Token
Stem
Lemma
Some stats
No of word count
Some kind insights

#### para="""

Reviewer 2 measured target lesion and made more reliable assessment.

reviewer 2 did not define any target lesions. reviewer 1 did - cervical lymph node left, which responded to therapy, in TP 2 less than 30% (SD) in TP3 more than 30% (PR).

At TP 4 there are new enlarged celiac and retroperitoneal lymph nodes that further enlarge at TP 5 - consistent with PD.

new hepatic lesion can indeed be detected in TP 3 (only detectable in arterial phase - difficult to see)

CR with resolution of visible esophageal lesion

not easy to decide. lesion is gone in ct at TP2 (condition for CR). on barium swallow esophagus is slightly rigid (slightly rigid is allowed for CR), no mucosal defects and barium passes smoothly (also condition for CR). what is missing in the barium swallow report is the ratio of upper esophagus part to narrow part. when measured by oneself the ratio in TP2 and 3 is more than 3:2 - what prevents CR. Hence right decision is NN At TP 6 right para tracheal lymph node that had decreased in size enlarges again, with further growth at TP 7 - consistent with PD.

I agree with reviewer 1 assessment of presence of tumor burden at TP 4 and also new equivocal lesion identified by reviewer1.

Reviewer 2 selection of lesion and measurement appears accurate.

PD not present at TP4...the retrotracheal LN marked as enlarging by reviewer 1 is smaller than BL

There is increased esophageal wall thickening at site of primary esophageal lesion on TP3, c/w PD

rev1

there are multiple new lung lesions at TP 2, therefore PD starting at TP 2 is correct not the same patient

Partial response is the correct evaluation - decreasing size of nodal lesions.

lung lesion measured by reviewer 2 is not necessarily a metastasis, therefore I would prefer review by reviewer 1.

Reviewer 2 measured to large and slightly different position to baseline correct is PD

bone lesion is growing but there is no real soft tissue component visible in the bone lesion.

Rev 2 market a bone lesion without a soft tissue component which should not be done according to study guidelines

PD at TP2 - significant increasing size of lesions

Retroperitoneal LNs are measurable at BL and can be chosen as a TL, therefore I would prefer the review by reviewer 2.

Reviewer measured the iliac vein in addition toi the lesion. The measurement of Reviewer 1 is exact

reviewer 1 is correct, there is a significant increase of the paraaortic LN at TP 4, and there seem to be new LN mets, PD at TP 4 is correct

there is a significant increase of the LNs iliac right, PD at TP 2 is correct more non specific fluid but not sure that pleural effusion is truly tumor related initial 130 mm TP4: 157 cm

there is a significant progression of multiple retroperitoneal LNs, clear PD at TP 2 there are no new liver lesions at TP 2. the hypodense area in liver segment 4 is due to fatty liver tissue, not a real lesion

There is progression as marked

new metastatic LNs since TP 3, review by reviewer 2 is preferable

progressive liver lesions already at tp 3

There is BL disease and the PD

lesion in the left adrenal gland is most likely benign

only non target lesion groups visible

Lesion in liver is calcified and seems NOT to be a NTL, therefore ND at BL and at the following TPs is correct

images are evaluable

it is not for sure that there are really new lesions in the liver at TP 2. At TP there is a good arterial phase which is missing at BL, at BL lesions might just not have been seen due to this missing arterial phase

I would not have marked the prostate but the pleural/soft tissue component of bony met is real

Incredibly subtle findings but present

no lesion out of the prostate

there is a target lesion like reviewer 2 described

PD at TP 2 due to increasing size of lesions

new lung lesions since TP 2, the decision is based on whether these are seen as new lung mets. As they persist at TP 3 it is more likely that they are new mets an not of inflammatory origin.

no clear disease according to RECST 1.1 visible

I don't primarily measure a intraprostatic lesions.

there is disease at Baseline - and a stable disease situation at TP 2.

lesions are progressive

At BL there is no clearly measurable lesion in the prostate, therefore I would prefer the evaluation of reviewer 1.

no clear disease visible

Fluid is no unequivocal lesion / therefore no progression

difference is due to different choice of TLs at BL, I think that the choice of reviewer 2 is more convincing and there is clear progression at TP 2

Has seen Prostate lesion earlier than Reviewer 2

Progression of T and NT

new nodal lesions iliacal left side and soft tissue component of bone lesionis Still PR at TP 3 is correct

Growth of retroperitoneal NTL LNs at TP 2 compared to BL is significant, therefore PD at TP 2 is correct.

there is a pelvic lesion like reviewer 2 stated

PD at TP 4 is correct

new fluid is no unequivocal lesion

Nothing that I would take as target. measured in long axis

PD at TP 2 is the correct evaluation due to increasing liver lesions

no disease. seminal vesicles

increasing size of bladder lesion

After carefully reviewing the imaging material I do not see a significant growthy of the liver lesions at TP 3 and 4. Review by reviewer 2 should be preferred.

nodal lesions are morphological suspect

increasing number and size of liver lesions

I think these are not unequivocal increase. I think there is likely an increase but I would have put NN. This is close and I understand the call.

TL% at tp2 do not warrant PD.

R2 provides appropriate response, unequivocal PD.

new adenopathy as described

TL chosen are more easily reproduced over time

Again TL are slightly better and more reproducible

TL are slightly better than R1

R1 measurements are more similar over time

reliable measures over all tp

No definite peritoneal disease

There is a new lucency in L1 at timepoint two which reviewer 1 marked as a new lesion and called PD. But, I would consider it an equivocal lesion as this could represent a benign compression fracture deformity. No other new lesions were seen. I would agree with partial response.

```
reviewer 2's target lesions are uniformly slightly over measured. Do not agree with
unequiv progression of disease
agree with PD at tp6
agree with pd at tp6
agree with pd at tp6
TL chosen by review 1 are slightly better than TL for reviewer 2.
the TL did increase over time compatible with Pd
TL are increasing overtime
PD at tp4 forward is correct "
the intramuscular metastases noted by reviewer 1 as a new lesion were present at BL
intramuscular metastasis was present at BL
intramuscular mets present at BL
intramuscular mets were present at BL
intramuscular mets present at BL
Agree with assessment of target lung and para-aortic nodal lesions and BOR of SD (based on
more consistent measurements by reader 1).
Agree with assessment of target lung and para-aortic nodal lesions and BOR of SD (based on
more consistent measurements by reader 1).
New lung lesions at TP 5, the bone lesions at TP 4 are equivocal
Sufficient evidence to declare PD at TP4, based on growth in target lesions.
agree with reviewer 1's more robust selection of measurable lesions
more robust selection of target lesions
continue to agree with reader 1 of PR as BOR
new adenopathy at tp2
Unequivocal PD based on progression in several mediastinal nodes.
PD at TP2 and 3 is reasonable as several nodes demonstrate 50+% growth (particularly
paraesophageal and retrocrural nodes).
R2 has more optimal target lesion selection
R2 has more optimal target lesion selection
BL-TP9: R1 has more optimal target lesion measurements; disagree with PD at TP9, as per R2
BL-TP14: R1 has more optimal target lesion measurements; disagree with PD at TP9, as per
R2 has more optimal detection of new lesions
R2 has more optimal detection of new lesions
BL-TP7: R2 has more optimal detection of NLs
BL-TP10: R2 has more optimal detection of NLs
R1 provides appropriate response. No unequivocal evidence of earlier PD.
reviewer 1's measurements are slightly more accurate at baseline
Do not agree with unequiv PD due to tiny nodules at tp9 (may be inflammatory)
overall agree with reviewer 1
overall agree with lesions measurements and analysis of reader 1
"overall agree with reader 1
agree with new adrenal met at tp3/PD
Agree with PD at tp3
R1 bases the decision to steer away from PD because the enlarging nodes which required it
later shrank. Of course there is the possibility that they were reactive (as he/she
claims), but the chest was full of metastatic nodes at baseline, and it's a stretch to
claim that the enlarged mediastinal nodes were due to two separate processes. We've all
seen tumor-bearing nodes (or non-nodal lesions, or new lesions) appear and then shrink
without thinking that the shrinking precluded tumor. They COULD have been reactive, of
course - we can never know for sure - but they were most likely malignant.
Same rationale as that in Reason for Selection for Review Period 1.
Same reasons as for selecting Reviewer 2 at Review Period 1 and Review Period 2.
the RP adenopathy has increased greater than 20% at TP2
Adenopathy has increased from baseline PD is correct BOR
Better assessment of non targets
```

text analysis - Jupyter Notebook Better assessment of non targets Better assessment of non targets Better assessment of non targets the right adrenal lesion is new from baseline. However, the R adrenal was present at TP3 should have been equivocal and then updated to PD at TP3. BOR would be SD. at TP2 I agree with reviewer 1's new lesions at timepoint 2 Reviewer 1's new lesions at timepoint 2 are real. As strange as it may seem to find a new liver lesion at timepoint 2 when all other lesions are shrinking, I think it's unequivocal, and therefore requires PD at timepoint 2 and thereafter. The reviewers differ with regard to when PD might have first occurred, and have revised their initial assessments. But there has been PD, and for overall assessments at global radiology review once PD has been assessed the overall assessments must remain PD. the lesion reviewer 2 marked as new at TP3 was not new and had been present at TP2 PD at TP4 PD at TP4 Impressive. I would not have measured the lesion at first - thinking it was intraprostatic. It is obviously growing. Well done. no convincing TLs, review by reviewer 2 should be preferred significant growth of soft tissue component of bone lesion new intrapulmonary lesions PD at TP3 is correct no clear disease visible pleural effusion and ascites since TP 2, PD is correct its SD in TP 2 and TP 3 - therefore reviewer 2 is more correct Yes, there are new liver mets since TP 2, PD at TP 2 All likely benign Measurements by reviewer 2 seems more precise, should be preferred Clear progression starts at TP 4. very small but still visible soft tissue component of bone lesion - therefore SD is the preferred valuation no clear disease visible leisions and PD is real it is correct that there are new metastatic mesenteric LNs starting at TP 4 stable disease at TP 2 is correct no clear measurable disease at BL agreed with reviewer 1 agreed with reviewer 2 rev2 I Agree With OQREV1 For RP1. I Agree with OQ REV1 for RP2. more accurate TL measurement/selection. Reviewer 2 selected ill-defined peritoneal caking more accurate TL measurement/selection there isn't clearly PD based on increased peritoneal disease at TP11 agree with lesion selection and measurements of reviewer 2 agree with measurements and lesion selection of reviewer 1 in retrospect, agree that their are three lung lesions, instead of 2 lung lesions plus a hilar node. R2 provides appropriate response. R2 provides appropriate response. R2 provides appropriate response. BL-TP17: R2 has more optimal NT assessment there is PD at TP 6 - newly enlarged R hilar LN there is a new lesion in the left lower lobe which looks similar to the other metastatic disease

Agree with consistency of measurements by reviewer 2 for solitary target lesion. SD is

localhost:8888/notebooks/text analysis.ipynb#

most appropriate at TP4.

Increased lung NT Agree with PD at tp5 Agree with consistency of measurements by reviewer 2 for solitary target lesion. SD is most appropriate at TP4.

Although SD is perhaps more appropriate early in the assessment, PR is clearly present from TP5 on and BOR of PR is appropriate from TP4 and beyond.

Although SD is perhaps more appropriate early in the assessment, PR is clearly present from TP5 on and BOR of PR is appropriate from TP4 and beyond. Bone lesion in question by reviewer 2 is overmeasured at later TPs.

insufficient # of new lung nodules to call it unequiv progression agree with PR

PD at TP2 is correct based on new brain mets at TP2 and accompanying new sites of disease at TP3.

differences related to techniques of measurement

Agree with PD identified by Reviewer 2

agree with BOR at tp3

in retrospect, agree with BOR at tp2

PD appropriate at TP3.

Periaortic nodes have been growing ever since baseline. I might not have called them PD until timepoint 3, but it's clear that PD occurred much closer to timepoint 2 than timepoint 7.

Agree with PD at timepoint 2

Agree with PD at tp2

agree with PD at tp7

left adrenal nodule present at baseline. agree with pd at tp7

agree with PD at tp7

both reviews are excellent; R1 has slightly better measurements.

TL measurements are slightly better

More targets selected and therefore more objective

More targets selected and therefore more objective

More targets selected and therefore more objective

the pleural lesion does increase at TP3 from nadir

new bone lesion best seen on bone scan and increase in the pleural lesion from nadir . PD is correct at TP3

given the totality of the case I would change my opinion from R2 to R1. the lesion that were TL are well chosen and show evidence of decrease over time. The new lesions chosen by R2 do appear but could represent fractures.

## In [9]:

import pandas as pd

# In [10]:

```
df=pd.read_csv("xyz2.csv",encoding='latin-1')
df
```

# Out[10]:

# i»¿Reviewer 2 measured target lesion and made more reliable assessment.

0	reviewer 2 did not define any target lesions
1	At TP 4 there are new enlarged celiac and retr
2	new hepatic lesion can indeed be detected in T
3	CR with resolution of visible esophageal lesion
4	not easy to decide. lesion is gone in ct at TP
195	More targets selected and therefore more objec
196	the pleural lesion does increase at TP3 from n
197	new bone lesion best seen on bone scan and inc
198	PD is correct at TP3
199	given the totality of the case I would change

## 200 rows × 1 columns

# In [11]:

```
df.head()
```

# Out[11]:

# $\label{eq:controller} \"{\text{"}}\textit{"}\textit{"}\textit{"}\textit{Reviewer 2 measured target lesion and made more reliable assessment.}$

0	reviewer 2 did not define any target lesions
1	At TP 4 there are new enlarged celiac and retr
2	new hepatic lesion can indeed be detected in T
3	CR with resolution of visible esophageal lesion
4	not easy to decide. lesion is gone in ct at TP

## In [12]:

df.head(7)

## Out[12]:

#### i»¿Reviewer 2 measured target lesion and made more reliable assessment.

0	reviewer 2 did not define any target lesions
1	At TP 4 there are new enlarged celiac and retr
2	new hepatic lesion can indeed be detected in T
3	CR with resolution of visible esophageal lesion
4	not easy to decide. lesion is gone in ct at TP
5	At TP 6 right para tracheal lymph node that ha
6	I agree with reviewer 1 assessment of presence

# In [14]:

df.rename(columns={'i»¿Reviewer 2 measured target lesion and made more reliable assessment.
df.head(5)

# Out[14]:

#### text

- **0** reviewer 2 did not define any target lesions. ...
- 1 At TP 4 there are new enlarged celiac and retr...
- 2 new hepatic lesion can indeed be detected in T...
- 3 CR with resolution of visible esophageal lesion
- 4 not easy to decide. lesion is gone in ct at TP...

## In [15]:

df.shape

## Out[15]:

(200, 1)

```
In [16]:
df.info()
<class 'pandas.core.frame.DataFrame'>
RangeIndex: 200 entries, 0 to 199
Data columns (total 1 columns):
     Column Non-Null Count Dtype
             -----
 0
     text
             200 non-null
                             object
dtypes: object(1)
memory usage: 1.7+ KB
In [17]:
# missing values
df.isnull().sum()
Out[17]:
text
dtype: int64
In [18]:
# check for duplicate values
df.duplicated().sum()
Out[18]:
19
In [19]:
# remove duplicates
df = df.drop_duplicates(keep='first')
In [21]:
df.duplicated().sum()
Out[21]:
0
In [22]:
df.shape
Out[22]:
(181, 1)
```

```
In [23]:
```

```
df['text'].value_counts()
```

## Out[23]:

reviewer 2 did not define any target lesions. reviewer 1 did - cervical lymp h node left, which responded to therapy, in TP 2 less than 30% (SD) in TP3 more than 30% (PR).

1

the RP adenopathy has increased greater than 20% at TP2

1

Better assessment of non targets

1

the right adrenal lesion is new from baseline. However, the R adrenal was p resent at TP3 should have been equivocal and then updated to PD at TP3. BOR would be SD. at TP2

1

I agree with reviewer 1's new lesions at timepoint 2

1

• •

increasing size of bladder lesion

1

After carefully reviewing the imaging material I do not see a significant growthy of the liver lesions at TP 3 and 4. Review by reviewer 2 should be preferred.

1

nodal lesions are morphological suspect

1

increasing number and size of liver lesions

1

given the totality of the case I would change my opinion from R2 to R1. the lesion that were TL are well chosen and show evidence of decrease over time. The new lesions chosen by R2 do appear but could represent fractures.

1 Name: text, Length: 181, dtype: int64

#### In [24]:

# df.describe()

#### Out[24]:

	text
count	181
unique	181
top	reviewer 2 did not define any target lesions
freq	1

```
In [30]:
```

```
df['num_sentences'] = df['text'].apply(lambda x:len(nltk.sent_tokenize(x)))
df['num_characters'] = df['text'].apply(len)
df['num_words'] = df['text'].apply(lambda x:len(nltk.word_tokenize(x)))
```

C:\Users\DELL\AppData\Local\Temp/ipykernel\_23880/2662147901.py:1: SettingWit
hCopyWarning:

A value is trying to be set on a copy of a slice from a DataFrame. Try using .loc[row\_indexer,col\_indexer] = value instead

See the caveats in the documentation: https://pandas.pydata.org/pandas-docs/stable/user\_guide/indexing.html#returning-a-view-versus-a-copy (https://pandas.pydata.org/pandas-docs/stable/user\_guide/indexing.html#returning-a-view-versus-a-copy)

df['num\_sentences'] = df['text'].apply(lambda x:len(nltk.sent\_tokenize
(x)))

C:\Users\DELL\AppData\Local\Temp/ipykernel\_23880/2662147901.py:2: SettingWit
hCopyWarning:

A value is trying to be set on a copy of a slice from a DataFrame. Try using .loc[row\_indexer,col\_indexer] = value instead

See the caveats in the documentation: https://pandas.pydata.org/pandas-docs/stable/user\_guide/indexing.html#returning-a-view-versus-a-copy (https://pandas.pydata.org/pandas-docs/stable/user\_guide/indexing.html#returning-a-view-versus-a-copy)

df['num\_characters'] = df['text'].apply(len)

C:\Users\DELL\AppData\Local\Temp/ipykernel\_23880/2662147901.py:3: SettingWit
hCopyWarning:

A value is trying to be set on a copy of a slice from a DataFrame. Try using .loc[row\_indexer,col\_indexer] = value instead

See the caveats in the documentation: https://pandas.pydata.org/pandas-docs/stable/user\_guide/indexing.html#returning-a-view-versus-a-copy (https://pandas.pydata.org/pandas-docs/stable/user\_guide/indexing.html#returning-a-view-versus-a-copy)

df['num\_words'] = df['text'].apply(lambda x:len(nltk.word\_tokenize(x)))

#### In [31]:

```
df.head()
```

## Out[31]:

	text	num_sentences	num_characters	num_words
0	reviewer 2 did not define any target lesions	2	172	43
1	At TP 4 there are new enlarged celiac and retr	1	120	23
2	new hepatic lesion can indeed be detected in T	1	105	21
3	CR with resolution of visible esophageal lesion	1	47	7
4	not easy to decide. lesion is gone in ct at TP	6	432	91

```
In [32]:
```

```
df[['num_characters','num_words','num_sentences']].describe()
```

## Out[32]:

	num_characters	num_words	num_sentences
count	181.000000	181.000000	181.000000
mean	71.143646	13.723757	1.193370
std	68.565632	13.843608	0.650778
min	4.000000	1.000000	1.000000
25%	31.000000	6.000000	1.000000
50%	51.000000	9.000000	1.000000
75%	86.000000	17.000000	1.000000
max	600.000000	115.000000	6.000000

# In [35]:

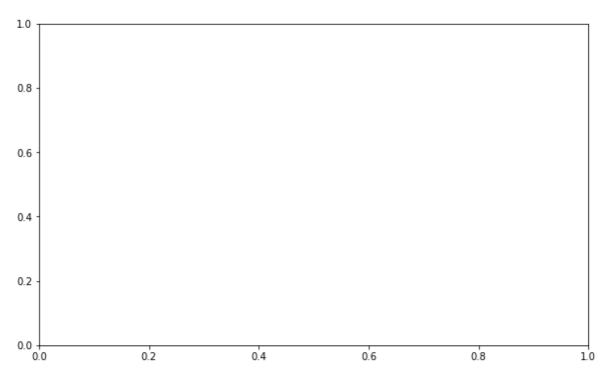
```
import seaborn as sns
import matplotlib.pyplot as plt
```

# In [41]:

```
plt.figure(figsize=(10,6))
sns.histplot(df[df['text'] == 0]['num_characters'])
sns.histplot(df[df['text'] == 1]['num_characters'])
```

## Out[41]:

# <AxesSubplot:>



#### In [42]:

```
c:\Users\DELL\Anaconda3\lib\site-packages\matplotlib\backends\backend_agg.p
y:240: RuntimeWarning: Glyph 13 missing from current font.
  font.set_text(s, 0.0, flags=flags)
C:\Users\DELL\Anaconda3\lib\site-packages\matplotlib\backends\backend_agg.p
y:203: RuntimeWarning: Glyph 13 missing from current font.
  font.set_text(s, 0, flags=flags)
```

#### Out[42]:

<seaborn.axisgrid.PairGrid at 0x1f32b2be3a0>

```
| Part |
```

## In [27]:

import nltk

## In [5]:

```
from nltk.tokenize import sent_tokenize

tokenized_sent=sent_tokenize(para)
print(tokenized_sent)
```

['\nReviewer 2 measured target lesion and made more reliable assessment.', 'reviewer 2 did not define any target lesions.', 'reviewer 1 did - cervica l lymph node left, which responded to therapy, in TP 2 less than 30% (SD) in TP3 more than 30% (PR).', 'At TP 4 there are new enlarged celiac and re troperitoneal lymph nodes that further enlarge at TP 5 - consistent with P D.', 'new hepatic lesion can indeed be detected in TP 3 (only detectable i n arterial phase - difficult to see) \nCR with resolution of visible esoph ageal lesion\nnot easy to decide.', 'lesion is gone in ct at TP2 (conditio n for CR).', 'on barium swallow esophagus is slightly rigid (slightly rigi d is allowed for CR), no mucosal defects and barium passes smoothly (also condition for CR).', 'what is missing in the barium swallow report is the ratio of upper esophagus part to narrow part.', 'when measured by oneself the ratio in TP2 and 3 is more than 3:2 - what prevents CR.', 'Hence right decision is NN \nAt TP 6 right para tracheal lymph node that had decrease d in size enlarges again, with further growth at TP 7 - consistent with P D.', 'I agree with reviewer 1 assessment of presence of tumor burden at TP 4 and also new equivocal lesion identified by reviewer1.', 'Reviewer 2 sel ection of lesion and measurement appears accurate.', 'PD not present at TP 4...the retrotracheal LN marked as enlarging by reviewer 1 is smaller than

#### In [6]:

from nltk.tokenize import word\_tokenize
tokenized\_word=word\_tokenize(para)
print(tokenized\_word)

```
['Reviewer', '2', 'measured', 'target', 'lesion', 'and', 'made', 'more', 'reliable', 'assessment', '.', 'reviewer', '2', 'did', 'not', 'define', 'a ny', 'target', 'lesions', '.', 'reviewer', '1', 'did', '-', 'cervical', 'l ymph', 'node', 'left', ',', 'which', 'responded', 'to', 'therapy', ',', 'i n', 'TP', '2', 'less', 'than', '30', '%', '(', 'SD', ')', 'in', 'TP3', 'mo re', 'than', '30', '%', '(', 'PR', ')', '.', 'At', 'TP', '4', 'there', 'ar e', 'new', 'enlarged', 'celiac', 'and', 'retroperitoneal', 'lymph', 'node s', 'that', 'further', 'enlarge', 'at', 'TP', '5', '-', 'consistent', 'wit h', 'PD', '.', 'new', 'hepatic', 'lesion', 'can', 'indeed', 'be', 'detecte d', 'in', 'TP', '3', '(', 'only', 'detectable', 'in', 'arterial', 'phase', '-', 'difficult', 'to', 'see', ')', 'CR', 'with', 'resolution', 'of', 'vis ible', 'esophageal', 'lesion', 'not', 'easy', 'to', 'decide', '.', 'lesio n', 'is', 'gone', 'in', 'ct', 'at', 'TP2', '(', 'condition', 'for', 'CR', ')', '.', 'on', 'barium', 'swallow', 'esophagus', 'is', 'slightly', 'rigid', 'is', 'allowed', 'for', 'CR', ')', ',', 'n o', 'mucosal', 'defects', 'and', 'barium', 'passes', 'smoothly', '(', 'als o', 'condition', 'for', 'CR', ')', '.', 'what', 'is', 'missing', 'in', 'the', 'barium', 'swallow', 'report', 'is', 'the', 'ratio', 'of', 'upper', 'e sophagus', 'part', 'to', 'narrow', 'part', '.', 'when', 'measured', 'by',
```

## In [7]:

```
#words count
from nltk.probability import FreqDist
fdist=FreqDist(tokenized_word)
print(fdist)
```

<FreqDist with 589 samples and 2629 outcomes>

# In [8]:

```
fdist.most_common(2)
```

#### Out[8]:

```
[('at', 97), ('.', 82)]
```

#### In [9]:

```
from nltk.corpus import stopwords
stop_words=set(stopwords.words("english"))
print(stop_words)
```

{'doesn', 'than', "shan't", 'which', 'couldn', 'yours', 'any', 'shan', 'hi
s', 'with', 'out', "should've", 'after', 'hers', "you've", 'same', 'too', 'w
on', 'below', 'own', 'don', 'into', "don't", 'what', 'been', 'had', 'most',
'not', 'mightn', 'and', 'before', 'me', 'these', 'll', 'm', 'y', 'again', "d
idn't", 'has', 'down', 'themselves', 'yourselves', 'further', "you'd", 'up',
'he', 'itself', 'that', 'be', 'being', 'have', 'i', "aren't", 'to', 'it', "n
eedn't", 'such', 'are', 'whom', 'hadn', 'few', "you're", 'doing', 'needn',
'where', 'by', 'against', "won't", "that'll", "mustn't", 'or', 'as', 've',
'wouldn', "shouldn't", 'ours', 'off', 'over', 'having', 'does', 'through',
'no', 'do', 's', 'if', 'them', 'is', "hasn't", "wouldn't", "doesn't", 'jus
t', 'the', 'during', 'because', 'on', 'now', 'in', 'mustn', 'here', 'how',
'd', 'haven', 'its', 'yourself', 'but', 'some', 'while', 'other', 'did', 'on
ly', 'him', 'once', 'this', 're', 'will', 'ourselves', "she's", 'they', 'wh
o', 'herself', 'didn', "isn't", "wasn't", 'why', 'at', 'a', 'hasn', 'until',
'haven't", 'ma', 'an', 'were', 'when', "it's", 'should', "you'll", 'so', 'sh
ouldn', 'their', 'there', 'o', 'those', 'all', 't', 'aren', "mightn't", 'was
n', "weren't", 'ain', 'under', "hadn't", 'of', 'you', 'we', 'each', 'our',
'myself', 'isn', 'more', 'theirs', 'weren', 'her', "couldn't", 'then', 'ca
n', 'himself', 'above', 'about', 'your', 'she', 'very', 'was', 'between', 'b
oth', 'nor', 'my', 'from', 'for', 'am'}

```
In [10]:
```

```
filtered_sent=[]
for w in tokenized_word:
    if w not in stop_words:
        filtered_sent.append(w)
print("ts:",tokenized_word)
print("fs:",filtered_sent)
```

```
ts: ['Reviewer', '2', 'measured', 'target', 'lesion', 'and', 'made', 'mor e', 'reliable', 'assessment', '.', 'reviewer', '2', 'did', 'not', 'defin e', 'any', 'target', 'lesions', '.', 'reviewer', '1', 'did', '-', 'cervica l', 'lymph', 'node', 'left', ',', 'which', 'responded', 'to', 'therapy', ',', 'in', 'TP', '2', 'less', 'than', '30', '%', '(', 'SD', ')', 'in', 'TP 3', 'more', 'than', '30', '%', '(', 'PR', ')', '.', 'At', 'TP', '4', 'ther e', 'are', 'new', 'enlarged', 'celiac', 'and', 'retroperitoneal', 'lymph', 'nodes', 'that', 'further', 'enlarge', 'at', 'TP', '5', '-', 'consistent', 'with', 'PD', '.', 'new', 'hepatic', 'lesion', 'can', 'indeed', 'be', 'det ected', 'in', 'TP', '3', '(', 'only', 'detectable', 'in', 'arterial', 'pha se', '-', 'difficult', 'to', 'see', ')', 'CR', 'with', 'resolution', 'of', 'visible', 'esophageal', 'lesion', 'not', 'easy', 'to', 'decide', '.', 'le sion', 'is', 'gone', 'in', 'ct', 'at', 'TP2', '(', 'condition', 'for', 'C R', ')', '.', 'on', 'barium', 'swallow', 'esophagus', 'is', 'slightly', 'rigid', '(', 'slightly', 'rigid', 'is', 'allowed', 'for', 'CR', ')', ',', 'no', 'mucosal', 'defects', 'and', 'barium', 'passes', 'smoothly', '(', 'a lso', 'condition', 'for', 'CR', ')', '.', 'what', 'is', 'missing', 'in', 'the', 'barium', 'swallow', 'report', 'is', 'the', 'ratio', 'of', 'upper', 'esophagus', 'part', 'to', 'narrow', 'part', '.', 'when', 'measured', 'b
```

#### In [1]:

```
from nltk.tokenize import sent_tokenize, word_tokenize
ps=PorterStemmer()
stemmed_words=[]
for w in filtered_sent:
    stemmed_words.append(ps.stem(para))
print(filtered_sent)
```

```
In [2]:
```

ls=WordNetLemmatizer()

from nltk.stem.wordnet import WordNetLemmatizer

```
#from nltk.stem import PorterStemmer
#stem =PorterStemmer()
word="flying"
print("l_w;",ls.lemmatize(word,"g"))
LookupError
                                         Traceback (most recent call last)
~\Anaconda3\lib\site-packages\nltk\corpus\util.py in load(self)
     83
                       try:
---> 84
                           root = nltk.data.find(f"{self.subdir}/{zip_nam
e}")
     85
                       except LookupError:
~\Anaconda3\lib\site-packages\nltk\data.py in find(resource_name, paths)
           resource_not_found = f"\n{sep}\n{msg}\n{sep}\n"
--> 583
            raise LookupError(resource_not_found)
    584
LookupError:
*************************
  Resource wordnet not found.
  Please use the NLTK Downloader to obtain the resource:
  >>> import nltk
  >>> nltk.download('wordnet')
  For more information see: https://www.nltk.org/data.html (https://www.nlt
k.org/data.html)
  Attempted to load corpora/wordnet.zip/wordnet/
  Searched in:
    - 'C:\\Users\\DELL/nltk_data'
    - 'C:\\Users\\DELL\\Anaconda3\\nltk data'
    - 'C:\\Users\\DELL\\Anaconda3\\share\\nltk_data'
    - 'C:\\Users\\DELL\\Anaconda3\\lib\\nltk data'
    - 'C:\\Users\\DELL\\AppData\\Roaming\\nltk_data'
    - 'C:\\nltk_data'
    - 'D:\\nltk_data'
    - 'E:\\nltk_data'
*******************************
During handling of the above exception, another exception occurred:
LookupError
                                         Traceback (most recent call last)
~\AppData\Local\Temp/ipykernel_20160/2970926820.py in <module>
     4 #stem =PorterStemmer()
      5 word="flying"
----> 6 print("1 w;",ls.lemmatize(word, "g"))
~\Anaconda3\lib\site-packages\nltk\stem\wordnet.py in lemmatize(self, word,
 pos)
     43
               :return: The lemma of `word`, for the given `pos`.
     44
```

#### In [13]:

```
from nltk.tokenize import sent_tokenize, word_tokenize
from nltk.stem.wordnet import WordNetLemmatizer
ls=WordNetLemmatizer()
lemm_words=[]
for w in filtered_sent:
    lemm_words.append(ls.lemmatize(para))
print(filtered_sent)
```

```
LookupError
                                       Traceback (most recent call las
t)
~\Anaconda3\lib\site-packages\nltk\corpus\util.py in __load(self)
    83
                      try:
---> 84
                          root = nltk.data.find(f"{self.subdir}/{zip_nam
e}")
    85
                      except LookupError:
~\Anaconda3\lib\site-packages\nltk\data.py in find(resource_name, paths)
           resource_not_found = f"\n{sep}\n{msg}\n{sep}\n"
--> 583
           raise LookupError(resource_not_found)
   584
LookupError:
************************
 Resource wordnet not found.
 Please use the NLTK Downloader to obtain the resource:
```

#### In [14]:

```
pip install wordnet
```

Requirement already satisfied: wordnet in c:\users\dell\anaconda3\lib\site-p ackages (0.0.1b2)Note: you may need to restart the kernel to use updated packages.

Requirement already satisfied: colorama==0.3.9 in c:\users\dell\anaconda3\lib\site-packages (from wordnet) (0.3.9)

## In [15]:

```
from nltk.tokenize import sent_tokenize, word_tokenize
from nltk.stem.wordnet import WordNetLemmatizer
ls=WordNetLemmatizer()
lemm_words=[]
for w in filtered_sent:
   lemm_words.append(ls.lemmatize(para))
print(filtered_sent)
LookupError
                                        Traceback (most recent call las
t)
~\Anaconda3\lib\site-packages\nltk\corpus\util.py in __load(self)
                       try:
---> 84
                           root = nltk.data.find(f"{self.subdir}/{zip_nam
e}")
    85
                       except LookupError:
~\Anaconda3\lib\site-packages\nltk\data.py in find(resource_name, paths)
           resource_not_found = f"\n{sep}\n{msg}\n{sep}\n"
    582
--> 583
           raise LookupError(resource_not_found)
    584
LookupError:
*************************
  Resource wordnet not found.
  Please use the NLTK Downloader to obtain the resource:
In [ ]:
```