Project_IAU

November 5, 2025

1 Import libraries

[750]:
```
import numpy as np
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt
import warnings
import scipy.stats as stats
from scipy.stats import shapiro, ttest_ind, mannwhitneyu, levene, pearsonr
from statsmodels.stats.power import TTestIndPower
from sklearn.model_selection import train_test_split
from pandas.api.types import CategoricalDtype
from sklearn.preprocessing import StandardScaler , RobustScaler ,␣
 ↪PowerTransformer, QuantileTransformer

#preprocesing
from sklearn.feature_selection import mutual_info_classif, chi2
from sklearn.ensemble import RandomForestClassifier
from sklearn.preprocessing import MinMaxScaler
```

[751]:
```
#warnings fix
warnings.filterwarnings("ignore", category=UserWarning, module="IPython")
plt.rcParams['font.family'] = 'DejaVu Sans'  # Change the font globally
plt.tight_layout()  # Ensure layout adjustments
```

```
<Figure size 640x480 with 0 Axes>
```

2 Load datasets

[752]:
```
patient_df = pd.read_csv('132/patient.csv', sep = '\t')
station_df = pd.read_csv('132/station.csv', sep= '\t')
observation_df = pd.read_csv('132/observation.csv', sep = '\t')
```

[753]:
```
patient_df.head()
```

[753]:
```
                        company  \
0
```

```
1  Gray, Cunningham and Morales
2                        Walter
3
4                Munari s.r.l.

                                  current_location              ssn  \
0      (Decimal('32.168477'), Decimal('9.804478'))      302-73-9054
1     (Decimal('-80.289857'), Decimal('2.308813'))      499-92-6793
2   (Decimal('63.5169555'), Decimal('-48.876252'))      925-81-9055
3   (Decimal('-71.015803'), Decimal('140.978474'))      175-19-6965
4  (Decimal('-58.1525245'), Decimal('-120.099037'))  FRSMRL26C50L167Z

               name                   mail  user_id  residence  \
0                    takuma47@gmail.com     1770        NaN
1  Nicholas Campbell    curtis06@yahoo.com      946        NaN
2       Raissa Rose  junkgisbert@yahoo.de     2010        NaN
3                  morimomoko@gmail.com     1100        NaN
4  Eraldo Anguillara   bpergolesi@poste.it     1247        NaN

            username   birthdate  \
0           snakajima  1937-03-07
1          benjamin02  1993-05-15
2            jkoehler         NaN
3       yamaguchikana  2000-08-20
4  guglielmomicheletti         NaN

                                       address          registration  \
0                                  27 2 7            2020-08-15
1  0515 Angela Run\r\nPort Thomasberg, GU 35535            2018-11-30
2     Louise-Stey-Platz 79\r\n88420 Bremervörde  05/09/2025, 00:00:00
3                       26 5 4     985  10/21/2022, 00:00:00
4                                        NaN            2024-11-26

  blood_group  station_ID
0          B-         464
1          A-         109
2          0-         462
3          B-         191
4          B+         588
```

[754]:
```
station_df.head()
```

[754]:
```
           QoS code  latitude              revision  longitude      station
0         good   JP  36.00000           2020-11-25  139.55722      Okegawa
1    excellent   IN  11.93381  05/24/2021, 00:00:00   79.82979   Puducherry
2  maintenance   DE  52.21099           2022-05-10    7.02238       Gronau
3    excellent   CN  41.09822           2018-01-23  120.74792      Nanpiao
```

```
4         good   US  33.54428           08 Jun 2024  -84.23381  Stockbridge
```

[755]:
```
observation_df.head()
```

[755]:
```
        SpO          HR         PI         RR      EtCO        FiO   \
0  97.538229  87.194745  11.225419  14.812012  42.113735  33.852538
1  97.933271  80.787303  11.730935  14.964972  39.537692  65.326035
2  98.209983  79.733895  12.839449  14.840668  39.758706  53.925230
3  98.202790  86.156903  11.204152  14.523288  43.448577  35.227704
4  97.951933  78.966258  13.691758  14.992054  39.722280  35.005559

          PRV          BP  Skin Temperature  Motion/Activity index  …  \
0  144.504405  100.455727         35.961920              10.302567  …
1  110.615787  102.133386         36.274352               8.975704  …
2  107.208040  104.036654         35.583851               7.653790  …
3  143.282224  105.723603         36.463180               8.795732  …
4  118.524021   98.996494         35.080937               8.388887  …

         CO  Blood Flow Index  PPG waveform features  Signal Quality Index  \
0  4.022852         58.317397              49.143701             42.399816
1  4.002043         70.127865              15.557799             46.078137
2  4.001451         76.139163              53.879956             41.525607
3  4.015162         49.461570              58.701159             36.535021
4  4.001110         47.065823              52.338305             29.506008

   Respiratory effort  O  extraction ratio        SNR  oximetry   latitude  \
0           46.497869             0.289012  39.334620       1.0  49.183239
1           53.351208             0.290879  26.006709       0.0  33.544280
2           52.124182             0.263171  31.890829       1.0 -27.505780
3           50.342830             0.256780  30.721375       1.0  37.656390
4           39.480811             0.276094  38.214856       0.0  51.202190

    longitude
0   15.454273
1  -84.233810
2  153.102360
3  126.835000
4    7.360270

[5 rows x 23 columns]
```

lets have a look at the attributes

[756]:
```
observation_df.columns
```

[756]:
```
Index(['SpO ', 'HR', 'PI', 'RR', 'EtCO ', 'FiO ', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
```

```
       'Signal Quality Index', 'Respiratory effort', 'O  extraction ratio',
       'SNR', 'oximetry', 'latitude', 'longitude'],
      dtype='object')
```

[757]:
```
station_df.columns
```

[757]:
```
Index(['QoS', 'code', 'latitude', 'revision', 'longitude', 'station'],
dtype='object')
```

[758]:
```
patient_df.columns
```

[758]:
```
Index(['company', 'current_location', 'ssn', 'name', 'mail', 'user_id',
       'residence', 'username', 'birthdate', 'address', 'registration',
       'blood_group', 'station_ID'],
      dtype='object')
```

[759]:
```
for column in observation_df.columns:
    if column in station_df.columns:
        print ('observation_df and station_df share column :' + str(column) )
    if column in patient_df.columns:
        print ('observation_df and patient_df share column :' + str(column) )

for column in patient_df.columns:
    if column in station_df.columns:
        print ('station_df and patient_df share column :' + str(column) )
```

```
observation_df and station_df share column :latitude
observation_df and station_df share column :longitude
```

we can see that observation and station dataframes might be joinable via coordinates but after manual revision we can see that patient.station_ID will probably map to station ids

before joining the datasets, lets perform some basic EDA

3 Fáza 1

3.1 1.1 Základný opis dát spolu s ich charakteristikami

3.1.1 A) Analyza struktur dat ako subory + zaznamy

station_df

[760]:
```
station_df.head()
```

[760]:

	QoS	code	latitude	revision	longitude	station
0	good	JP	36.00000	2020-11-25	139.55722	Okegawa
1	excellent	IN	11.93381	05/24/2021, 00:00:00	79.82979	Puducherry
2	maintenance	DE	52.21099	2022-05-10	7.02238	Gronau
3	excellent	CN	41.09822	2018-01-23	120.74792	Nanpiao
4	good	US	33.54428	08 Jun 2024	-84.23381	Stockbridge

[761]:
```
station_df.shape
```

[761]:
```
(703, 6)
```

[762]:
```
station_df.columns
```

[762]:
```
Index(['QoS', 'code', 'latitude', 'revision', 'longitude', 'station'],
dtype='object')
```

[763]:
```
station_df.dtypes
```

[763]:
```
QoS           object
code          object
latitude     float64
revision      object
longitude    float64
station       object
dtype: object
```

[764]:
```
#2 nas, not bad
station_df.isna().sum()
```

[764]:
```
QoS          0
code         2
latitude     0
revision     0
longitude    0
station      0
dtype: int64
```

[765]:
```
station_df[station_df['code'].isna()]
```

[765]:

	QoS	code	latitude	revision	longitude	station
274	maintenance	NaN	-21.98333	02/23/2016, 00:00:00	16.91667	Okahandja
318	good	NaN	-21.98333	06 Dec 2021	16.91667	Okahandja

[766]:
```
#mby there is valid code for Okahandja?
station_df[station_df['station'] == 'Okahandja']
#there is not :(
```

[766]:

	QoS	code	latitude	revision	longitude	station
274	maintenance	NaN	-21.98333	02/23/2016, 00:00:00	16.91667	Okahandja
318	good	NaN	-21.98333	06 Dec 2021	16.91667	Okahandja

[767]:
```
station_df.nunique()
```

[767]:
```
QoS           4
code         98
```

```
latitude     498
revision     683
longitude    497
station      498
dtype: int64
```

[768]:
```
#prob useless
station_df.describe()
```

[768]:
```
          latitude   longitude
count  703.000000  703.000000
mean    28.699220   16.946846
std     24.406067   70.122555
min    -44.396720 -156.474320
25%     14.354040  -14.410810
50%     36.650000   13.321270
75%     47.432685   71.552920
max     65.848110  171.253640
```

patient df

[769]:
```
patient_df.head()
```

[769]:
```
                        company  \
0
1  Gray, Cunningham and Morales
2                        Walter
3
4                Munari s.r.l.

                                  current_location              ssn  \
0       (Decimal('32.168477'), Decimal('9.804478'))      302-73-9054
1      (Decimal('-80.289857'), Decimal('2.308813'))      499-92-6793
2    (Decimal('63.5169555'), Decimal('-48.876252'))      925-81-9055
3    (Decimal('-71.015803'), Decimal('140.978474'))      175-19-6965
4  (Decimal('-58.1525245'), Decimal('-120.099037'))  FRSMRL26C50L167Z

               name                   mail  user_id  residence  \
0                    takuma47@gmail.com     1770        NaN
1  Nicholas Campbell    curtis06@yahoo.com      946        NaN
2       Raissa Rose  junkgisbert@yahoo.de     2010        NaN
3                   morimomoko@gmail.com     1100        NaN
4  Eraldo Anguillara   bpergolesi@poste.it     1247        NaN

             username   birthdate  \
0           snakajima  1937-03-07
1          benjamin02  1993-05-15
2            jkoehler         NaN
```

```
3        yamaguchikana  2000-08-20
4  guglielmomicheletti         NaN

                                      address          registration  \
0                                27 2 7            2020-08-15
1  0515 Angela Run\r\nPort Thomasberg, GU 35535           2018-11-30
2     Louise-Stey-Platz 79\r\n88420 Bremervörde  05/09/2025, 00:00:00
3                      26 5 4     985  10/21/2022, 00:00:00
4                                         NaN           2024-11-26

  blood_group  station_ID
0          B-         464
1          A-         109
2          O-         462
3          B-         191
4          B+         588
```

[770]:
```
patient_df.shape
```

[770]:
```
(2197, 13)
```

[771]:
```
patient_df.columns
```

[771]:
```
Index(['company', 'current_location', 'ssn', 'name', 'mail', 'user_id',
       'residence', 'username', 'birthdate', 'address', 'registration',
       'blood_group', 'station_ID'],
      dtype='object')
```

[772]:
```
patient_df.dtypes
```

[772]:
```
company              object
current_location     object
ssn                  object
name                 object
mail                 object
user_id               int64
residence           float64
username             object
birthdate            object
address              object
registration         object
blood_group          object
station_ID            int64
dtype: object
```

[773]:
```
patient_df.isna().sum()
#residence is full NaN so it will be dropped, birthdate and address probably␣
 ↪too,
```

```
#current location needs to be processed
```

[773]:
```
company               0
current_location    110
ssn                   0
name                  0
mail                  0
user_id               0
residence          2197
username              0
birthdate           989
address             330
registration          0
blood_group           0
station_ID            0
dtype: int64
```

[774]:
```
patient_df.nunique()
#many unique values
```

[774]:
```
company             1989
current_location    2087
ssn                 2197
name                2131
mail                2192
user_id             1407
residence              0
username            2175
birthdate           1189
address             1867
registration        1987
blood_group            8
station_ID           677
dtype: int64
```

[775]:
```
patient_df.describe()
#also useless
```

[775]:

	user_id	residence	station_ID
count	2197.000000	0.0	2197.000000
mean	1105.308603	NaN	348.892126
std	644.728055	NaN	205.367454
min	0.000000	NaN	0.000000
25%	547.000000	NaN	172.000000
50%	1118.000000	NaN	340.000000
75%	1668.000000	NaN	536.000000
max	2196.000000	NaN	702.000000

observation df

[776]:
```
observation_df.head()
```

[776]:
```
        SpO          HR         PI         RR      EtCO        FiO  \
0  97.538229  87.194745  11.225419  14.812012  42.113735  33.852538
1  97.933271  80.787303  11.730935  14.964972  39.537692  65.326035
2  98.209983  79.733895  12.839449  14.840668  39.758706  53.925230
3  98.202790  86.156903  11.204152  14.523288  43.448577  35.227704
4  97.951933  78.966258  13.691758  14.992054  39.722280  35.005559

          PRV          BP  Skin Temperature  Motion/Activity index  …  \
0  144.504405  100.455727         35.961920              10.302567  …
1  110.615787  102.133386         36.274352               8.975704  …
2  107.208040  104.036654         35.583851               7.653790  …
3  143.282224  105.723603         36.463180               8.795732  …
4  118.524021   98.996494         35.080937               8.388887  …

         CO  Blood Flow Index  PPG waveform features  Signal Quality Index  \
0  4.022852          58.317397              49.143701             42.399816
1  4.002043          70.127865              15.557799             46.078137
2  4.001451          76.139163              53.879956             41.525607
3  4.015162          49.461570              58.701159             36.535021
4  4.001110          47.065823              52.338305             29.506008

   Respiratory effort  O  extraction ratio        SNR  oximetry   latitude  \
0           46.497869             0.289012  39.334620       1.0  49.183239
1           53.351208             0.290879  26.006709       0.0  33.544280
2           52.124182             0.263171  31.890829       1.0 -27.505780
3           50.342830             0.256780  30.721375       1.0  37.656390
4           39.480811             0.276094  38.214856       0.0  51.202190

    longitude
0   15.454273
1  -84.233810
2  153.102360
3  126.835000
4    7.360270

[5 rows x 23 columns]
```

[777]:
```
observation_df.shape
```

[777]:
```
(12177, 23)
```

[778]:
```
observation_df.columns
```

[778]:
```
Index(['SpO ', 'HR', 'PI', 'RR', 'EtCO ', 'FiO ', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
       'Signal Quality Index', 'Respiratory effort', 'O  extraction ratio',
       'SNR', 'oximetry', 'latitude', 'longitude'],
      dtype='object')
```

[779]:
```
observation_df.dtypes
```

[779]:
```
SpO                       float64
HR                        float64
PI                        float64
RR                        float64
EtCO                      float64
FiO                       float64
PRV                       float64
BP                        float64
Skin Temperature          float64
Motion/Activity index     float64
PVI                       float64
Hb level                  float64
SV                        float64
CO                        float64
Blood Flow Index          float64
PPG waveform features     float64
Signal Quality Index      float64
Respiratory effort        float64
O  extraction ratio       float64
SNR                       float64
oximetry                  float64
latitude                  float64
longitude                 float64
dtype: object
```

[780]:
```
observation_df.isna().sum()
#great
```

[780]:
```
SpO                       0
HR                        0
PI                        0
RR                        0
EtCO                      0
FiO                       0
PRV                       0
BP                        0
Skin Temperature          0
Motion/Activity index     0
```

```
PVI                       0
Hb level                  0
SV                        0
CO                        0
Blood Flow Index          0
PPG waveform features     0
Signal Quality Index      0
Respiratory effort        0
O  extraction ratio       0
SNR                       0
oximetry                  0
latitude                  0
longitude                 0
dtype: int64
```

[781]:
```
observation_df.nunique()
#makes sense, only oximetry, latitude and longitude are not completely unique
```

[781]:
```
SpO                      11997
HR                       11997
PI                       11997
RR                       11997
EtCO                     11997
FiO                      11997
PRV                      11997
BP                       11997
Skin Temperature         11997
Motion/Activity index    11997
PVI                      11997
Hb level                 11997
SV                       11997
CO                       11997
Blood Flow Index         11997
PPG waveform features    11997
Signal Quality Index     11997
Respiratory effort       11997
O  extraction ratio      11997
SNR                      11997
oximetry                     2
latitude                   498
longitude                  497
dtype: int64
```

[782]:
```
observation_df.describe()
```

[782]:
```
                SpO            HR            PI            RR         EtCO   \
count  12177.000000  12177.000000  12177.000000  12177.000000  12177.000000
```

mean	97.336001	83.397242	10.360642	16.158948	40.235152
std	0.657577	7.609815	2.417855	1.398210	1.715679
min	95.000000	60.000000	0.200000	12.000000	35.000000
25%	96.895657	77.573676	8.800161	15.044708	38.749846
50%	97.332851	84.323757	10.365288	15.979021	40.531056
75%	97.776356	89.763115	11.893853	17.393451	41.590048
max	100.000000	100.000000	20.000000	20.000000	45.000000

	FiO	PRV	BP	Skin Temperature	\
count	12177.000000	12177.000000	12177.000000	12177.000000	
mean	58.821759	117.675964	104.591413	35.426048	
std	12.119443	21.841513	4.088282	0.619283	
min	21.000000	20.000000	90.000000	33.000000	
25%	49.328258	103.101631	101.857377	35.007462	
50%	59.402365	117.695370	104.604980	35.424762	
75%	68.437184	132.088845	107.321633	35.845164	
max	100.000000	200.000000	120.000000	38.000000	

	Motion/Activity index	…	CO	Blood Flow Index	\
count	12177.000000	…	12177.000000	12177.000000	
mean	9.436818	…	4.078126	52.640590	
std	0.998907	…	0.222547	13.127091	
min	5.652322	…	4.000000	0.000000	
25%	8.766904	…	4.000762	43.996467	
50%	9.432476	…	4.007395	52.709829	
75%	10.105665	…	4.064212	61.440013	
max	13.997052	…	8.000000	100.000000	

	PPG waveform features	Signal Quality Index	Respiratory effort	\
count	12177.000000	12177.000000	12177.000000	
mean	46.734247	47.572094	49.788653	
std	13.374194	13.487056	13.006681	
min	0.000000	0.000000	0.000000	
25%	37.650597	38.396055	40.992562	
50%	46.733166	47.810036	49.742898	
75%	55.671672	56.736043	58.630016	
max	100.000000	100.000000	100.000000	

	O extraction ratio	SNR	oximetry	latitude	\
count	12177.000000	12177.000000	12177.000000	12177.000000	
mean	0.249557	29.994576	0.597602	28.701943	
std	0.028947	5.765251	0.490401	24.402668	
min	0.200000	20.000000	0.000000	-44.396720	
25%	0.224466	24.978691	0.000000	14.082300	
50%	0.249072	30.094799	1.000000	36.650000	
75%	0.274733	34.961220	1.000000	47.484440	
max	0.300000	40.000000	1.000000	65.848110	

```
          longitude
count  12177.000000
mean      17.028640
std       70.059129
min     -156.474320
25%      -13.235600
50%       13.321270
75%       71.577370
max      171.253640

[8 rows x 23 columns]
```

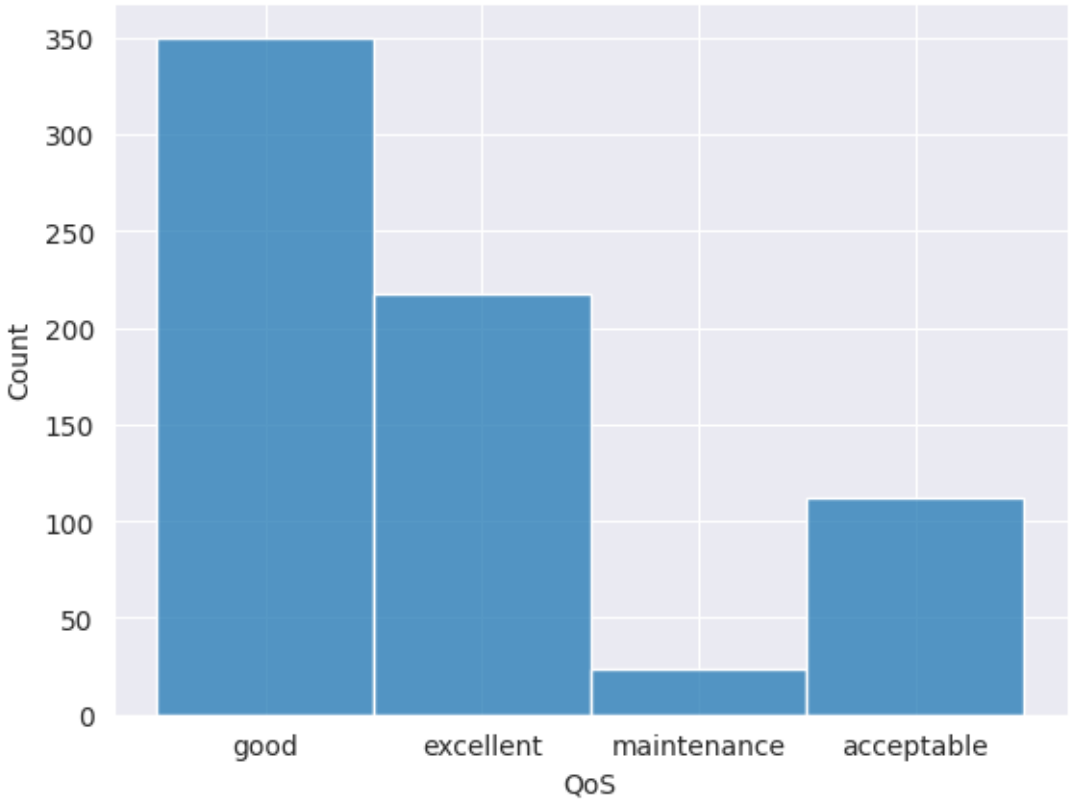
3.1.2 B) Analýza jednotlivých atribútov

[783]:
```
#Qos(station_df)
station_df['QoS'].unique().tolist()
```

[783]:
```
['good', 'excellent', 'maintenance', 'acceptable']
```

[784]:
```
sns.histplot(station_df['QoS'])
```

[784]:
```
<Axes: xlabel='QoS', ylabel='Count'>
```



[785]:
```
#for exact nums
station_df['QoS'].value_counts()
```

[785]:
```
QoS
good           350
excellent      217
acceptable     112
maintenance     24
Name: count, dtype: int64
```

[786]:
```
station_df['QoS'].isna().sum()
```

[786]:
```
np.int64(0)
```

[787]:
```
#latitude (from both station_df and observation_df)
print (station_df['latitude'].min(), observation_df['latitude'].min())
print (station_df['latitude'].max(), observation_df['latitude'].max())
#the min and the max are both realistic values, from -180 to 180

set(station_df['latitude']) == set(observation_df['latitude'])
#every station is included in observation by latitude
```

```
-44.39672 -44.39672
65.84811 65.84811
```

[787]:
```
True
```

[788]:
```
#longitude (from both station_df and observation_df)
print (station_df['longitude'].min(), observation_df['longitude'].min())
print (station_df['longitude'].max(), observation_df['longitude'].max())
#the min and the max are both realistic values, from -180 to 180

set(station_df['longitude']) == set(observation_df['longitude'])
#every station is included by longitude also

#this could be checked with patient -> current location but that needs␣
 ↪preprocessing in further steps
```

```
-156.47432 -156.47432
171.25364 171.25364
```

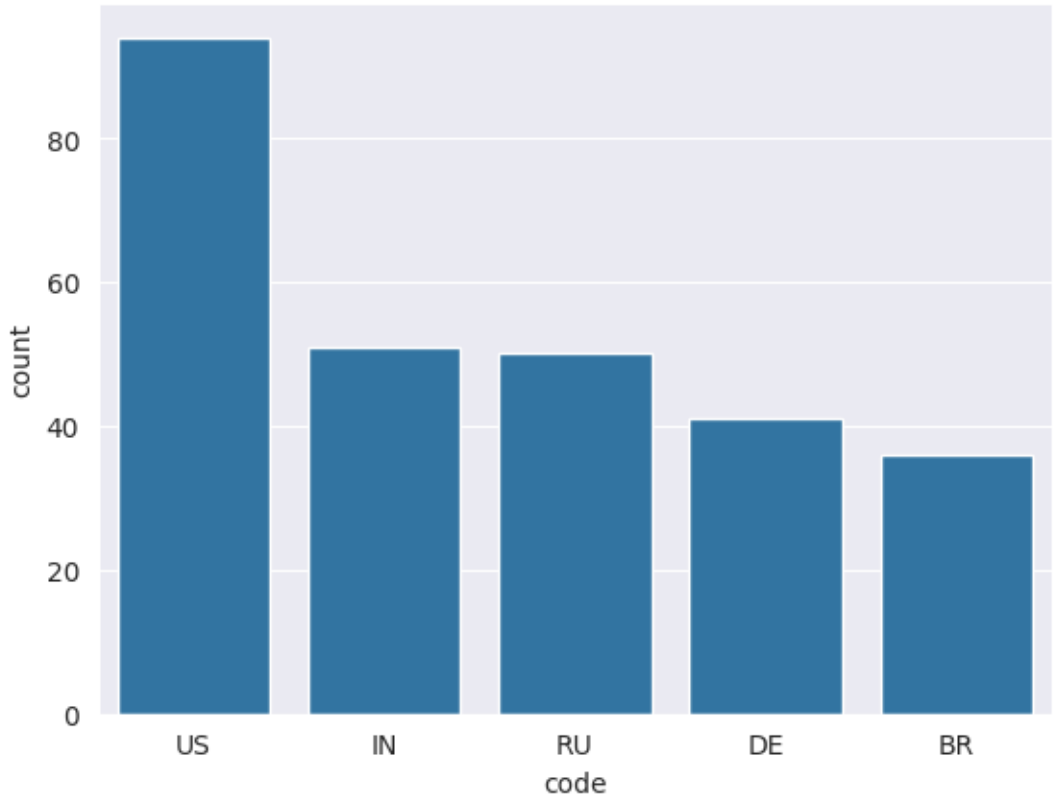
[788]:
```
True
```

[789]:
```
#Code (station_df)
station_df['code'].value_counts()
```

[789]:
```
code
US    94
IN    51
RU    50
DE    41
BR    36
      ..
DK     1
CU     1
GH     1
AF     1
AT     1
Name: count, Length: 98, dtype: int64
```

[790]:
```
#top 5
top5 = station_df['code'].value_counts().head(5).reset_index()
top5.columns = ['code', 'count']

sns.barplot(x='code', y='count', data=top5)
```

[790]:
```
<Axes: xlabel='code', ylabel='count'>
```

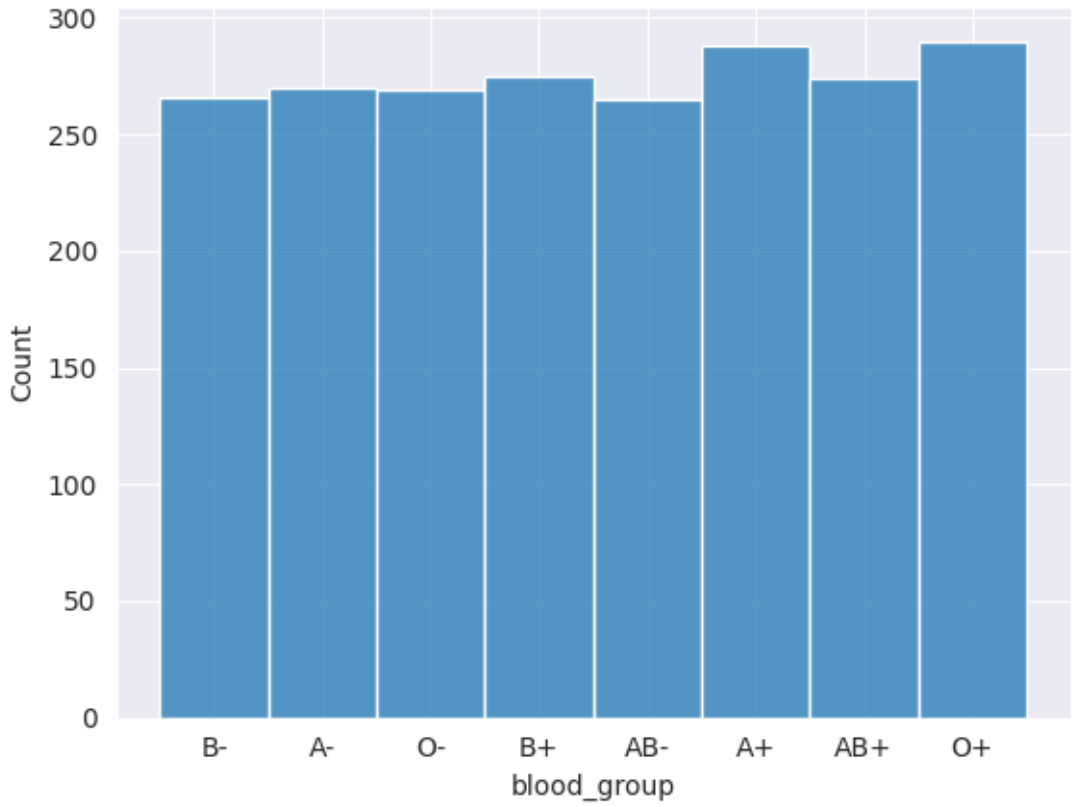


[791]:
```
#Blood group (patient_df)
patient_df['blood_group'].unique().tolist()
#All blood groups are represented
```

[791]:
```
['B-', 'A-', 'O-', 'B+', 'AB-', 'A+', 'AB+', 'O+']
```

[792]:
```
sns.histplot(patient_df['blood_group'])
#not that big of a range
```

[792]:
```
<Axes: xlabel='blood_group', ylabel='Count'>
```



[793]:
```
patient_df['blood_group'].value_counts()
```

[793]:
```
blood_group
O+     290
A+     288
B+     275
AB+    274
A-     270
O-     269
B-     266
```

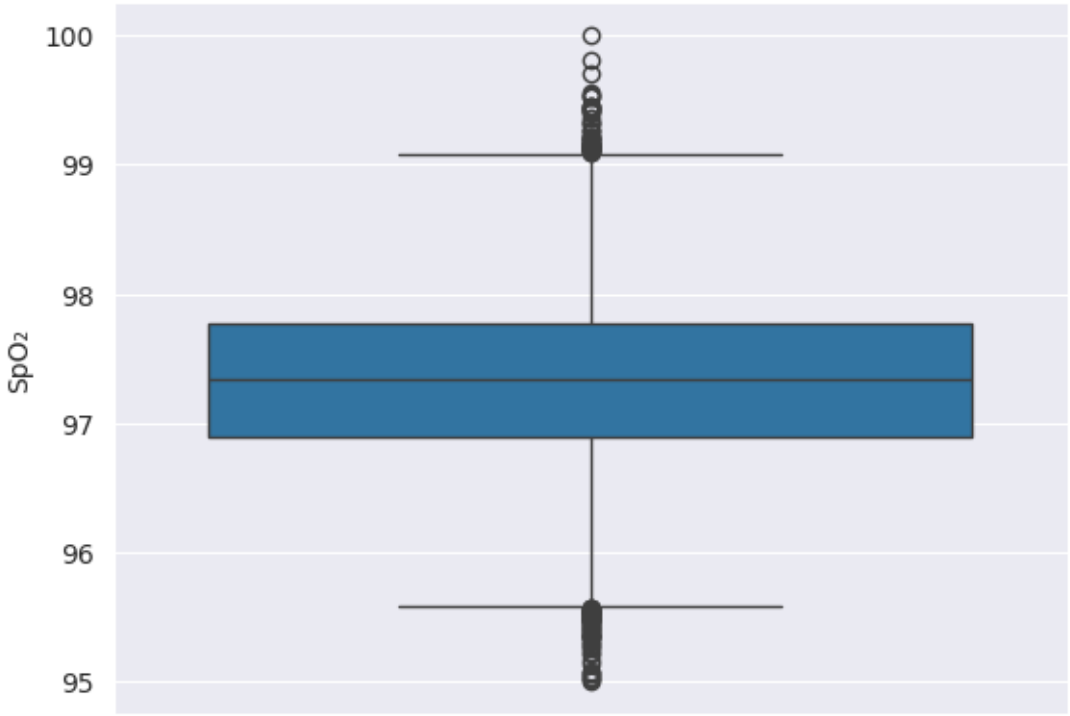
```
AB-    265
Name: count, dtype: int64
```

[794]:
```
patient_df['blood_group'].isna().sum()
#great
```

[794]:
```
np.int64(0)
```

[795]:
```
# SpO2
sns.boxplot(y = observation_df['SpO '])
```

[795]:
```
<Axes: ylabel='SpO '>
```

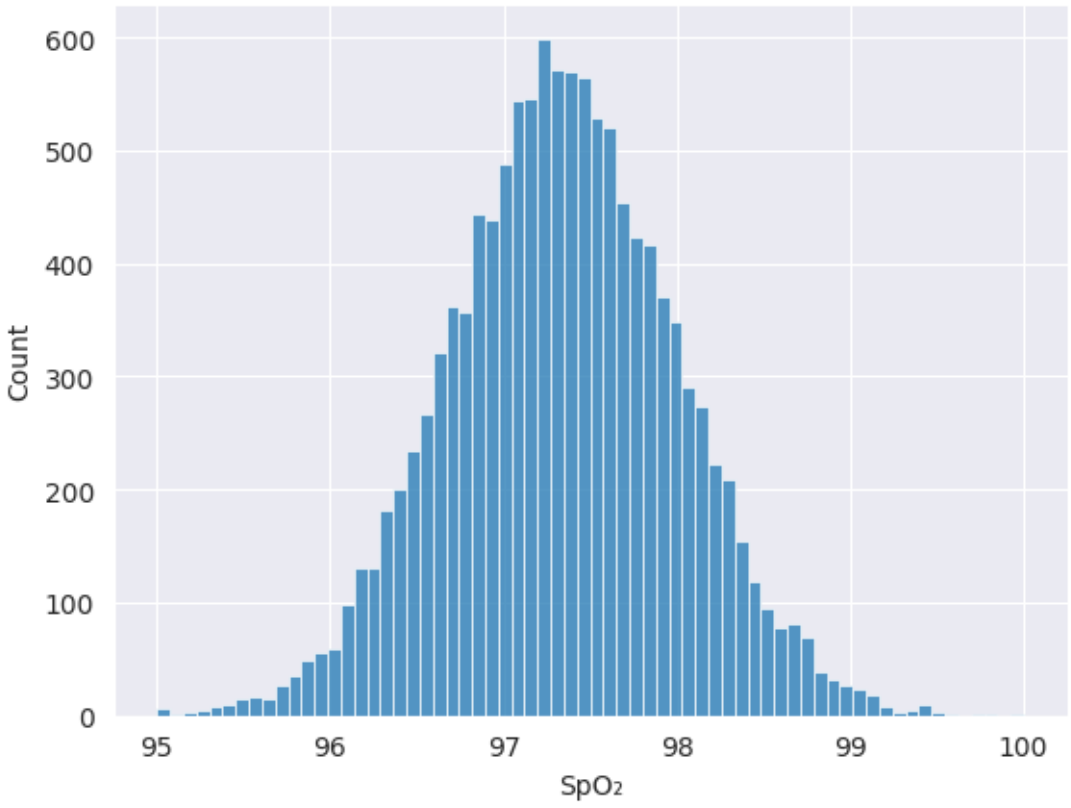


[796]:
```
#just to have the exact range
observation_df['SpO '].min(), observation_df['SpO '].max()
#realistic values
```

[796]:
```
(np.float64(95.0), np.float64(100.0))
```

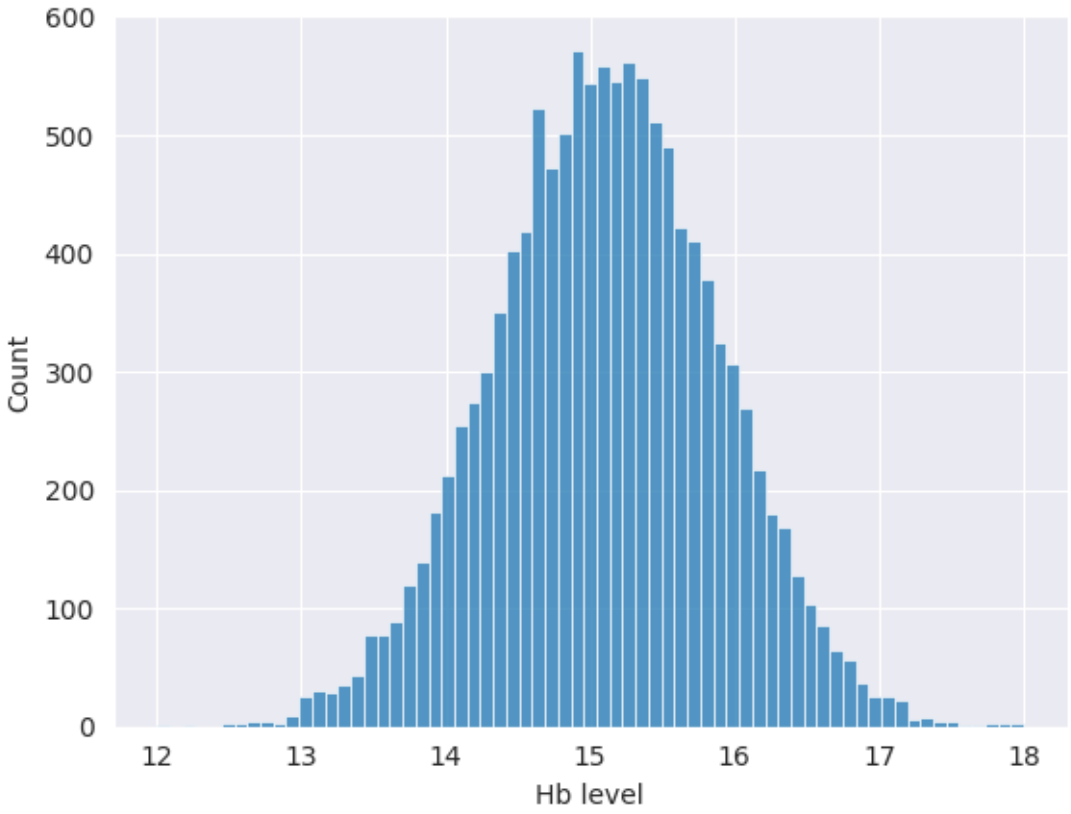
[797]:
```
sns.histplot(observation_df['SpO '])
#normal distribution
```

[797]:
```
<Axes: xlabel='SpO ', ylabel='Count'>
```



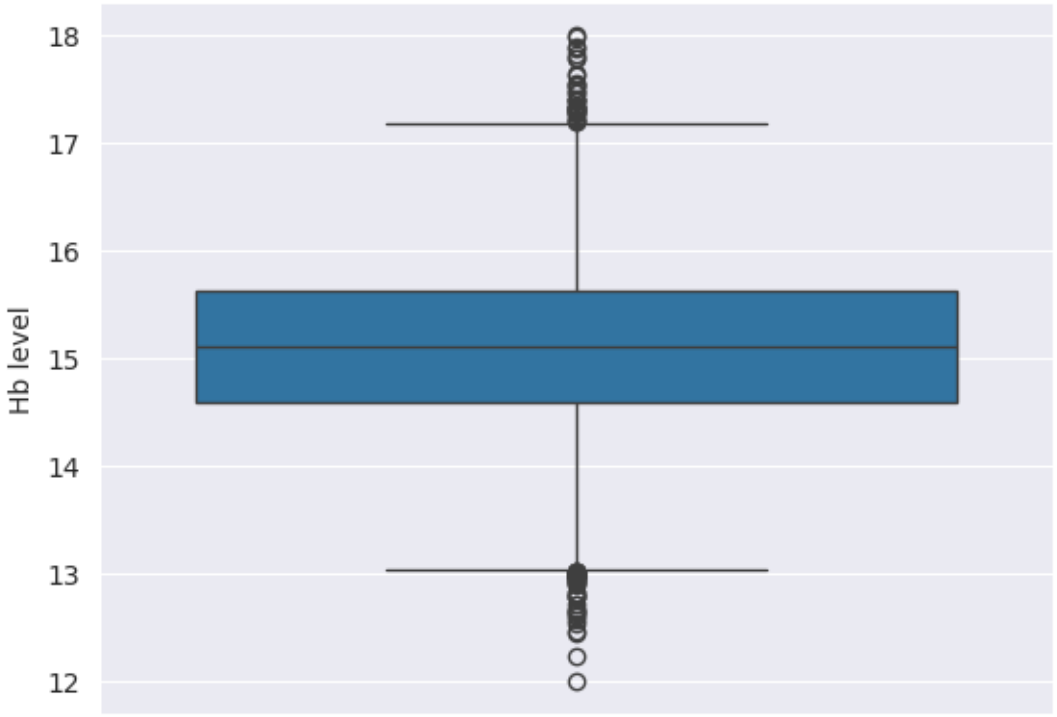
[798]:
```
#Hemoglobin (Observation_df)
sns.histplot(observation_df['Hb level'])
#pretty normal, realistic values
```

[798]: <Axes: xlabel='Hb level', ylabel='Count'>



[799]:
```
sns.boxplot(y = observation_df['Hb level'])
```

[799]: <Axes: ylabel='Hb level'>

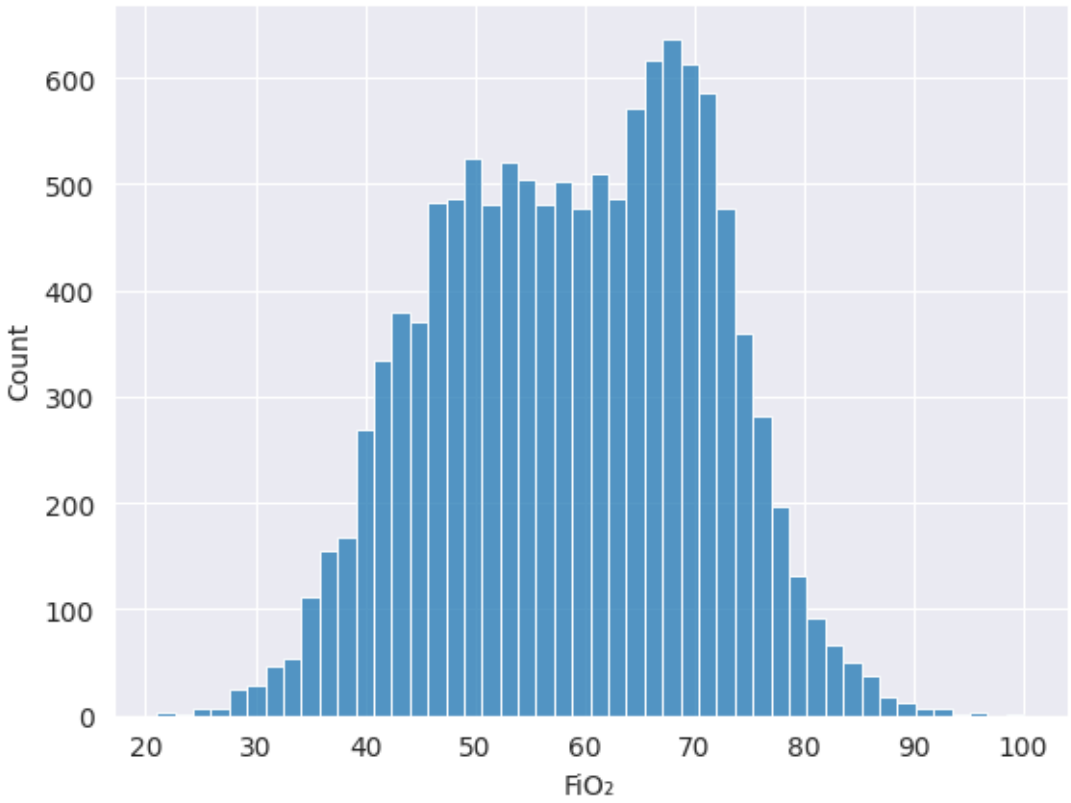


[800]:
```
#exact range
observation_df['Hb level'].min(), observation_df['Hb level'].max()
```

[800]:
```
(np.float64(12.0), np.float64(18.0))
```

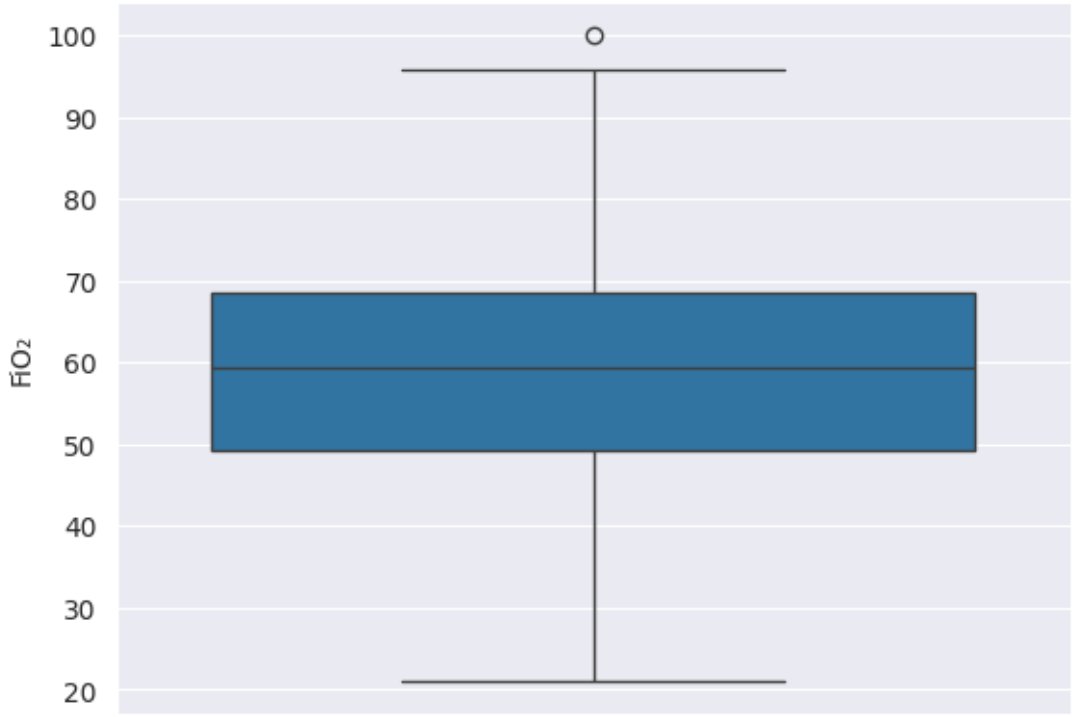
[801]:
```
#Fio (Observation_df)
sns.histplot(observation_df['FiO '])
#not that normal, slight skew to the right at the peak
#also values are not realistic for common people, probably on oxygen therapy or␣
 ↪something
```

[801]:
```
<Axes: xlabel='FiO ', ylabel='Count'>
```



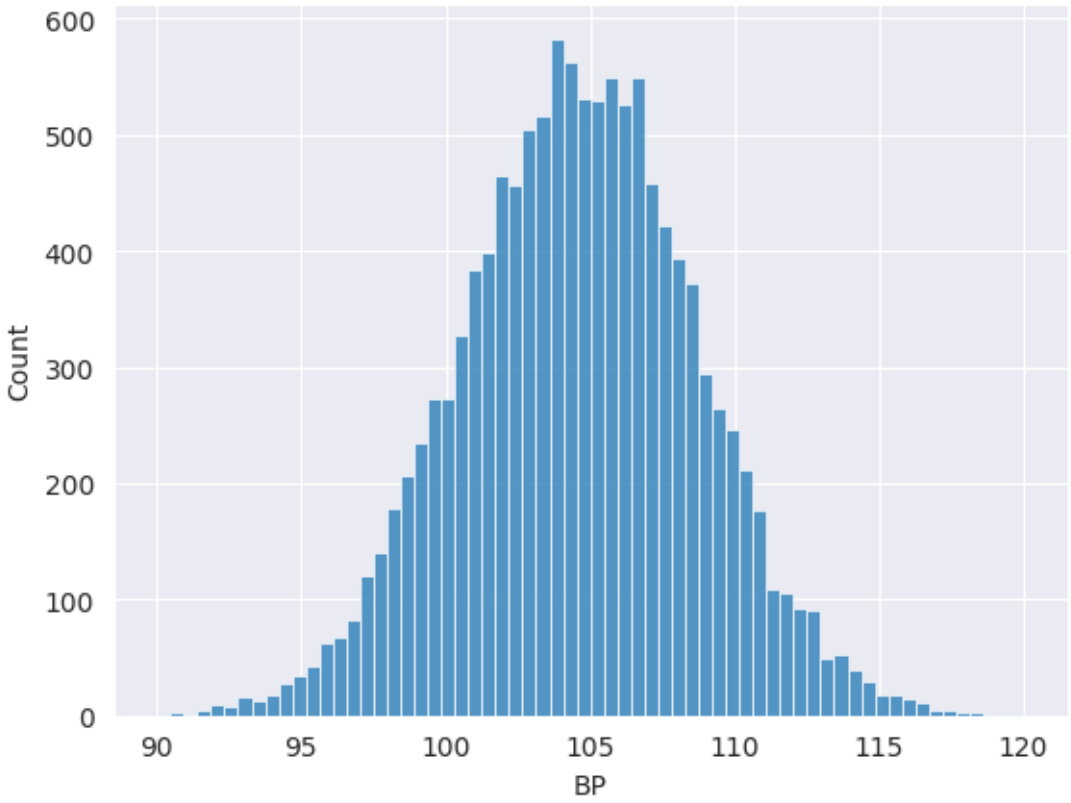
[802]:
```
sns.boxplot(observation_df['FiO '])
```

[802]:
```
<Axes: ylabel='FiO '>
```



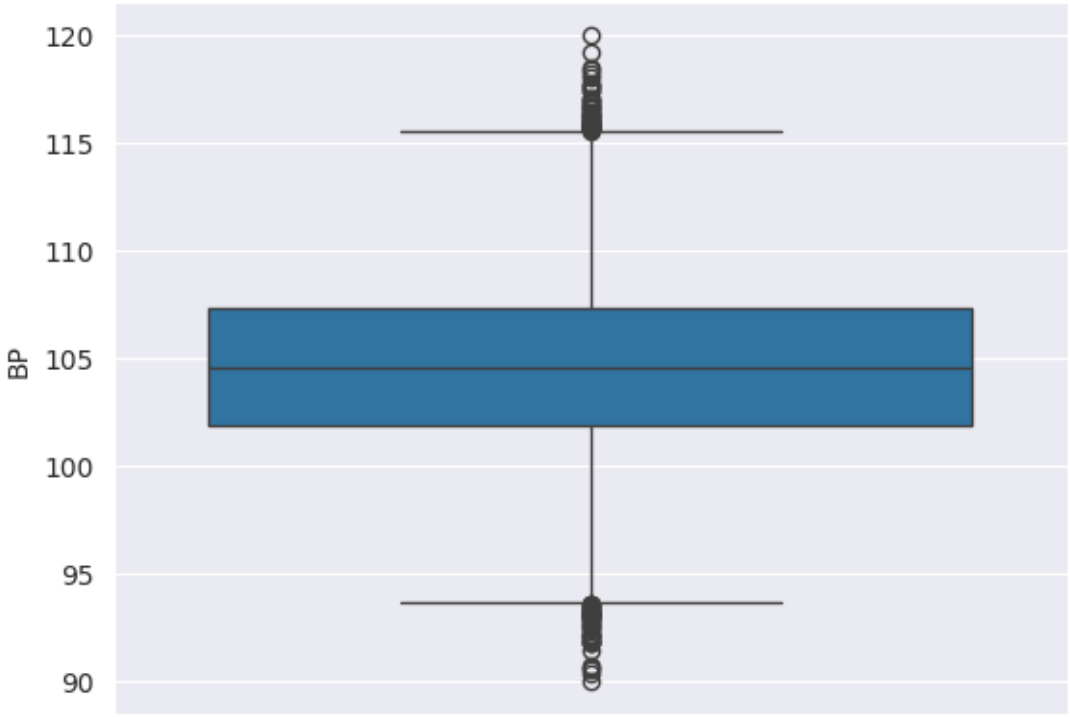
[803]:
```
#blood pressure (observation_df)
sns.histplot(observation_df['BP'])
#normal distribution with usual values
```

[803]: <Axes: xlabel='BP', ylabel='Count'>



[804]:
```
sns.boxplot(y = observation_df['BP'])
```

[804]:
```
<Axes: ylabel='BP'>
```

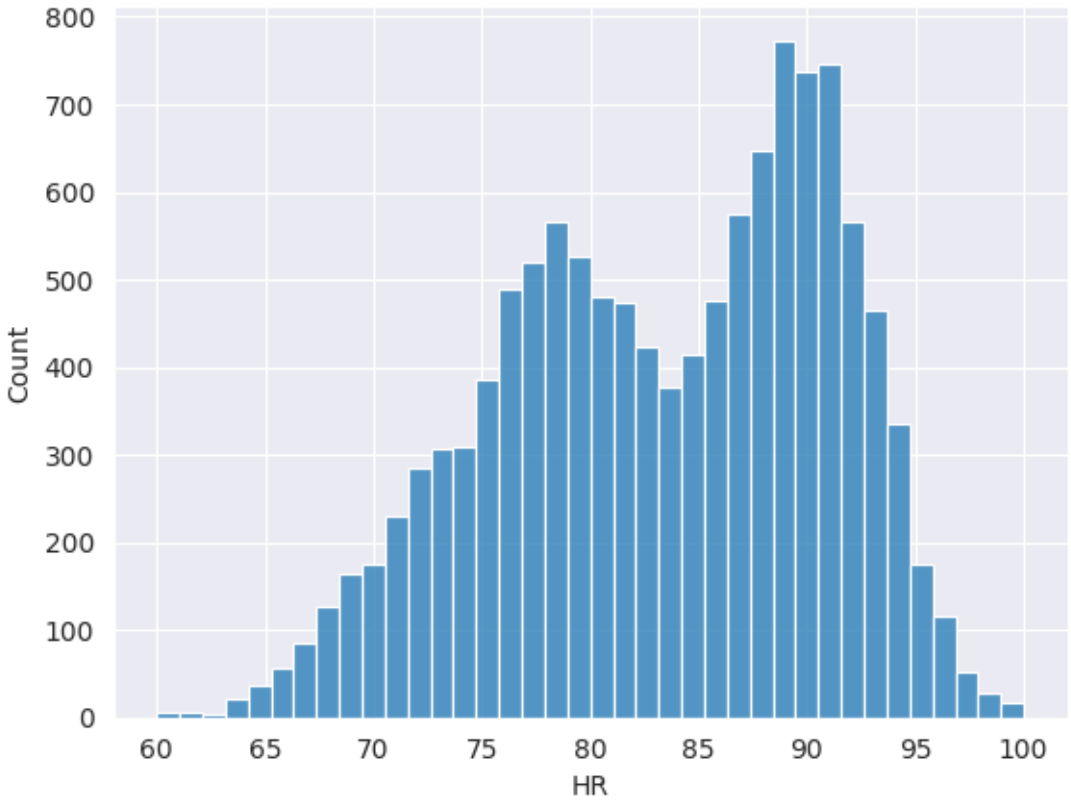


[805]:
```
observation_df['BP'].min(), observation_df['BP'].max()
```

[805]:
```
(np.float64(90.0), np.float64(120.0))
```

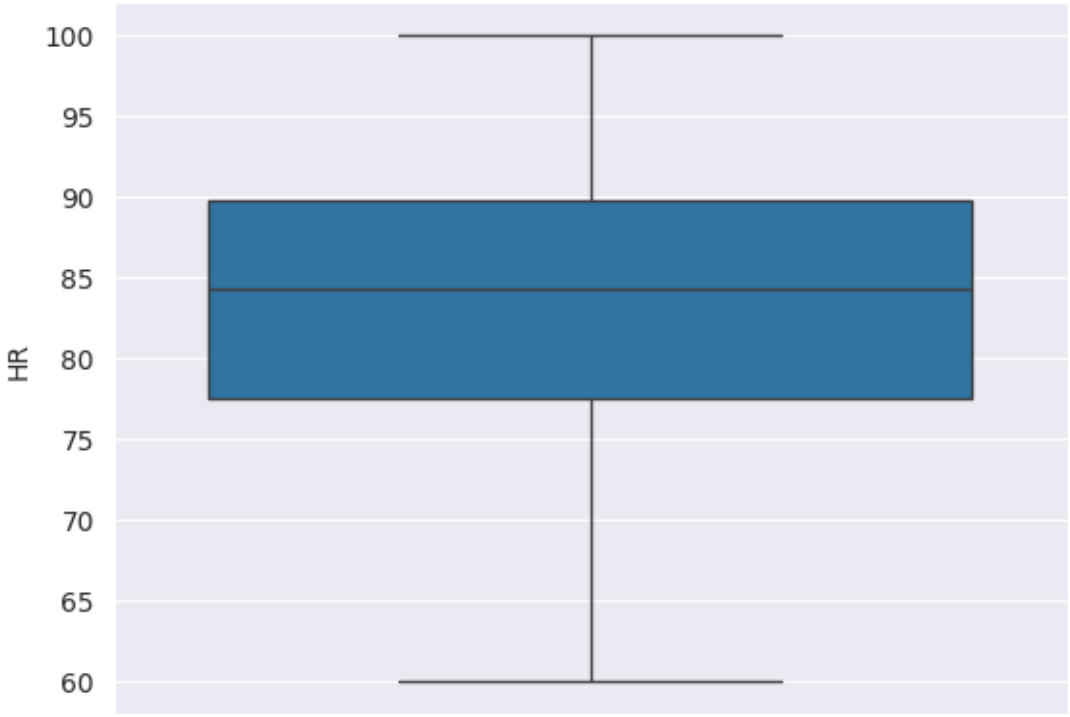
[806]:
```
#Heart rate
sns.histplot(observation_df['HR'])
#pretty bimodal, but realistic values
```

[806]:
```
<Axes: xlabel='HR', ylabel='Count'>
```



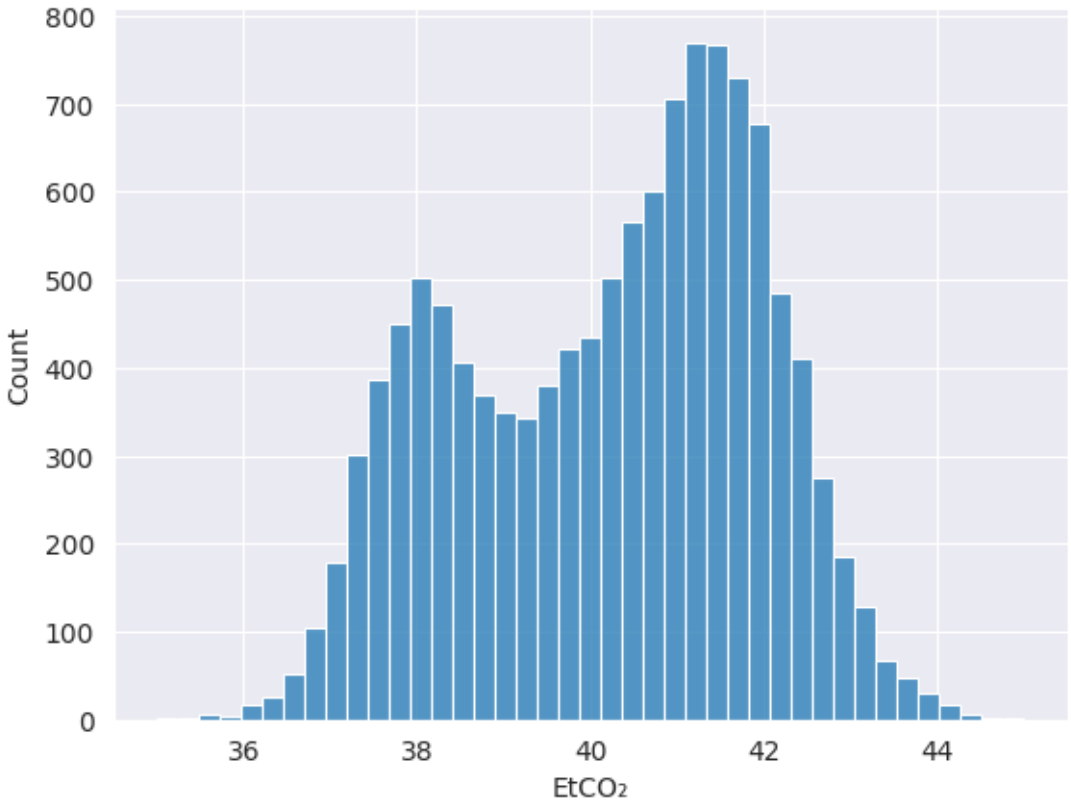
[807]:
```
sns.boxplot(y = observation_df['HR'])
```

[807]:
```
<Axes: ylabel='HR'>
```



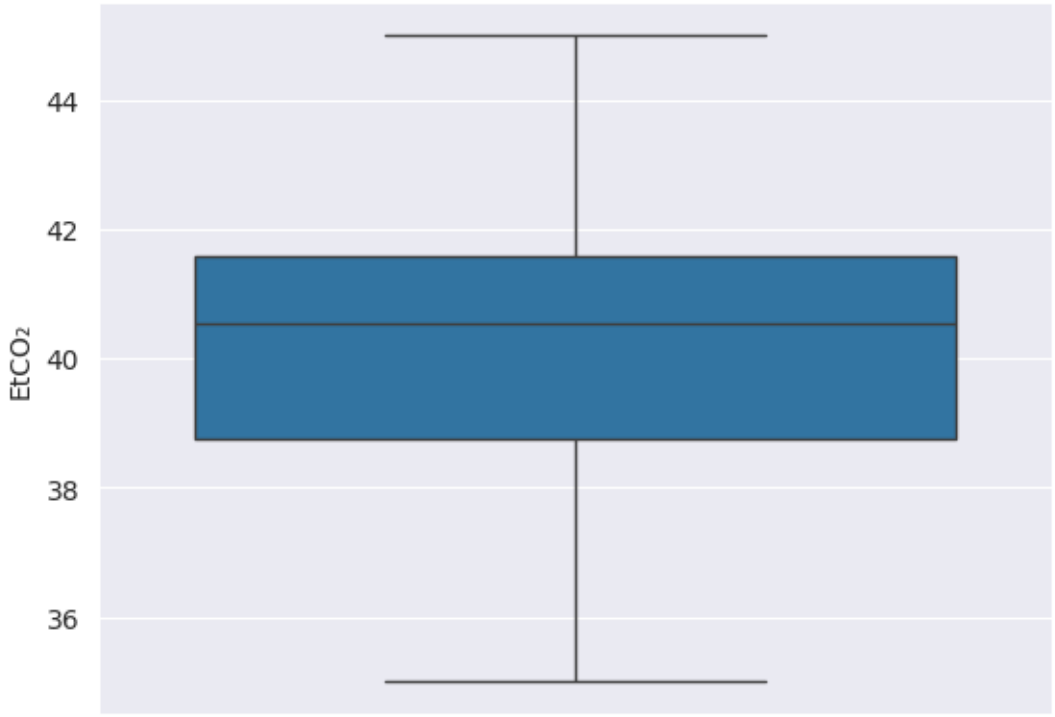
[808]:
```
#ETCO2
sns.histplot(observation_df['EtCO '])
#also bimodal
```

[808]: <Axes: xlabel='EtCO ', ylabel='Count'>



[809]:
```
sns.boxplot(observation_df['EtCO '])
```

[809]: <Axes: ylabel='EtCO '>



[810]:
```
observation_df['EtCO '].min(), observation_df['EtCO '].max()
#also usual values
```

[810]:
```
(np.float64(35.0), np.float64(45.0))
```

[811]:
```
#just checking if this is relevant
patient_df['company'].nunique(), patient_df.shape
#too many unique vals, irrelevant
```

[811]:
```
(1989, (2197, 13))
```

3.1.3 C) Párová analýza dát: Identifikujte vzťahy a závislostí medzi dvojicami atribútov.

But first lets create a dataset with merged tables with only the attributes we consider as necessary

Creation of the joined df

[812]:
```
#This is a list of attributes that will make up the new merged dataset

#QoS might be of significance with Fio and other atts,
#some codes may also have a touch (for example US station are healthier... idk)
#latitude and longitude will be kept for potentional current location␣
 ↪(patient_df) relation
```

```
station_attributes = ['QoS', 'code', 'latitude', 'longitude', ]

#current location has been mentioned in station atts, may be dropped -> later␣
 ↪comment : it was dropped
#blood group may be relevant, may be not we will see
#user id just so we have some unique user identifier since we are dropping name␣
 ↪and everything
patient_attributes = ['user_id', 'blood_group', 'station_ID']

#everything apart from longitude and latitude since that is already kept from␣
 ↪station_attributes and we mentioned before that all stations are mentioned␣
 ↪in obs_df
observation_attributes = ['SpO ', 'HR', 'PI', 'RR', 'EtCO ', 'FiO ', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
       'Signal Quality Index', 'Respiratory effort', 'O  extraction ratio',
       'SNR', 'oximetry']

#for check, lets count the atts
len(station_attributes) + len(patient_attributes) + len(observation_attributes)
```

[812]: 28

[813]:
```
station_df2 = station_df[station_attributes].copy()
patient_df2 = patient_df[patient_attributes].copy()
observation_df2 = observation_df[observation_attributes + ['latitude',␣
 ↪'longitude']].copy()
#this is needed to convert the ID into a normal attribute
station_df2 = station_df2.reset_index().rename(columns={'index': 'station_ID'})
```

[814]:
```
observation_df.shape
```

[814]: (12177, 23)

[815]:
```
df_obs_stat = observation_df2.merge(
    station_df2,
    left_on=['latitude', 'longitude'],   # from merged patiens
    right_on=['latitude', 'longitude'],  # from observation_df
    how='inner'
)
```

[816]:
```
#the shape should be the same as observation_df.shape
df_obs_stat.shape
#it is not because in the following cell we can see that some stations have the␣
 ↪same coordinates
```

[816]: (21385, 26)

[817]:
```
station_df2[['latitude', 'longitude']].duplicated().sum()
```

[817]:
```
np.int64(205)
```

[818]:
```
df_obs_stat.head(5)
```

[818]:

	SpO	HR	PI	RR	EtCO	FiO	\
0	97.538229	87.194745	11.225419	14.812012	42.113735	33.852538	
1	97.933271	80.787303	11.730935	14.964972	39.537692	65.326035	
2	97.933271	80.787303	11.730935	14.964972	39.537692	65.326035	
3	98.209983	79.733895	12.839449	14.840668	39.758706	53.925230	
4	98.202790	86.156903	11.204152	14.523288	43.448577	35.227704	

	PRV	BP	Skin Temperature	Motion/Activity index	…	\
0	144.504405	100.455727	35.961920	10.302567	…	
1	110.615787	102.133386	36.274352	8.975704	…	
2	110.615787	102.133386	36.274352	8.975704	…	
3	107.208040	104.036654	35.583851	7.653790	…	
4	143.282224	105.723603	36.463180	8.795732	…	

	Signal Quality Index	Respiratory effort	O	extraction ratio	SNR	\
0	42.399816	46.497869		0.289012	39.334620	
1	46.078137	53.351208		0.290879	26.006709	
2	46.078137	53.351208		0.290879	26.006709	
3	41.525607	52.124182		0.263171	31.890829	
4	36.535021	50.342830		0.256780	30.721375	

	oximetry	latitude	longitude	station_ID	QoS	code
0	1.0	49.183239	15.454273	403	good	CZ
1	0.0	33.544280	-84.233810	4	good	US
2	0.0	33.544280	-84.233810	426	excellent	US
3	1.0	-27.505780	153.102360	219	excellent	AU
4	1.0	37.656390	126.835000	319	excellent	KR

[5 rows x 26 columns]

[819]:
```
#now we need to join the df_obs_stat with patient_df
df = df_obs_stat.merge(
    patient_df2,
    left_on=['station_ID'],
    right_on=['station_ID'],
    how='inner'
)
```

[820]:
```
df.shape
#now we have many more entries since there are many patients sharing the same␣
 ↪station as we can see in the next cell
```

[820]: (66973, 28)

[821]:
```
patient_df2[['station_ID']].duplicated().sum()
```

[821]: np.int64(1520)

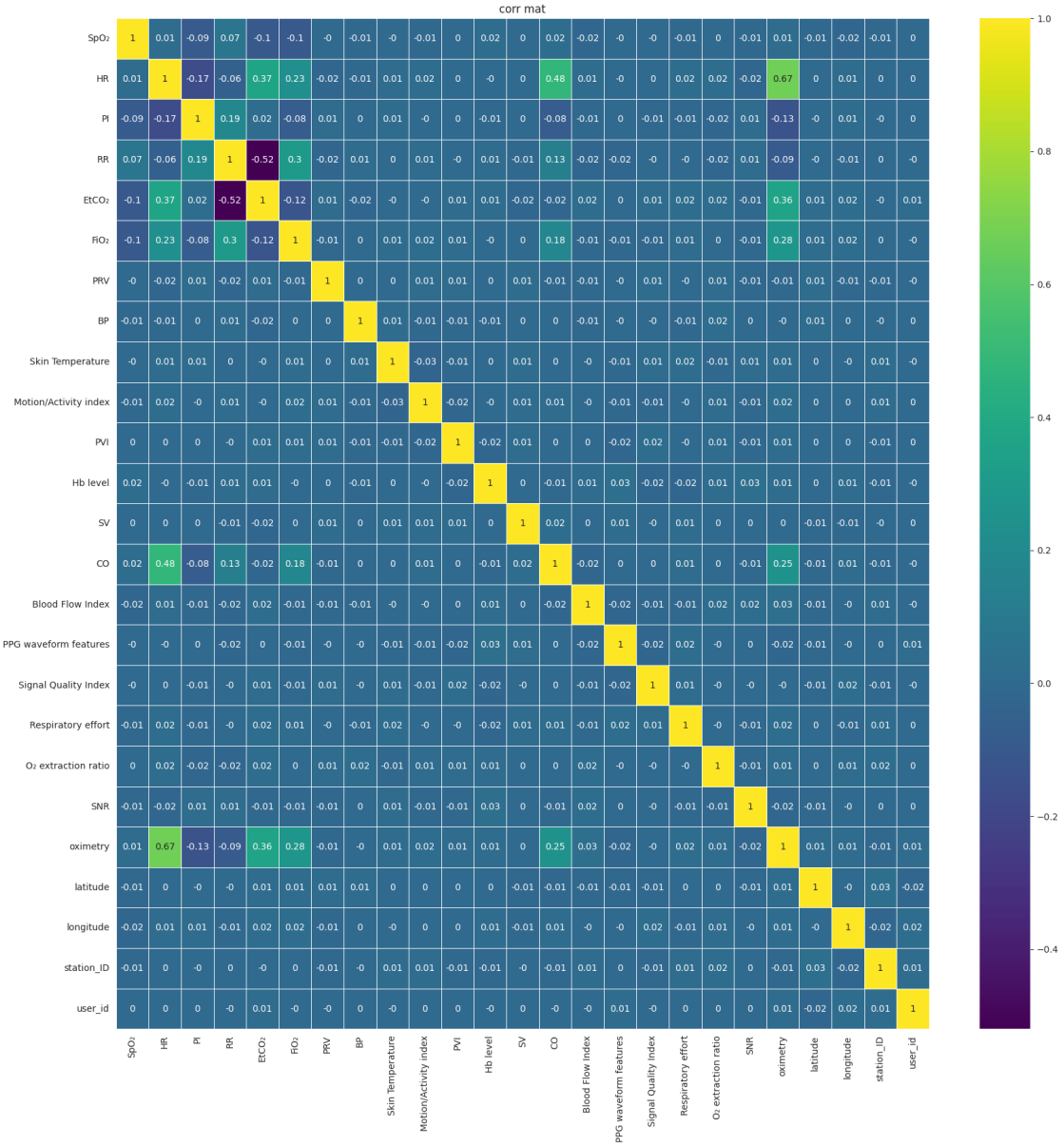
[822]:
```
df.columns
```

[822]:
```
Index(['SpO ', 'HR', 'PI', 'RR', 'EtCO ', 'FiO ', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
       'Signal Quality Index', 'Respiratory effort', 'O  extraction ratio',
       'SNR', 'oximetry', 'latitude', 'longitude', 'station_ID', 'QoS', 'code',
       'user_id', 'blood_group'],
      dtype='object')
```

correlation matrix with heatmap

[823]:
```
numeric_df = df.select_dtypes(include=['float64', 'int64'])

corr_matrix = numeric_df.corr().round(2)
```

[824]:
```
plt.figure(figsize=(20, 20))
sns.heatmap(corr_matrix, cmap='viridis', annot=True,linewidths=0.5)
plt.title("corr mat")
plt.show()
```



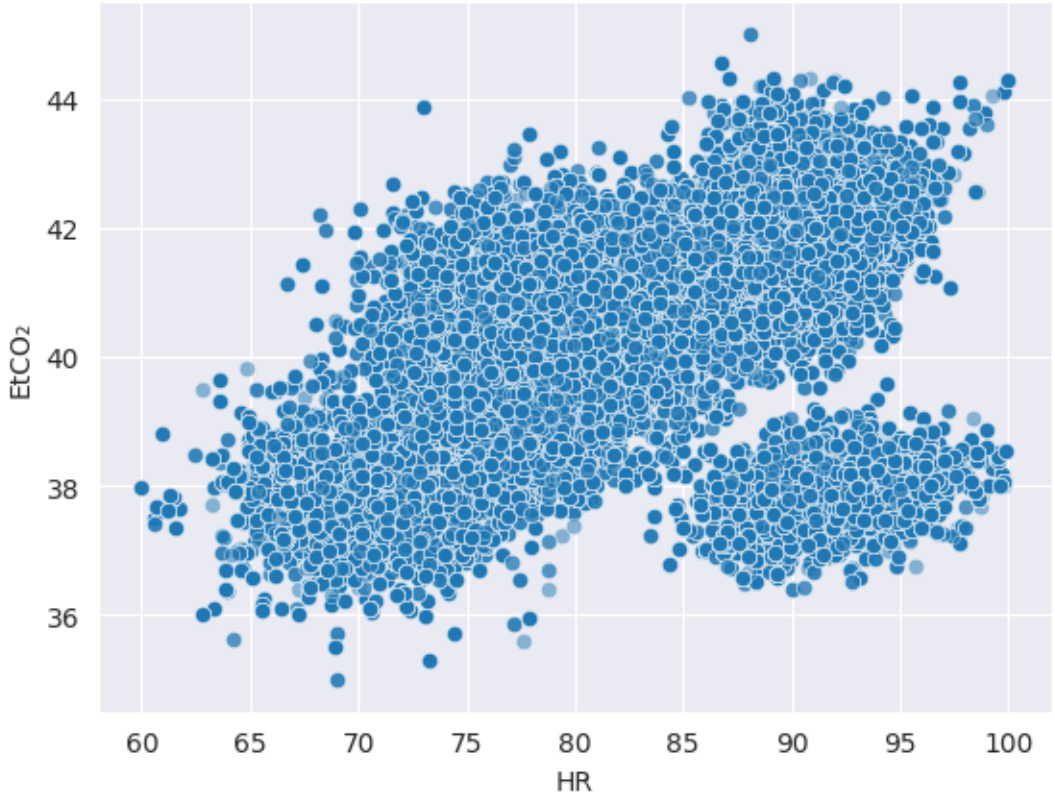
[825]:
```
#based on this heatmap lets write out all the correlations so we can take a␣
 ↪look at them later
#I only included minimum 0,2 corr

#GROUP 1 (WITHOUT OXIMETRY):              -- used in this step

#HR - EtCO2(weak), FiO2(weak), CO(mid)
#RR - Etco2(mid), FiO2(weak),

#Group 2(OXIMETRY):
```

```
#OX - HR(strong),EtCO2(weak) ,FiO2(weak), CO(weak)       -- used in 1D
```

solution for 1.1C) here are the comparisons between the attributes we got from the correlation heatmap

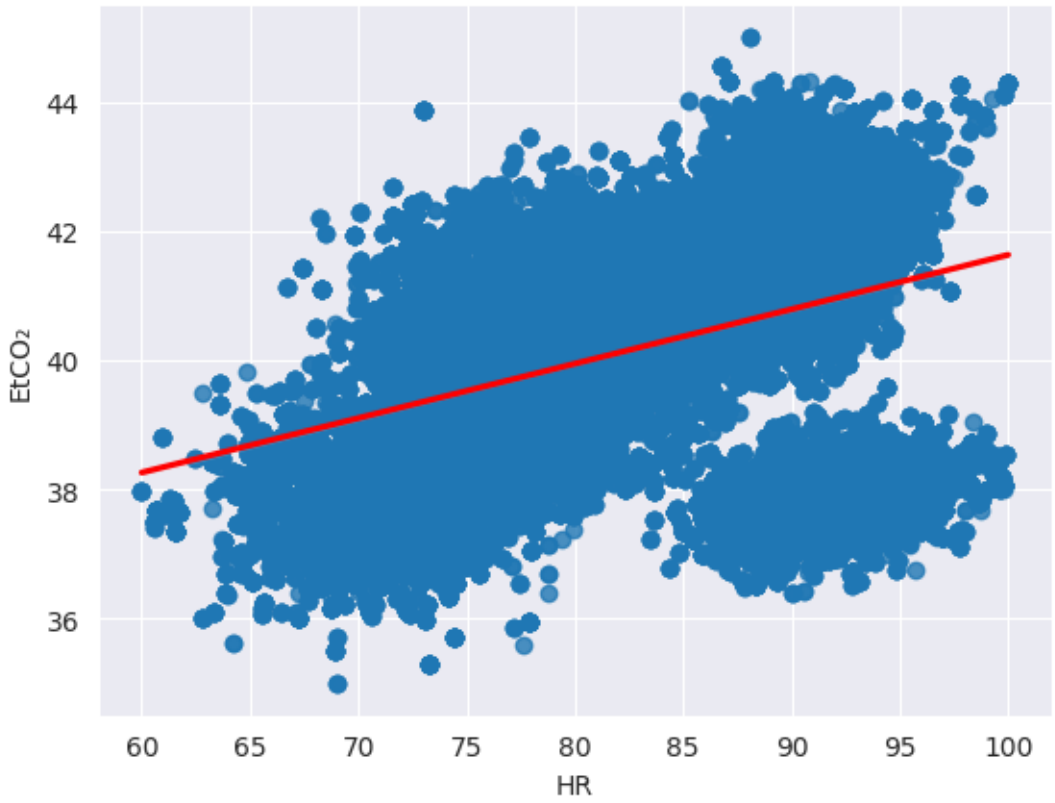
[826]:
```
#HR-EtCO2
sns.scatterplot(data=df, x='HR', y='EtCO ', alpha=0.5)
```

[826]: <Axes: xlabel='HR', ylabel='EtCO '>



[827]:
```
sns.regplot(data=df, x='HR', y='EtCO ', line_kws={'color':'red'})
```

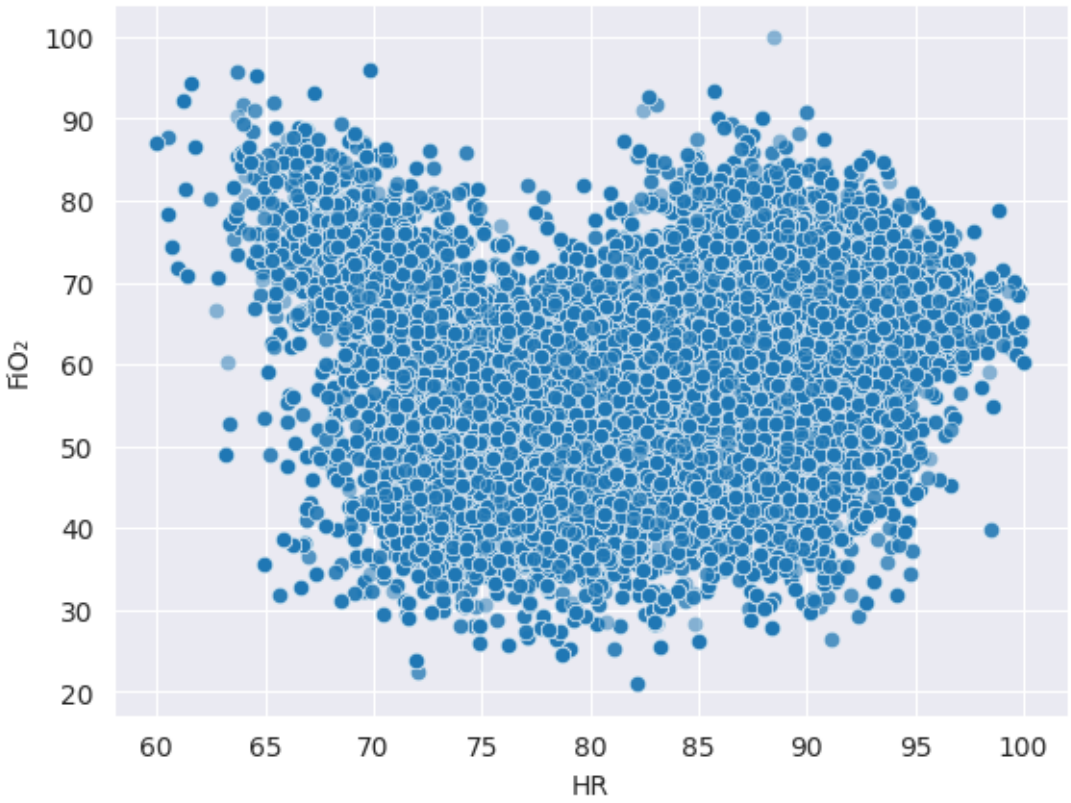
[827]: <Axes: xlabel='HR', ylabel='EtCO '>



[828]:
```
#this is not enough to determine that these two variables correlate, maybe␣
↪without the cluster around [92.5 , 38]
```

[829]:
```
#HR-FiO2
sns.scatterplot(data=df, x='HR', y='FiO ', alpha=0.5)
```

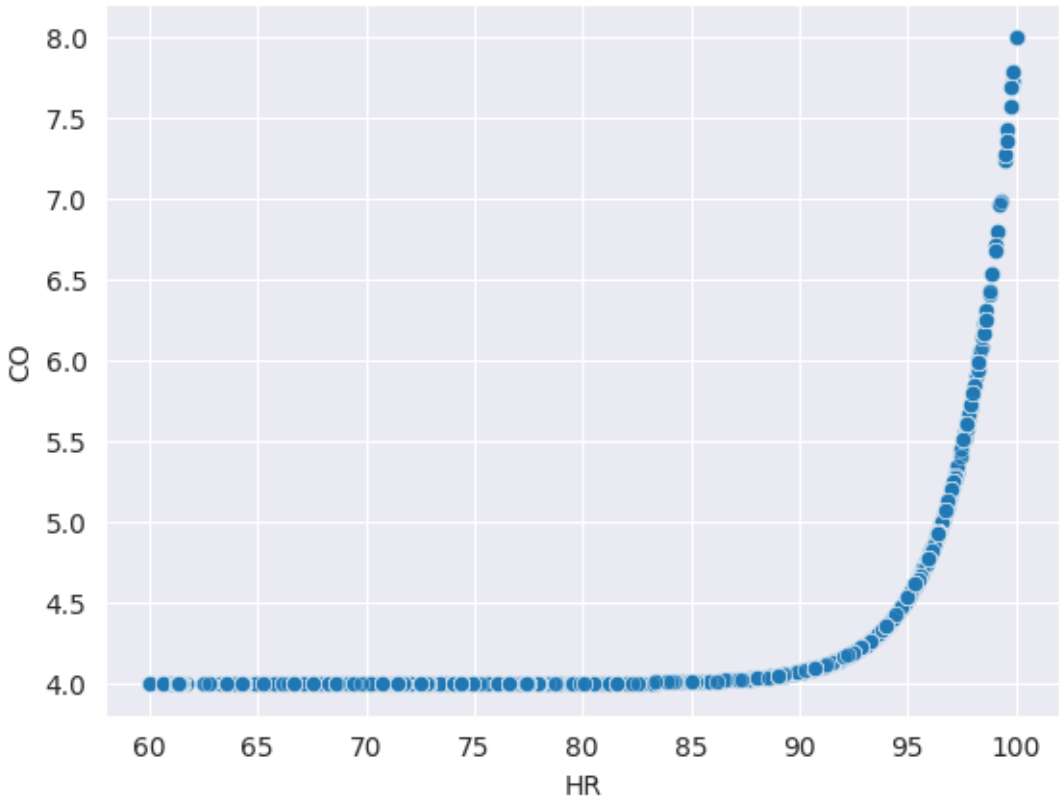
[829]: <Axes: xlabel='HR', ylabel='FiO '>



[830]:
```
#way too spread out, insignificant
```

[831]:
```
#HR-CO
sns.scatterplot(data=df, x='HR', y='CO', alpha=0.5)
```

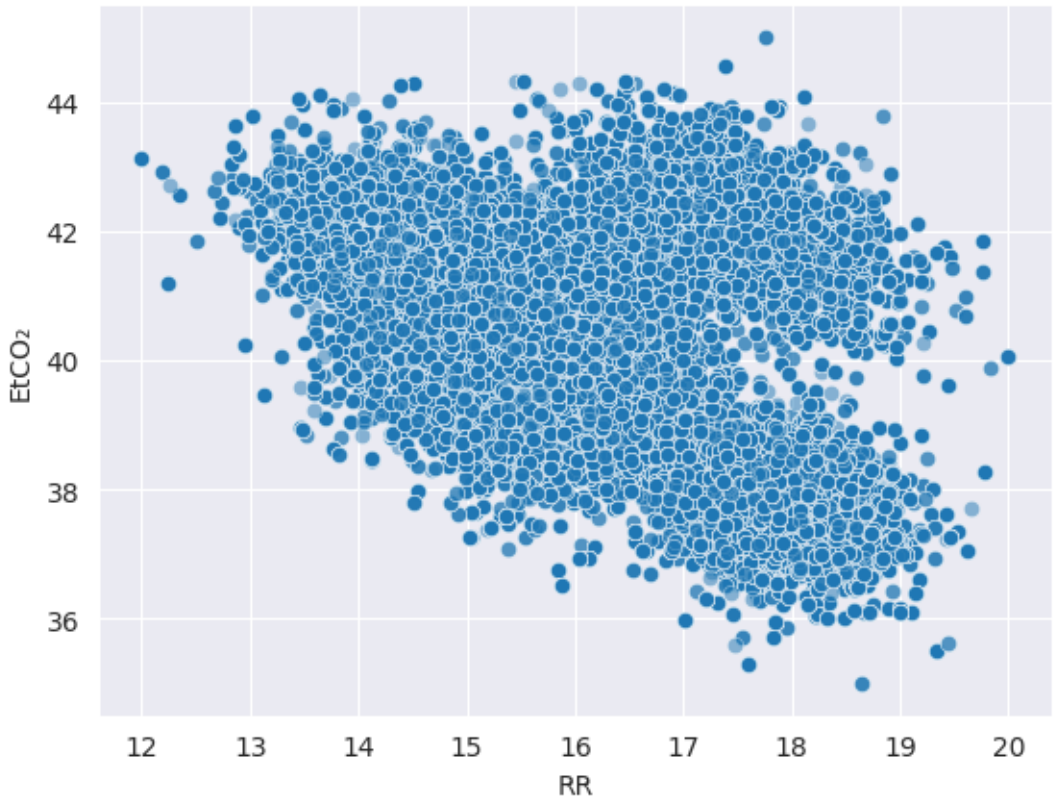
[831]: <Axes: xlabel='HR', ylabel='CO'>



[832]:
```
#This is definitely a significant correlation, but not linear since CO values␣
↪are cut off at 4.0
```

[833]:
```
#RR - Etco2
sns.scatterplot(data=df, x='RR', y='EtCO ', alpha=0.5)
```

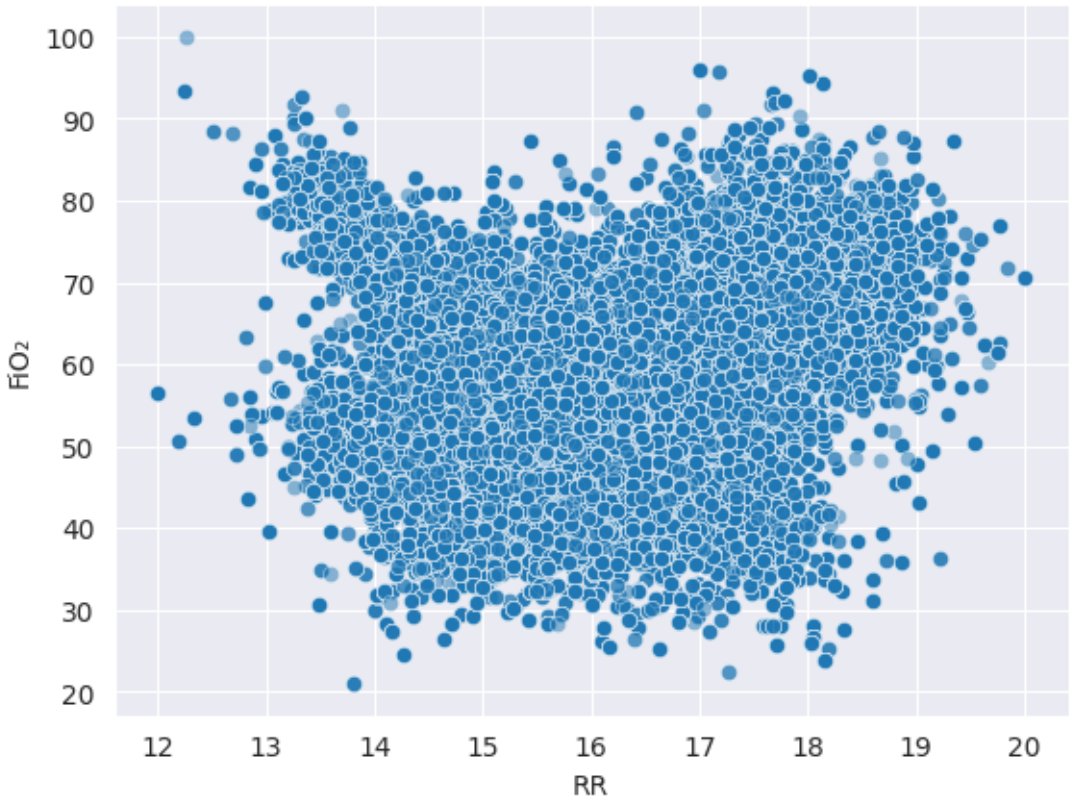
[833]:
```
<Axes: xlabel='RR', ylabel='EtCO '>
```



[834]:
```
#slight show of negative correlation, but ruined by the upper right cluster
```

[835]:
```
#RR FiO2
sns.scatterplot(data=df, x='RR', y='FiO ', alpha=0.5)
```

[835]:
```
<Axes: xlabel='RR', ylabel='FiO '>
```



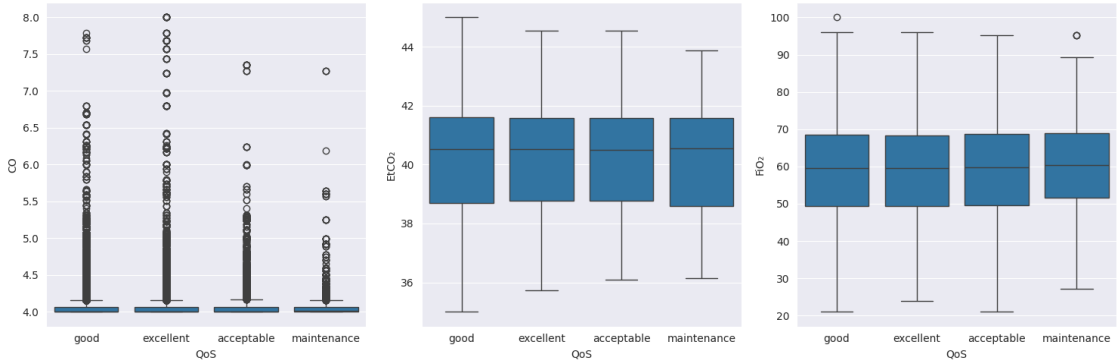
[836]:
```
#no correlation
```

these were only the numeric attributes we took from the heatmap, now lets take at look at non numeric correlations aswell

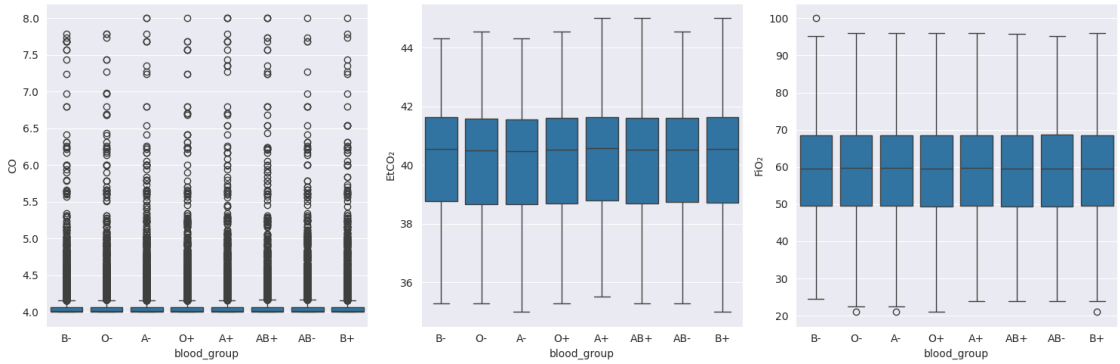
[837]:
```
non_numeric = df.select_dtypes(exclude=['float64', 'int64']).columns
print(non_numeric)
#these will be tested for correlation with CO,FIO2 and EtCO2 since these seem␣
 ↪to be the most important attributes
```

```
Index(['QoS', 'code', 'blood_group'], dtype='object')
```

[838]:
```
#QoS
fig, axes = plt.subplots(1, 3, figsize=(15, 5))

sns.boxplot(data=df, x='QoS', y='CO', ax=axes[0])
sns.boxplot(data=df, x='QoS', y='EtCO ', ax=axes[1])
sns.boxplot(data=df, x='QoS', y='FiO ', ax=axes[2])

plt.tight_layout()
plt.show()
#no real significance
```

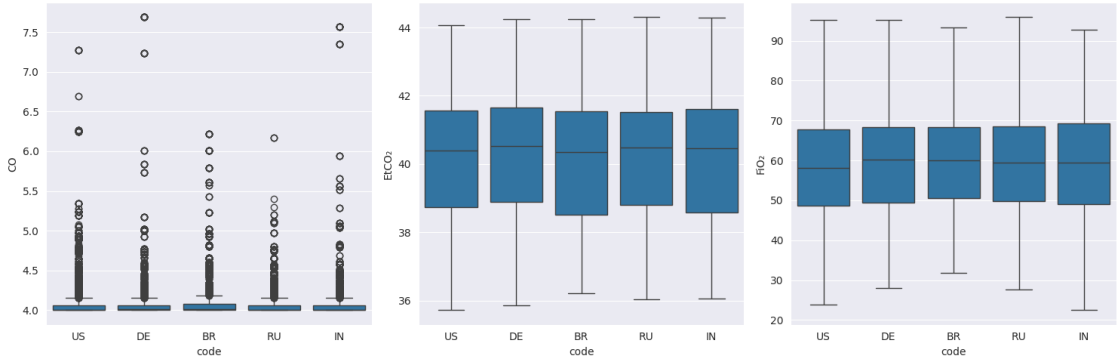


[839]:
```
#QoS
fig, axes = plt.subplots(1, 3, figsize=(15, 5))

sns.boxplot(data=df, x='blood_group', y='CO', ax=axes[0])
sns.boxplot(data=df, x='blood_group', y='EtCO ', ax=axes[1])
sns.boxplot(data=df, x='blood_group', y='FiO ', ax=axes[2])

plt.tight_layout()
plt.show()
```



[840]:
```
#code - top 5
top_codes = df['code'].value_counts().head(5).index
filtered_df = df[df['code'].isin(top_codes)]

fig, axes = plt.subplots(1, 3, figsize=(15, 5))

sns.boxplot(data=filtered_df, x='code', y='CO', ax=axes[0])
sns.boxplot(data=filtered_df, x='code', y='EtCO ', ax=axes[1])
sns.boxplot(data=filtered_df, x='code', y='FiO ', ax=axes[2])
```

```
plt.tight_layout()
plt.show()
```

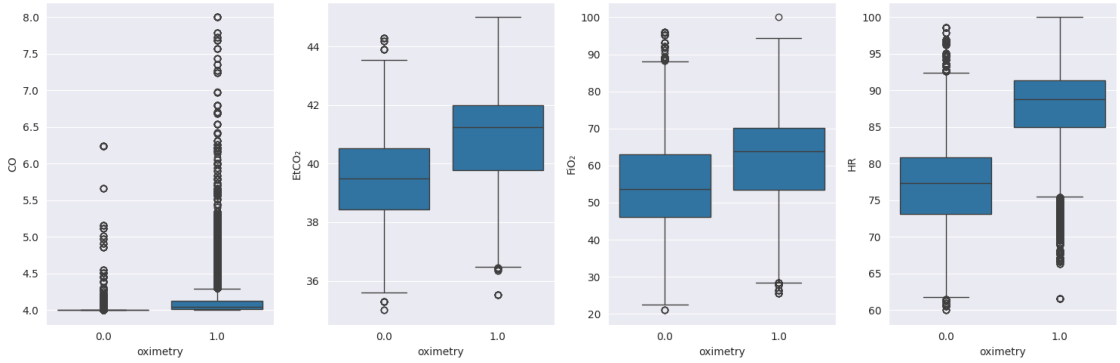


3.1.4 D)Párová analýza dát: Identifikujte závislosti medzi predikovanou premennou a ostatnými premennými (potenciálnymi prediktormi)

from the correlation heatmap we have these attributes that seem to have some correlation with the predicted attribute

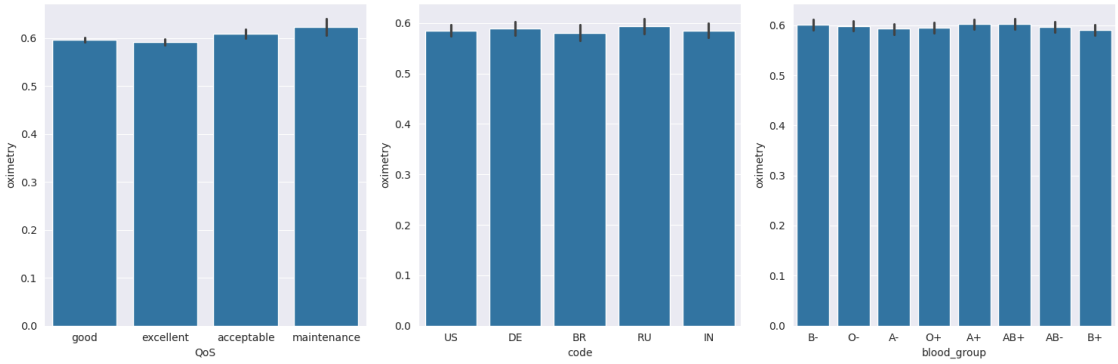
HR(strong),EtCO2(weak) ,FiO2(weak), CO(weak)

[841]:
```
fig, axes = plt.subplots(1, 4, figsize=(15, 5))

sns.boxplot(data=df, x='oximetry', y='CO', ax=axes[0])
sns.boxplot(data=df, x='oximetry', y='EtCO ', ax=axes[1])
sns.boxplot(data=df, x='oximetry', y='FiO ', ax=axes[2])
sns.boxplot(data=df, x='oximetry', y='HR', ax=axes[3])

plt.tight_layout()
plt.show()
```



[842]:
```
#CO is hard to determine, the IQRs are close but the outliers may mean a lot␣
↪(will probably see more after normalization), the EtCO2 and FiO2 seem to be␣
↪higher when oximetry is set to 1, HR seems to have a really significant␣
↪correlation with oximetry, as suggested by the correlation heatmap
```

[843]:
```
#correlation with non numerical values
fig, axes = plt.subplots(1, 3, figsize=(15, 5))

# QoS
sns.barplot(ax=axes[0], x=df['QoS'], y=df['oximetry'])
# code (top 5)
top5 = df['code'].value_counts().head(5).index
sns.barplot(ax=axes[1], x=df[df['code'].isin(top5)]['code'], y=df[df['code'].
↪isin(top5)]['oximetry'])
# blood_group
sns.barplot(ax=axes[2], x=df['blood_group'], y=df['oximetry'])

plt.tight_layout()
plt.show()
```



[844]:
```
#this graphs show the oximetry 1/0 ratio per each QoS, Code and blood group. So␣
↪rather than correlation, this shows that the data are evenly distributed
```

3.1.5 E) Findings, thought ect.

all of our thoughs and findings are thoroughly documented throughout the operations, but here is the summary:

Firstly we analysed each table and gave a quick look for every attribute, for dataset station, we used the coordinations to merge it with observation_df, we ended up not using the station name as it is insignificant as well as revision. The only missing values were 2 codes which will probably be filled later with a newly created code. The patient dataframe is full of insignificant attributes, we end up only using the blood group and station ID for merge with station df. From observation we keep every value, although some seem to be more signicicant than others, for example FiO2, CO,RR, EtCO2. We take some time to look at individual distributions of attributes in B), but we dont

really find any abnormalities that would pose a threat to our models precition, most distributions are either normal or bimodal or uniform. In C) we needed to take a look at correlation between the attributes, so we needed to create a correlation heatmap which was created from a joint dataframe consisting of all three datasets, but only the attributes that we deemed as important. We discarded the useless features and we will not work with them from that point onward. However we find out minor correlation between HR and EtCO2 as well as a negative one between HR and FiO2 and a RR and EtCO2. HR and CO seem to have a strong positive correlation but since CO is capped at 4.0, it is not shown that nicely. In D) we see that the predicted value oximetry strongly correlates with HR, and FiO2 and EtCO2 also show medium signs of correlation. Correlation with CO is harder to determine without normalising the data first.

3.2 1.2 Identifikácia problémov, integrácia a čistenie dát

3.2.1 A) nevhodná štruktúra, nejednotne formaty ,duplikáty, chýbajúce hodnoty, vychýlene hodnoty, abnormalne hodnoty, nelogické vzťahy

[845]:
```
df.columns
```

[845]:
```
Index(['SpO ', 'HR', 'PI', 'RR', 'EtCO ', 'FiO ', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
       'Signal Quality Index', 'Respiratory effort', 'O  extraction ratio',
       'SNR', 'oximetry', 'latitude', 'longitude', 'station_ID', 'QoS', 'code',
       'user_id', 'blood_group'],
      dtype='object')
```

[846]:
```
# I see many collumn name that are corectly name but it is so anoying to copy␣
 ↪the same ` ` symbol over and over again so that's why im gonna rename it just␣
 ↪for now

rename_map = {
    'SpO ': 'SpO2',
    'EtCO ': 'EtCO2',
    'FiO ': 'FiO2',
    'O  extraction ratio': 'O2 extraction ratio',
}
df = df.rename(columns={k:v for k,v in rename_map.items() if k in df.columns})
```

[847]:
```
for c in ['QoS','code','blood_group']:
    if c in df.columns:
        df[c] = df[c].astype('category')
```

[848]:
```
#checking if every value in oximetry col is type int so it does not make any␣
 ↪issues in the future but most of this is already done in previous cells in 1.
 ↪1

if 'oximetry' in df.columns:
    df['oximetry'] = df['oximetry'].astype(int)
```

[849]:
```
df.dtypes.value_counts()
```

[849]:
```
float64     22
int64        3
category     1
category     1
category     1
Name: count, dtype: int64
```

[850]:
```
df.duplicated().sum()
```

[850]:
```
np.int64(0)
```

[851]:
```
df.duplicated(subset=['HR','RR','BP']).sum()
```

[851]:
```
np.int64(55244)
```

[852]:
```
duplicates_output = df[df.duplicated(subset=[col for col in df.columns if col !
 ↪= 'user_id'], keep=False)]
duplicates_output
```

[852]:

	SpO2	HR	PI	RR	EtCO2	FiO2	\
6	97.933271	80.787303	11.730935	14.964972	39.537692	65.326035	
7	97.933271	80.787303	11.730935	14.964972	39.537692	65.326035	
9	97.933271	80.787303	11.730935	14.964972	39.537692	65.326035	
30	96.851933	91.008225	11.323571	15.148349	42.056062	58.615353	
32	96.851933	91.008225	11.323571	15.148349	42.056062	58.615353	
…	…	…	…	…	…	…	
66965	96.698981	71.499309	13.427977	15.751513	41.201634	55.177636	
66966	96.698981	71.499309	13.427977	15.751513	41.201634	55.177636	
66967	96.698981	71.499309	13.427977	15.751513	41.201634	55.177636	
66968	96.698981	71.499309	13.427977	15.751513	41.201634	55.177636	
66970	96.698981	71.499309	13.427977	15.751513	41.201634	55.177636	

	PRV	BP	Skin Temperature	Motion/Activity index	…	\
6	110.615787	102.133386	36.274352	8.975704	…	
7	110.615787	102.133386	36.274352	8.975704	…	
9	110.615787	102.133386	36.274352	8.975704	…	
30	77.660061	102.695264	35.163246	11.823341	…	
32	77.660061	102.695264	35.163246	11.823341	…	
…	…	…	…	…	…	
66965	103.386006	102.693239	34.895596	11.429244	…	
66966	103.386006	102.693239	34.895596	11.429244	…	
66967	103.386006	102.693239	34.895596	11.429244	…	
66968	103.386006	102.693239	34.895596	11.429244	…	
66970	103.386006	102.693239	34.895596	11.429244	…	

	O2 extraction ratio	SNR	oximetry	latitude	longitude	\

```
6                 0.290879  26.006709         0  33.54428  -84.23381
7                 0.290879  26.006709         0  33.54428  -84.23381
9                 0.290879  26.006709         0  33.54428  -84.23381
30                0.285955  32.768112         1  10.29085  105.75635
32                0.285955  32.768112         1  10.29085  105.75635
…                    …         …        …        …         …
66965             0.283565  28.887543         1  51.04962   12.13690
66966             0.283565  28.887543         1  51.04962   12.13690
66967             0.283565  28.887543         1  51.04962   12.13690
66968             0.283565  28.887543         1  51.04962   12.13690
66970             0.283565  28.887543         1  51.04962   12.13690

       station_ID        QoS  code  user_id  blood_group
6             426  excellent    US      398           O+
7             426  excellent    US      988           O+
9             426  excellent    US     2033           O+
30            663  excellent    VN     1225           A-
32            663  excellent    VN     1454           A-
…           …        …   …       …          …
66965         223  excellent    DE      575           B-
66966         223  excellent    DE      468           O+
66967         223  excellent    DE     1212           B-
66968         223  excellent    DE      928           B-
66970         223  excellent    DE       51           B-

[21334 rows x 28 columns]
```

[853]:
```
duplicates_output['user_id'].nunique(), len(duplicates_output)
```

[853]:
```
(599, 21334)
```

[854]:
```
duplicates_output['station_ID'].value_counts().head()
```

[854]:
```
station_ID
186    468
8      414
350    352
208    318
223    318
Name: count, dtype: int64
```

After checking for duplicates, we found that some patients shared identical measurements. It is normal but not if we found out that 21 336 row that are similar. This was caused by merging patient_df and observation_df using station_ID.

[855]:
```
na_count = df.isna().sum().sort_values(ascending=False)
na_count
```

[855]:
```
code                      170
SpO2                        0
PI                          0
HR                          0
EtCO2                       0
FiO2                        0
PRV                         0
BP                          0
Skin Temperature            0
Motion/Activity index       0
PVI                         0
RR                          0
Hb level                    0
SV                          0
Blood Flow Index            0
CO                          0
Signal Quality Index        0
Respiratory effort          0
O2 extraction ratio         0
PPG waveform features       0
SNR                         0
oximetry                    0
longitude                   0
latitude                    0
station_ID                  0
QoS                         0
user_id                     0
blood_group                 0
dtype: int64
```

3.2.2 B) Kontrola správnosť v dátach

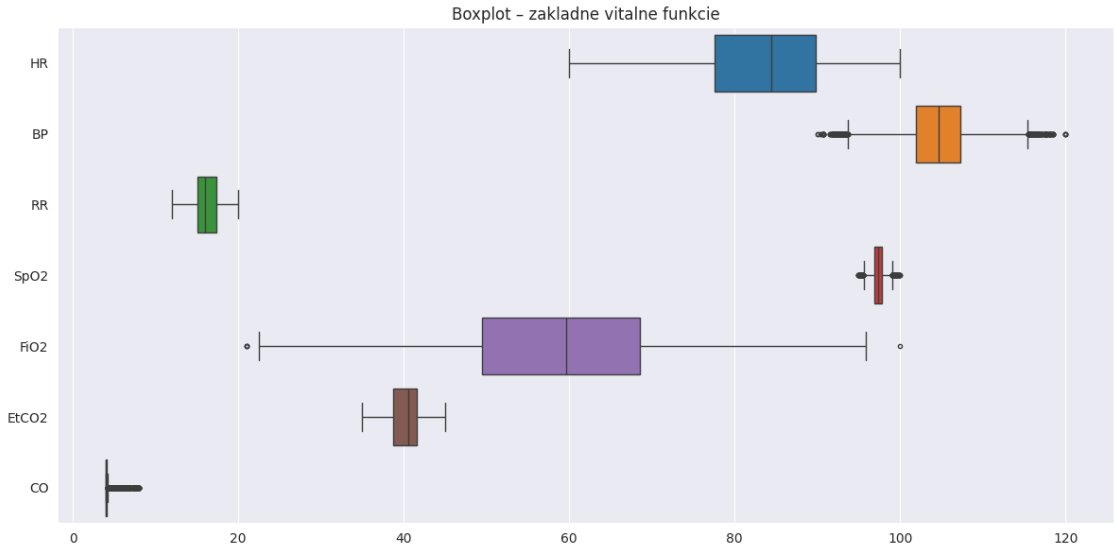
Most of the analyzing the atributes was done in part 1.1, individual findings were documented along the way.

Some of the most important findings that were captured along the way about the abnormality of the data could be divided into 3 groups. 1) Value-range-wise, most of the attributes have real values, for example the latitude and longitude do range from -180 to 180, SPO2 and blood groups are also in check and so are many more atributes. The only attribute where this was not the case would be FIO2, where the patients must have been altered from normal people, for example on oxygen therapy. 2) Distribution-wise, no attributes are abnormaly different. Some distributions were normal, e.g. SPO2, Hb level. Some were uniform, for example blood group. Some were bimodal, for example HR or EtCO2 which does show quite a skew to abnormality. FIO2 is slightly abnormal, but in most values it would pass as a normal distribution. 3) Shape-wise, patients current location is mapped as an object, but we did not waste time with asserting this issue as we dropped the column alltogether. The station's revision attribute is also an anomaly, since the shape is inconsistent, sometimes having only the date, sometimes also the time… The residence attribute is empty, having no entries

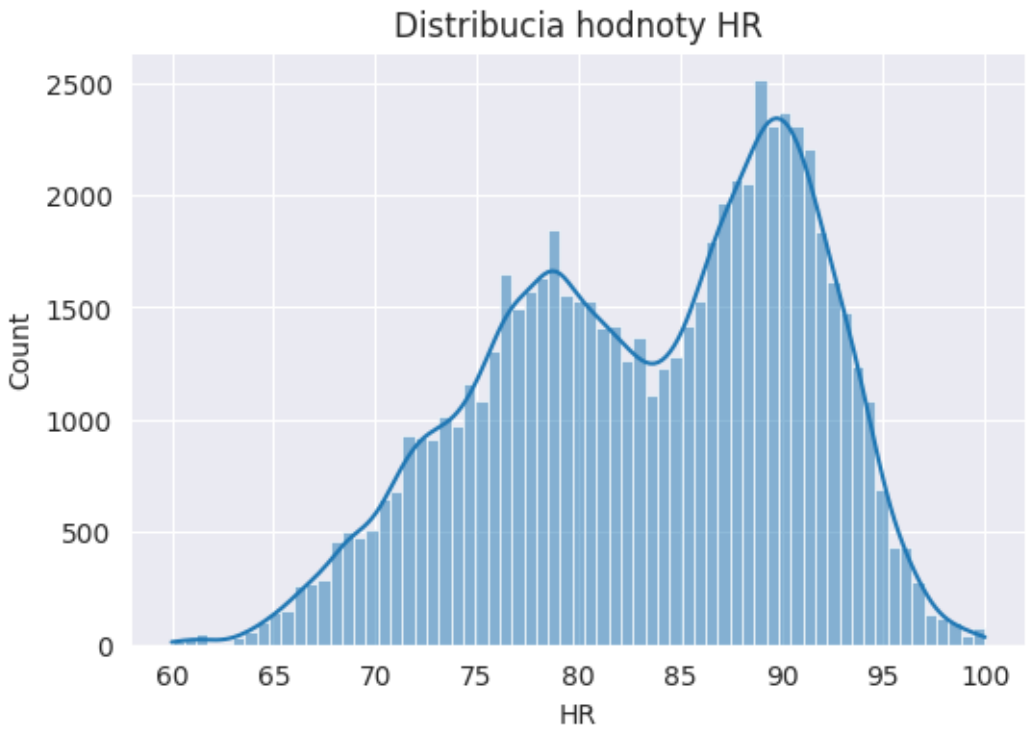
Illogical relationships would also be the division of the data into three tables. What struck me the most was the inability to find a patients observation, due to the fact that after merging patients and stations, we find out that multiple patient are linked to a singular station, and no further link to observation is provided apart than a 1:many relationship between a station and an observation with coordinates.

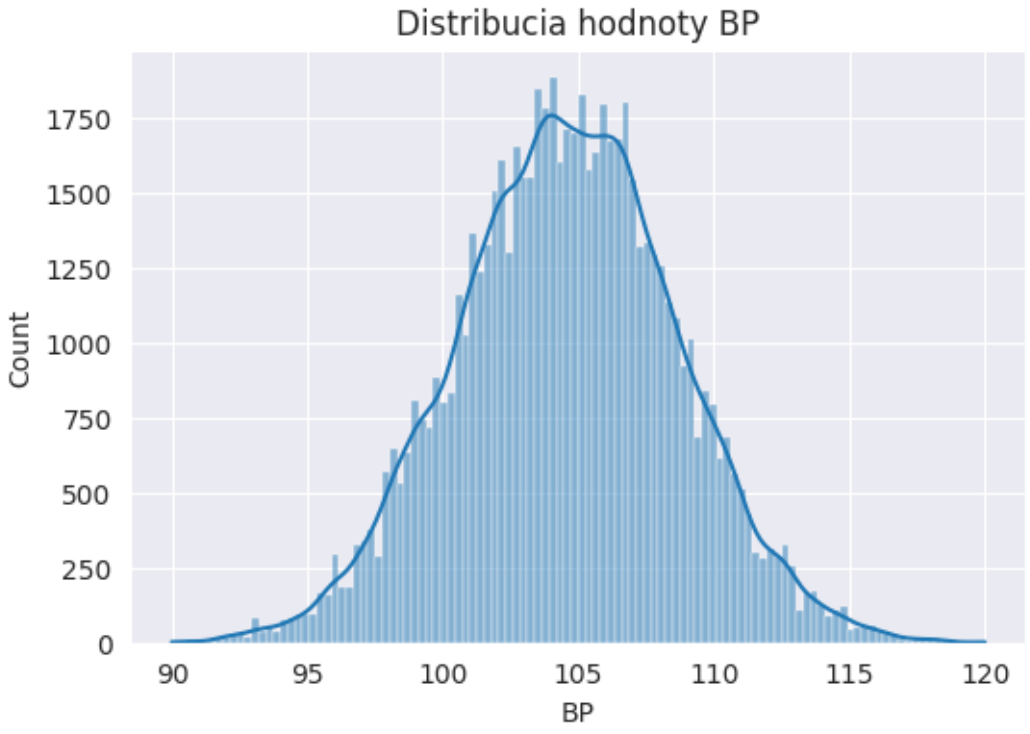
3.2.3 C) Outlier detection

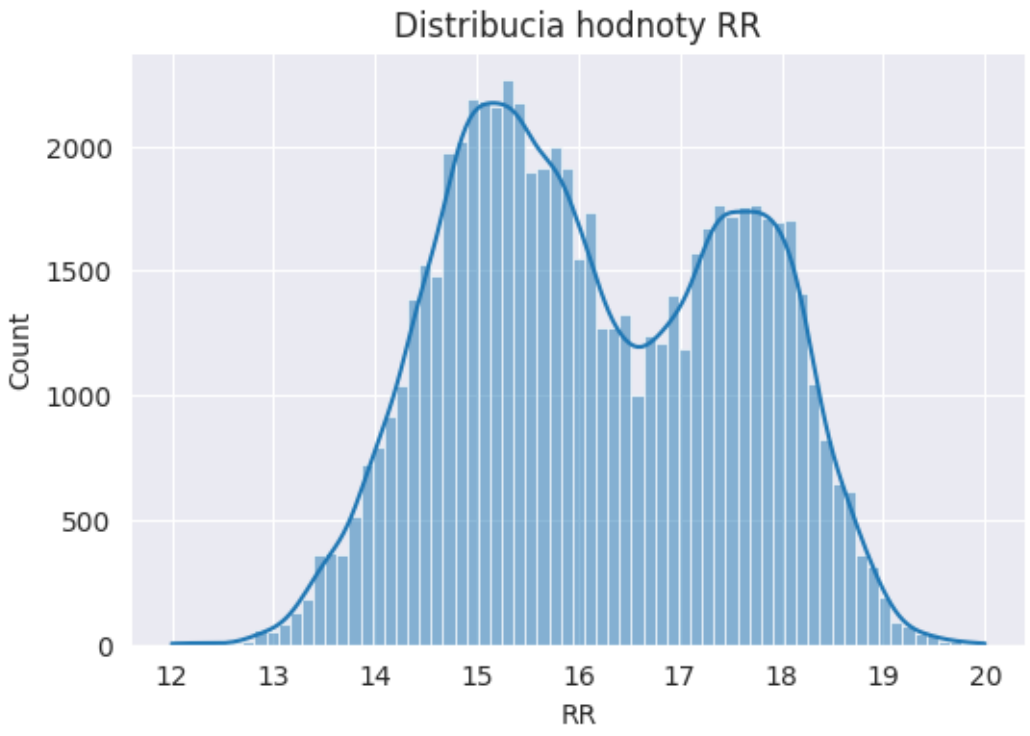
[856]:
```
cols1 = ['HR', 'BP', 'RR', 'SpO2', 'FiO2', 'EtCO2',"CO"]
plt.figure(figsize=(12, 6))
sns.boxplot(data=df[cols1], orient='h', fliersize=3)
plt.title('Boxplot - zakladne vitalne funkcie')
plt.tight_layout()
plt.show()
```

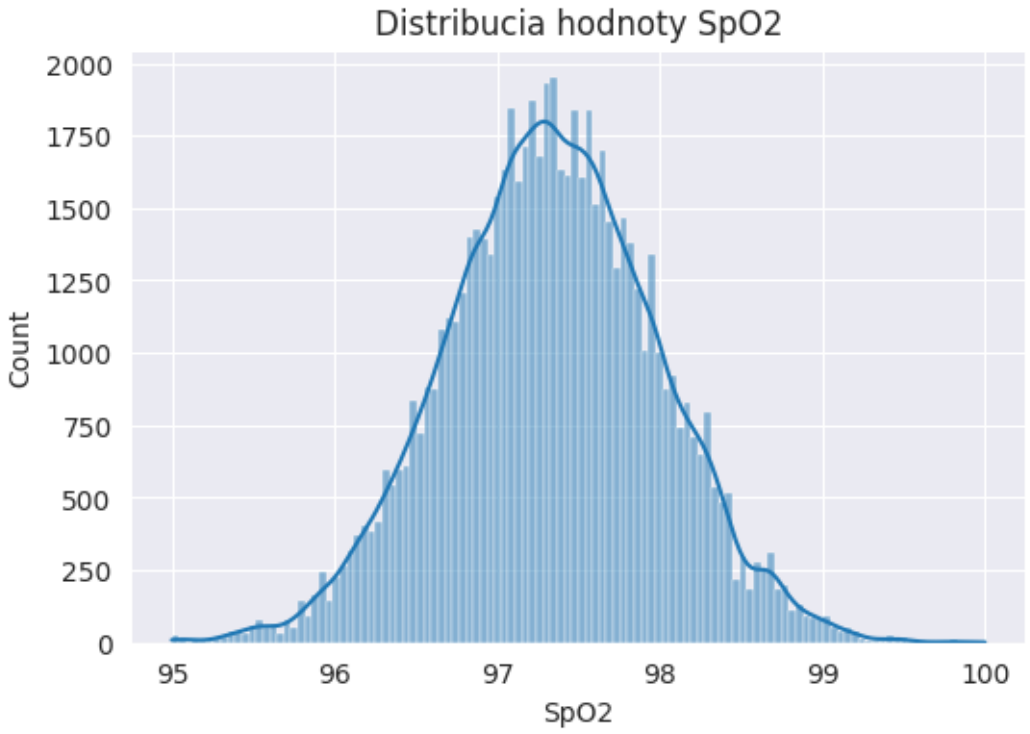


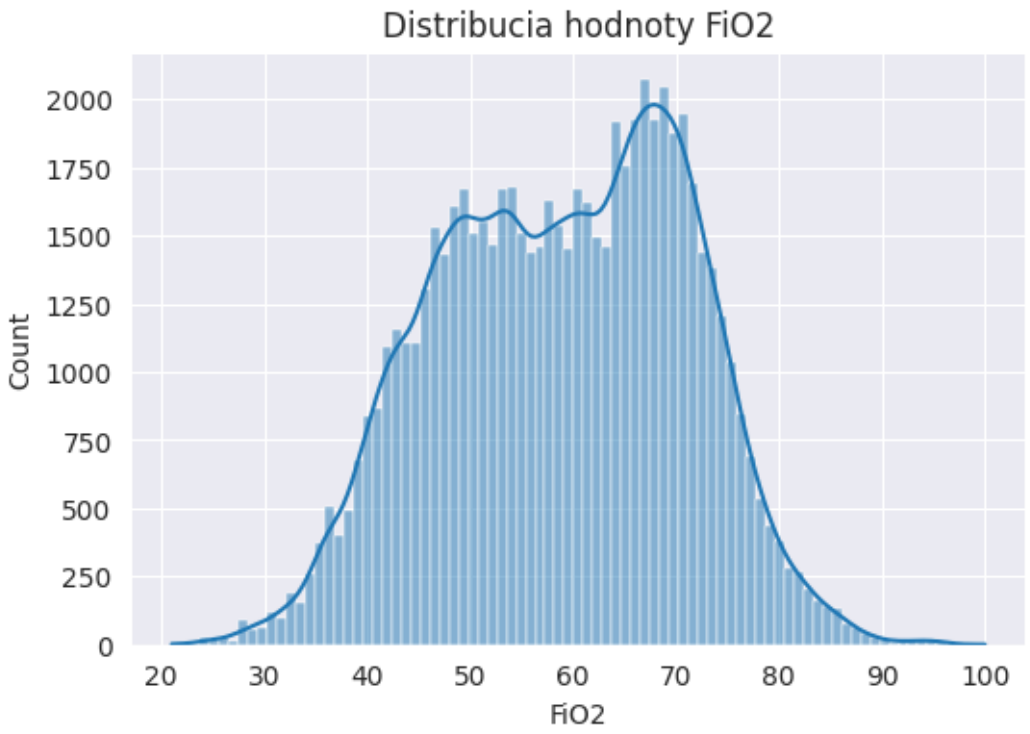
[857]:
```
for col in cols1:
    plt.figure(figsize=(6, 4))
    sns.histplot(df[col], kde=True)
    plt.title(f'Distribucia hodnoty {col}')
    plt.show()
```

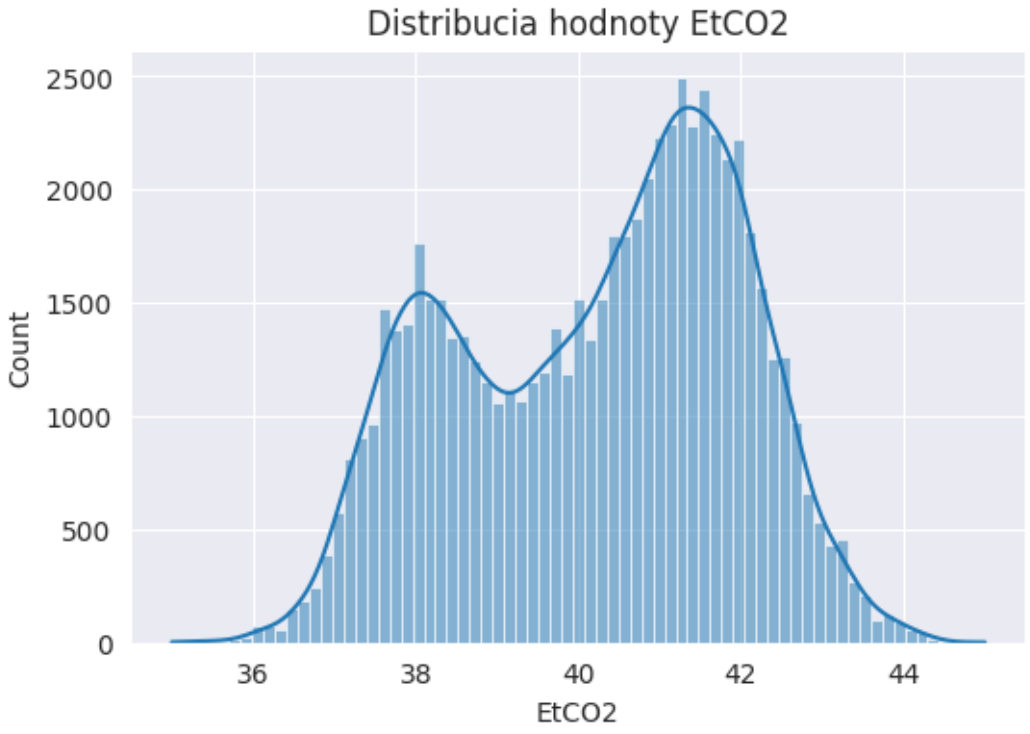

Distribucia hodnoty HR
Count
2500
2000
1500
1000
500
0
60
65
70
75
80
85
90
95
100
HR

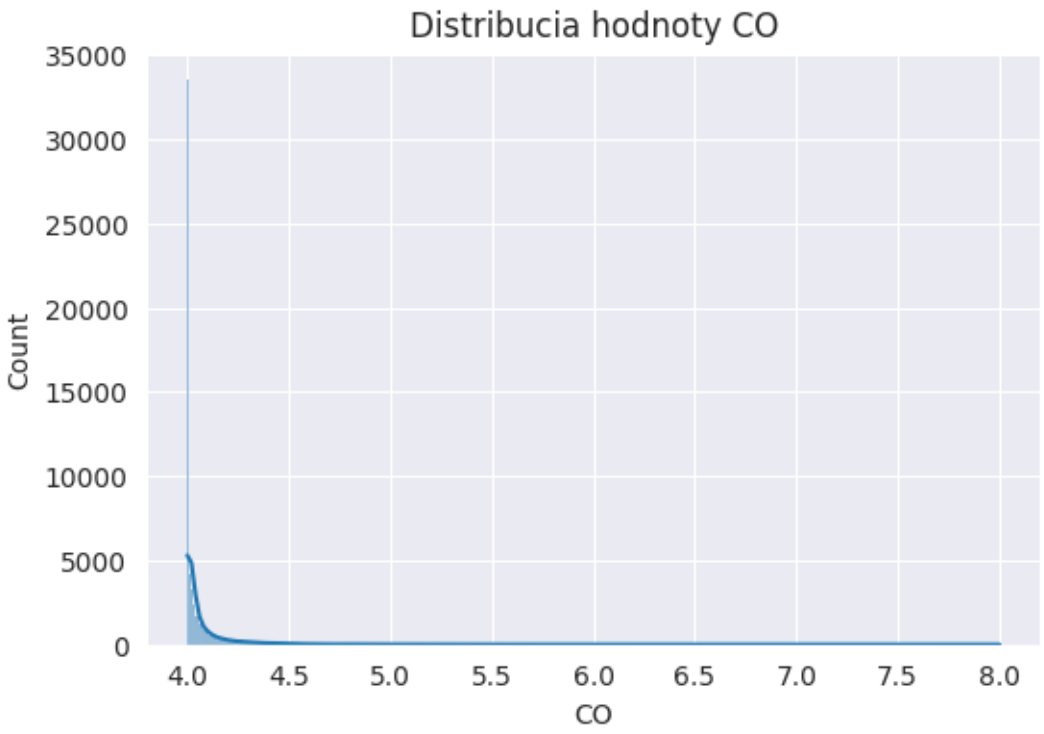

Distribucia hodnoty BP
Count
1750
1500
1250
1000
750
500
250
0
90
95
100
105
110
115
120
BP


Distribucia hodnoty RR
Count
2000
1500
1000
500
0
12
13
14
15
16
17
18
19
20
RR


Distribucia hodnoty SpO2
Count
2000
1750
1500
1250
1000
750
500
250
0
95
96
97
98
99
100
SpO2

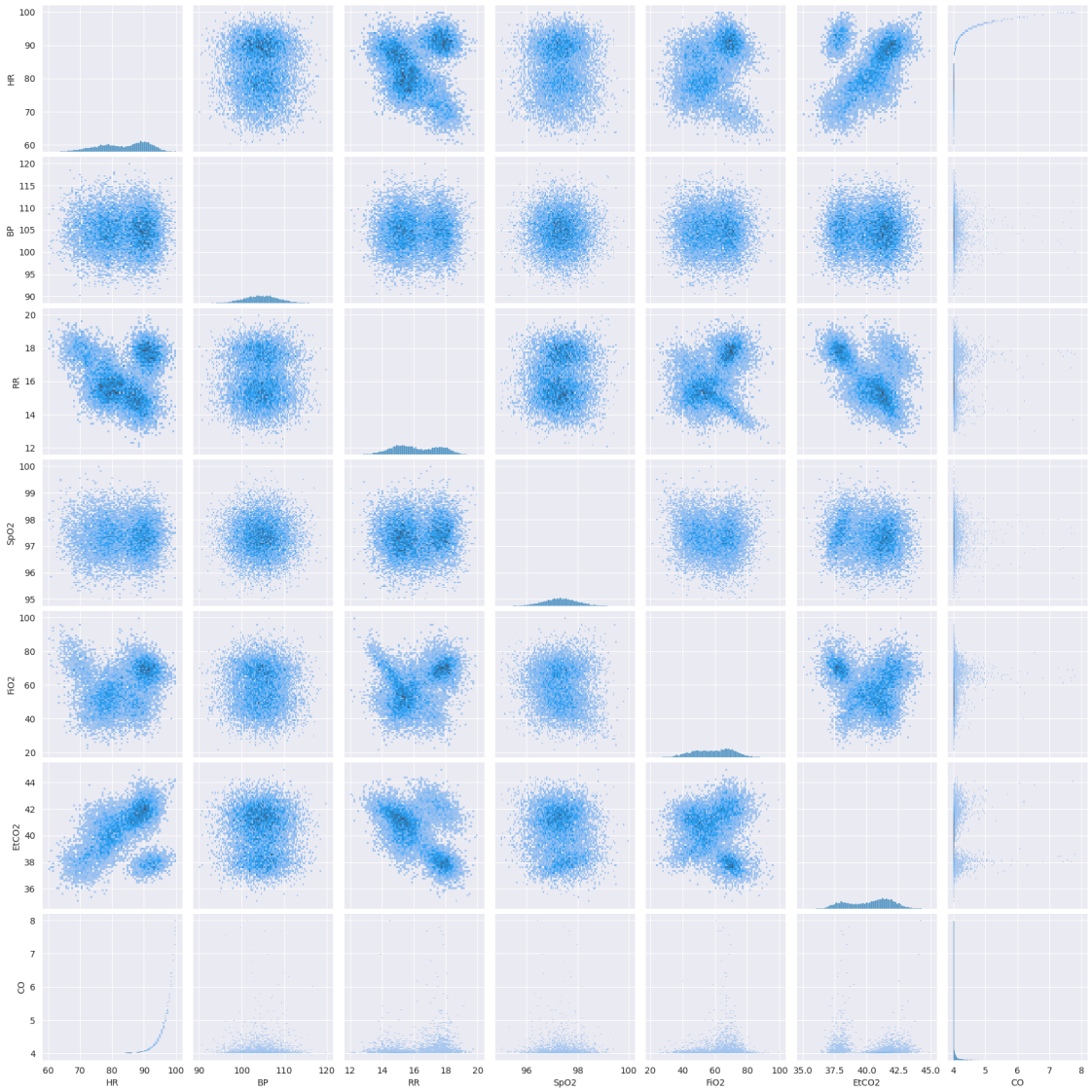

Distribucia hodnoty FiO2
Count
2000
1750
1500
1250
1000
750
500
250
0
20
30
40
50
60
70
80
90
100
FiO2


Distribucia hodnoty EtCO2
Count
2500
2000
1500
1000
500
0
36
38
40
42
44
EtCO2

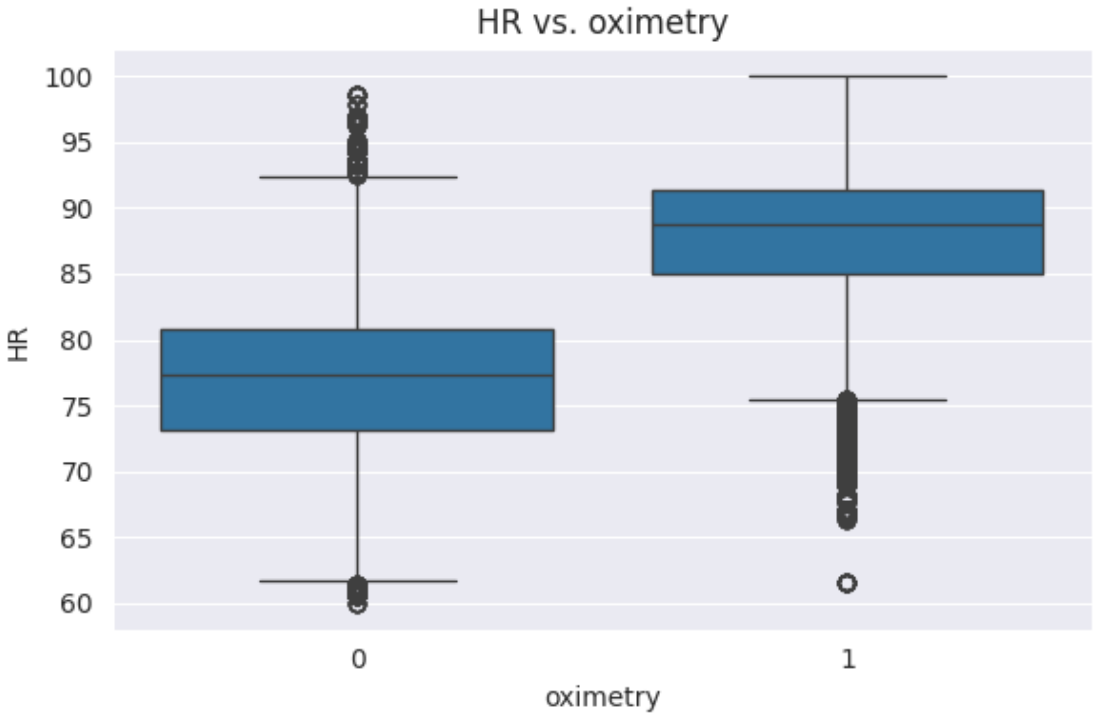


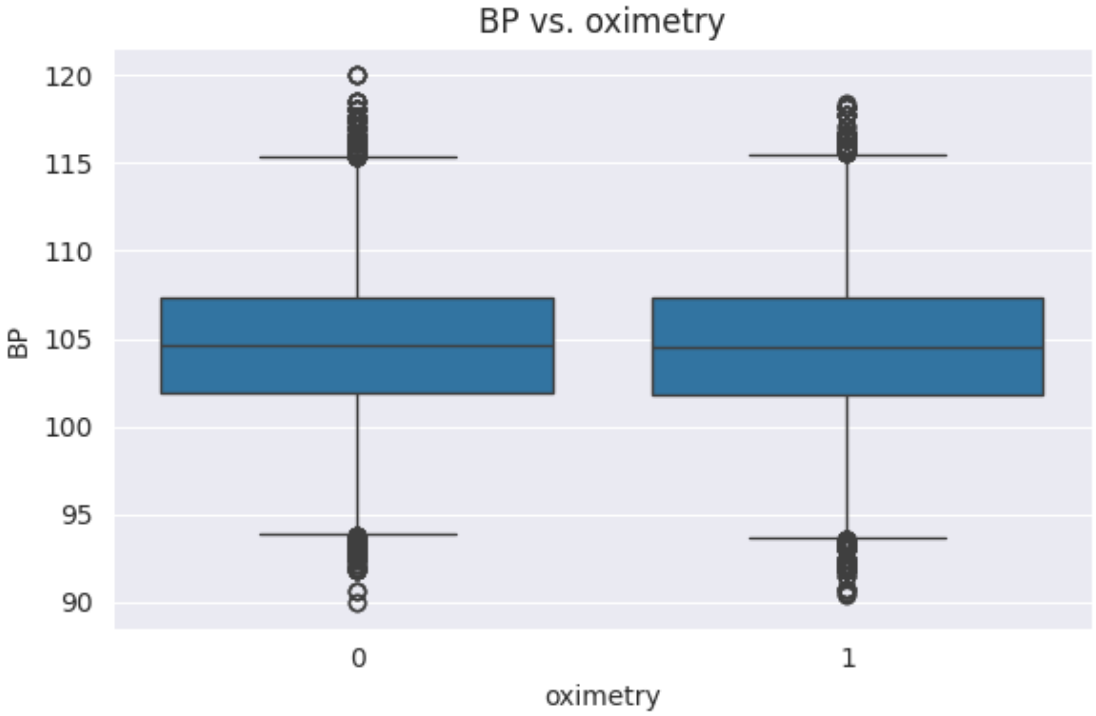
[858]:
```
sns.pairplot(data=df[cols1], kind='hist')
```

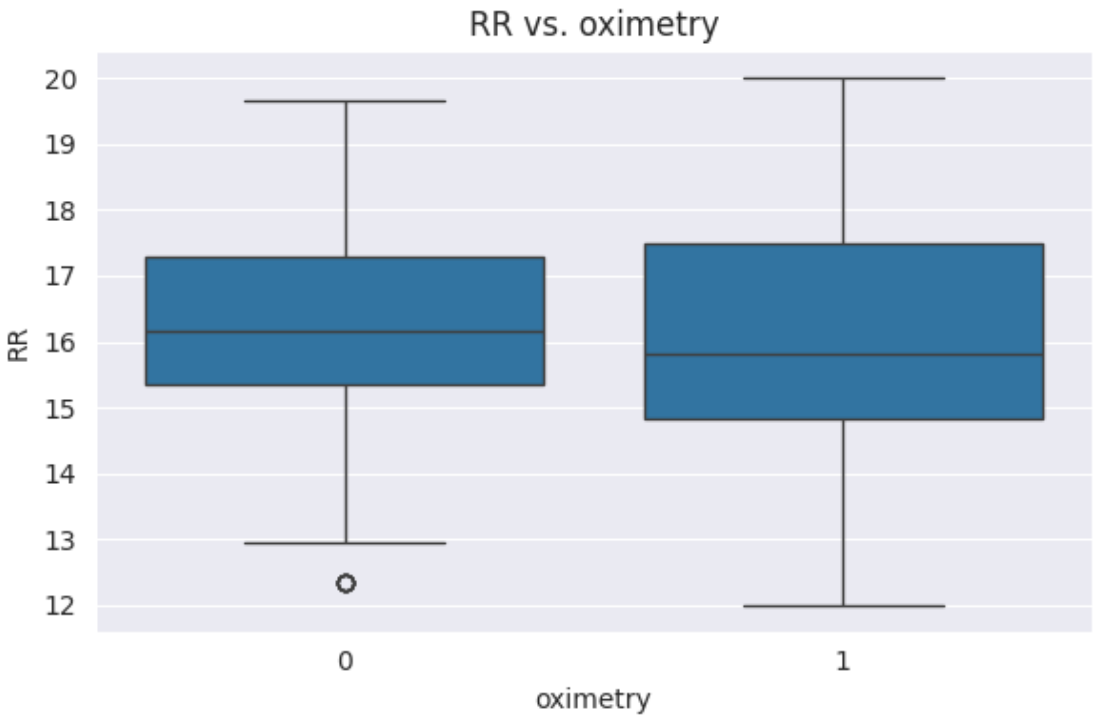
[858]: <seaborn.axisgrid.PairGrid at 0x2fb0f4f96e0>

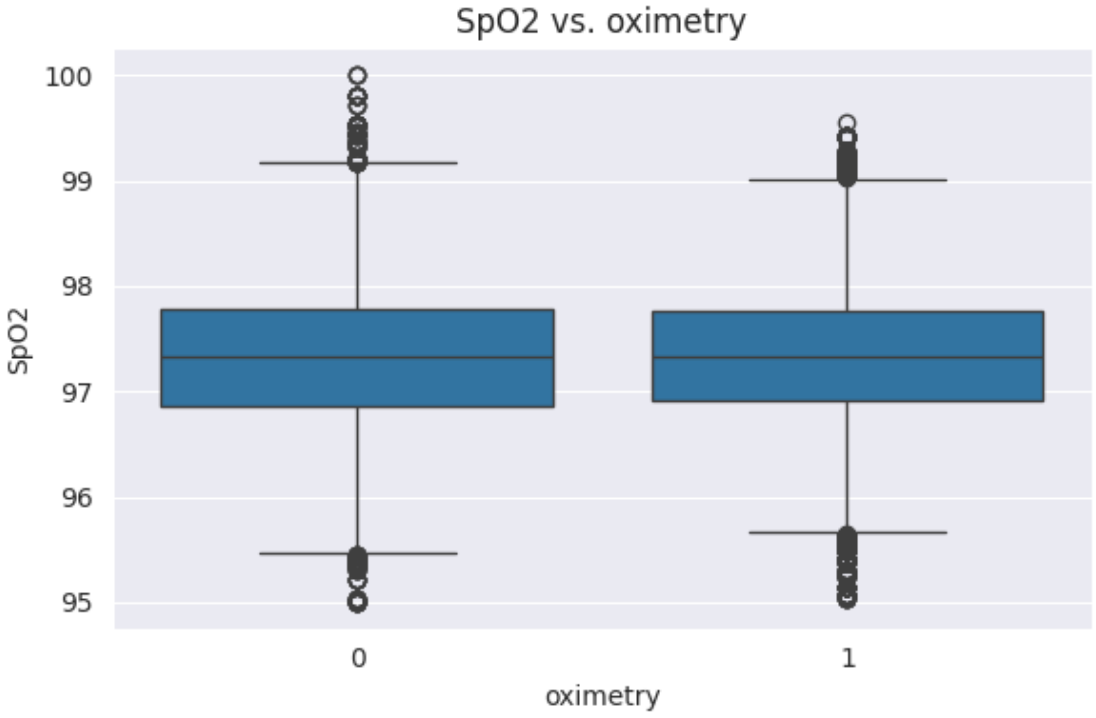


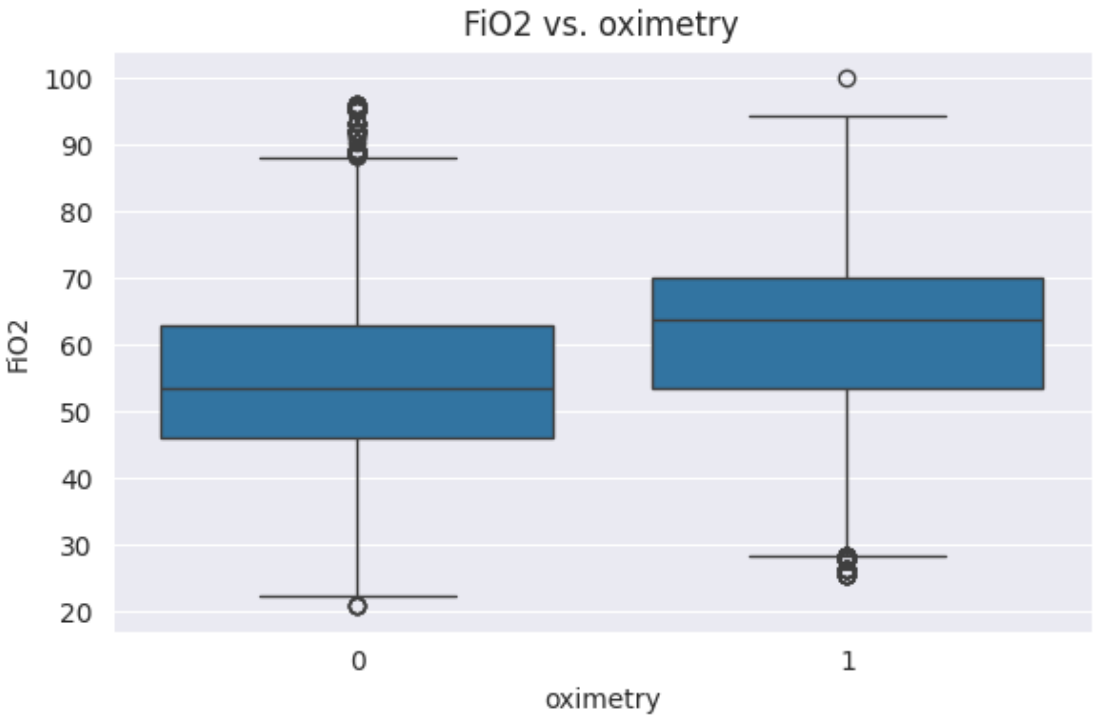
[859]:
```
for col in cols1:
    plt.figure(figsize=(6,4))
    sns.boxplot(data=df, x='oximetry', y=col)
    plt.title(f'{col} vs. oximetry')
    plt.tight_layout()
    plt.show()
```

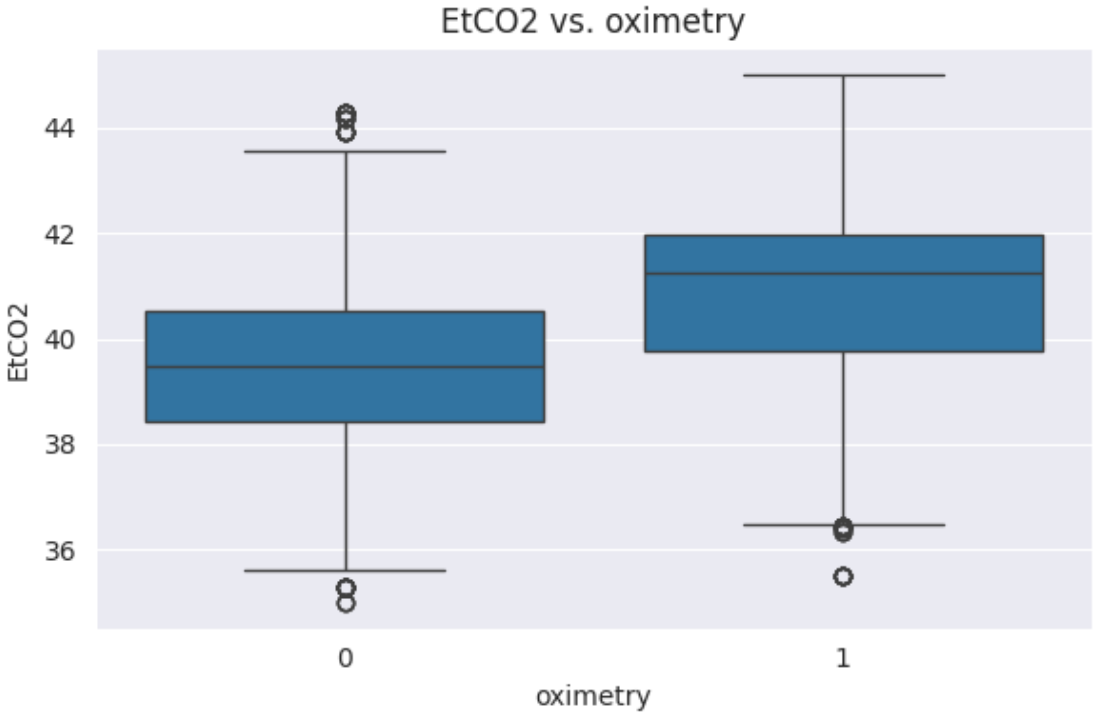

HR vs. oximetry
HR
100
95
90
85
80
75
70
65
60
0
1
oximetry

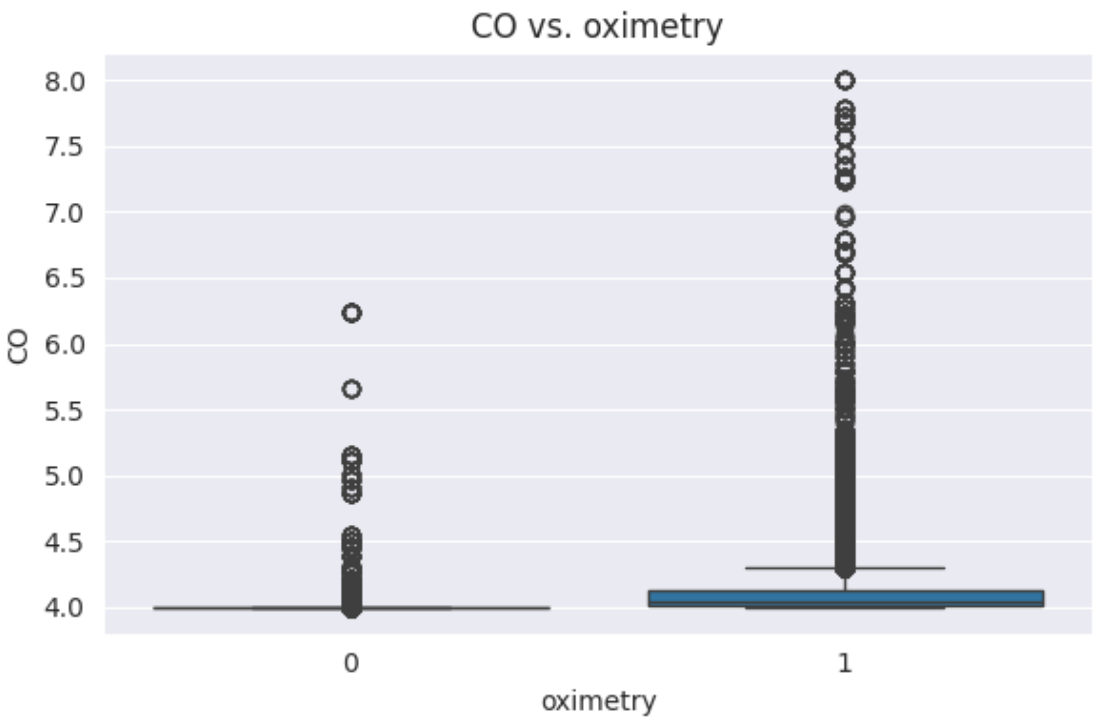

BP vs. oximetry
BP
120
115
110
105
100
95
90
0
1
oximetry


RR vs. oximetry
RR
20
19
18
17
16
15
14
13
12
0
1
oximetry


SpO2 vs. oximetry
SpO2
100
99
98
97
96
95
0
1
oximetry


FiO2 vs. oximetry
FiO2
100
90
80
70
60
50
40
30
20
0
1
oximetry


EtCO2 vs. oximetry
EtCO2
44
42
40
38
36
0
1
oximetry



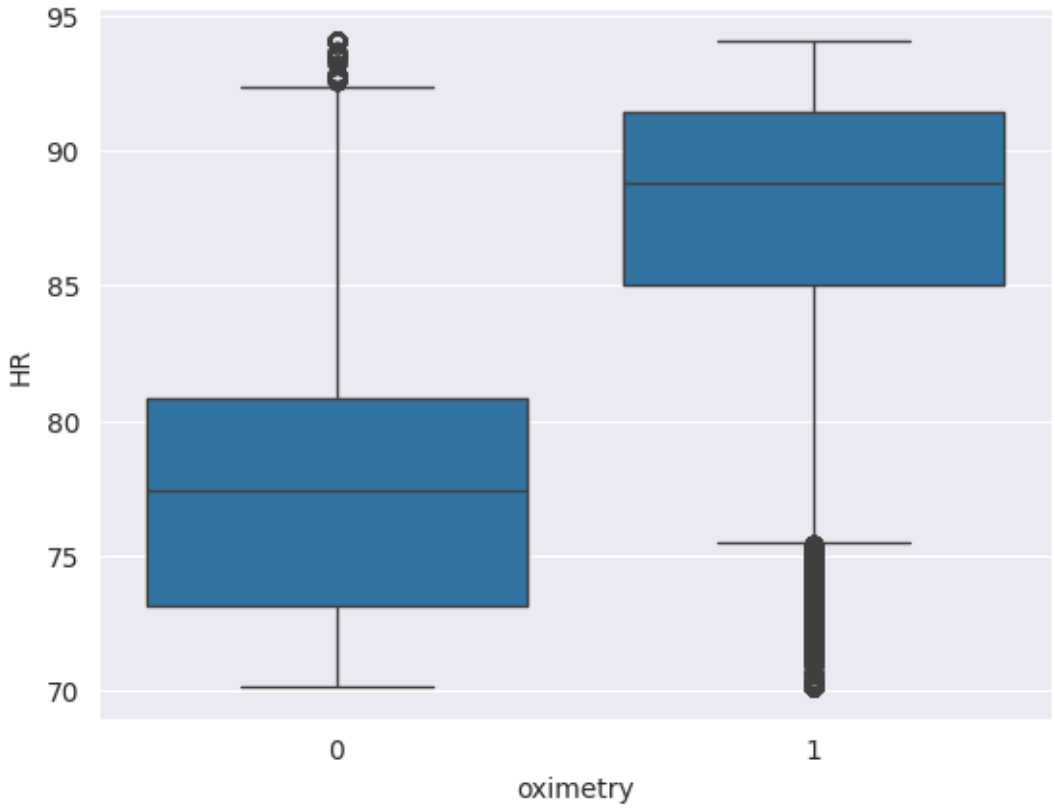
[860]:
```
# HR (Heart Rate) showed several extreme values .Very high for oximetry = 0 and very low for oximetry = 1.
#These outliers can affect model accuracy, so in the next phase we will clean them using the IQR method and winsorization (5th-95th percentile) to keep only realistic heart rate values.

#The CO variable shows a highly right-skewed distribution with a long upper tail, indicating a large number of potential outliers compared to other attributes.
#These extreme values are likely to distort the model, so CO will require normalization or outlier treatment (IQR filtering or winsorization).
```

[861]:
```
# Calculate 5th and 95th percentiles
low, high = df['HR'].quantile([0.05, 0.95])

# before winsorization
before = ((df['HR'] < low) | (df['HR'] > high)).sum()

# apply wins.
df_win = df.copy()
df_win['HR'] = df_win['HR'].clip(lower=low, upper=high)

# after winsorization
```

```
after = ((df_win['HR'] < low) | (df_win['HR'] > high)).sum()

print(f"Before winsorization: {before} outliers")
print(f"After winsorization: {after} outliers")

sns.boxplot(data = df_win, x='oximetry', y='HR')
```

```
Before winsorization: 6696 outliers
After winsorization: 0 outliers
```

[861]: <Axes: xlabel='oximetry', ylabel='HR'>



Now I can see that there is not such a values that are higher or lower to quntile range from 0.05 to 0.95 that's why we did not get any after winsortization value on the other hand we caught some outliers with the IQR method on df["CO"] atribute because he included many abnormal values

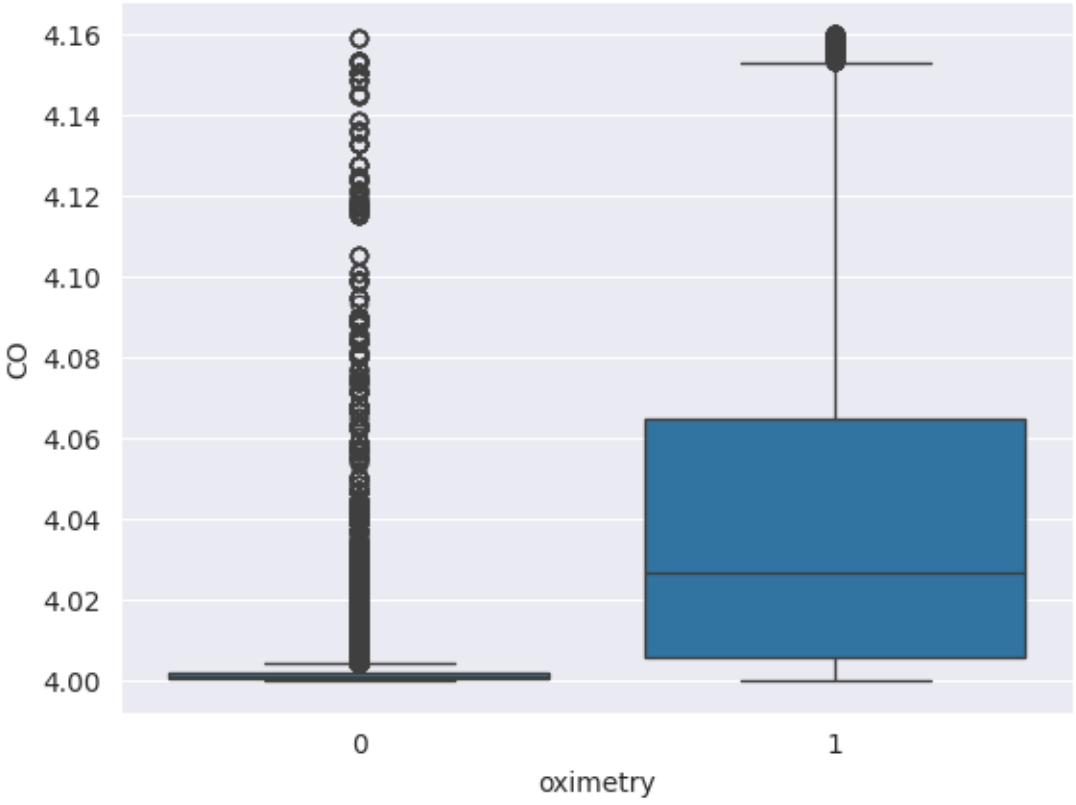
[862]:
```
# IQR bounders
Q1 = df['CO'].quantile(0.25)
Q3 = df['CO'].quantile(0.75)
IQR = Q3 - Q1

lower_bound = Q1 - 1.5 * IQR
```

```
upper_bound = Q3 + 1.5 * IQR

print(f"CO bounds (IQR): {lower_bound:.2f} - {upper_bound:.2f}")

# delete values out of range
df_iqr = df[(df['CO'] >= lower_bound) & (df['CO'] <= upper_bound)]

print(f"Removed {len(df) - len(df_iqr)} rows")
print(len(df_iqr))
```

```
CO bounds (IQR): 3.91 - 4.16
Removed 8460 rows
58513
```

[863]:
```
sns.boxplot(data=df_iqr, x='oximetry', y='CO')
```

[863]: <Axes: xlabel='oximetry', ylabel='CO'>



On the graph above we can see extreme difference on outlier values because the previous graph had values close to 9 and this has maximum values 4.16

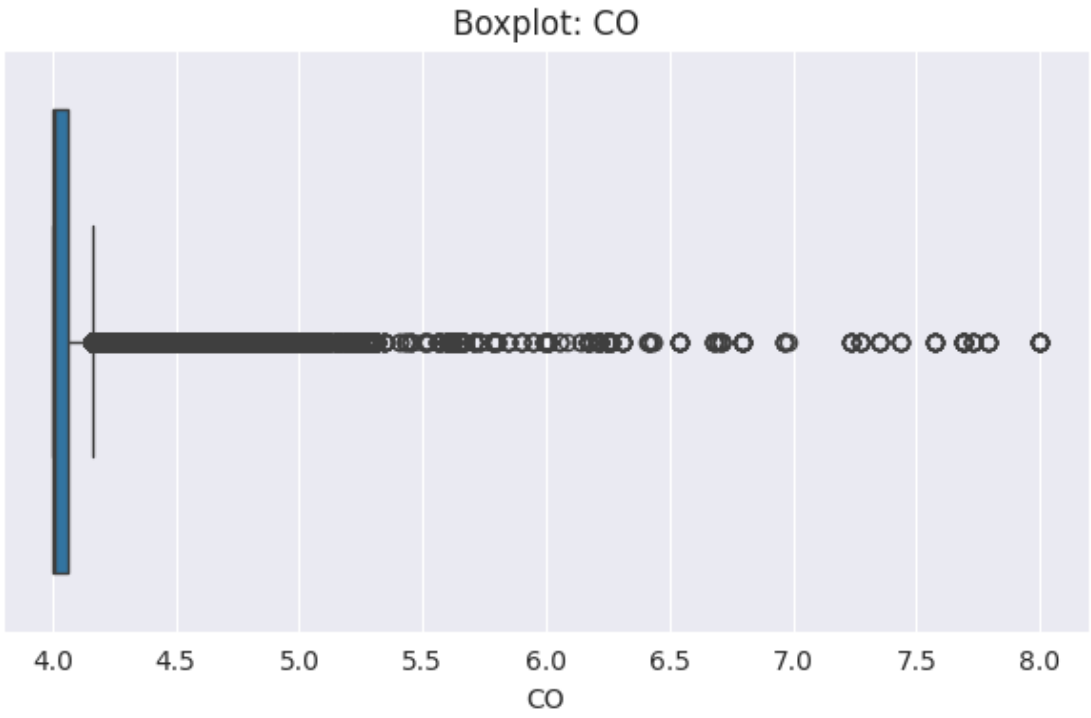
[864]:
```
# all number collumns exept oximetry
num_cols = df.select_dtypes(include='number').columns.drop('oximetry',␣
 ↪errors='ignore')

#add collumns if in range of upper or lower bounder
iqr_summary = []
for col in num_cols:
    q1, q3 = df[col].quantile([0.25, 0.75])
    iqr = q3 - q1
    lower, upper = q1 - 1.5 * iqr, q3 + 1.5 * iqr
    outliers = ((df[col] < lower) | (df[col] > upper)).sum()
    iqr_summary.append([col, outliers, round(outliers / len(df) * 100, 2)])

# Display as a small DataFrame
iqr_df = pd.DataFrame(iqr_summary, columns=['Column', 'Outliers', 'Percent'])
iqr_df.sort_values('Percent', ascending=False, inplace=True)
iqr_df
```

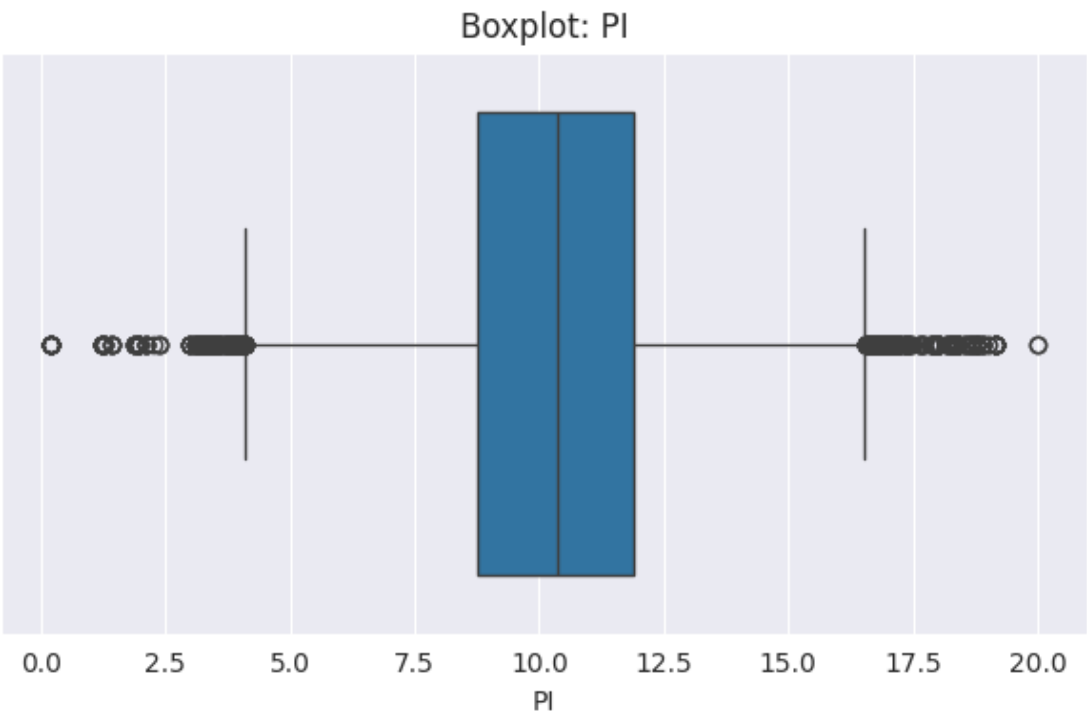
[864]:

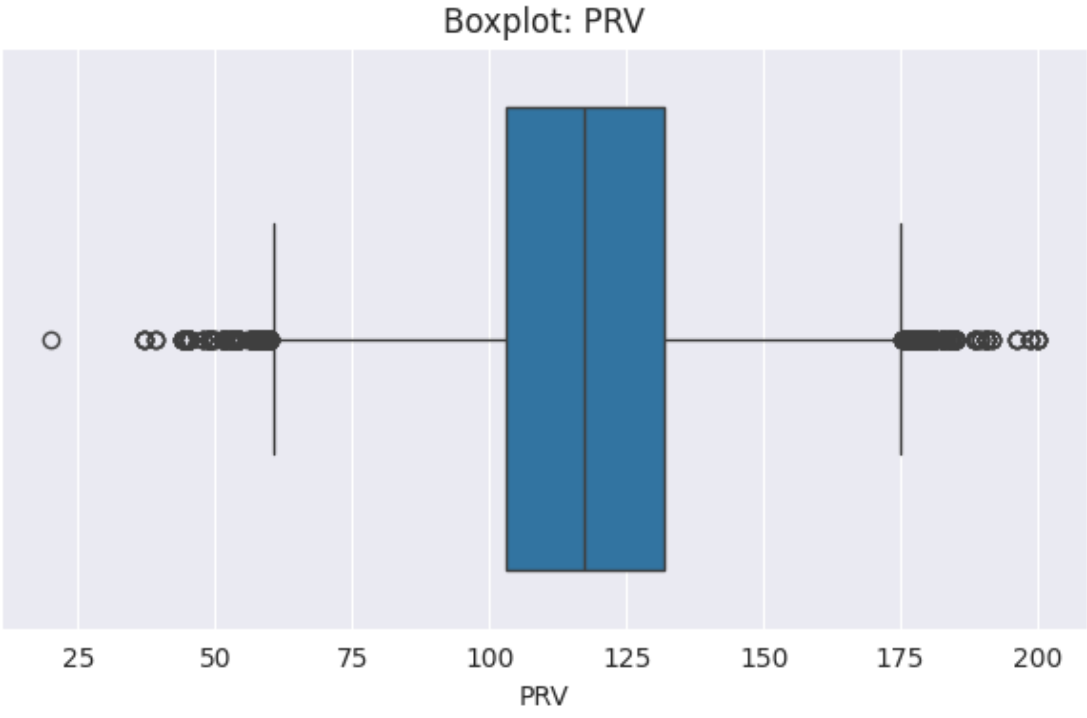
	Column	Outliers	Percent
13	CO	8460	12.63
2	PI	785	1.17
6	PRV	633	0.95
20	latitude	627	0.94
14	Blood Flow Index	602	0.90
0	SpO2	587	0.88
7	BP	560	0.84
15	PPG waveform features	549	0.82
21	longitude	544	0.81
17	Respiratory effort	468	0.70
9	Motion/Activity index	472	0.70
12	SV	439	0.66
8	Skin Temperature	432	0.65
11	Hb level	352	0.53
16	Signal Quality Index	351	0.52
10	PVI	344	0.51
5	FiO2	5	0.01
4	EtCO2	0	0.00
1	HR	0	0.00
3	RR	0	0.00
19	SNR	0	0.00
18	O2 extraction ratio	0	0.00
22	station_ID	0	0.00
23	user_id	0	0.00

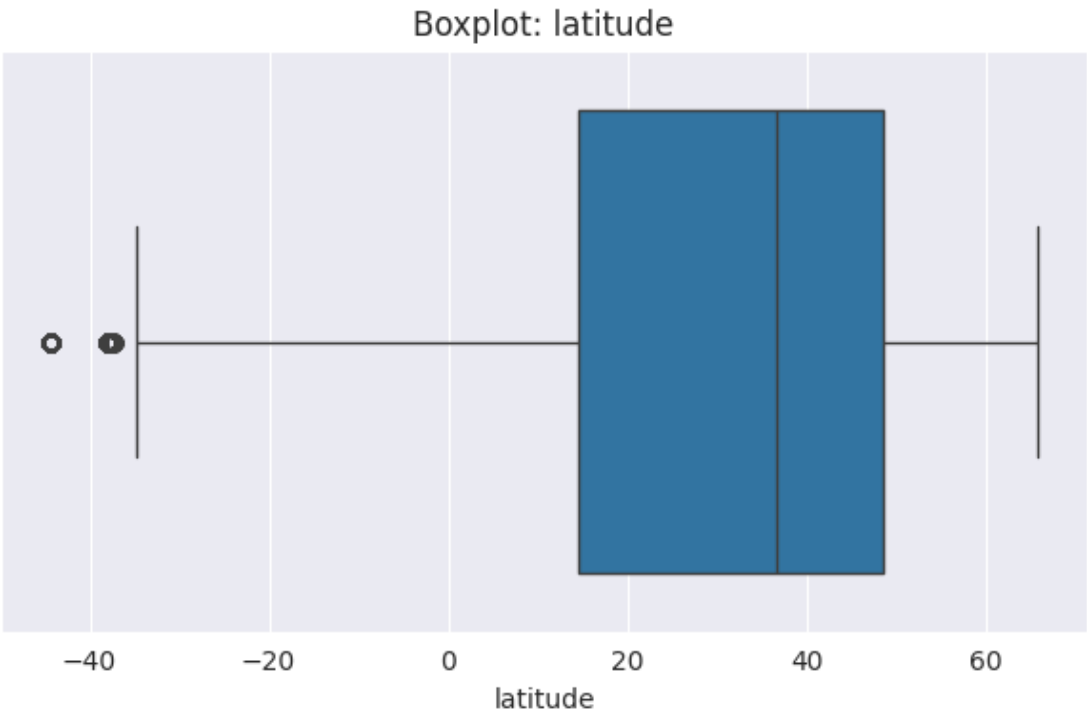
It's bad practice to analyze a atribute one by one so I created a list of numeric values that I can analyze at once. I calculated they're IQR based on dividing they're quantile (0.25 and 0.75) .If any value is less than lower value (<0.25) that it is added to list of outliers this process also include calculating if value is greater that upper bounder (>0.75).

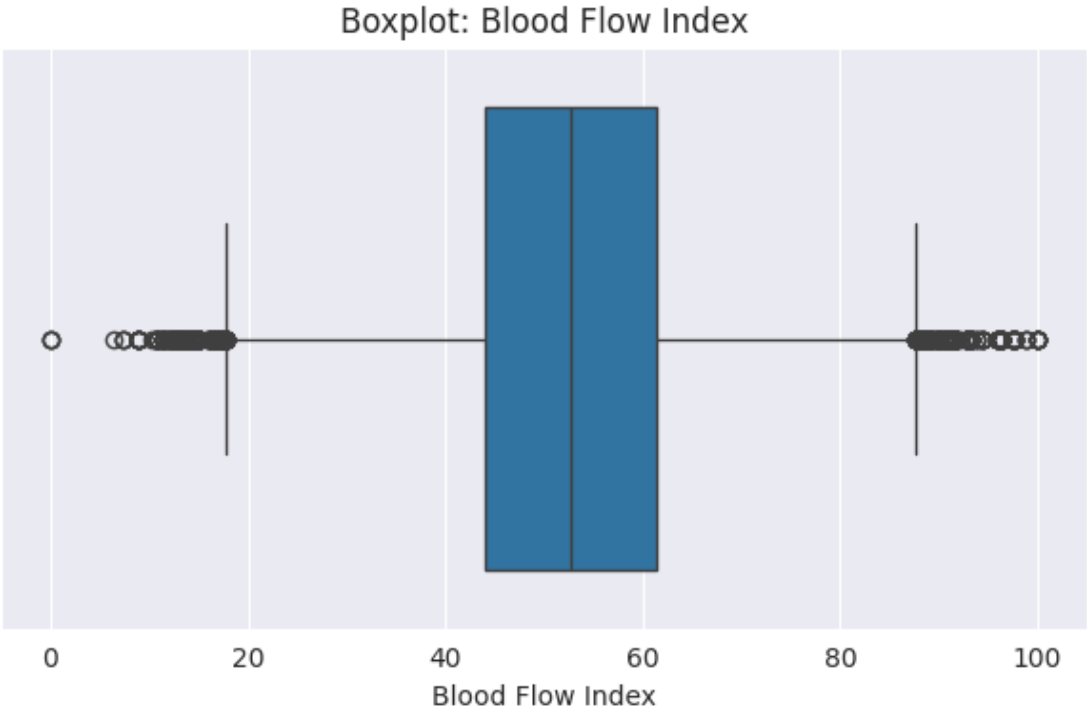
[865]:
```
suspects = iqr_df[ iqr_df['Outliers'] > 0 ]['Column'].head(6).tolist()

for col in suspects:
    plt.figure(figsize=(6,4))
    sns.boxplot(x=df[col])
    plt.title(f'Boxplot: {col}')
    plt.tight_layout()
    plt.show()
```

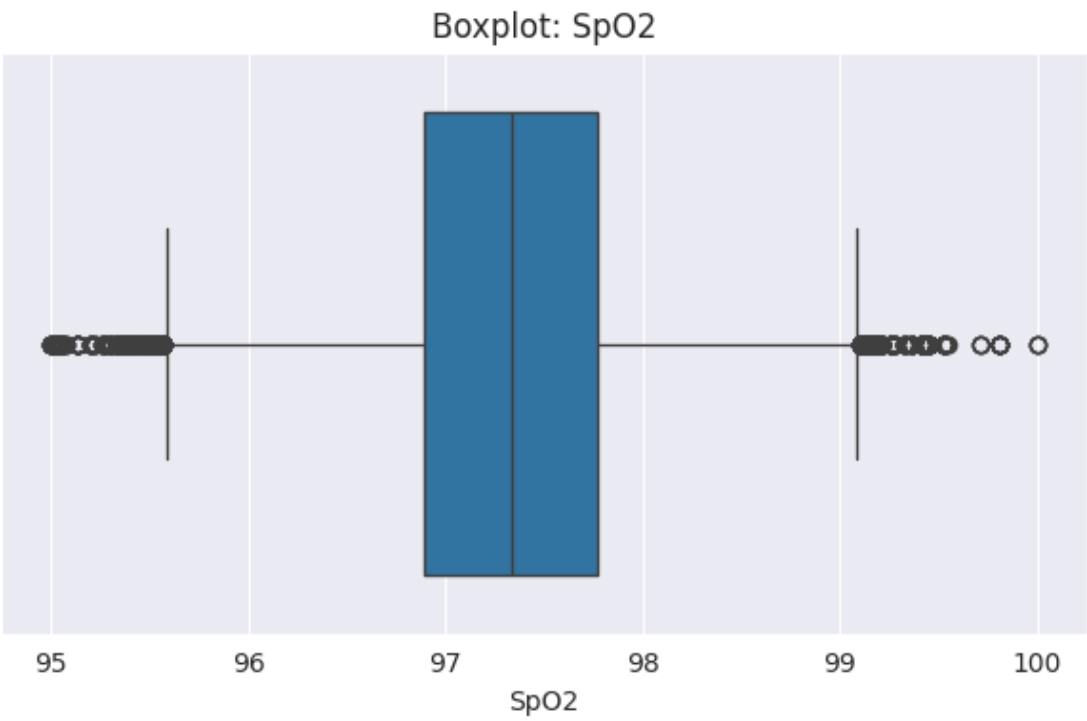



Boxplot: PI
0.0
2.5
5.0
7.5
10.0
12.5
15.0
17.5
20.0
PI


Boxplot: PRV
25
50
75
100
125
150
175
200
PRV


Boxplot: latitude
−40
−20
0
20
40
60
latitude


Boxplot: Blood Flow Index
0
20
40
60
80
100
Blood Flow Index



3.3 1.3 Formulácia a štatistické overenie hypotéz o dátach

3.3.1 Zle hypotezy

Hypotéza 1 – rozdiel v tepovej frekvencii podľa stavu saturácie

H (Null Hypothesis): The mean heart rate (HR) is the same for both groups — patients with oximetry = 0 and oximetry = 1.

H (Alternative Hypothesis): The mean heart rate (HR) differs between the groups — patients with oximetry = 0 and oximetry = 1.

[866]:
```
const_shapiro = 5000
# I will create 2 datasets hro0 (HE and oximtery = 0) and hro1 (HE column and␣
 ↪oximetry = 1)
hr0 = df[df['oximetry'] == 0]['HR']
hr1 = df[df['oximetry'] == 1]['HR']


sample_hr0 = hr0.sample(const_shapiro, random_state=42)
sample_hr1 = hr1.sample(const_shapiro, random_state=42)

print(shapiro(sample_hr0))
print(shapiro(sample_hr1))
```

```
ShapiroResult(statistic=np.float64(0.9964622127654921),
pvalue=np.float64(1.50860565376863e-09))
ShapiroResult(statistic=np.float64(0.9381229280127523),
pvalue=np.float64(1.7803348598149635e-41))
```

[867]:
```
#we can see that the pvalues from both samples are lower than 0.05 which␣
 ↪indicates to not normal distributed data
#based on that we need to perform Mann-whitney U test not basic T-test that␣
 ↪need normal distribution

stat, p = stats.mannwhitneyu(sample_hr0, sample_hr1)
print(f"Mann-Whitney U-test p-value: {p:.3e}")

# the value is super close to 0 that means that we denied H0 => there is a␣
 ↪statistically significant difference
# in heart rate (HR) between the groups (oximetry = 0 and oximetry = 1).
```

```
Mann-Whitney U-test p-value: 0.000e+00
```

Hypothesis 2 – Difference in EtCO based on oxygen saturation (oximetry)

H (Null Hypothesis): The mean EtCO level is the same for both groups — patients with oximetry = 0 and oximetry = 1.

H (Alternative Hypothesis): The mean EtCO level differs between the groups — patients with oximetry = 0 and oximetry = 1.

[868]:
```
df.columns
```

[868]:
```
Index(['SpO2', 'HR', 'PI', 'RR', 'EtCO2', 'FiO2', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
       'Signal Quality Index', 'Respiratory effort', 'O2 extraction ratio',
       'SNR', 'oximetry', 'latitude', 'longitude', 'station_ID', 'QoS', 'code',
       'user_id', 'blood_group'],
      dtype='object')
```

[869]:
```
etco0 = df[df['oximetry'] == 0]['EtCO2']
etco1 = df[df['oximetry'] == 1]['EtCO2']

#sample for max 5000
sample_etco0 = etco0.sample(5000, random_state=42)
sample_etco1 = etco1.sample(5000, random_state=42)

# normality check
print(shapiro(sample_etco0))
print(shapiro(sample_etco1))
```

```
# if both p > 0.05 → data are normally distributed
# then perform Student's T-test
stat, p = stats.mannwhitneyu(etco0, etco1)
print(f"U-test p-value: {p:.3e}")
```

```
ShapiroResult(statistic=np.float64(0.9938690077488066),
pvalue=np.float64(8.316180710196192e-14))
ShapiroResult(statistic=np.float64(0.9191799162237433),
pvalue=np.float64(1.0762260080747649e-45))
U-test p-value: 0.000e+00
```

there is a statistically significant difference in mean EtCO between patients with oximetry = 0 and oximetry = 1.

3.3.2 A) Hypothesis 1 – Difference in SPO based on oxygen saturation (oximetry)

H (Null Hypothesis): The mean SPO level is the same for both groups — patients with oximetry = 0 and oximetry = 1.

H (Alternative Hypothesis): The mean SPO level differs between the groups — patients with oximetry = 0 and oximetry = 1.

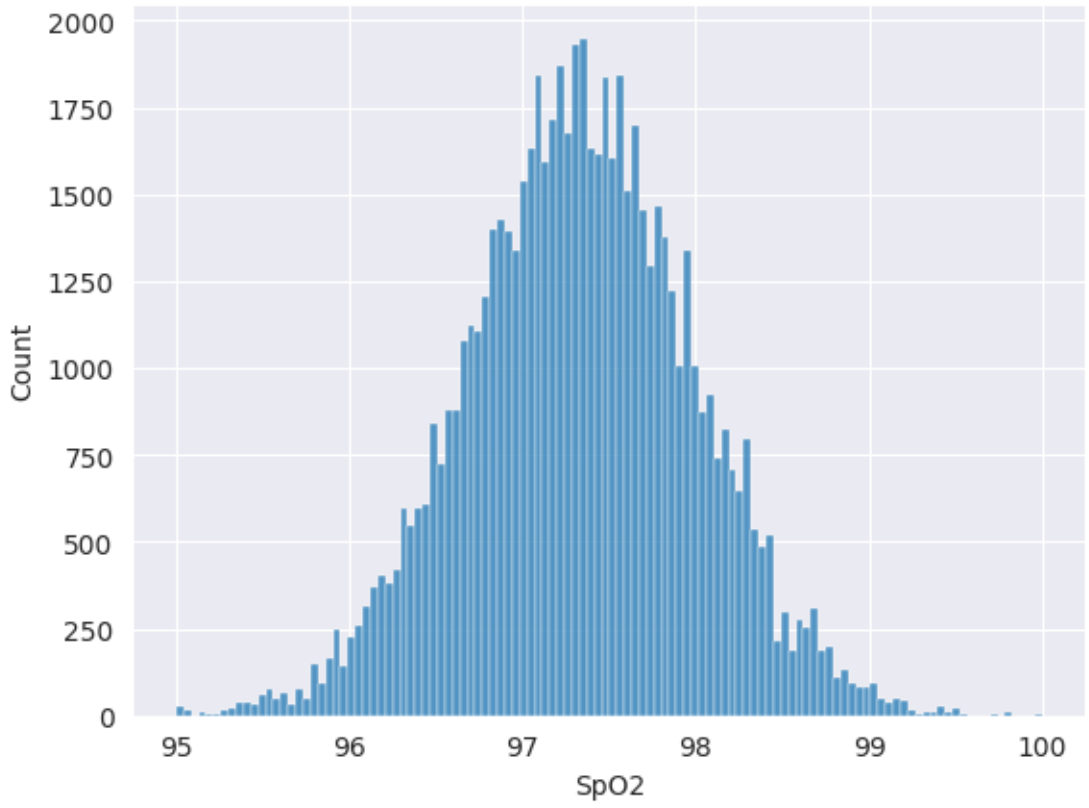
[870]:
```
#division of the data into oximetry == 1 and oximetry == 0 groups
g1 = df[df['oximetry'] == 1]['SpO2']
g0 = df[df['oximetry'] == 0]['SpO2']

#doing the shapiro test -> pvalue for both values
p1 = shapiro(g1).pvalue
p0 = shapiro(g0).pvalue
print('shapiro:', p1, p0)
# even if the p values do not indicate normal division of the data, we will␣
 ↪continue with levenne and ttest since from the histogram we can see that the␣
 ↪data is indeed rather normally distributed, this error is probably caused by␣
 ↪the table creation
```

```
shapiro: 9.472557968212563e-07 0.0018560650823116259
```

```
C:\Users\mudry\AppData\Local\Programs\Python\Python313\Lib\site-
packages\scipy\stats\_axis_nan_policy.py:579: UserWarning: scipy.stats.shapiro:
For N > 5000, computed p-value may not be accurate. Current N is 40029.
  res = hypotest_fun_out(*samples, **kwds)
C:\Users\mudry\AppData\Local\Programs\Python\Python313\Lib\site-
packages\scipy\stats\_axis_nan_policy.py:579: UserWarning: scipy.stats.shapiro:
For N > 5000, computed p-value may not be accurate. Current N is 26944.
  res = hypotest_fun_out(*samples, **kwds)
```

[871]:
```
sns.histplot(df['SpO2'])
```

[871]:
```
<Axes: xlabel='SpO2', ylabel='Count'>
```

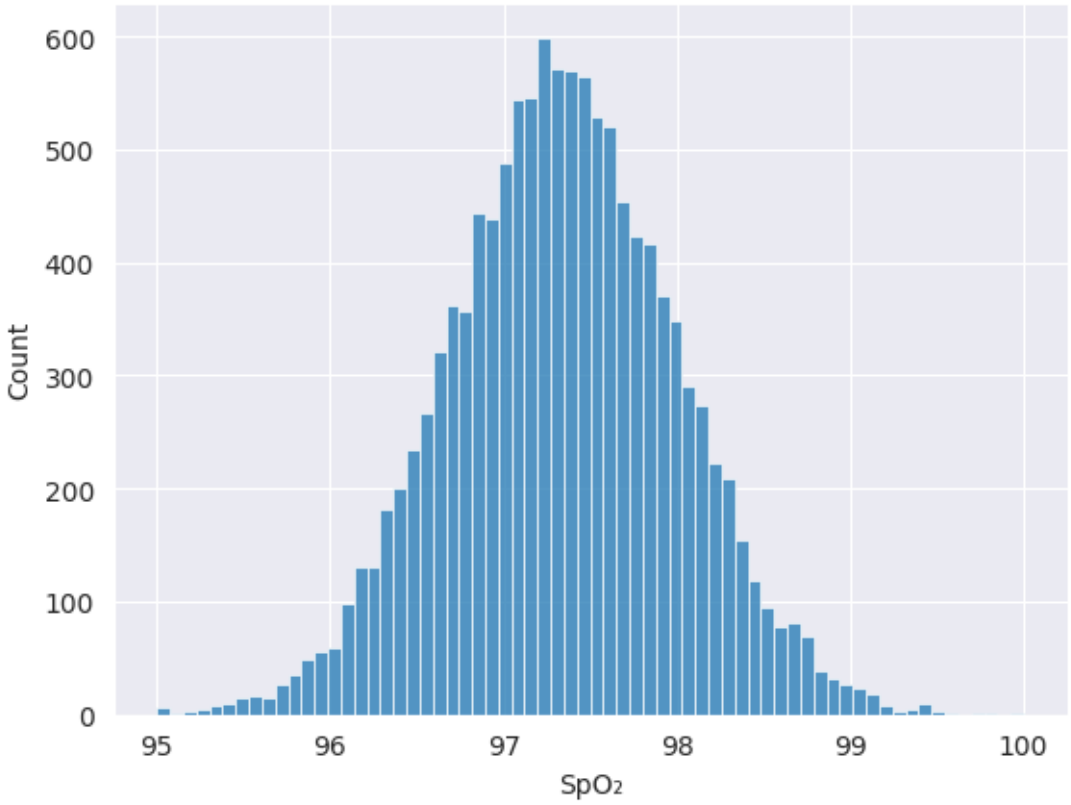


[872]:
```
#due to the findings in the previous cell, we found out the merging of the␣
 ↪tables reshaped the distribution of the data into not so normally␣
 ↪distributed, this however could be fixed by performing the tests on the␣
 ↪observation_df (non merged table).
g1 = observation_df[observation_df['oximetry'] == 1]['SpO ']
g0 = observation_df[observation_df['oximetry'] == 0]['SpO ']

#we need to set sample to max 5000 so the data are in range of the recommended␣
 ↪sample value
sample_g1 = g1.sample(3000, random_state=42)
sample_g0= g0.sample(3000, random_state=42)

#doing the shapiro test -> pvalue for both values
p1 = shapiro(sample_g1).pvalue
p0 = shapiro(sample_g0).pvalue
print('shapiro:', p1, p0)
#after performing the shapiro test we can see that the p values from both the␣
 ↪grous are larger than 0.05 which proves that the normal distribution is␣
 ↪present which leads us to perform the levene and t test
```

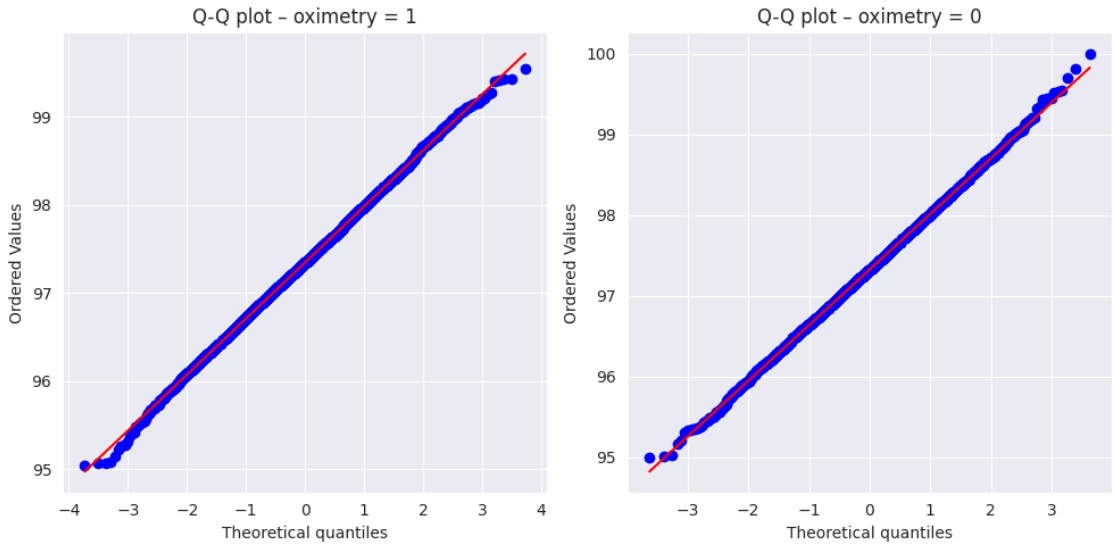
```
shapiro: 0.10880973933264598 0.6169824476877354
```

[873]:
```
sns.histplot(observation_df['SpO '])
```

[873]:
```
<Axes: xlabel='SpO ', ylabel='Count'>
```



[874]:
```
#lets verify the distributions of both the subsets via qqplot
fig, ax = plt.subplots(1, 2, figsize=(10, 5))

stats.probplot(g1, dist="norm", plot=ax[0])
ax[0].set_title("Q-Q plot - oximetry = 1")

stats.probplot(g0, dist="norm", plot=ax[1])
ax[1].set_title("Q-Q plot - oximetry = 0")

plt.tight_layout()
plt.show()
#they indeed are
```



[875]:
```
#levenne tests, this whole block is executed since the data are normally␣
 ↪distributed
p_lev = stats.levene(sample_g1,sample_g0).pvalue
print('levene:', p_lev)

if p_lev > 0.05:
    # since the variance is the same we can perfrom the usual t-test
    t, p = stats.ttest_ind(sample_g1, sample_g0)
    print('t-test', t, p)
else:
    # variances differ, use Welch's t-test - we know this wasnt in our class␣
 ↪notebooks, but we needed to find out if the two groups differ even though␣
 ↪they have different variance, so we found this method
    t, p = stats.ttest_ind(sample_g1, sample_g0, equal_var=False)
    print('Welch t-test (unequal variances):', t, p)
```

```
levene: 1.253188761462382e-05
Welch t-test (unequal variances): 0.09803193034222664 0.9219102349142181
```

[876]:
```
# Both groups are normally distributed (Shapiro > 0.05), but their variances␣
 ↪differ (Levene < 0.05). The Mann-Whitney U test (p = 0.0984 > 0.05) shows no␣
 ↪statistically significant difference in SpO  levels between oximetry = 0 and␣
 ↪oximetry = 1.
```

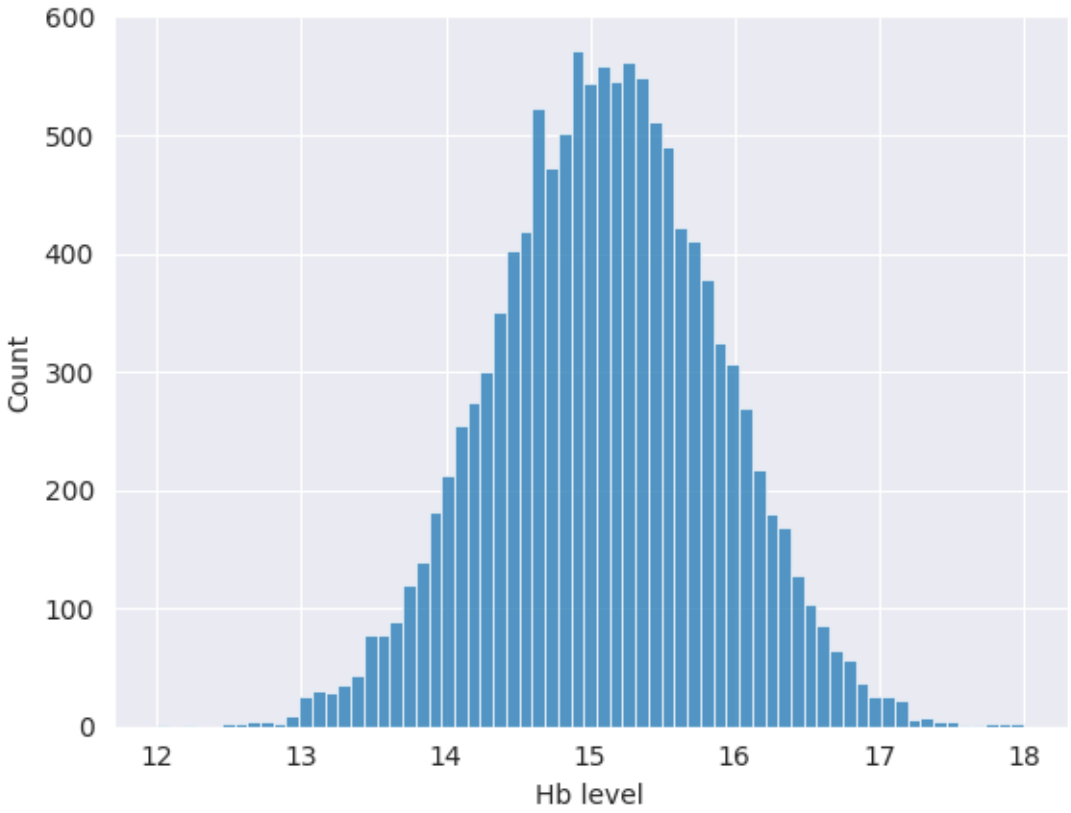
3.3.3 A) Hypothesis 2 - Comparison of Hemoglobin (Hb) Levels Between Oximetry Groups

H (Null Hypothesis): The mean hemoglobin (Hb) level is the same for both groups — patients with oximetry = 0 and oximetry = 1.

H (Alternative Hypothesis): The mean hemoglobin (Hb) level differs between the groups — patients with oximetry = 0 and oximetry = 1.

[877]:
```
sns.histplot(observation_df['Hb level'])
```

[877]:
```
<Axes: xlabel='Hb level', ylabel='Count'>
```

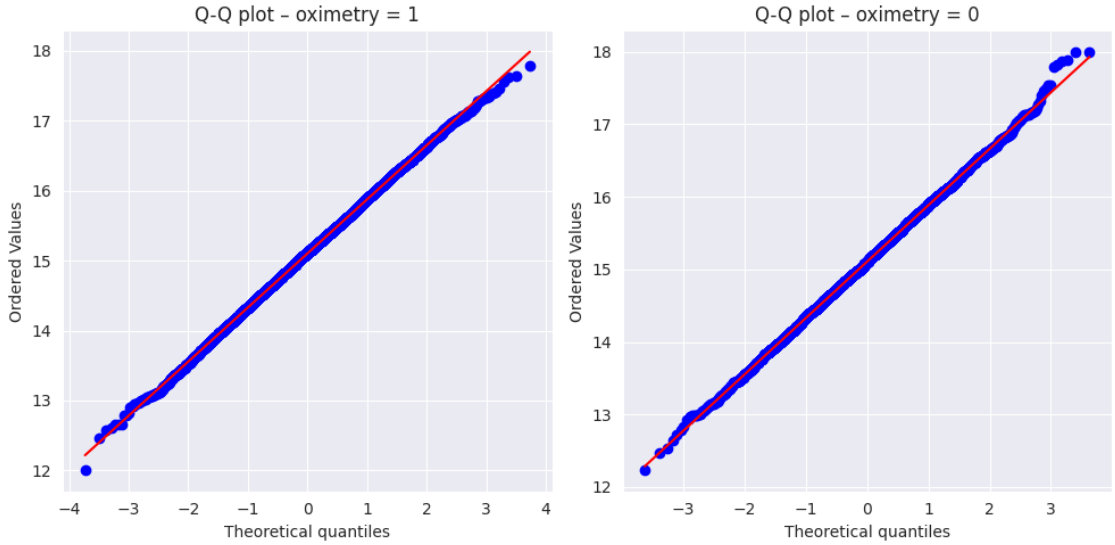


[878]:
```
hb0 = observation_df[observation_df['oximetry'] == 0]['Hb level']
hb1 = observation_df[observation_df['oximetry'] == 1]['Hb level']

#we need to set sample to max 5000 so the data are in range of the recommended␣
 ↪sample value
sample_hb0= hb0.sample(2000, random_state=42)
sample_hb1 = hb1.sample(2000, random_state=42)

#doing the shapiro test -> pvalue for both values
p0 = shapiro(sample_hb0).pvalue
p1 = shapiro(sample_hb1).pvalue
print('shapiro:\np0:{} \np1:{}'.format(p0,p1))
```

```
shapiro:
p0:0.11878125043102106
```

p1:0.85391674151546

[879]:
```
#lets verify the distributions of both the subsets via qqplot
fig, ax = plt.subplots(1, 2, figsize=(10, 5))

stats.probplot(hb1, dist="norm", plot=ax[0])
ax[0].set_title("Q-Q plot - oximetry = 1")

stats.probplot(hb0, dist="norm", plot=ax[1])
ax[1].set_title("Q-Q plot - oximetry = 0")

plt.tight_layout()
plt.show()
#they indeed are
```



[880]:
```
#same as H0
p_lev = stats.levene(sample_hb1,sample_hb0).pvalue
print('levene:', p_lev)

if p_lev > 0.05:
    # this was explained in H0
    t, p = stats.ttest_ind(sample_hb1, sample_hb0)
    print('t-test ', t, p)
else:
    # this was explained in H0
    t, p = stats.ttest_ind(sample_hb1, sample_hb0, equal_var=False)
    print('Welch t-test (unequal variances):', t, p)
```

levene: 0.6624508281537517

```
t-test  -1.588383168023951 0.11227879476272101
```

[881]:
```
# Both groups are normally distributed (Shapiro > 0.05), with their variance␣
↪being very similar as well. Students t test shows that the difference of the␣
↪means was not statistically significant.
```

3.3.4 B) Overte či Vaše štatistické testy majú dostatok podpory z dát, teda či majú dostatočne silnú štatistickú silu.

[882]:
```
# SpO2 attribute
y0 = sample_g0.values
y1 = sample_g1.values

# base stats
m0, m1 = np.mean(y0), np.mean(y1)
s0, s1 = np.std(y0, ddof=1), np.std(y1, ddof=1)
n0, n1 = len(y0), len(y1)

# CI
se = np.sqrt(s0**2/n0 + s1**2/n1)
df_temp = (s0**2/n0 + s1**2/n1)**2 / ((s0**4)/((n0**2)*(n0-1)) + (s1**4)/
↪((n1**2)*(n1-1)))
ci = stats.t.interval(0.95, df_temp, loc=(m1 - m0), scale=se)
print(f"95% CI for delta means: {ci[0]:.3f} to {ci[1]:.3f}")
#very little difference -> strong statistical power
```

```
95% CI for delta means: -0.032 to 0.035
```

[883]:
```
#Cohen
sp = np.sqrt(((n0 - 1)*s0**2 + (n1 - 1)*s1**2) / (n0 + n1 - 2))
d = (m1 - m0) / sp
print(f"Cohen's d: {d:.3f}")
#almost 0 difference -> strong statistical power
```

```
Cohen's d: 0.003
```

CI ranging from -0.032 to 0.035 and cohens'd being 0.003 shows us that there is strong statistical power and that the tests have enough support from the data to be legitimate

[884]:
```
# HB level attribute
y0 = sample_hb0.values
y1 = sample_hb1.values

# base stats
m0, m1 = np.mean(y0), np.mean(y1)
s0, s1 = np.std(y0, ddof=1), np.std(y1, ddof=1)
n0, n1 = len(y0), len(y1)

# CI
```

```
se = np.sqrt(s0**2/n0 + s1**2/n1)
df_temp = (s0**2/n0 + s1**2/n1)**2 / ((s0**4)/((n0**2)*(n0-1)) + (s1**4)/
 ↪((n1**2)*(n1-1)))
ci = stats.t.interval(0.95, df_temp, loc=(m1 - m0), scale=se)
print(f"95% CI for delta means: {ci[0]:.3f} to {ci[1]:.3f}")
#same result as with SPO2
```

```
95% CI for delta means: -0.087 to 0.009
```

[885]:
```
#Cohen
sp = np.sqrt(((n0 - 1)*s0**2 + (n1 - 1)*s1**2) / (n0 + n1 - 2))
d = (m1 - m0) / sp
print(f"Cohen's d: {d:.3f}")
#almost 0 difference -> strong statistical power
```

```
Cohen's d: -0.050
```

CI ranging from -0.087 to 0.009 and cohens'd being -0.050 shows us that there is strong statistical power and that the tests have enough support from the data to be legitimate

4 Fáza 2

4.1 2.1 Realizácia predspracovania dát

4.1.1 A)

Before doing the magic with spliting data we need to set them up by filtering only atributes we need and are usefull in future analysis

[886]:
```
df_2 = df.copy()

# we want to work with the data that has some connectivity to "health"␣
 ↪specification not anything like position ...
drop_cols = [c for c in ['user_id','station_ID','latitude','longitude'] if c in␣
 ↪df_2.columns]
df_2 = df_2.drop(columns=drop_cols)

df_2 = df_2.drop_duplicates()
```

[887]:
```
target = "oximetry"

#split featires and target
X = df_2.drop(columns=[target])
y = df_2[target]
```

[888]:
```
X_train, X_test, y_train, y_test = train_test_split(
    X, y,
    test_size=0.2,
```

```
    random_state=42,
    stratify=y
)

print("Train dataframe:", X_train.shape)
print("Test dataframe:", X_test.shape)
print("Oximetry=1/total for train:", y_train.mean().round(3))
print("Oximetry=1/total for test:", y_test.mean().round(3))
#we can see the division of the original df into train and test subdatasets, we can also see the stratification focused on the oximetry attribute worked
```

```
Train dataframe: (41501, 23)
Test dataframe: (10376, 23)
Oximetry=1/total for train: 0.597
Oximetry=1/total for test: 0.597
```

4.1.2 B)

[889]:
```
X_test.head()
```

[889]:

	SpO2	HR	PI	RR	EtCO2	FiO2	\
49445	96.601411	80.344377	8.884552	15.344047	38.825321	55.089020	
30741	97.993688	83.684406	11.323195	15.208839	41.193087	63.886566	
52230	96.805991	89.893353	7.491702	14.655980	42.801724	62.273550	
50948	97.293984	77.630562	11.833415	16.148954	42.182432	65.501698	
58059	97.263038	84.972915	10.198301	14.049168	39.057607	59.162991	

	PRV	BP	Skin Temperature	Motion/Activity index	…	\
49445	84.643454	105.944792	34.263087	10.094809	…	
30741	111.221376	106.803949	34.890998	9.349818	…	
52230	108.099974	101.153579	36.156039	10.064168	…	
50948	130.012206	100.104509	35.036113	8.346491	…	
58059	129.506948	99.689616	36.135584	9.809290	…	

	CO	Blood Flow Index	PPG waveform features	\
49445	4.001692	67.333233	38.870747	
30741	4.005741	42.993997	33.119117	
52230	4.067662	50.017832	37.370695	
50948	4.000921	49.619197	60.764507	
58059	4.009634	60.365075	58.944290	

	Signal Quality Index	Respiratory effort	O2 extraction ratio	\
49445	76.138614	40.726619	0.268550	
30741	24.206459	33.059317	0.256908	
52230	59.558249	44.561747	0.210238	
50948	42.742079	46.305055	0.219876	
58059	55.162309	69.790015	0.245487	

```
              SNR          QoS  code  blood_group
49445  25.952849         good    AM           AB+
30741  31.628228    excellent    PH            O-
52230  28.975340    excellent    AU            O+
50948  29.580558  maintenance    FR            O+
58059  22.505126    excellent    IN            A+

[5 rows x 23 columns]
```

[890]:
```
X_train.columns
```

[890]:
```
Index(['SpO2', 'HR', 'PI', 'RR', 'EtCO2', 'FiO2', 'PRV', 'BP',
       'Skin Temperature', 'Motion/Activity index', 'PVI', 'Hb level', 'SV',
       'CO', 'Blood Flow Index', 'PPG waveform features',
       'Signal Quality Index', 'Respiratory effort', 'O2 extraction ratio',
       'SNR', 'QoS', 'code', 'blood_group'],
      dtype='object')
```

[891]:
```
print(X_train.dtypes)
#every attribute is either category or float64, however this was not the case␣
 ↪when we started working with the dataset, QoS, code and blood group were␣
 ↪changed into category datatype in 1.2 A), 3rd cell -> we now have to turn␣
 ↪category attributes into numberic attributes
```

```
SpO2                      float64
HR                        float64
PI                        float64
RR                        float64
EtCO2                     float64
FiO2                      float64
PRV                       float64
BP                        float64
Skin Temperature          float64
Motion/Activity index     float64
PVI                       float64
Hb level                  float64
SV                        float64
CO                        float64
Blood Flow Index          float64
PPG waveform features     float64
Signal Quality Index      float64
Respiratory effort        float64
O2 extraction ratio       float64
SNR                       float64
QoS                      category
code                     category
blood_group              category
```

```
dtype: object
```

[892]:
```
X_train['QoS'].unique()
```

[892]:
```
['good', 'excellent', 'acceptable', 'maintenance']
Categories (4, object): ['acceptable', 'excellent', 'good', 'maintenance']
```

[893]:
```
#It makes sense to use ordinal encoding on QoS, since we can set which is the␣
 ↪best(excellent) and which is the worst(maintenece)
qos_order = CategoricalDtype(
    categories=["maintenance", "acceptable", "good", "excellent"],
    ordered=True
)

X_train["QoS"] = X_train["QoS"].astype(qos_order)

X_train["QoS_ord"] = X_train["QoS"].cat.codes
X_train = X_train.drop(columns=["QoS"])
```

[894]:
```
X_train['blood_group'].unique()
```

[894]:
```
['A-', 'B+', 'AB+', 'A+', 'AB-', 'O+', 'O-', 'B-']
Categories (8, object): ['A+', 'A-', 'AB+', 'AB-', 'B+', 'B-', 'O+', 'O-']
```

[895]:
```
X_train = pd.get_dummies(X_train, columns=["blood_group"], prefix="bg")
```

[896]:
```
X_train['code'].unique()
```

[896]:
```
['KR', 'CN', 'BJ', 'AR', 'US', …, 'EC', 'HK', 'CH', 'MK', 'CF']
Length: 97
Categories (96, object): ['AF', 'AM', 'AO', 'AR', …, 'VE', 'VN', 'VU', 'ZA']
```

[897]:
```
#we also remember that the code attribute is the only attribute that has a␣
 ↪missing value which belonged to Okinawa station, we will simply drop these␣
 ↪rows
#I did think about imputing the station with a new code for Japan, but its␣
 ↪really only 100 rows and we cant use an imputing strategy that could be␣
 ↪later replicated as this is Okinawa only...
X_train = X_train.dropna()
```

[898]:
```
#after further analysis we decided to drop code column alltogether, as the only␣
 ↪viable encoding option was frequency encoding but that would eventually make␣
 ↪no sense...
#We know we should not change te test dataset in this part but this is not␣
 ↪really a preprocessing method
X_train = X_train.drop(columns=["code"])
X_test = X_test.drop(columns=["code"])
```

[899]:
```
print(X_test.dtypes)
#We can see that there are no more non numeric attributes!
```

```
SpO2                       float64
HR                         float64
PI                         float64
RR                         float64
EtCO2                      float64
FiO2                       float64
PRV                        float64
BP                         float64
Skin Temperature           float64
Motion/Activity index      float64
PVI                        float64
Hb level                   float64
SV                         float64
CO                         float64
Blood Flow Index           float64
PPG waveform features      float64
Signal Quality Index       float64
Respiratory effort         float64
O2 extraction ratio        float64
SNR                        float64
QoS                       category
blood_group               category
dtype: object
```

4.1.3 C)

[900]:
```
X_train.dtypes.head(30)
```

[900]:
```
SpO2                       float64
HR                         float64
PI                         float64
RR                         float64
EtCO2                      float64
FiO2                       float64
PRV                        float64
BP                         float64
Skin Temperature           float64
Motion/Activity index      float64
PVI                        float64
Hb level                   float64
SV                         float64
CO                         float64
Blood Flow Index           float64
PPG waveform features      float64
```

```
Signal Quality Index     float64
Respiratory effort       float64
O2 extraction ratio      float64
SNR                      float64
QoS_ord                     int8
bg_A+                       bool
bg_A-                       bool
bg_AB+                      bool
bg_AB-                      bool
bg_B+                       bool
bg_B-                       bool
bg_O+                       bool
bg_O-                       bool
dtype: object
```

[901]:
```
num_cols = X_train.select_dtypes(include=["float64", "int64","int8"]).columns.
 ↪tolist()
num_cols
```

[901]:
```
['SpO2',
 'HR',
 'PI',
 'RR',
 'EtCO2',
 'FiO2',
 'PRV',
 'BP',
 'Skin Temperature',
 'Motion/Activity index',
 'PVI',
 'Hb level',
 'SV',
 'CO',
 'Blood Flow Index',
 'PPG waveform features',
 'Signal Quality Index',
 'Respiratory effort',
 'O2 extraction ratio',
 'SNR',
 'QoS_ord']
```

[902]:
```
# Scaniling #1 - Standart Scaler

std_scaler = StandardScaler()

X_train_std = pd.DataFrame(
    std_scaler.fit_transform(X_train[num_cols]),
```

```
    columns=num_cols,
    index=X_train.index
)
# values check
check_means = X_train_std[num_cols].mean().round(3)
check_stds  = X_train_std[num_cols].std(ddof=0).round(3)

print("Means (should be ~0):")
print(check_means.head(10))
print("\nStds (should be ~1):")
print(check_stds.head(10))
```

```
Means (should be ~0):
SpO2                     0.0
HR                      -0.0
PI                      -0.0
RR                       0.0
EtCO2                   -0.0
FiO2                    -0.0
PRV                     -0.0
BP                      -0.0
Skin Temperature        -0.0
Motion/Activity index    0.0
dtype: float64

Stds (should be ~1):
SpO2                     1.0
HR                       1.0
PI                       1.0
RR                       1.0
EtCO2                    1.0
FiO2                     1.0
PRV                      1.0
BP                       1.0
Skin Temperature         1.0
Motion/Activity index    1.0
dtype: float64
```

[903]:
```
#Scaling #2 - RobustScaler (outlierproof; median=0, IQR=1)

rob_scaler = RobustScaler(with_centering=True,with_scaling=True)

X_train_robust = pd.DataFrame(
    rob_scaler.fit_transform(X_train[num_cols]),
    columns=num_cols,
    index=X_train.index
)
```

```
#check if the values are fine
medians = X_train_robust[num_cols].median().round(3)
q75 = X_train_robust[num_cols].quantile(0.75)
q25 = X_train_robust[num_cols].quantile(0.25)
iqr = (q75 - q25).round(3)

print("Medians (should be ~0):")
print(medians.head(10))
print("\nIQR (should be ~1):")
print(iqr.head(10))
```

```
Medians (should be ~0):
SpO2                      0.0
HR                        0.0
PI                        0.0
RR                        0.0
EtCO2                     0.0
FiO2                      0.0
PRV                       0.0
BP                        0.0
Skin Temperature         -0.0
Motion/Activity index     0.0
dtype: float64

IQR (should be ~1):
SpO2                      1.0
HR                        1.0
PI                        1.0
RR                        1.0
EtCO2                     1.0
FiO2                      1.0
PRV                       1.0
BP                        1.0
Skin Temperature          1.0
Motion/Activity index     1.0
dtype: float64
```

[904]:
```
# Transformer #1 PowerTransformer
power_tx = PowerTransformer(method="yeo-johnson", standardize=True)

X_train_power = X_train.copy()
X_train_power[num_cols] = power_tx.fit_transform(X_train[num_cols])

# quick checks
means = X_train_power[num_cols].mean().round(3).head(10)
stds  = X_train_power[num_cols].std(ddof=0).round(3).head(10)
```

```
print("PowerTransformer -> means ~0:\n", means)
print("\nPowerTransformer -> stds ~1:\n", stds)
```

```
PowerTransformer -> means ~0:
 SpO2                      0.0
HR                        -0.0
PI                        -0.0
RR                        -0.0
EtCO2                     -0.0
FiO2                      -0.0
PRV                        0.0
BP                        -0.0
Skin Temperature           0.0
Motion/Activity index     -0.0
dtype: float64

PowerTransformer -> stds ~1:
 SpO2                      1.0
HR                        1.0
PI                        1.0
RR                        1.0
EtCO2                     1.0
FiO2                      1.0
PRV                       1.0
BP                        1.0
Skin Temperature          1.0
Motion/Activity index     1.0
dtype: float64
```

[905]:
```
nq = min(1000, len(X_train))  # practical cap
quant_tx = QuantileTransformer(
    n_quantiles=nq,
    output_distribution="normal",
    random_state=42
)

X_train_quant = X_train.copy()
X_train_quant[num_cols] = quant_tx.fit_transform(X_train[num_cols])

# quick checks
means_q = X_train_quant[num_cols].mean().round(3).head(10)
stds_q  = X_train_quant[num_cols].std(ddof=0).round(3).head(10)

print("QuantileTransformer -> means ~0:\n", means_q)
print("\nQuantileTransformer -> stds ~1:\n", stds_q)
```

```
QuantileTransformer -> means ~0:
```

```
 SpO2                      0.021
HR                        -0.002
PI                        -0.001
RR                        -0.007
EtCO2                      0.007
FiO2                      -0.002
PRV                        0.009
BP                        -0.005
Skin Temperature          -0.006
Motion/Activity index     -0.005
dtype: float64

QuantileTransformer -> stds ~1:
 SpO2                       0.998
HR                         1.003
PI                         1.003
RR                         0.999
EtCO2                      0.998
FiO2                       0.997
PRV                        1.002
BP                         0.998
Skin Temperature           1.015
Motion/Activity index      1.000
dtype: float64
```

[906]:
```
# we already did more techniques such as Outlier ( winsortizatoin , IQR ) ,␣
↪encoding
```

4.1.4 D)

As part of data preprocessing (section 2.1(C)), four commonly used transformation methods were applied to numerical attributes — two **scaling** and two **distribution transformation** techniques.

Scaling techniques

1. **StandardScaler**
   - Chosen as the baseline normalization method.
   - Scales each feature to have mean 0 and standard deviation 1.
   - Improves model convergence and training stability for algorithms sensitive to feature magnitude (e.g., Logistic Regression, SVM).
2. **RobustScaler**
   - Used as a more resilient alternative to handle outliers.
   - Scales data using the median and interquartile range (IQR), reducing the influence of extreme values (e.g., in the `CO` feature).

Transformation techniques

3. **PowerTransformer (Yeo–Johnson)**
   - Applied to stabilize variance and reduce skewness in feature distributions.

- Helps make distributions more symmetric and supports linear model assumptions.

4. **QuantileTransformer (normal)**
   - Maps the feature quantiles to a normal distribution, effectively reducing the impact of outliers and non-Gaussian shapes.
   - Useful for attributes with irregular or multimodal distributions.

Summary These four techniques cover the most relevant preprocessing needs for numerical data: - **StandardScaler** – baseline normalization, - **RobustScaler** – robust to outliers, - **PowerTransformer** – reduces skewness and stabilizes variance, - **QuantileTransformer** – enforces normal-like distribution.

This combination allows comparison of preprocessing effects in later phases (2.3 and 3), ensuring a replicable, consistent, and model-friendly data preparation workflow.

4.2 2.2 Výber atribútov pre strojové učenie

4.2.1 A)

[907]:
```
X = X_train.copy()
y = y_train.copy()

# 1) align indices first (very important if X was dropna'd earlier)
common_idx = X.index.intersection(y.index)
X = X.loc[common_idx]
y = y.loc[common_idx]

# 2) drop rows with any NaN in X (apply the SAME mask to both X and y)
mask = ~X.isna().any(axis=1)
X = X.loc[mask]
y = y.loc[mask]

print("Shapes after align -> X:", X.shape, "| y:", y.shape)
```

```
Shapes after align -> X: (41392, 29) | y: (41392,)
```

[908]:
```
scaler_minmax = MinMaxScaler()
X_chi = pd.DataFrame(
    scaler_minmax.fit_transform(X),
    columns=X.columns,
    index=X.index
)
chi_scores, p_values = chi2(X_chi, y)
chi_results = pd.Series(chi_scores, index=X.columns).
 ↪sort_values(ascending=False)

print("Top 10 features by Chi-Square:\n")
print(chi_results.head(10))
```

```
Top 10 features by Chi-Square:
```

```
HR                      1164.230502
CO                       435.709035
EtCO2                    305.397141
FiO2                     154.531106
PI                        21.104691
RR                        17.934760
O2 extraction ratio        1.538853
bg_B-                      1.303170
QoS_ord                    1.243664
bg_O+                      1.183909
dtype: float64
```

[909]:
```
X_chi[X_chi.columns]
```

[909]:

	SpO2	HR	PI	RR	EtCO2	FiO2	PRV	\
6475	0.525146	0.787473	0.513921	0.715944	0.326645	0.566474	0.530721	
12274	0.506686	0.857688	0.723126	0.796223	0.532064	0.647200	0.518096	
47746	0.386994	0.478045	0.546312	0.553884	0.475704	0.347002	0.626112	
56807	0.445673	0.256708	0.625327	0.712715	0.307308	0.588409	0.687110	
46687	0.474444	0.713392	0.461103	0.341468	0.683023	0.541257	0.390535	
…	…	…	…	…	…	…	…	
32793	0.460962	0.548346	0.474521	0.466746	0.580586	0.381199	0.677896	
52966	0.264648	0.781487	0.534218	0.640457	0.617411	0.557990	0.635862	
26667	0.477546	0.363955	0.680007	0.510529	0.509821	0.361080	0.791034	
6446	0.466543	0.523679	0.486441	0.692979	0.301562	0.321686	0.402102	
42302	0.403658	0.599102	0.467683	0.409394	0.466401	0.428401	0.404867	

	BP	Skin Temperature	Motion/Activity index	…	SNR	\
6475	0.328978	0.259424	0.548962	…	0.581740	
12274	0.263212	0.671844	0.273192	…	0.324382	
47746	0.387173	0.243384	0.378412	…	0.681468	
56807	0.580476	0.379736	0.410406	…	0.342466	
46687	0.556896	0.454367	0.623607	…	0.524398	
…	…	…	… …	…		
32793	0.406224	0.354757	0.692835	…	0.704693	
52966	0.688202	0.509013	0.420458	…	0.889518	
26667	0.544325	0.440188	0.309792	…	0.316030	
6446	0.436908	0.409603	0.415853	…	0.394871	
42302	0.459714	0.463596	0.241395	…	0.700864	

	QoS_ord	bg_A+	bg_A-	bg_AB+	bg_AB-	bg_B+	bg_B-	bg_O+	bg_O-
6475	0.666667	0.0	1.0	0.0	0.0	0.0	0.0	0.0	0.0
12274	1.000000	0.0	0.0	0.0	0.0	1.0	0.0	0.0	0.0
47746	1.000000	0.0	0.0	1.0	0.0	0.0	0.0	0.0	0.0
56807	0.666667	1.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
46687	0.666667	0.0	0.0	0.0	0.0	1.0	0.0	0.0	0.0

```
…          …    …    …    …    …    …    …    …    …
32793  1.000000  0.0  0.0  0.0  0.0  0.0  0.0  0.0  1.0
52966  0.666667  0.0  1.0  0.0  0.0  0.0  0.0  0.0  0.0
26667  0.333333  0.0  0.0  0.0  0.0  0.0  1.0  0.0  0.0
6446   0.666667  0.0  0.0  0.0  0.0  1.0  0.0  0.0  0.0
42302  0.666667  0.0  0.0  0.0  0.0  0.0  0.0  1.0  0.0

[41392 rows x 29 columns]
```

[910]:
```
X_mi = X.copy()
mi_scores = mutual_info_classif(X, y, random_state=42)
mi_results = pd.Series(mi_scores, index=X.columns).sort_values(ascending=False)

print("Top 10 features by Mutual Information:\n")
print(mi_results.head(10))
```

```
Top 10 features by Mutual Information:

HR                        0.559387
CO                        0.554434
EtCO2                     0.511240
RR                        0.489521
FiO2                      0.484673
PPG waveform features     0.482610
Blood Flow Index          0.482178
PVI                       0.479729
SV                        0.478260
PI                        0.477896
dtype: float64
```

[ ]:
```
rf = RandomForestClassifier(n_estimators=200, random_state=42)
rf.fit(X, y)

rf_importances = pd.Series(rf.feature_importances_, index=X.columns).
 ↪sort_values(ascending=False)

print("Top 10 features by Random Forest Importance:\n")
print(rf_importances.head(10))
```

[748]:
```
#compare 3 methods
feature_scores = pd.concat([
    mi_results.rename("Mutual Information"),
    chi_results.rename("Chi2"),
    rf_importances.rename("Random Forest")
], axis=1)

# normalize to 0-1 scale for easier comparison
```

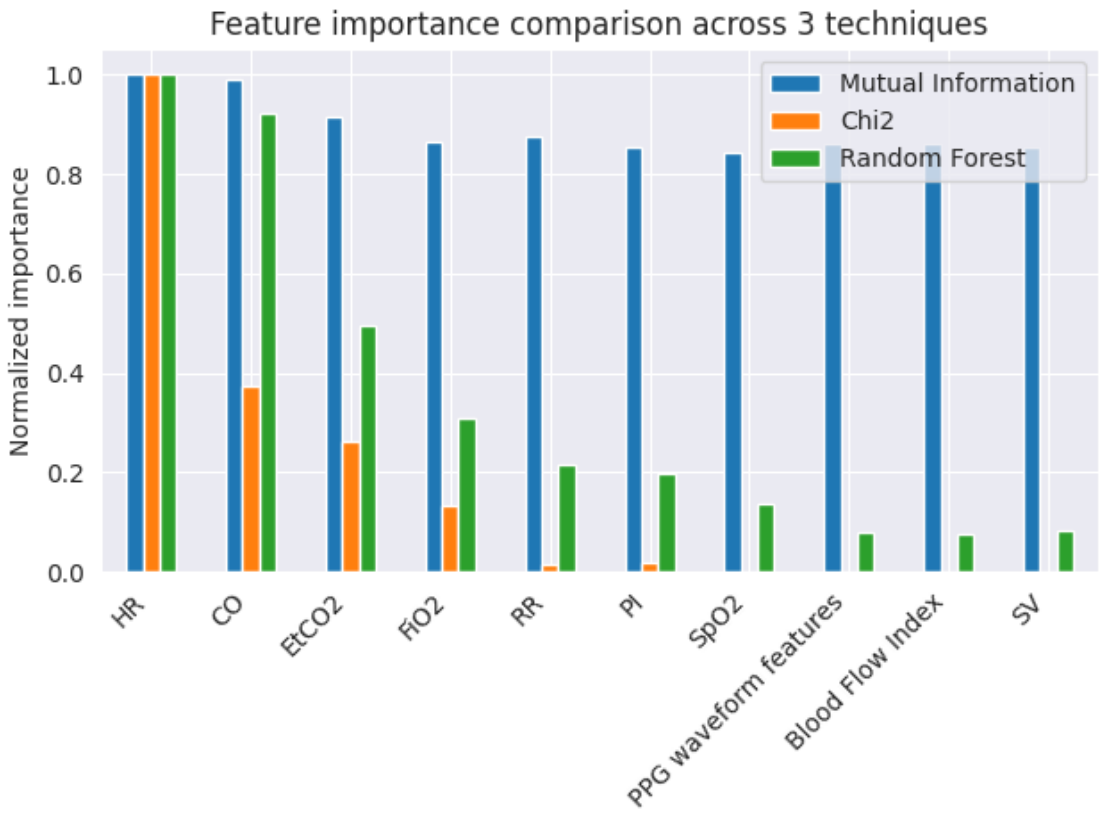
```
feature_scores = feature_scores.apply(lambda x: (x - x.min()) / (x.max() - x.
 ↪min()))

# compute average score
feature_scores["Average"] = feature_scores.mean(axis=1)
feature_scores.sort_values("Average", ascending=False).head(10)
```

[748]:

	Mutual Information	Chi2	Random Forest	Average
HR	1.000000	1.000000	1.000000	1.000000
CO	0.991147	0.374244	0.921206	0.762199
EtCO2	0.913930	0.262314	0.494068	0.556771
FiO2	0.866436	0.132730	0.308987	0.436051
RR	0.875103	0.015402	0.215498	0.368668
PI	0.854322	0.018125	0.197009	0.356485
SpO2	0.843700	0.000513	0.137522	0.327245
PPG waveform features	0.862748	0.000470	0.077891	0.313703
Blood Flow Index	0.861977	0.000793	0.076666	0.313145
SV	0.854972	0.000000	0.081849	0.312274

[749]:
```
#visualization
top10 = feature_scores.sort_values("Average", ascending=False).head(10)

plt.figure(figsize=(10,6))
top10[["Mutual Information", "Chi2", "Random Forest"]].plot(kind="bar")
plt.title("Feature importance comparison across 3 techniques")
plt.ylabel("Normalized importance")
plt.xticks(rotation=45, ha="right")
plt.tight_layout()
plt.show()
```

```
<Figure size 1000x600 with 0 Axes>
```



4.2.2 4.2.1 A) Findings & decisions (feature selection)

Setup. Informativeness of features for the target `oximetry` was evaluated by three independent methods: (1) Mutual Information (MI), (2) Chi-Square (^{2}; after MinMax scaling to ensure non-negativity), and (3) model-based Random Forest importance. Scores were normalized to 0,1 and averaged for comparison.

Consistent signals across methods. - **HR** and **CO** ranked **#1–#2** in both MI and Random Forest and stayed on top in the averaged ranking → strongest, method-agnostic predictors. - **EtCO2** and **FiO2** showed **high to medium** importance in MI and RF (respiratory physiology) → robust, clinically plausible predictors. - **RR** and **PI** were **medium**; **SpO2** medium-to-lower (still relevant). - Other waveform/flow indices (PPG waveform features, Blood Flow Index, SV) were **lower**, yet non-zero, suggesting auxiliary value.

Why ^{2} looks conservative. - ^{2} assumes independence on **binned/positive** data; with continuous variables (scaled to 0–1) it is less sensitive to nuanced, non-linear relations. - MI and RF capture **non-linear/monotone** effects better, therefore we weigh them more when resolving ties.

Decision – feature shortlist for ML (to carry forward). - **Core set (high confidence):** `HR`, `CO`, `EtCO2`, `FiO2`. - **Support set (keep / test with regularization):** `RR`, `PI`, `SpO2`. - **Optional (use if helpful in CV, otherwise drop to reduce complexity):** `PPG waveform features`, `Blood Flow Index`, `SV`.

Rationale. - Chosen features are consistently ranked by **2 methods** and align with earlier EDA (HR oximetry, respiratory parameters oxygenation). - Keeping a compact core reduces multicollinearity and speeds training; the support/optional groups allow controlled expansion if cross-validated performance improves.

Next steps (planned in 2.2/2.3). - Compare *Core* vs. *Core+Support* (and optionally +PCA) in stratified CV with the final preprocessing choice (scaling/transformers). - Use model regularization (e.g., L1/L2) or tree-based models to manage redundancy.

4.2.3 C)

4.3 2.3 Replikovateľnosť predspracovania

4.3.1 A)

4.3.2 B)