.7 / 46.9 | 34.7 / 53.7 | 53.4 / 32.0 | 39.6 / 37.0 |
| PubmedBert + BRAN         | 46.5 / 70.9 | 61.1 / 31.6 | 52.8 / 43.8 | 53.0 / 50.6 | 45.8 / 69.8 | 59.0 / 32.5 | 47.0 / 40.5 |

Table 6: ‘PubmedBert + BRAN’ model metrics for each relation type in the curated corpus, sorted on decreasing relation frequency in the training data.

| Relation Type | F1   |
|---------------|------|
| Chemical-Disease : marker/mechanism | 54.1 |
| Chemical-Disease : therapeutic | 45.5 |
| Chemical-Gene : expression - increases | 58.2 |
| Chemical-Gene : expression - decreases | 61.6 |
| Gene-Disease : marker/mechanism | 47.1 |
| Chemical-Gene : activity - increases | 52.4 |
| Chemical-Gene : activity - decreases | 56.3 |
| Chemical-Gene : metabolic_processing - increases | 36.4 |
| Chemical-Gene : binding - affects | 58.1 |
| Chemical-Gene : transport - increases | 36.1 |
| Chemical-Gene : metabolic_processing - decreases | 34.4 |
| Chemical-Gene : localization - affects | 48.9 |
| Chemical-Gene : expression - affects | 0.4 |
| Gene-Disease : therapeutic | 28.6 |

6. Conclusion

We introduced ChemDisGene, a new dataset of research abstracts labeled with biomedical entity mentions and distance-labeled with biomedical relationships, for training and evaluating multi-type multi-label biomedical RE models. The dataset includes a large automatically derived corpus with noisy relationship labels (~ 22% noise based on manual curation), and a cleaner manually curated dataset of 523 abstracts. We also provided three baseline ML models for RE, trained and evaluated on the ChemDisGene dataset. We believe this is the first dataset for biomedical relation extraction tasks that addresses multiple entity (more than 2) and relation types, and includes both a large automatically derived corpus (useful for model training), as well as a smaller corpus labeled by human experts.

Manually annotating raw text with biomedical relationships is a hard and time consuming task, even for domain experts. We facilitated the curation with high quality models for entity recognition. Future refinements to this dataset could include verifying the linked entities in the curated corpus, and adding Protein-Protein interactions, useful for understanding disease mechanisms and drug repurposing.

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