

Appendix - COVID-19 test fraud detection: Findings from a pilot study comparing conventional and statistical approaches

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- ⁵ Hasso Plattner Institute, Department Digital Global Public Health

Zitieren

Bosnjak M, Dahm S, Kuhnert R, Weihrauch D, Schaffrath Rosario A, Hurraß J, Schmich P, Wieler L und Nießen J (2024): Appendix - COVID-19 test fraud detection: Findings from a pilot study comparing conventional and statistical approaches. [Dataset] Zenodo. DOI:10.5281/zenodo.10608926.

The methods and results of the publication "COVID-19 test fraud detection: Findings from a pilot study comparing conventional and statistical approaches" are described in more detail in this appendix. The R-syntax for the calculation is provided, as well as a pseudo data set with which the syntax can also be tested.

Organisational and administrative information

The publicaton "COVID-19 test fraud detection: Findings from a pilot study comparing conventional and statistical approaches", is a joined projekt of the Department for Psychological Research Methods - Trier University, Department 2 | Epidemiology and Health Monitoring - Robert Koch Institute, the Department Infectious and Environmental Hygieneare - Health Authority of the City of Cologne and the Department Digital Global Public Health - Hasso Plattner Institute. The appendix presented here provides additional results and data for the publication and was currated by Department 2 | Epidemiology and Health Monitoring of the Robert Koch Institute. Questions regarding the content of the data can be addressed directly to the correspondending author Michael Bosnjak (bosnjak@uni-trier.de).

The publication of the data as well as the quality management of the (meta-)data is done by the department MF 4 | Research Data and Infromation Management. Questions regarding data management and the publication infrastructure can be directed to the Open Data Team of the Department MF4 at OpenData@rki.de.

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Data

We used data on claims for COVID-19 antigen tests submitted for reimbursement by 907 test centers operating in a German city with approximately one million residents for the timespan April 8, 2021 through August 28, 2022.

The data were transmitted on a daily basis via an online portal provided for this purpose by the ministry of a federal German state. Transmission was mandatory by law for the test centers by "CoronaTeststrukturVO" from 2021-03-09, regulated in §5.

For each claim, the following information was provided: test center category (pharmacy, doctor's or dentist's office, private test center), date of testing, number of tests performed per day, number of positive tests per day. The detailed data schmea of the analysed data can be found in section data schema of the simulated data, as we provide simulated data in the same format.

Methods and Results

We used four statistical methods to detect fraud, which are described in the following sections. All results shown are based on the original data.

- Outlier identification from the mean number of tests per day invoiced (high number of tests)
- Low positive rates identified by Poisson regression (low positive rate)
- Deviations from Benford's Law
- Deviations from the assumption of equally distributed last digits

Outlier identification from the mean number of tests per day invoiced (high number of tests)

In our first statistical approach aimed at identifying disproportionately high test volumes invoiced, the numbers of tests invoiced per day are classified as conspicuous if they fall outside the 90% percentile in terms of the mean number of tests performed per day within a test center category.

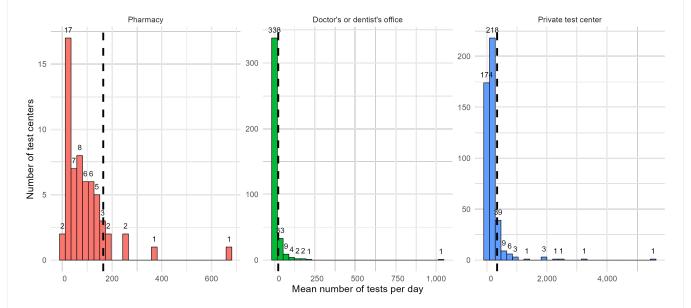
Analysis script

The analysis on outlier identification from the mean number of tests per day invoiced was performed using an R script. The content of the script is provided as R file:

supporting_material/scripts/Fraud_Methods_Description.r

Figure 1 shows the corresponding distributions resulting form the analysis.

Figure 1: Histograms of the mean number of tests performed per day (x-axis) by test center type (pharmacies, doctor's or dentist's offices, private test centers). The dashed vertical line indicates the 90% percentile of each distribution. Test centers falling on the right sides of these lines are considered statistically conspicuous. The numbers above the bars indicate the number of test centers within each bar.



A total of 91 testing centers (6 pharmacies, 39 physician practices/dentists, and 46 private testing sites) were classified as suspicious using this approach. Table 1 shows the basic statistics of the tests performed per day, divided into conspicuous and non-conspicuous test centers according to the statistical method.

Table 1: Basic statistics of the mean number of tests per day and test centers by facility type, statistically conspicuous and statistically not conspicuous of fraud

| For ellitary to many | Statistically conspicuous | | | Statistically not conspicuous | | | Total | | |
|------------------------------|---------------------------|--------|---------|-------------------------------|--------|-------|-------|--------|---------|
| Facility type | N | Median | Max | N | Median | Max | N | Median | Max |
| Pharmacy | 6 | 252.2 | 679.3 | 54 | 64.1 | 161.5 | 60 | 71.0 | 679.3 |
| Doctor's or dentist's office | 39 | 51.7 | 1,032.0 | 351 | 3.5 | 23.0 | 390 | 3.8 | 1,032.0 |
| Private test center | 46 | 515.4 | 5,520.0 | 411 | 106.6 | 342.2 | 457 | 116.3 | 5,520.0 |

Low positive rates identified by Poisson regression (low positive rate)

The positive rates were modeled by a Poisson regression model with random effects using the logarithms of the number of positive tests per day and per test center as dependent variable and the logarithms of the respective total number of tests as offset. The variability between the test centers were modeled by random intercepts for test centers. In addition, to account for changes in the positive rates over time for example induced by changing incidence, calendar week specific random intercepts were introduced in the model. Differences in positive rates between the facility types were controlled by fixed effects.

$$\log(pos_{ijkl}) = \log(test_{ijkl}) + A + \beta_j + \gamma_k + x_l \cdot typ_l + \varepsilon_{ijkl}$$

 pos_{ijkl} : Number of positive tests at day i in test center $j(j=1,\ldots,907)$ the week $k(k=1,\ldots,73)$ and in facility type l(l=1,2,3)

 $test_{ijkl}$: Number of tests at day i in test center j in week k in facility type l

A: Global intercept (fixed effect)

 β_i : Test center-specific deviation from the global intercept (random effect)

 γ_k : Calendar week specific deviation from the global intercept (random effect)

 typ_l : Facility type: Pharmacy, doctor's or dentist's office or private test center (fixed effect)

 x_l : Factor accounting for differences in positive rates by facility type (fixed)

 ε_{ijkl} : Residual at day i in test center j in week k and facility type l not explained by the regression.

Analysis script

The analysis to identify low positive rates by Poisson regression was performed with an R script. The content of the script is provided as R file:

supporting_material/scripts/Fraud_Methods_Poisson_Regression.r

Corona tests were performed in the time span April 8, 2021 through August 28, 2022 on 73 weeks respective 508 days in 907 test centers, but not all centers operated for the entire period. This resulted in a total of N = 118,908 positive rates.

Table 2: Statistics of estimated fixed effects

| Variable | Estimate | Standard Error | P-value |
|---|----------|----------------|----------|
| A (global intercept) | -5.004 | 0.257 | < 0.0001 |
| typ_2 (Facility type: doctor's or dentist's office) | 0.149 | 0.192 | 0.45 |
| typ_3 (Facility type: private test center) | -0.477 | 0.182 | 0.0086 |

The differences between the mean positive rates by facility type (Table 2) were significant in the regression at P = 0.009. In particular, the rate of positive tests was lower for the private test centers than for the pharmacies or the physicians' practices. The estimated random intercepts for the 907 test centers (β_j) were centered by 0 and had a variance of 1.65. The variance of the 73 week specific random intercepts (γ_k) was estimated to be 2.67. Both variables (β and γ) were significant with P < 0.0001

A low center specific random intercept j indicates a low mean positive rate for the tests in resp. center. Therefore, the reporting of tests conducted by a test center was considered conspicuous if its estimated random intercept was significantly low. The estimated test center intercepts (β_j) and their standard deviations $sd(\beta_j)$ were used to generate test values comparable to the t-values of the t-test:

$$r_j = rac{eta_j}{sd(eta_j)}, j = 1, \ldots, 907$$

The test values r_j ranged from -23.0 to 49.0 corresponding to positive rates of 0.5% resp. 10.6%. A value of r_j < -6 was regarded as significant. According to this criterion, the 907 test centers could be classified to 88 conspicuous and 819 not conspicuous test centers (s. Table 3), where the mean positive rate in conspicuous test centers amounted to 0.6% and the not conspicuous test centers had a mean positive rate of 2.3%.

Table 3: Summary of classifications into statistical conspicuous versus not conspicuous test centers according to the Poisson regression model used

| Facility type | Statisticall | Statistically conspicuous | | not conspicuous | Total | | |
|---------------|--------------|---------------------------|--------|-----------------|--------|---------------|--|
| Facility type | Number | Positive rate | Number | Positive rate | Number | Positive rate | |
| Pharmacy | 11 | 0.88 | 49 | 2.82 | 60 | 2.44 | |

| Doctor's or dentist's office | 16 | 0.59 | 374 | 3.95 | 390 | 2.75 |
|------------------------------|----|------|-----|------|-----|------|
| Private test center | 61 | 0.54 | 396 | 2.24 | 457 | 1.95 |
| Total | 88 | 0.58 | 819 | 2.33 | 907 | 2.02 |

Deviations from Benford's Law

Requirements for Benford Analysis:

For data to undergo Benford analysis, it must exhibit a specific range and be reported over a certain number of days. The Benford principle may be compromised if test centers frequently operate at maximum capacity, leading to the reporting of similar test numbers. This can lead to a false positive result for a test center according to Benford's Law. Test centers reporting tests on only a few days are ineligible for Benford analysis because the distribution of the first digit lacks the necessary variability for evaluation. To address this, we stipulate that the number of tests must be reported over a minimum of 30 days.

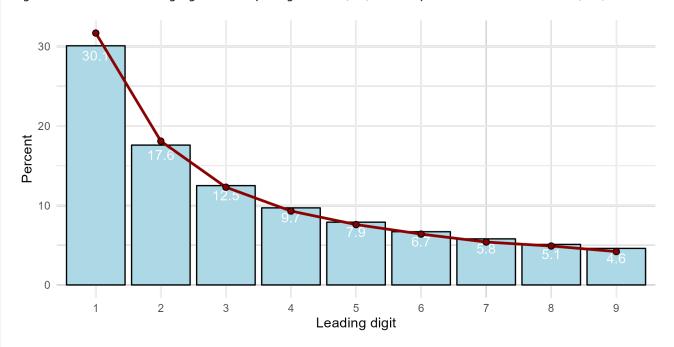
Analysis script

The analysis for deviations from Benford's Law was performed using an R script. The content of the script is provided as R file:

supporting_material/scripts/Fraud_Methods_Benford.r

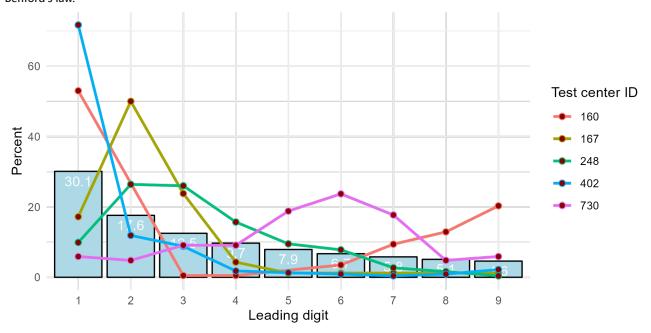
Figure 2 shows the distribution of the leading digit according to Benford's law and the distribution over all data available for the observation period. Overall, there is good agreement. The leading 1 as well as the 2 occur slightly disproportionately according to Benford's law.

Figure 2: Distribution of leading digit of total reporting numbers (line) versus expected values of Benford's law (bars).



A chi-square test is calculated for each of these 665 test centers. The chi-square test value determines the degree of deviation.

Figure 3: Illustration of the distribution of the leading digit of the five test centers (lines) with the largest deviations from Benford's law.



In table 4, we have summarized the number of test centers classified by conventional methods and Benford's Law. The threshold for test centers considered to be conspicuous according to Benford's Law was set to those 10% with the largest chi-square test value.

Table 4: Number of test centers by facility type, (non) suspected of fraud by the conventional approach, and (non) suspected of fraud by the statistical approach focusing on the deviation from Benford's law.

| Facility to the | Suspected of fraud by the health authorities (conventional approach) | | | Not suspected of fraud by the health authorities (conventional approach) | | | | |
|---------------------------------|--|-------------------------------|-------|--|-------------------------------|-------|--|--|
| Facility type | Statistically conspicuous | Statistically not conspicuous | Total | Statistically conspicuous | Statistically not conspicuous | Total | | |
| Pharmacy | 0 | 0 | 0 | 5 | 55 | 60 | | |
| Doctor's or dentist's office | 0 | 4 | 4 | 8 | 197 | 205 | | |
| Private test center | 10 | 65 | 75 | 44 | 227 | 321 | | |
| Total | 10 | 69 | 79 | 57 | 529 | 586 | | |

Based on table 4, the percentage of positive overlap between the traditional and Benford's law-based methods amounts to 12.7% (10/79), the percentage of negative overlap 90.3% (529/586), and the share of incrementally identified potentially fraudulent test centers identified by Benford's law which were undetected by traditional approaches amounts to 9.7% (57/586) related to all test centers.

In contrast to the publication, only the test sites that could also be evaluated by the method are considered here. This means that the positive, negative overlap and the proportion of undetected by traditional approaches are higher than in the results in the publication.

Deviations from the assumption of equally distributed last digits

Requirements for Last Digit Analysis:

Similar to the Benford approach, the last digit method requires data with a sufficient range for evaluating the distribution of the last digit. A minimum number of daily reports (\geq 30) is essential for this evaluation. Unlike Benford's law, the distribution of the last digit includes zero. However, in the single-digit range 0-9, testing centers were not obligated to report "0" on days without testing, leading to a systematic underrepresentation of zero. To mitigate this bias, only test reports with more than 9 tests per day are considered for the distribution. In summary, for the last digit method evaluation, a test center must have reported more than 9 tests per day for at least 30 days. This method could only be applied to 512 of the 907 test sites.

Analysis script

The analysis on deviations from the assumption of equally distributed last digits was performed using an R script. The content of the script is provided as R file:

supporting_material/scripts/Fraud_Methods_Last_Digit.r

0

1

2

3

Figure 4 shows the distribution of the last digit comparing the expected distribution versus all reported data in the observation period. Overall, there is good agreement. The last digits 1 and 2 seem to be slightly overrepresented.

15 Second of the second of the

Figure 4: Distribution of the last digit of the total number of reports (line) versus the expected distribution (bar).

A chi-square test is calculated for each of the 512 test centers. The chi-square test value determines the degree of deviation.

The five test centers with the greatest deviation from the expected distribution can be seen in Figure 5. It seems that the zero and the number 5 were deliberately avoided as the last digit in the test center 48 and 68. Test center 486 reported only an average of 11 tests per day and all numbers below 10 are not considered. Therefore, the distribution has only limited informative value.

5

Last digit

6

7

8

9

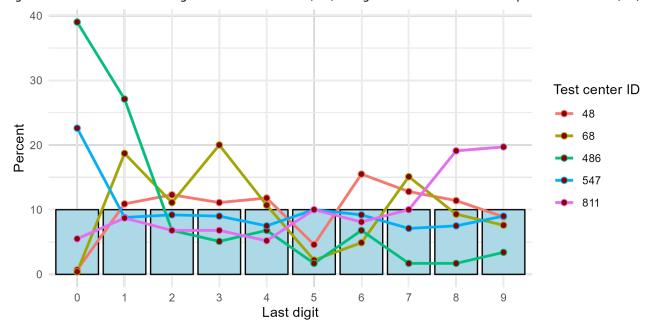


Figure 5: Distribution of the last digit of the five test centes (line) with greatest deviation from the expected distribution (bar).

The threshold for test centers considered to be conspicuous according to the assumption about the last digit distribution was set to those 10% with the largest chi-square test value, yielding 52 test centers. In Table 5, we have summarized the number of test centers classified by traditional approach and the Last Digit method.

Table 5: Number of test centers by facility type, (non) suspected of fraud by the conventional approach, and (non) suspected of fraud by the statistical approach focusing on the deviation from the law of equally distributed last digits.

| Facility as an | Suspected of fraud by the health authorities (conventional approach) | | | Not suspected of fraud by the health authorities (conventional approach) | | | | |
|------------------------------|--|-------------------------------|-------|--|-------------------------------|-------|--|--|
| Facility type | Statistically conspicuous | Statistically not conspicuous | Total | Statistically conspicuous | Statistically not conspicuous | Total | | |
| Pharmacy | 0 | 0 | 0 | 1 | 58 | 59 | | |
| Doctor's or dentist's office | 1 | 3 | 4 | 14 | 48 | 62 | | |
| Private test center | 7 | 68 | 75 | 29 | 283 | 312 | | |
| Total | 8 | 71 | 79 | 44 | 389 | 433 | | |

Based on table 5, the percentage of positive overlap between the traditional and Last-Digit-assumption-based methods amounts to 10.1% (8/79), the percentage of negative overlap 89.8% (389/433), and the share of incrementally identified potentially fraudulent test centers identified by the last digit assumption which were undetected by traditional approaches amounts to 10.2% (44/433).

As with Benford, only the test sites that could also be assessed by the method are considered here. This means that the positive, negative overlap and the proportion of undetected by traditional approaches are higher than in the results in the publication.

Comparison and combination of the four statistical methods used

The positive and negative overlap give the ratio betweet the number of identified test centers by our analysis and the number given by health authorities. If all test sites that are conspicuous by at least one statistical method are considered, the positive overlap also increases by 50.5% (47 test centers of the 93 test centers conspicuous by health authorities were classified also as conspicuous by at least one of the statistical methods). The negative overlap decreases to 78.1% (table 6). If test sites are considered to be conspicuous in the combination of at least 2 methods, the positive and negative overlap is comparable to that of the individual methods.

A more detailed descrition of the comparison can be found in the "Outcomes" section of our corsponding publication:

Bosnjak M, Dahm S, Kuhnert R, Weihrauch D, Schaffrath Rosario A, Hurraß J, Schmich P, Wieler L und Nießen J (2024): COVID-19 test fraud detection: Findings from a pilot study comparing conventional and statistical approaches.

Table 6: Positive and negative overlap and share of incrementally identified potentially fraudulent test centers by statistical approaches

| Statistical approach | Positive overlap | Negative overlap | Share of incrementally identified potentially fraudulent test centers |
|------------------------------------|------------------|---------------------|---|
| At least identified by one method | 50.5% | 78.1% | 21.9% |
| At least identified by two methods | 9.7% | 93.6% | 6.4% |

Simulated data

Based on the data schema of the analysed data, we have simulated data for overall 807 test centers for which we assumed that they operated within the time period 2021-03-01 till 2021-12-31, containing 136,192 observations. Furthermore, we assumed that 15% of the test centers invoiced for fraudulent test results. For these 'criminal' test centers are simulated high numbers of tests, low positive rates, deviations from Benford's Law and deviations from the assumption of equally distributed last digits. The resulting characteristics of simulated data are shown in table 7. The simulated data does not contain any information from the original data and can be used to test the R-scripts we provided for the analysis.

Table 7: Characteristic of simulated data

| Facility type | test centers | test centers suspected | Tosts [N] | Tests per day | | | |
|---------------|--------------|------------------------|------------|---------------|-------|--------|------|
| raciity type | [N] | of fraud [N] | iests [iv] | Mean | Max | median | [%] |
| Pharmacy | 290 | 4 | 6,795,707 | 137.5 | 1,387 | 98 | 1.42 |

| Doctor's or dentist's office | 357 | 6 | 8,312,270 | 141.4 | 1,663 | 110 | 1.75 |
|------------------------------|-----|----|------------|-------|-------|-----|------|
| Private test center | 160 | 3 | 4,457,176 | 159.3 | 2,019 | 113 | 1.16 |
| Total | 807 | 13 | 19,565,153 | 143.7 | 2,019 | 106 | 1.50 |

Content and structure of the appendix data

The tables, figures and their data, R-scripts and simulated data described in the section Methods and Results are made available in the appendix as open data. The following section describes the structure of the dataset in more detail.

Data Tables

The results of the analyses conducted are provided as data tables. The files are named as the corresponding tables in the Methods and Results sextion as Table_1.tsv , etc.

| File | Description | Download |
|-------------|---|----------|
| Table_1.tsv | Basic statistics of the mean number of tests per day and test centers by facility type, statistically conspicuous and statistically not conspicuous of fraud | |
| Table_2.tsv | Statistics of estimated fixed effects | |
| Table_3.tsv | Summary of classifications into statistical conspicuous versus not conspicuous test centers according to the Poisson regression model used | |
| Table_4.tsv | Number of test centers by facility type, (non) suspected of fraud by the conventional approach, and (non) suspected of fraud by the statistical approach focusing on the deviation from Benford's law. | |
| Table_5.tsv | Number of test centers by facility type, (non) suspected of fraud by the conventional approach, and (non) suspected of fraud by the statistical approach focusing on the deviation from the law of equally distributed last digits. | |
| Table_6.tsv | Positive and negative overlap and share of incrementally identified potentially fraudulent test centers by statistical approaches | |
| Table_7.tsv | Characteristic of simulated data | |

Supporting Materials

The Supporting Materials folder contains

- All figures used in the appendix
- The underlying data for the figures
- The R-scripts used for the analysis
- Sample data for testing the analysis scripts

Figures and Figure Data

The figures listed in the appendix and their underlying data are provided in the "Figures" subfolder as .png and .tsv files.

supporting_material/figures

The figures are named according to the numbering in the appendix as figure_1.png, etc. The underlying data for the figures are provided following the naming of the figures as figure_1_data.tsv, etc. The variables and values of the figure data are explained below.

Variables and valuests of the figure data

| Variable | Туре | Variations | Description |
|---------------------|---------|------------|-------------------|
| testcenter_id / tnr | integer | 1 807 | ID of test center |

| Variable | Туре | Variations | Description |
|---------------------|---------|--|--|
| typ_l | string | Pharmacy, Doctors or dentists office, Private test center | Facility type |
| mean | float | ≥0 | mean number of tests per day |
| p90 | float | ≥0 | 90% percentil |
| leading_figure | integer | 1 9 | first digit |
| last_figure | integer | 0 9 | last digit |
| theoretical_percent | float | ≥0 | theoretical percentage of digit |
| observed_percent | float | ≥0 | observed percentage of digit |
| suspicious_des | integer | 0, 1 | suspicious yes or no (1 = yes, 0 = no) |

R Scripts

The analysis scripts for the individual methods are provided as .r-files in the following subfolder:

supporting_material/scripts

The naming of the scripts is based on the methods used and is linked in the "Analysis script" sections.

Simulated data

Simulated data for testing the evaluation R-scripts are provided. These correspond to the data schema described below.

Data schema of the simulated data

| Variable | Туре | Values | Description |
|---------------|----------|--|--|
| typ | string | Pharmacy, Doctors or dentists office, Private test center | Facility type (pharmacy, doctor's or dentist's office or private test center) |
| tnr | interger | 1 999 | Testcenter identification number |
| date | date | yyyy-mm-dd | Date of the tests in ISO 8601 format |
| week_yr | string | ww_yyyy | Identifer for the calendar week (ww) of the Year (yyyy) |
| getestet | interger | ≥0 | Number of tests per day |
| positiv | interger | ≥0 | Number of positiv tests per day |
| investigation | interger | 0,1 | Indicator for investigation by the health authorities (1 = yes, θ = no) |

The simulated data are available in the folder supporting_material in the file

simulated_data.csv

Further information on the simulated data can be found in the corresponding section.

Metadata

To increase findability, the provided data are described with metadata. The Metadata are distributed to the relevant platforms via GitHub Actions. There is a specific metadata file for each platform; these are stored in the metadata folder:

Metadata/

Versioning and DOI assignment are performed via <u>Zenodo.org</u>. The metadata prepared for import into Zenodo are stored in the <u>zenodo.json</u>. Documentation of the individual metadata variables can be found at https://developers.zenodo.org/representation.

Metadata/zenodo.json

The zenodo.json includes the publication date ("publication_date") and the date of the data status:

```
"dates": [
{
    "start": "2023-09-11T15:00:21+02:00",
    "end": "2023-09-11T15:00:21+02:00",
    "type": "Collected",
    "description": "Date when the Dataset was created"
}
],
```

Guidelines for Reuse of the Data

Open data from the RKI are available on Zenodo.org, GitHub.com, OpenCoDE, and Edoc.rki.de:

- https://zenodo.org/communities/robertkochinstitut
- https://github.com/robert-koch-institut
- https://gitlab.opencode.de/robert-koch-institut
- https://edoc.rki.de/

License

The "Appendix - COVID-19 test fraud detection: Findings from a pilot study comparing conventional and statistical approaches" dataset is licensed under the <u>Creative Commons Attribution 4.0 International Public License</u> | <u>CC-BY.</u>

The data provided in the dataset are freely available, with the condition of attributing the Robert Koch Institute as the source, for anyone to process and modify, create derivatives of the dataset and use them for commercial and non-commercial purposes. Further information about the license can be found in the <u>LICENSE</u> or <u>LIZENZ</u> file of the dataset.