

예비프로젝트 #6 : Pubmed 논문지 저널별 분석

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목차

1. File Generator

- 2015~2019 Journal 별 논문 수 확인
- 2015~2019 5년간 Journal 별 논문 추출

2. Analysis

- wordcloud
- keyword 분포
- GNI와 유사한 논문지 찾기
- + 12개의 저널별 논문 accepted 기간 분석

3. K-Means Clustering

- 논문지 clustering

1. File Generator

- 2015~2019 Journal 별 논문 수 확인
- 2015~2019 5년간 Journal 별 논문 추출

2015~2019 Journal 별 논문 수

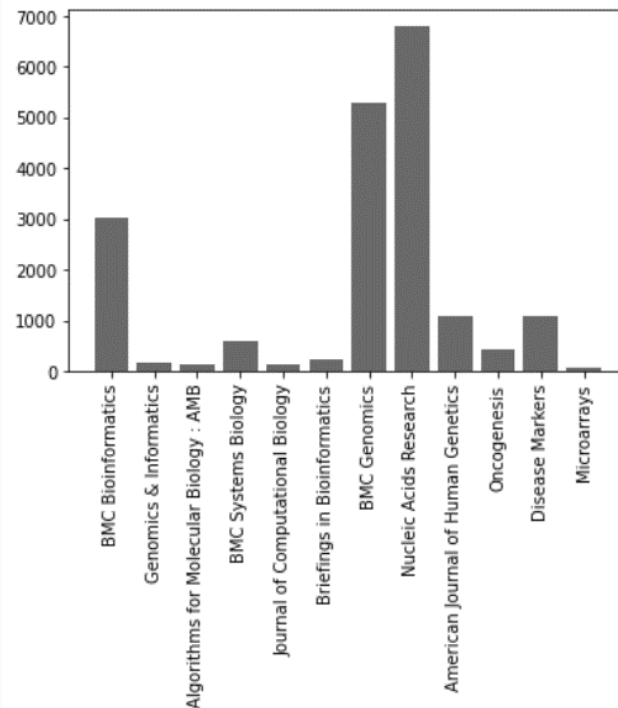
["BMC Bioinformatics", "Genomics & Informatics", "Algorithms for Molecular Biology : AMB", "BMC Systems Biology", "Journal of Computational Biology", "Briefings in Bioinformatics", "BMC Genomics", "Nucleic Acids Research", "American Journal of Human Genetics", "Oncogenesis", "Disease Markers", "Microarrays"] 저널에 대해,

최근 5년 간의 12개의 Journal 별 발행 논문수를 살펴보면 다음과 같다.

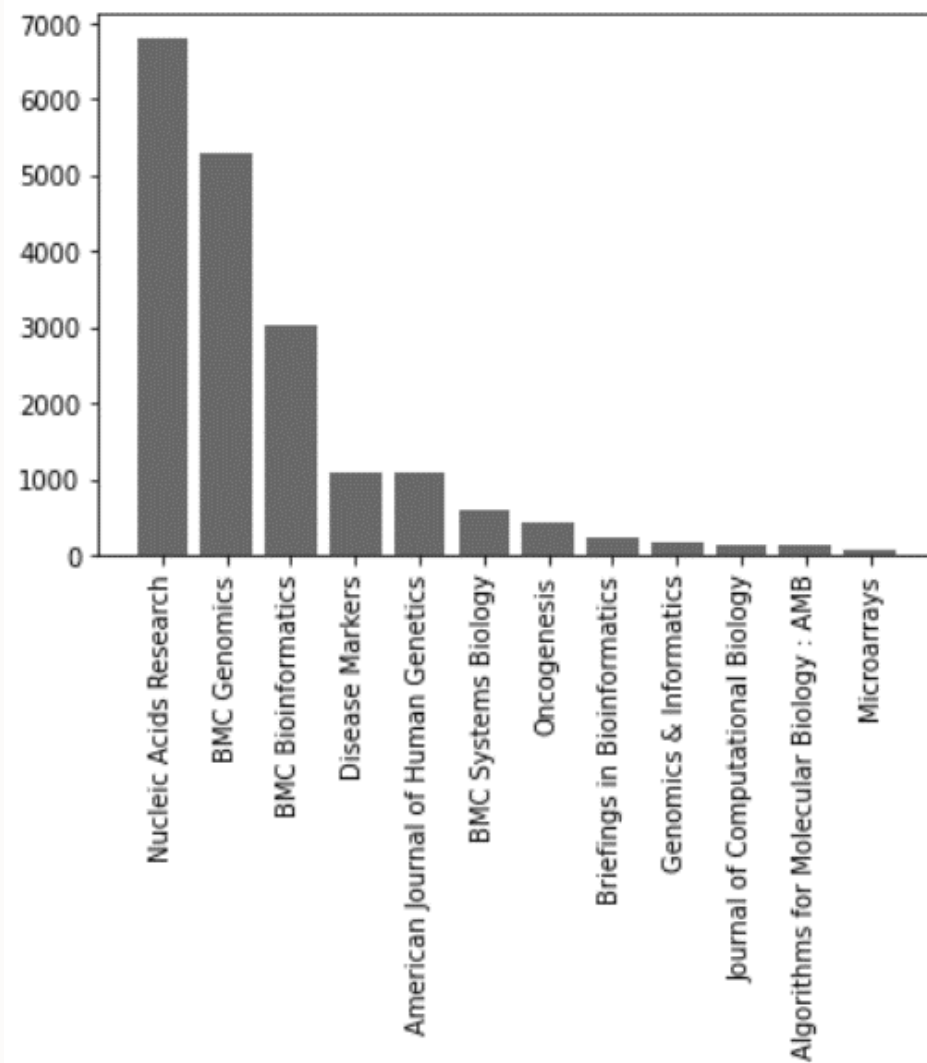
```
import pandas as pd
import numpy as np
```

```
journalList = ["BMC Bioinformatics", "Genomics & Informatics", "Algorithms for Molecular Biology : AMB", "BMC Systems Biology", "Journal of Computational Biology", "Briefings in Bioinformatics", "BMC Genomics", "Nucleic Acids Research", "American Journal of Human Genetics", "Oncogenesis", "Disease Markers", "Microarrays"]
journalList_count = [3017, 181, 128, 596, 130, 241, 5294, 6803, 1072, 416, 1081, 79]
journalList2 = pd.DataFrame({'Name': journalList, 'Count': journalList_count})
journalList2 = journalList2.sort_values(by='Count', ascending=False)
journalList2
```

	Name	Count
0	BMC Bioinformatics	3017
1	Genomics & Informatics	181
2	Algorithms for Molecular Biology : AMB	128
3	BMC Systems Biology	596
4	Journal of Computational Biology	130
5	Briefings in Bioinformatics	241
6	BMC Genomics	5294
7	Nucleic Acids Research	6803
8	American Journal of Human Genetics	1072
9	Oncogenesis	416
10	Disease Markers	1081
11	Microarrays	79



2015~2019 Journal 별 논문 수



2015~2019년 5년간 저널별 논문 발행 수는 Nucleic Acids Research가 약 6000여개, BMC Genomics가 약 5000여개, BMC Bioinformatics 약 3000여개로 높았다.

논문의 개수가 너무 많아 파일 관리하기가 어려우므로, 2015,2016,2017,2018,2019년 각각 abstract 만 따로 떼어내어 파일을 저장하였다.

2015~2019 Journal 별 논문 수 - Code

[2015~2019 5년간]

12개의 Journal List별로 파일을 생성하기 전, 5년간 발행된 논문의 개수를 확인하기위한 코드이다.

```
[ ] # -*- encoding: utf-8 -*-
from Bio import Entrez
from Bio import Medline

Entrez.email = "neo.ewha@gmail.com"
journalList = ["BMC Bioinformatics", "Genomics & Informatics", "Algorithms for Molecular Biology : AMB", "BMC Systems Biology", "Journal of Computational Biology", "Briefings in Bioinformatics", "BMC Genomics", "Nucleic Acids Research", "American J

curdir = "/content/gdrive/My Drive/인공지능/5년간꺼번에/"
for journal in journalList:
    keyword = "(#" + journal + "#"[Journal]) AND (#"2015/01/01#[Publication Date] : #"2019/12/31#[Publication Date]) "
    handle1 = Entrez.esearch(db='pmc', term=keyword, retmax=10000)
    record = Entrez.read(handle1)
    idlist = record['IdList']
    handle1.close()
    #print(journal, ":", idlist)

    handle2 = Entrez.efetch(db='pmc', id=idlist, rettype='medline', retmode='text')
    records = Medline.parse(handle2)
    records = list(records)
    #print(records[0].keys())
    i = 0
    if idlist: # if journal list is not empty
        for record in records:
            #print(idlist[i] + ": " + record.get("TI", "?"))
            #f = open(curdir + journal + "_" + str(idlist[i]) + ".txt", "w", encoding="utf-8")
            i += 1
            #f.write("Title: " + ''.join(record.get("TI", "?")) + "\n")
            #f.write("Authors: " + ''.join(record.get("AU", "?")) + "\n")
            #f.write("Source: " + ''.join(record.get("SO", "?")) + "\n")
            #f.write("Abstract: " + ''.join(record.get("AB", "?")) + "\n")
            #f.close()
    else:
        print(journal + ": list empty!!!")
print(i)
```

3017
181
128
596
130
241
5294
6803
1072
416
1081
79

File Generator with Bio Python

[2015년]

12개의 Journal List별로 논문의 abstract만을 추출하여 파일 생성

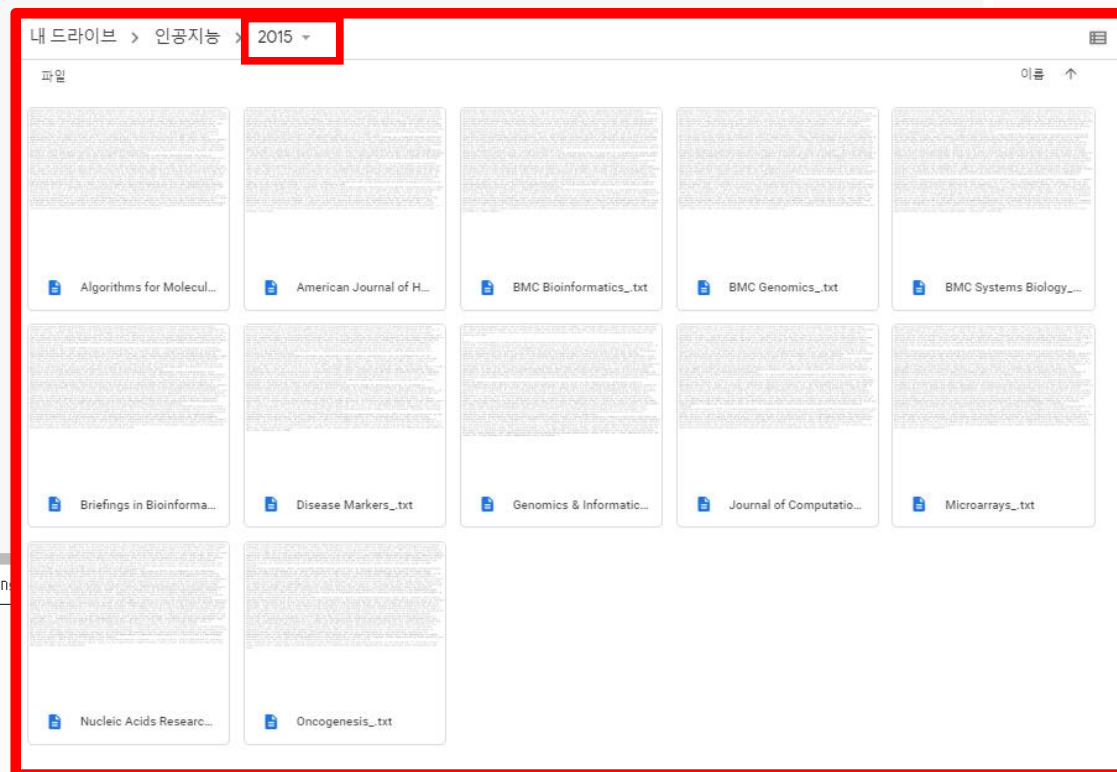
```
# -*- encoding: utf-8 -*-
from Bio import Entrez
from Bio import Medline

Entrez.email = "neo.ewha@gmail.com"
journalList = ["BMC Bioinformatics", "Genomics & Informatics", "Algorithms for Molecular Biology : AMB", "BMC Systems Biology", "Journal of Computational Biology", "Briefings in Bioinformatics", "BMC Genomics", "Nucleic Acids Research", "American Journal of Human Genetics"]
# keyword = ("BMC Bioinformatics"[Journal]) AND ("2015/01/01"[Publication Date] : "2020/05/31"[Publication Date]) "

curdir = "/content/gdrive/My Drive/인공지능/2015년/"
for journal in journalList:
    f = open(curdir + journal + "_" + ".txt", "w", encoding="utf-8")
    keyword = ("BMC Bioinformatics"[Journal]) AND ("2015/01/01"[Publication Date] : "2015/12/31"[Publication Date]) "
    handle1 = Entrez.esearch(db="pmc", term=keyword, retmax=20)
    record = Entrez.read(handle1)
    idlist = record['IdList']
    handle1.close()
    #print(journal, ":", idlist)

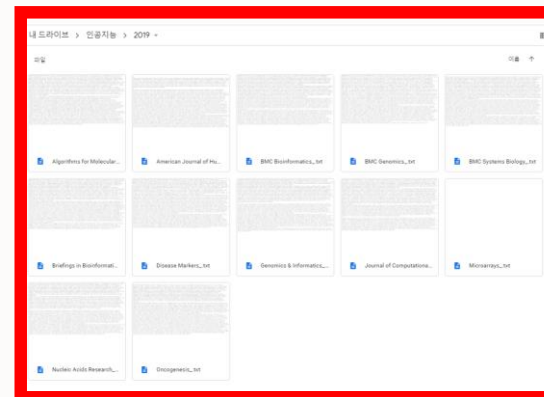
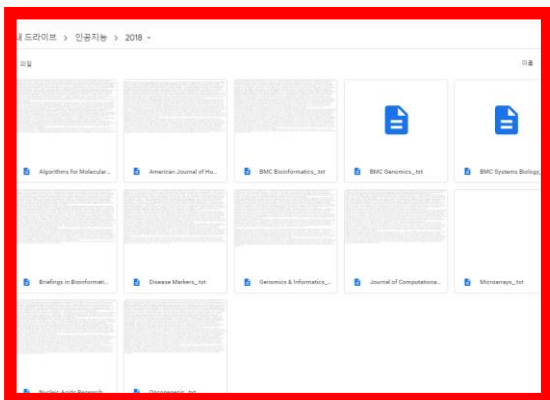
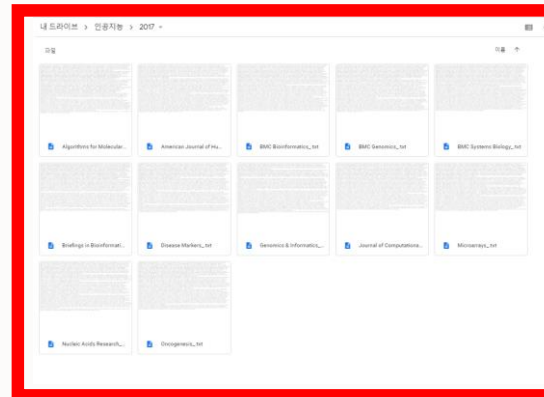
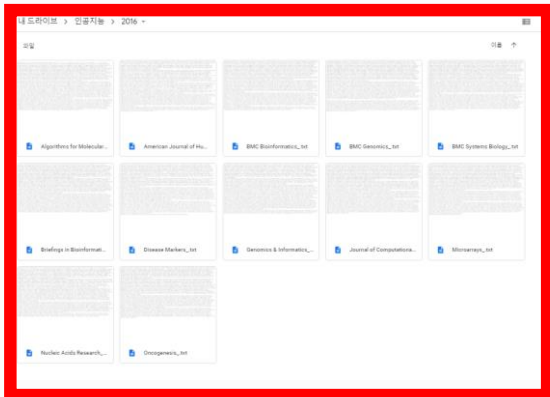
    handle2 = Entrez.efetch(db="pmc", id=idlist, rettype="medline", retmode="text")
    records = Medline.parse(handle2)
    records = list(records)
    #print(records[0].keys())
    i = 0
    if idlist: # if journal list is not empty
        for record in records:
            print(idlist[i] + ":", record.get("TI", "?"))
            #f = open(curdir + journal + "_" + str(idlist[i]) + ".txt", "w", encoding="utf-8")
            i += 1
            f.write(":" + record.get("AB", "?"))
    else:
        print(journal + ": list empty!!!")
    f.close()
```

4697333: Multi-label multi-instance transfer learning for simultaneous reconstruction and cross-talk modeling of multiple human signaling pathways
4595314: BioPy: a MySQL-based framework to handle genomic variants in 12k2



File Generator with Bio Python

[2016년]/[2017년]/[2018년]/[2019년] 도 마찬가지로
12개의 Journal List별로 논문의 abstract만을 추출하여 파일 생성

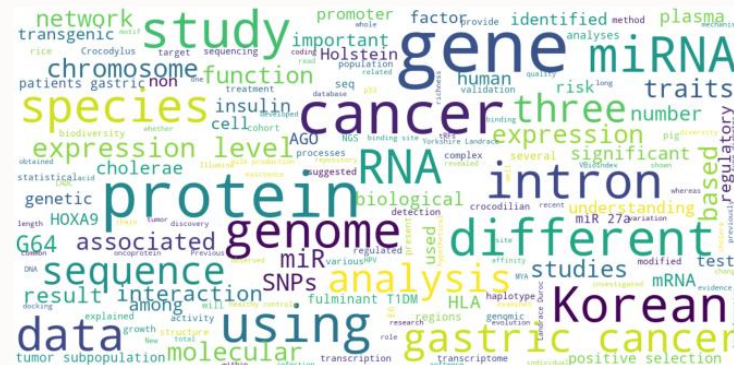


2. Analysis

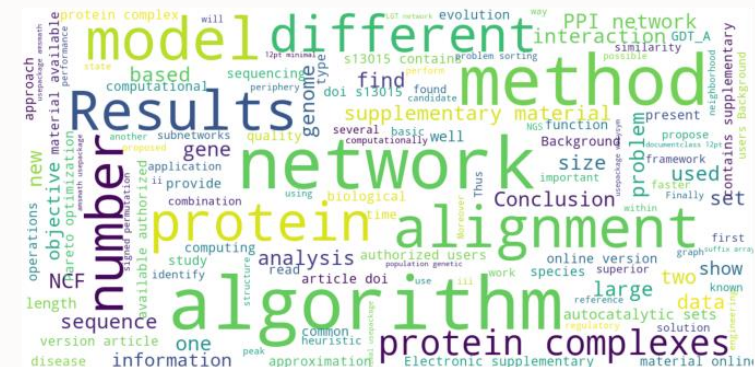
- wordcloud
- keyword 분포
- GNI와 유사한 논문지 찾기
- + 12개의 저널별 논문 accepted 기간 분석



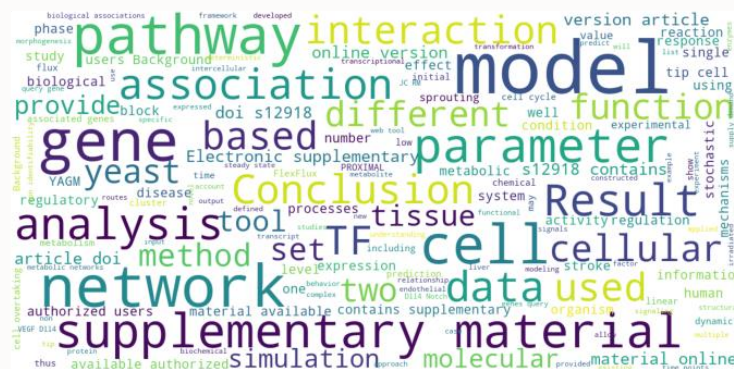
BMC Bioinformatics



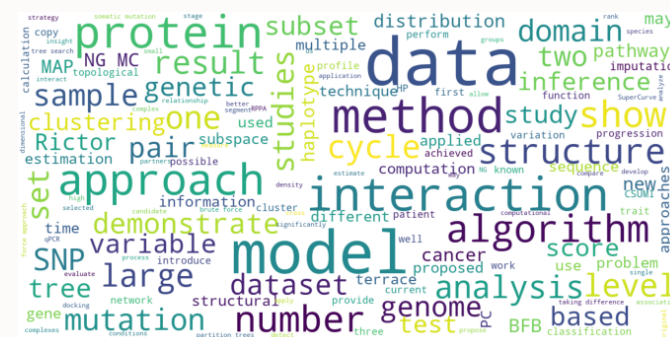
Genomics & Informatics



Algorithms for Molecular Biology : AMB



BMC Systems Biology



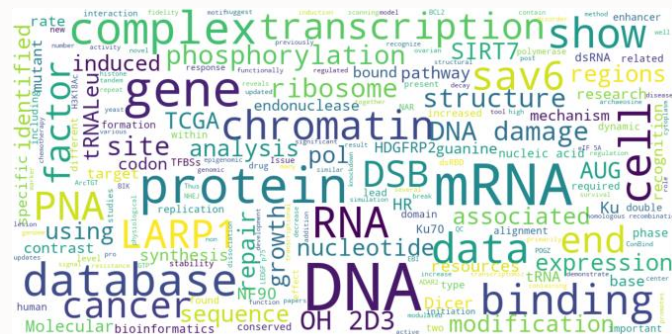
journal of Computational Biology



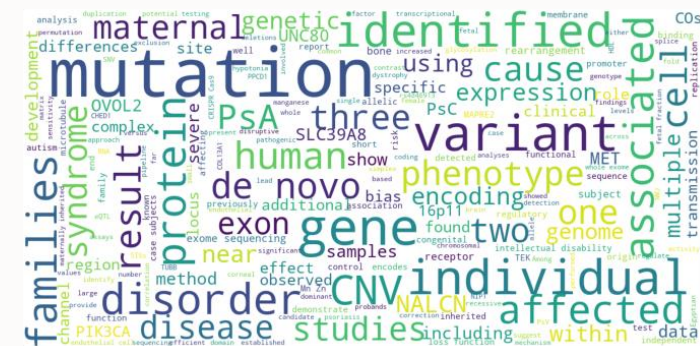
Briefings in Bioinformatics



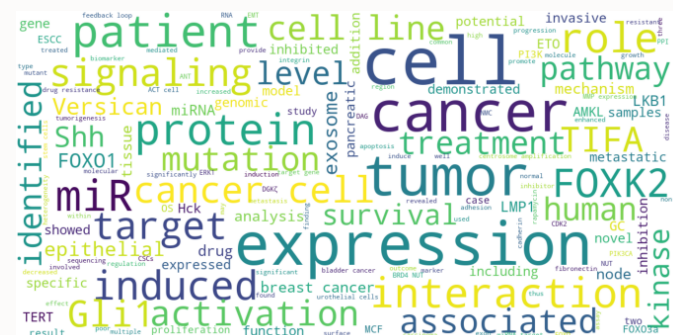
BMC Genomics



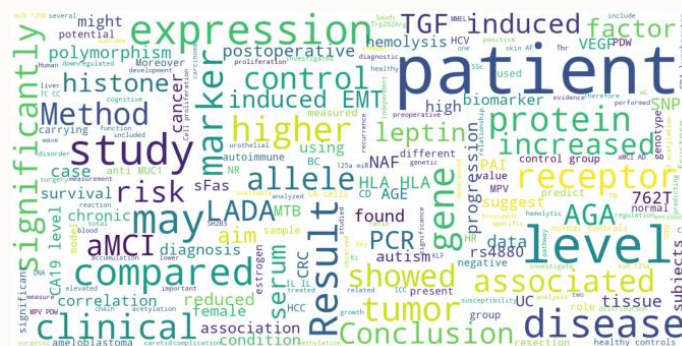
Nucleic Acids Research



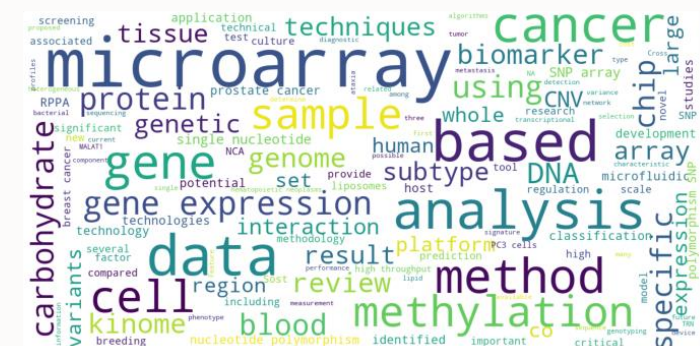
American Journal of Human Genetics



Oncogenesis



Disease Markers



Microarrays

저널별 워드클라우드 - Code

```
[9] curdir = "/content/gdrive/My Drive/인공지능/2015/"
journalList = ["BMC Bioinformatics", "Genomics & Informatics", "Algorithms for Molecular Biology : AMB", "BMC Systems Biology", "Journal of Computational Biology", "Briefings in Bioinformatics", "BMC Genomics", "Nucleic Acids Research", "American Journal c
j = []
for journal in journalList:
    f = open(curdir + journal + "_ " + ".txt", "r", encoding="utf-8")
    f = f.read()
    j.append(f)
```

```
[11] !pip install wordcloud
```

Requirement already satisfied: wordcloud in /usr/local/lib/python3.6/dist-packages (1.5.0)
Requirement already satisfied: numpy>=1.6.1 in /usr/local/lib/python3.6/dist-packages (from wordcloud) (1.18.5)
Requirement already satisfied: pillow in /usr/local/lib/python3.6/dist-packages (from wordcloud) (7.0.0)

```
from wordcloud import WordCloud, STOPWORDS
import matplotlib.pyplot as plt
stopwords = set(STOPWORDS)

def show_wordcloud(data, title = None):
    wordcloud = WordCloud(
        background_color='white',
        stopwords=stopwords,
        max_words=200,
        max_font_size=40,
        scale=3,
        random_state=1 # chosen at random by flipping a coin: it was heads
    ).generate(str(data))

    fig = plt.figure(1, figsize=(12, 12))
    plt.axis('off')
    if title:
        fig.suptitle(title, fontsize=20)
        fig.subplots_adjust(top=2.3)

    plt.imshow(wordcloud)
    plt.show()

#txt = 'Background: During evolution, global mutations may alter the order and the orientation of the genes in a genome. Such mutations are referred to as rearrangement events, or simply operations. In unichromosomal genomes, the most common operations are
for i in range(12):
    print(journalList[i])
    show_wordcloud(j[i])
```

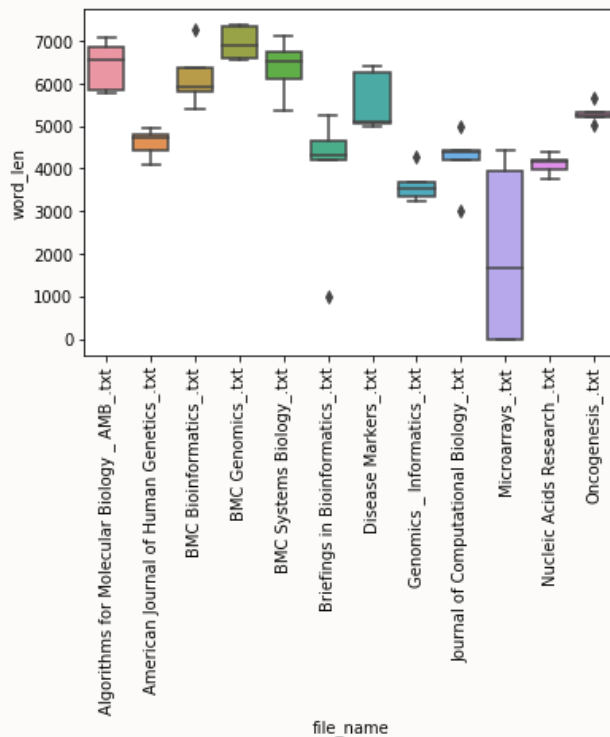
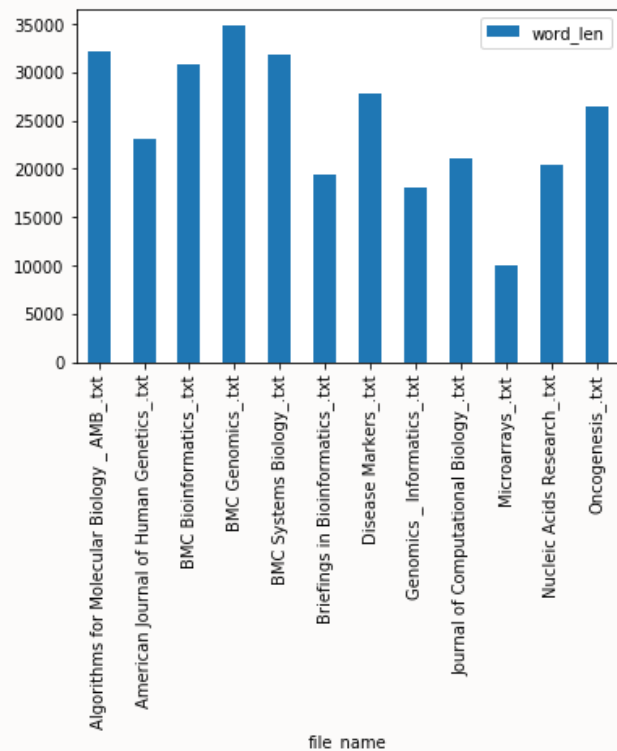
저널별 단어분포

앞에서 살펴본 워드클라우드를 통해, 자주 등장하는 단어를 정리해보면 다음과 같다.

BMC Bioinformatics	- data, sequence, protein, genome
Genomics & Informatics	- algorithm, network, method, alignment
Algorithms for Molecular Biology : AMB	- protein, cancer, genome, gene, RNA, Korean
BMC Systems Biology	- model, gene, network, material, analysis, pathway, association
Journal of Computational Biology	- data, method, model, algorithm, approach, interaction, structure
Briefings in Bioinformatics	- data, gene, genome, method, cancer, analysis
BMC Genomics	- gene, genome, analysis, pretein infection
Nucleic Acids Research	- DNA, mRNA, database, gene, data, binding
American Journal of Human Genetics	- mutation, variant, gene, individual, disorder
Oncogenesis	- cell, cancer, expression, tumor
Disease Markers	- patient, study, level, result, disease
Microarrays	- microarray,based, data, analysis, method, cell

저널별 단어분포

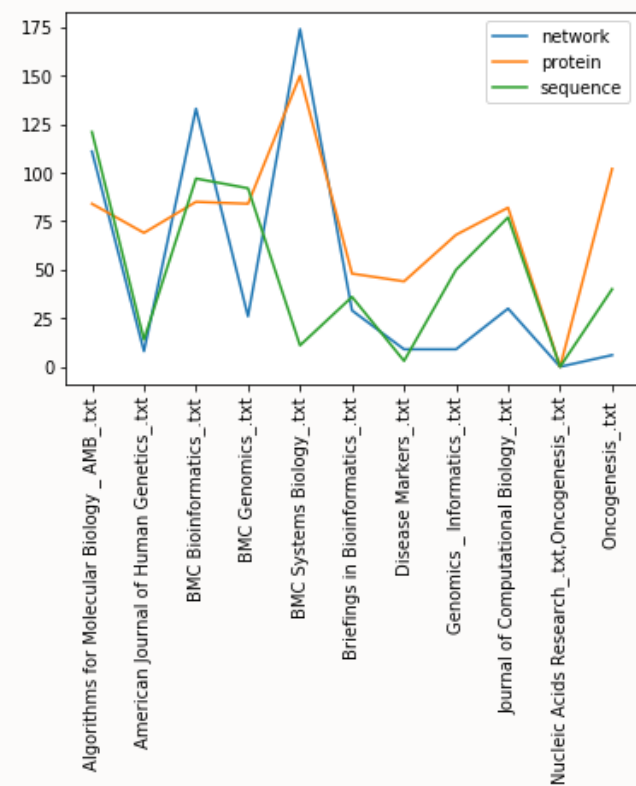
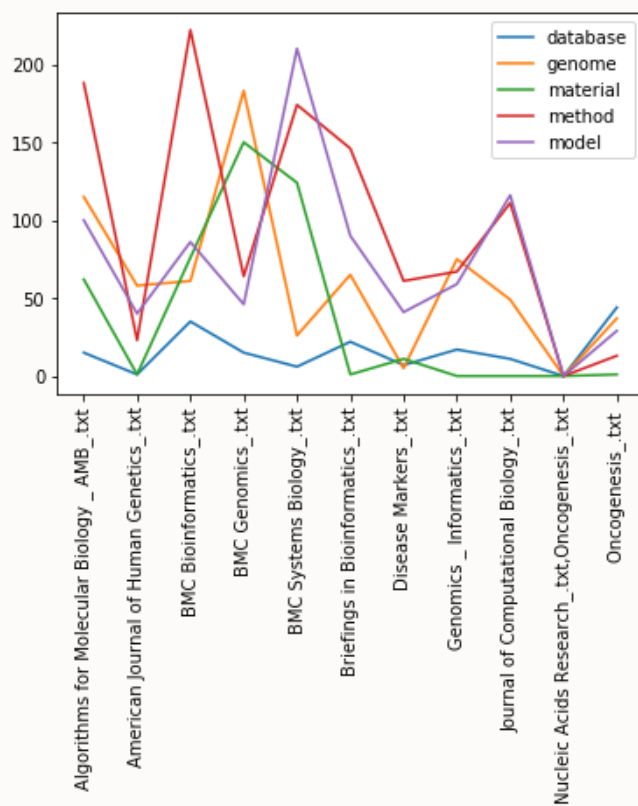
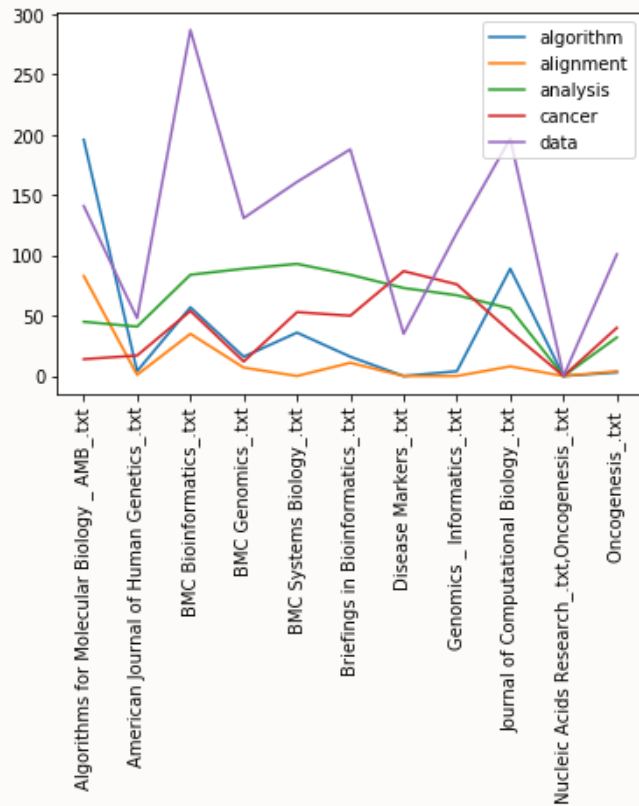
단어분포를 살펴보기 전에 우선 저널별 단어의 길이를 살펴보면 다음과 같다.
비슷한 수의 파일을 선택했음에도 불구하고 약간의 차이가 존재하여 이를 고려하여 앞으로 분석을 할 것이다.



*Microarray 저널은 표본의 수가 너무 부족하기 때문에 왜곡이 될 가능성이 있어 우선 제외하고 분석함.

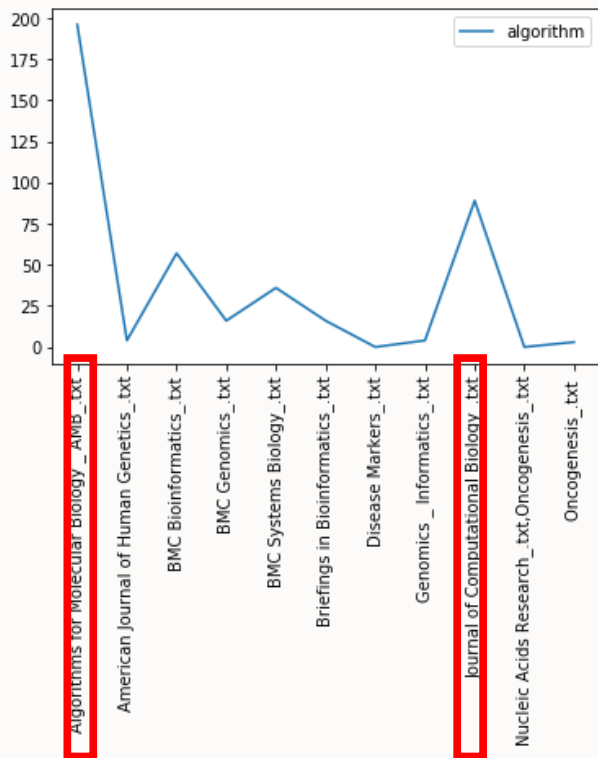
저널별 단어분포 - Code

앞에서 살펴본 자주 등장하는 단어들에 대해 시각화하여 저널별로 비교를 해보면 다음과 같다.

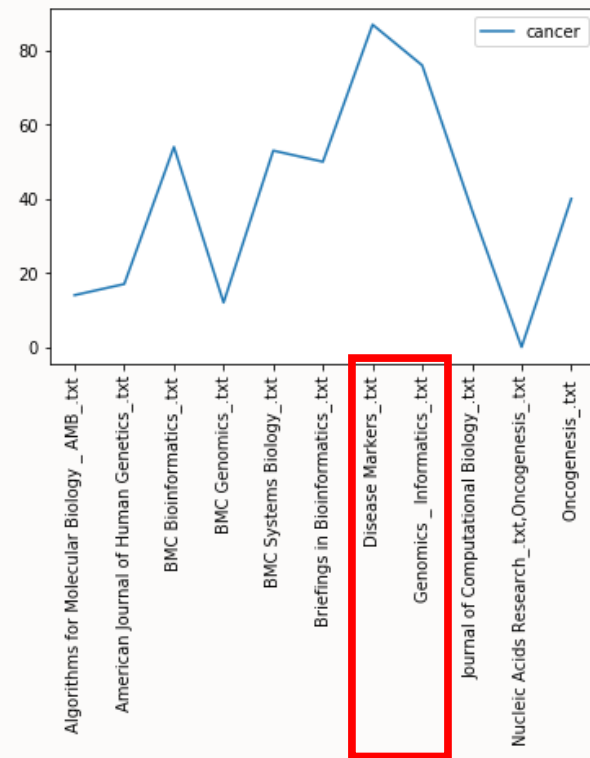


저널별 단어분포 - Code

앞의 그래프를 통해 의미있는 결과를 간단히 정리해보자.



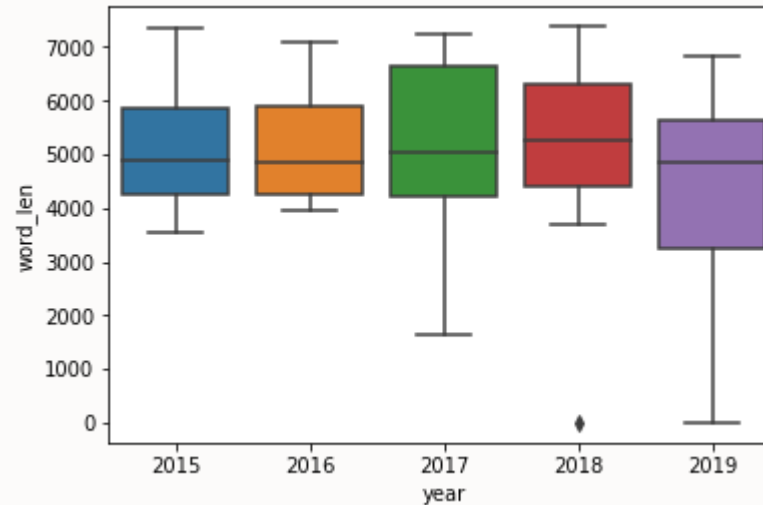
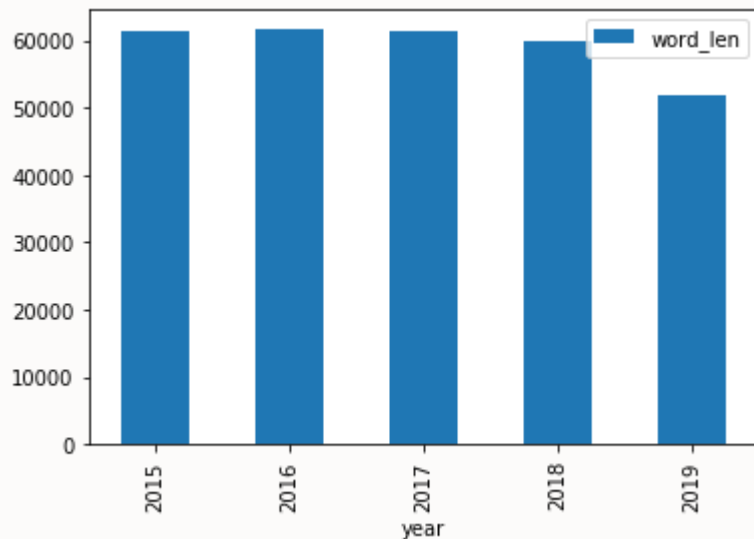
-> algorithm 단어는 Algorithms for Molecular Biology _AMB_저널에서 압도적으로 많이 등장하였으며, Journal of Computational Biology 저널에서도 꽤 많이 등장하였다. 이는 알고리즘, 컴퓨터학문에 관련한 저널이기 때문이라고 유추할 수 있다.



-> cancer 단어는 Disease Markers 저널, Genomics_Informatics에서 자주 등장하였다.

2015~2019 시간에 따른 단어분포 변화

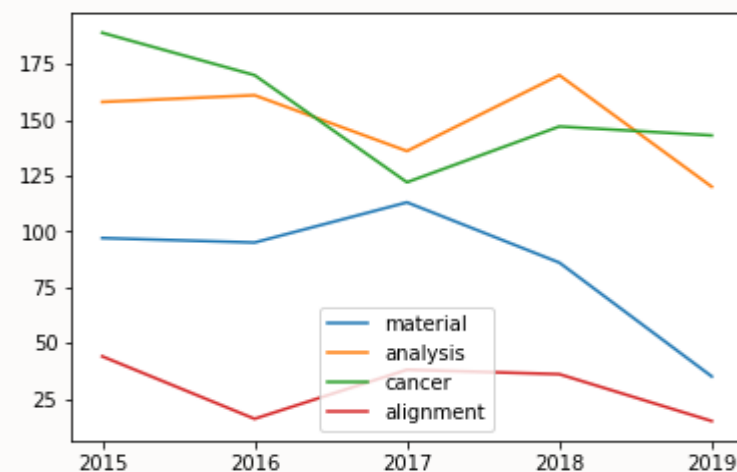
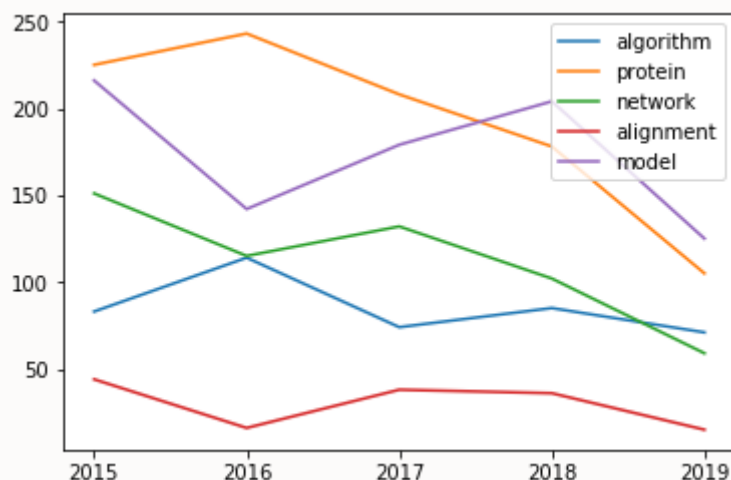
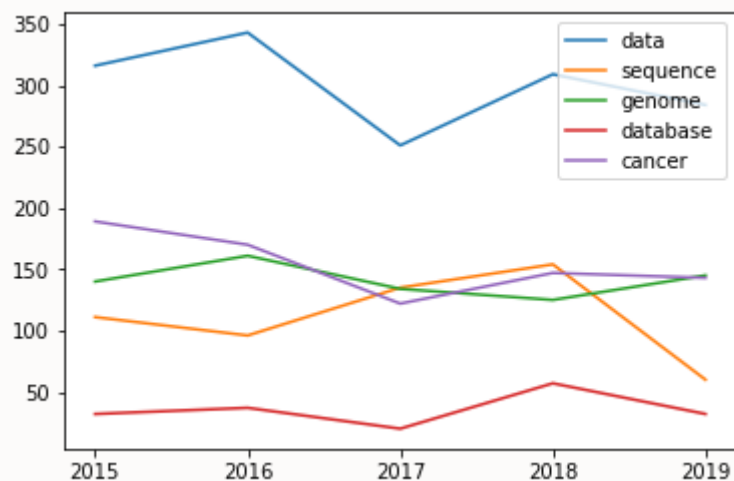
단어분포를 살펴보기 전에 우선 연도별 단어의 길이를 살펴보았다.



논문의 수를 비슷하게 뽑아왔기 때문에, 데이터 분포에 큰 치우침이 없어보인다.
따라서 이 데이터를 그대로 분석에 이용해도 크게 문제 없을 것 같다.
실제로 총단어갯수를 고려한 그래프를 따로 그려보았는데 기존 그래프와 큰 차이가 없었다.

2015~2019 시간에 따른 단어분포 변화

pubmed data의 2015~2019년 abstract의 단어 분포를 살펴본 결과이다.



대체적으로 data 단어가 자주 사용되는 것을 확인할 수 있고 model이라는 단어가 2018년에 많이 사용된 것을 확인할 수 있다. 이외에도 각 단어별 시간에 따른 분포 변화를 확인할 수 있다.

+ 12개의 저널별 논문 accepted 기간 분석

received ~ accepted 기간

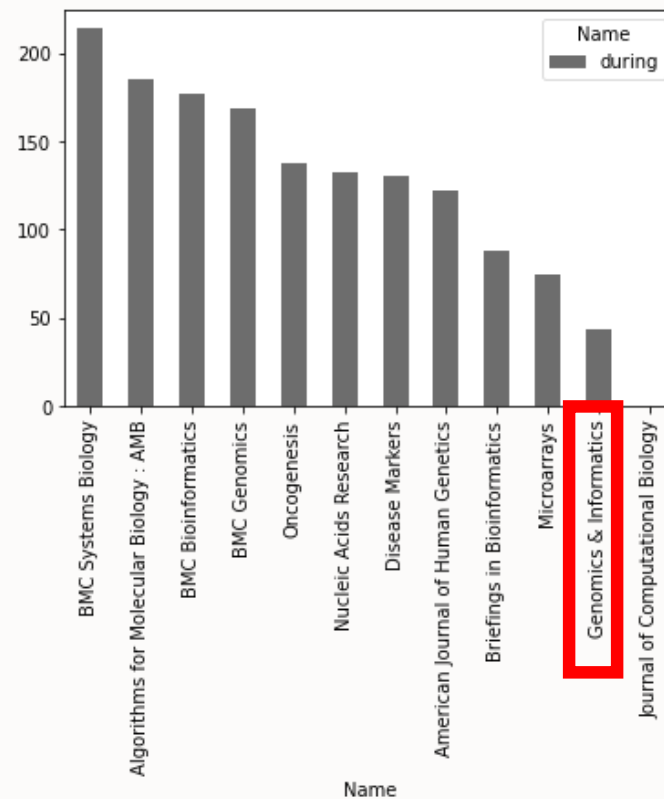
	Name	received	accepted	during
0	BMC Bioinformatics	2020-04-17	2020-05-11	24.0
1	BMC Bioinformatics	2019-06-05	2020-04-30	330.0
2	BMC Bioinformatics	2019-11-03	2020-05-11	190.0
3	BMC Bioinformatics	2019-11-19	2020-04-15	148.0
4	BMC Bioinformatics	2019-08-06	2020-03-31	238.0
...
595	Microarrays	2015-08-19	2015-11-04	77.0
596	Microarrays	2015-08-14	2015-10-16	63.0
597	Microarrays	2015-09-19	2015-10-20	31.0
598	Microarrays	2015-09-15	2015-10-22	37.0
599	Microarrays	2015-07-30	2015-10-15	77.0

600 rows × 4 columns

저널별 received ~ accepted 기간 평균

Name	
Algorithms for Molecular Biology : AMB	185.220000
American Journal of Human Genetics	122.441860
BMC Bioinformatics	176.812500
BMC Genomics	168.659091
BMC Systems Biology	214.391304
Briefings in Bioinformatics	87.933333
Disease Markers	130.040000
Genomics & Informatics	43.347826
Journal of Computational Biology	NaN
Microarrays	74.645833
Nucleic Acids Research	132.720000
Oncogenesis	137.708333

저널별 received ~ accepted 기간 평균



-> genomics & informatics의 논문들은
다른 논문들에 비해 received 하고 accepted 되는데에 걸리는 기간이 굉장히 짧은 편이었다.

+ 12개의 저널별 논문 accepted 기간 분석 - Code

*자세한 코드는 생략

```
[ ] # -*- encoding: utf-8 -*-
from Bio import Entrez
from Bio import Medline

Entrez.email = "neo.esha@gmail.com"
JournalList = ["BMC Bioinformatics", "Genomics & Informatics", "Algorithms for Molecular Biology : AMB", "BMC Systems Biology", "Journal of Computational Biology", "Briefin

PHST_list = []

for journal in JournalList:
    keyword = "(" + journal + ")" AND ("2015/01/01"[Publication Date] : "2020/05/31"[Publication Date]) "
    handle1 = Entrez.esearch(db="pub", term=keyword, retmax=50)
    record = Entrez.read(handle1)
    idlist = record["idlist"]
    handle1.close()

    handle2 = Entrez.efetch(db="pub", id=idlist, rettype="medline", retmode="text")
    records = Medline.parse(handle2)
    records = list(records)
    i = 0
    if idlist: # If Journal list is not empty
        for record in records:
            PHST_list.append(record.get("PHST", ""))
            #f = open(cwdir + journal + "_" + str(idlist[i]) + ".txt", "w", encoding="utf-8")
            i += 1
            #f.write("Title: " + ''.join(record.get('TI', "")) + "\n")
            #f.close()
        else:
            print(journal + ": list empty!!!")


[ ] len(PHST_list)
PHST_list
JournalList
name = []
for i in range(12):
    for j in range(50):
        name.append(JournalList[i])
```

```
[ ] import pandas as pd
PHST_dataframe = pd.DataFrame({'Name': name, 'received':received, 'accepted':accepted})
PHST_dataframe['received'] = pd.to_datetime(PHST_dataframe['received'],errors='coerce')
PHST_dataframe['accepted'] = pd.to_datetime(PHST_dataframe['accepted'],errors='coerce')
```

```
[ ] PHST_dataframe['during'] = abs(PHST_dataframe['accepted']- PHST_dataframe['received'])
```

```
[ ] PHST_dataframe['during'] = pd.to_numeric(PHST_dataframe['during'].dt.days, downcast='integer')
```

```
[ ] PHST_dataframe
```

	Name	received	accepted	during
0	BMC Bioinforma	 name [néim] 이름,명성,명명하다,성명,명칭		복사
1	BMC Bioinformatics	2019-06-05	2020-04-30	330.0
2	BMC Bioinformatics	2019-11-03	2020-05-11	190.0
3	BMC Bioinformatics	2019-11-19	2020-04-15	148.0
4	BMC Bioinformatics	2019-08-06	2020-03-31	238.0
...
595	Microarrays	2015-08-19	2015-11-04	77.0
596	Microarrays	2015-08-14	2015-10-16	63.0
597	Microarrays	2015-09-19	2015-10-20	31.0
598	Microarrays	2015-09-15	2015-10-22	37.0
599	Microarrays	2015-07-30	2015-10-15	77.0

600 rows × 4 columns

3. Clustering

- 논문지 Kmeans clustering

Clustering

앞의 12가지 저널에 대해 유사한 저널을 찾아보기 위해 clustering을 수행해본 결과 (k=3일때)

Top terms per cluster:

Cluster 0 words: b'patient', b'p', b'tumor', b'mutation', b'mrna', b'mirnas',

Cluster 0 titles: American Journal of Human Genetics_.txt, Disease Markers_.txt, Genomics _ Informatics_.txt, Nucleic Acids Research_.txt, Oncogenesis_.txt,

Cluster 1 words: b'supplementary', b'supplementary', b'material', b'conclusion', b'background', b'algorithm',

Cluster 1 titles: Algorithms for Molecular Biology _ AMB_.txt, BMC Bioinformatics_.txt, BMC Genomics_.txt, BMC Systems Biology_.txt,

Cluster 2 words: b'microarray', b'review', b'set', b'methylation', b'algorithm', b'array',

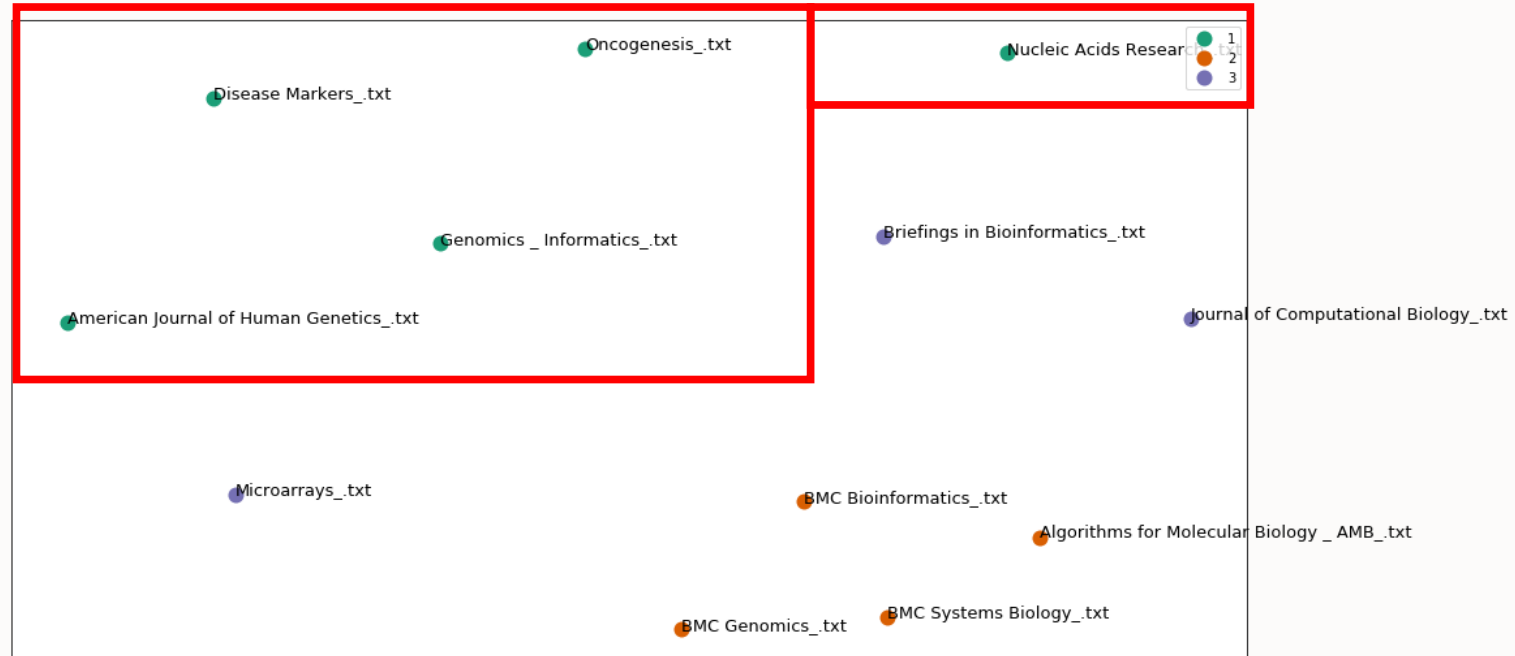
Cluster 2 titles: Briefings in Bioinformatics_.txt, Journal of Computational Biology_.txt, Microarrays_.txt,

총 12가지 저널에 대해 GNI와 유사한 저널을 찾아보기 위하여 tf-idf를 이용하여 Kmeans Clustering을 수행해보았다.
먼저 k=3이었을 때 GNI는 patient, tumor, mutation, mran, miranas 와 같은 단어들로 이루어진 0번 클러스터로 분류가 되었다.

	title	cluster
1	Algorithms for Molecular Biology _ AMB_.txt	1
0	American Journal of Human Genetics_.txt	0
1	BMC Bioinformatics_.txt	1
1	BMC Genomics_.txt	1
1	BMC Systems Biology_.txt	1
2	Briefings in Bioinformatics_.txt	2
0	Disease Markers_.txt	0
0	Genomics _ Informatics_.txt	0
2	Journal of Computational Biology_.txt	2
2	Microarrays_.txt	2
0	Nucleic Acids Research_.txt	0
0	Oncogenesis_.txt	0

Clustering

앞의 12가지 저널에 대해 유사한 저널을 찾아보기 위해 clustering을 수행해본 결과 (k=3일때)



GNI와 같은 클러스터로는 American journal of Human Genetics, Disease Markers, Nucleic Acids Research 였다.
이는 앞에서 주요 키워드로만 유사도를 측정 한 결과와 같았다.

Clustering

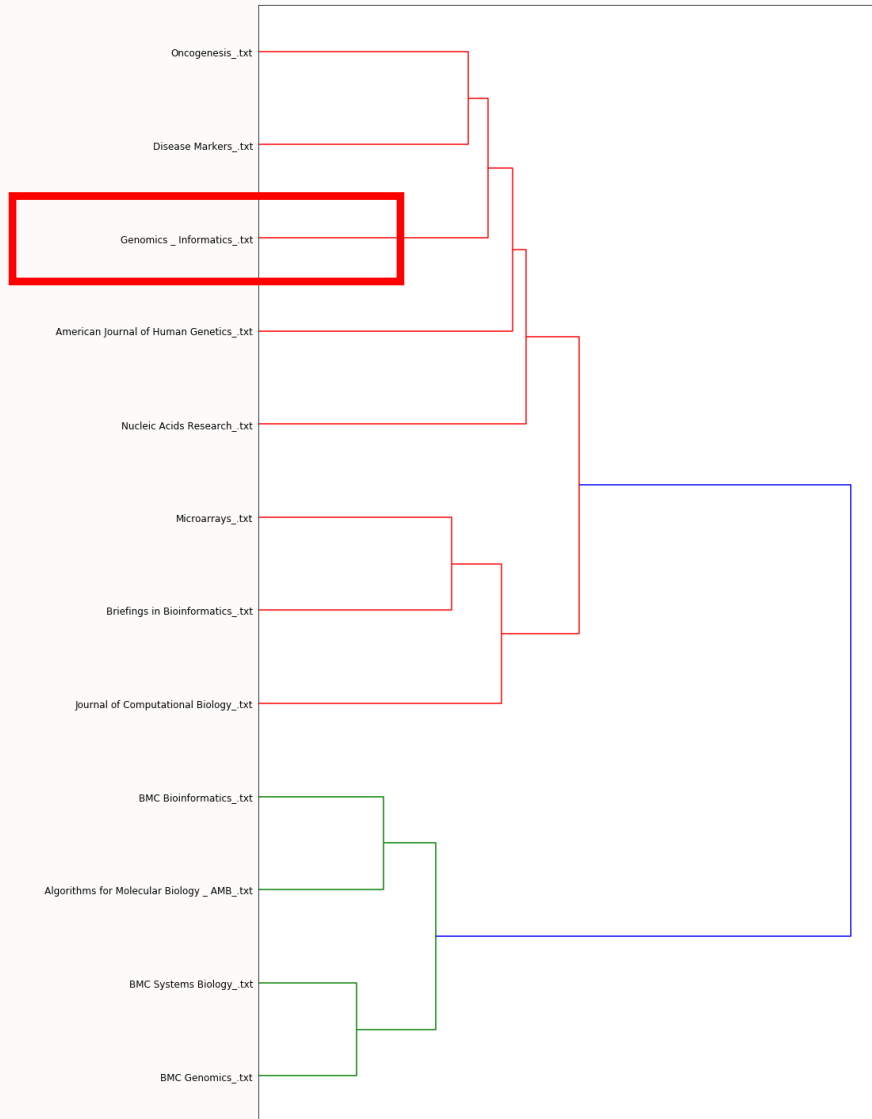
앞의 12가지 저널에 대해 유사한 저널을 찾아보기 위해 clustering을 수행해본 결과 (k=5일때)

	title	cluster
1	Algorithms for Molecular Biology _AMB_.txt	1
3	American Journal of Human Genetics_.txt	3
1	BMC Bioinformatics_.txt	1
1	BMC Genomics_.txt	1
1	BMC Systems Biology_.txt	1
0	Briefings in Bioinformatics_.txt	0
3	Disease Markers_.txt	3
4	Genomics _ Informatics_.txt	4
2	Journal of Computational Biology_.txt	2
0	Microarrays_.txt	0
4	Nucleic Acids Research_.txt	4
4	Oncogenesis_.txt	4



다음으로 k=5이었을 때 GNI는 4번 cluster로, Nucleic Acids Research, Oncogenesis와 같은 클러스터로 분류되었다.

Clustering - Hierarchical document clustering



다음은 Hierarchical document clustering 결과이다.

이전에 계산한 거리인 ward clustering을 사용하여 linkage_marix를 정의하고 dendrogram을 만들어 이를 시각화 하였다.

-> 그 결과 Genomics&Informatics 논문지는 Oncogenesis, Disease Markers, American journal of Human Genetics, Nucleic Acids Research 등과 유사한 것으로 확인됐다.

Clustering Code – (1)

```
In [1]: import numpy as np
import pandas as pd
import nltk
from nltk.corpus import *
from bs4 import BeautifulSoup
import re
import os
import codecs
from sklearn import feature_extraction
import nlp3

os.getcwd()
```

```
Out[1]: 'C:\\Users\\KimMinyoung'
```

```
In [4]: path = 'C:/Temp/2015'
filelist = os.listdir(path)
print(filelist)
stopwords = nltk.corpus.stopwords.words('english')

['Algorithms for Molecular Biology _ AMB_.txt', 'American Journal of Human Genetics_.txt', 'BMC Bioinformatics_.txt', 'BMC Genomics_.txt',
'BMC Systems Biology_.txt', 'Briefings in Bioinformatics_.txt', 'Disease Markers_.txt', 'Genomics _ Informatics_.txt', 'Journal of Computat
ional Biology_.txt', 'Microarrays_.txt', 'Nucleic Acids Research_.txt', 'Oncogenesis_.txt']
```

```
In [5]: pubmed = PlaintextCorpusReader(path, filelist, encoding='utf-8')
```

```
In [7]: raw=[]
for i in range(len(filelist)):
    item = pubmed.raw(pubmed.fileids()[i])
    raw.append(item)
```

```
In [10]: from nltk.stem.snowball import SnowballStemmer
stemmer = SnowballStemmer("english")
```

```
In [12]: def tokenize_and_stem(text):
# first tokenize by sentence, then by word to ensure that punctuation is caught as it's own token
tokens = [word for sent in nltk.sent_tokenize(text) for word in nltk.word_tokenize(sent)]
filtered_tokens = []
# filter out any tokens not containing letters (e.g., numeric tokens, raw punctuation)
for token in tokens:
    if re.search('[a-zA-Z]', token):
        filtered_tokens.append(token)
stems = [stemmer.stem(t) for t in filtered_tokens]
return stems
```

```
def tokenize_only(text):
# first tokenize by sentence, then by word to ensure that punctuation is caught as it's own token
tokens = [word.lower() for sent in nltk.sent_tokenize(text) for word in nltk.word_tokenize(sent)]
filtered_tokens = []
# filter out any tokens not containing letters (e.g., numeric tokens, raw punctuation)
for token in tokens:
    if re.search('[a-zA-Z]', token):
        filtered_tokens.append(token)
return filtered_tokens
```

```
In [13]: totalvocab_stemmed = []
totalvocab_tokenized = []
for i in raw:
    allwords_stemmed = tokenize_and_stem(i)
    totalvocab_stemmed.extend(allwords_stemmed)

    allwords_tokenized = tokenize_only(i)
    totalvocab_tokenized.extend(allwords_tokenized)
```

```
In [16]: vocab_frame = pd.DataFrame({'words': total_vocab_tokenized}, index = total_vocab_stemmed)
vocab_frame
```

	words
background	background
markov	markov
chain	chains
are	are
a	a
...	...
exosom	exosome
research	research
in	in
urotheli	urothelial
cell	cells

49800 rows × 1 columns

```
In [18]: from sklearn.feature_extraction.text import TfidfVectorizer

tfidf_vectorizer = TfidfVectorizer(max_df=0.8, max_features=200000,
                                   min_df=0.2, stop_words='english',
                                   use_idf=True, tokenizer=tokenize_and_stem, ngram_range=(1,3))

%time tfidf_matrix = tfidf_vectorizer.fit_transform(raw)

print(tfidf_matrix.shape)
```

```
In [19]: terms = tfidf_vectorizer.get_feature_names()
         terms
```

'absent',
'absent',
'abund',
'aceler',
'access',
'accompani',
'accord',
'account',
'accumul',
'accur',
'accuraci',
'achiev',
'acid',
'acquir',
'acquisit',
'act',
'action',
'acut',
'acut myeloid',

```
In [20]: from sklearn.metrics.pairwise import cosine_similarity
dist = 1 - cosine_similarity(tfidf_matrix)
```

```
[In [40]: from sklearn.cluster import KMeans

num_clusters = 3

km = KMeans(n_clusters=num_clusters)

time km.fit(tfidf_matrix)

clusters = km.labels_.tolist()
```

Wall time: 301 ms

Clustering Code – (3)

```
In [42]: import pandas as pd

gni = { 'title': filelist, 'text': raw, 'cluster': clusters}

frame = pd.DataFrame(gni, index = [clusters] , columns = ['title', 'cluster'])
frame
```

```
Out[42]:
```

	title	cluster
1	Algorithms for Molecular Biology _AMB_bd	1
0	American Journal of Human Genetics _bd	0
1	BMC Bioinformatics _bd	1
1	BMC Genomics _bd	1
1	BMC Systems Biology _bd	1
2	Briefings in Bioinformatics _bd	2
0	Disease Markers _bd	0
0	Genomics _ Informatics _bd	0
2	Journal of Computational Biology _bd	2
2	Microarrays _bd	2
0	Nucleic Acids Research _bd	0
0	Oncogenesis _bd	0

```
In [43]: from __future__ import print_function

print("Top terms per cluster:")
print()
order_centroids = km.cluster_centers_.argsort()[:, :-1]
for i in range(num_clusters):
    print("Cluster %d words:" % i, end='')
    for ind in order_centroids[i, :6]:
        print(' %s' % vocab_frame.loc[terms[ind].split(' ')].values.tolist()[0][0].encode('utf-8', 'ignore'), end=',')
    print()
    print()
    print("Cluster %d titles:" % i, end='')
    for title in frame.loc[i]['title'].values.tolist():
        print(' %s,' % title, end='')
    print()
    print()
```

Top terms per cluster:

Cluster 0 words: b'patient', b'p', b'tumor', b'mutation', b'mrna', b'mirnas',

Cluster 0 titles: American Journal of Human Genetics .txt, Disease Markers .txt, Genomics _ Informatics .txt, Nucleic Acids Research .txt, Oncogenesis .txt,

```
In [44]: import os # for os.path.basename

import matplotlib.pyplot as plt
import matplotlib as mpl

from sklearn.manifold import MDS

MDS()

# two components as we're plotting points in a two-dimensional plane
# "precomputed" because we provide a distance matrix
# we will also specify 'random_state' so the plot is reproducible.
mds = MDS(n_components=2, dissimilarity="precomputed", random_state=1)

pos = mds.fit_transform(dist) # shape (n_components, n_samples)

xs, ys = pos[:, 0], pos[:, 1]
```

```
In [45]: #strip any proper nouns (NNP) or plural proper nouns (NNPS) from a text
from nltk.tag import pos_tag

def strip_propprs_POS(text):
    tagged = pos_tag(text.split()) #use NLTK's part of speech tagger
    non_proper nouns = [word for word,pos in tagged if pos != 'NNP' and pos != 'NNPS']
    return non_proper nouns
```

```
In [46]: #set up colors per clusters using a dict
cluster_colors = {0: '#1b9e77', 1: '#d95f02', 2: '#7570b3', 3: '#e7298a', 4: '#66a61e', 5: '#00ff00'}

#set up cluster names using a dict
cluster_names = {0: '1',
                  1: '2',
                  2: '3',
                  3: '4',
                  4: '5',
                  5: '6',
                  }
```

Clustering Code – (4)

```
In [48]: df = pd.DataFrame(dict(xs=xs, ys=ys, label=clusters, title=filelist))
```

```
In [52]: #group by cluster
groups = df.groupby('label')

# set up plot
fig, ax = plt.subplots(figsize=(17, 9)) # set size
ax.margins(0.05) # Optional, just adds 5% padding to the autoscaling

#iterate through groups to layer the plot
#note that I use the cluster_name and cluster_color dicts with the 'name' lookup to return the appropriate color/label
for name, group in groups:
    ax.plot(group.x, group.y, marker='o', linestyle='', ms=12, label=cluster_names[name], color=cluster_colors[name], mec='none')
    ax.set_aspect('auto')
    ax.tick_params(
        axis='x',          # changes apply to the x-axis
        which='both',      # both major and minor ticks are affected
        bottom='off',      # ticks along the bottom edge are off
        top='off',         # ticks along the top edge are off
        labelbottom='off')
    ax.tick_params(
        axis='y',          # changes apply to the y-axis
        which='both',      # both major and minor ticks are affected
        left='off',        # ticks along the bottom edge are off
        top='off',         # ticks along the top edge are off
        labelleft='off')

ax.legend(numpoints=1) #show legend with only 1 point

#add label in x,y position with the label as the film title
for i in range(len(df)):
    ax.text(df.loc[i]['x'], df.loc[i]['y'], df.loc[i]['title'], size=13)

plt.show() #show the plot

#uncomment the below to save the plot if need be
#plt.savefig('olusters_small_noaxes.png', dpi=200)
```

```
In [50]: from scipy.cluster.hierarchy import ward, dendrogram

linkage_matrix = ward(dist) #define the linkage_matrix using ward clustering pre-computed distances

fig, ax = plt.subplots(figsize=(15, 20)) # set size
ax = dendrogram(linkage_matrix, orientation="right", labels=filelist);

plt.tick_params(
    axis='x',          # changes apply to the x-axis
    which='both',      # both major and minor ticks are affected
    bottom='off',      # ticks along the bottom edge are off
    top='off',         # ticks along the top edge are off
    labelbottom='off')

plt.tight_layout() #show plot with tight layout

#uncomment below to save figure
plt.savefig('ward_clusters.png', dpi=200) #save figure as ward_clusters
```

