aracne.networks, a data package containing gene regulatory networks assembled from TCGA data by the ARACNe algorithm

Federico M. Giorgi^{1,2}, Mariano J. Alvarez^{1,3}, and Andrea Califano¹

¹Department of Systems Biology, Columbia University, New York, USA
²CRUK, Cambridge University, Cambridge, UK
³DarwinHealth Inc., New York, USA

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1 Overview of aracne.networks data package

The *aracne.networks* data package provides context-specific transcriptional regulatory networks (also called interactomes or regulators) reverse engineered by the ARACNe algorithm from The Cancer Genome Atlas (TCGA) RNAseq expression profiles.

ARACNe networks This package contains 24 Mutual Information-based networks assembled by ARACNe-AP [1] with default parameters (MI p-value = 10^{-8} , 100 bootstraps and permutation seed = 1). ARACNe is a network inference algorithm based on an Adaptive Partioning (AP) Mutual Information (MI) approach [1]. In short, ARACNe-AP estimates all pairwise Mutual Information scores between gene expression profiles, then assesses the significance of such Mutual Information by comparison to a null dataset. ARACNe then draws network edges between centroid genes (Transcription Factors and Signaling Proteins) and genes significantly associated with them (i.e. with significant MI). It then calculates Data Processing Inequality (DPI) to reduce the number of indirect connections.

ARACNe-AP was run on RNA-Seq datasets normalized using Variance-Stabilizing Transformation [2]. The raw data was downloaded on April 15th, 2015 from the TCGA official website [3]. We follow the TCGA naming convention (e.g. BRCA = Breast Carcinoma) to name the individual context-specific networks.

```
> library(aracne.networks)
> data(package="aracne.networks")$results[, "Item"]
```

```
[1] "regulonblca" "regulonbrca" "reguloncesc" "reguloncoad" "regulonesca" [6] "regulongbm" "regulonhnsc" "regulonkirc" "regulonkirp" "regulonlaml" [11] "regulonlihc" "regulonluad" "regulonlusc" "regulonov" "regulonpaad" [16] "regulonpcpg" "regulonprad" "regulonread" "regulonsarc" "regulonstad" [21] "regulontgct" "regulonthca" "regulonthym" "regulonucec"
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Write a network to file The package contains a function to print individual networks into a file. Four columns will be printed: the Regulator id, the Target id, the Mode of Action (MoA, inferred by Spearman correlation analysis [4]) that indicates the sign of the association between regulator and target gene and ranges between -1 and +1, the Likelihood (essentially an edge weight that indicates how strong the mutual information for an edge is when compared to the maximum observed MI in the network, it ranges between 0 and 1). Further details about the regulon object as a model for transcriptional regulation are present in the manuscript [4].

In the following example, we print the first 10 interactions from the bladder carcinoma (blca) network. The network genes are identified by Entrez Gene ids.

- > data(regulonblca)
- > write.regulon(regulonblca, n = 10)

Regulator	Target	MoA	likelihood
10002	2648	0.99468959127046	0.886774633189913
10002	677827	0.116175345640	0.707841406455471
10002	80152	0.9997704370156	0.950286744281199
10002	284382	-0.03684243335	0.0419762049859333
10002	9866	0.97206659815444	18 0.442238853411591
10002	283422	-0.57408492938	35018 0.260828476620346
10002	221613	-0.09592426018	320319 0.717904706549976
10002	348174	0.953943934091	1558 0.814491117578869
10002	373509	0.704691385719	9852 0.244337186726846
10002	8803	-0.9591656560869	0.831653033754096

The user may want to analyze all the connections of a particular regulator (E.g. "399", the RHOH gene).

- > data(regulonblca)
- > write.regulon(regulonblca, regulator="399")

Regulator	Targe	t	MoA	likelihood
399	9595	1	0.9999994	39751274
399	54440	1	0.999999	439753891
399	5788	1	0.9999936	91255193
399	2124	1	0.9999939	72431349
399	10563	0.9999	9999999987	0.999880973084544
399	80342	1	0.999979	237947268
399	1840	0.99999	9959099145	0.994240739975982
399	8875	0.99999	999999397	0.999602389369848
399	6689	0.99999	9999998723	0.999531614767901
399	200186	0.154	40359065400	8 0.948828817305409
399	165631	0.999	9999995056	5 0.998777586463862
399	54509	0.9999	99981560018	0.997883918024065
399	171389	0.999	99999482404	4 0.996800613785205
399	147929	-0.99	91545345527	0.985197674740525
399	23416	0.9999	29331217517	0.96812145442081
399	26015	-0.992	83846636841	2 0.834785111763068
399	10148	0.9999	9999999872	0.999729153685544
399	4951	-0.0504	64773052601	5 0.544073601564966
399	57003	-0.075	17089290228	0.714920200879607

References

- [1] Giorgi, F.M. et al. (2016) ARACNe-AP: Gene Network Reverse Engineering through Adaptive Partitioning inference of Mutual Information. Bioinformatics doi: 10.1093/bioinformatics/btw216.
- [2] Anders, S and Huber W. (2010) Differential expression analysis for sequence count data. Genome Biol 2010;11(10):R106
- [3] Weinstein J.N. et al. (2013) The cancer genome atlas pan-cancer analysis project. Nature Genetics 45, 1113-1120 2013

[4]	Alvarez M.J. et al. (2016) Functional characterization of somatic mutations in cancer using network-based inference of protein activity. Nature Genetics $in\ press.$					