CT Findings in Patients with COVID-19-Compatible Symptoms but Initially Negative qPCR Test

CT-Manifestationen in Patienten mit COVID-19-typischen Symptomen trotz initial negativem qPCR-Test

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ZUSAMMENFASSUNG
Ziel
Mittels dieser Studie soll evaluiert werden, ob Patienten mit starkem Verdacht auf eine SARS-CoV-2 Infektion trotz initial negativer qPCR- und anschließend durchgeführter Thorax-CT verlässig diagnostiziert werden können.

Material und Methoden
In dieser retrospektiven Studie wurden 437 Patienten mit Verdacht auf COVID-19, aber initial negativem qPCR-Test und anschließend durchgeführter Thorax-CT zwischen dem 13. März und 30. November 2020 eingeschlossen. Der Referenzstandard zu den CT-Befunden war die qPCR (mindestens 3 aufeinander folgende qPCR-Tests, im Falle eines Infektionsverdachts durch die CT, um die Sensitivität, Spezifität, den positiven prädiktiven Wert (PPV) und den negativen prädiktiven Wert (NPV) des CT zu bestimmen.

Ergebnisse
Die CT erzielte eine Sensitivität von 100 % (95 % Konfidenzintervall [KI]: 65–100), eine Spezifität von 88 % (95 % KI: 84–90), einen PPV von 12 % (95 % KI: 6–22), einen NPV von 100 % (95 % KI: 99–100) und eine diagnostische Genauigkeit von 88 % (95 % KI: 84–91).

Schlussfolgerung
In diesem speziellen Studiensetting kann die CT trotz initial negativer qPCR eine SARS-CoV-2 Infektion detektieren. Alle Patienten mit einem positiven CT-Befund zeigten einen fortgeschrittenen pulmonalen Befall trotz aktueller negativer qPCR. Die CT kann als diagnostische Methode bei weiterbestehendem klinischem Verdacht auf eine SARS-CoV-2 Infektion trotz negativem qPCR-Test eingesetzt werden und eine Infektion sicher ausschließen.

Kernaussagen:
▪ Die Low-Dose-Thorax-CT kann infizierte Patienten trotz vorliegendem negativen qPCR-Test erkennen und eignet sich aus diesem Grund, besonders im frühen Krankheitsstadium, als additive diagnostische Methode.
▪ Die Low-Dose-Thorax-CT kann eine SARS-CoV-2 Infektion in einer Pandemie verlässlich ausschließen.
▪ Eine zuverlässige Differenzierung zu anderen viralen Parenchymveränderungen ist schwierig.

ABSTRACT
Purpose
To assess whether it is possible to reliably detect patients with strong suspicion of COVID-19 despite initially negative quantitative polymerase-chain-reaction (qPCR) tests by means of computed tomography (CT).

Materials and Methods
437 patients with suspected COVID-19 but initially negative qPCR and subsequent chest CT between March 13 and November 30, 2020 were included in this retrospective study. CT findings were compared to results of successive qPCR tests (minimum of 3 qPCR tests if
CT suggested infection) to determine the sensitivity, specificity, positive predictive value (PPV), and negative predictive value (NPV) of CT for diagnosing COVID-19.

**Results** COVID-19 was diagnosed correctly with a sensitivity of 100 % [95 % confidence interval (CI): 65–100] and a specificity of 88 % [95 % CI: 84–90]. A PPV of 12 % [95 % CI: 6–22] and an NPV of 100 % [95 % CI: 99–100] were determined.

**Conclusion** CT is able to detect COVID-19 before qPCR in initially negative patients in this special study setting. Similar CT findings in COVID-19 and other atypical pneumonias might be challenging, especially in a large hospital and in the case of a high patient volume. Here, it is of utmost importance to identify patients with CT findings typical for COVID-19 pneumonia to a) initiate adequate patient treatment and b) to prevent transmission of infections between patients and between patients and staff. The purpose of this retrospective study was therefore to assess whether low-dose chest CT allows the diagnosis of COVID-19 in symptomatic patients with initially negative qPCR.

**Key Points:**
- Low-dose chest CT is able to diagnose COVID-19 in symptomatic patients even in cases of an initially negative quantitative PCR result and therefore is a fast support method to detect COVID-19, especially in early disease.
- Low-dose chest CT can reliably exclude COVID-19 in a pandemic setting.
- CT does not always ensure a reliable differentiation from other viral diseases.

**Citation Format**

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**Introduction**

The coronavirus disease 2019 (COVID-19) pandemic caused by the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has kept the entire world in suspense. The gold standard of COVID-19 diagnosis is quantitative polymerase chain reaction (qPCR). However, the sensitivity of qPCR depends on different factors, such as disease stage, the method of sample acquisition, and the type of test in use [1]. For this reason, the Fleischner society recommends additionally performing a low-dose chest CT examination in patients with clinical presentation and/or medical history indicative of COVID-19 or in patients experiencing an aggravation of symptoms [2]. CT examinations may indicate COVID-19 pneumonia earlier than qPCR in specific cases since they allow independent assessment of the involvement of the lung parenchyma [3, 4]. Typical CT findings of COVID-19 pneumonia are ground-glass opacities, infiltrations, crazy paving pattern, and predominant involvement of the lower lobes and the periphery [3, 5–11]. However, these imaging findings are non-specific and may appear in a similar way in other diseases [9, 12], like other viral pneumonias. In an early stage of a viral infection, imaging patterns caused by different viruses are similar and cannot be distinguished easily by means of CT [13]. Besides in a SARS-CoV-2 infection, ground-glass opacities occur in 75 % of patients with cytomegalovirus, 50–75 % with adenovirus, and up to 25 % with influenza [14]. Consolidations can also occur in 50–75 % of patients with adenovirus and up to 25 % with influenza [14]. Hence, ground-glass opacities and consolidation are typical but not specific for COVID-19.

Since qPCR tests might provide false-negative results and since CT might not allow differentiation between COVID-19 pneumonia and other pneumonias, infection control in a pandemic situation might be challenging, especially in a large hospital and in the case of a high patient volume. Here, it is of utmost importance to identify patients with CT findings typical for COVID-19 pneumonia to a) initiate adequate patient treatment and b) to prevent transmission of infections between patients and between patients and staff. The purpose of this retrospective study was therefore to determine the sensitivity, specificity, positive predictive value (PPV), and negative predictive value (NPV) of CT for diagnosing COVID-19.

**Materials and methods**

**Patient cohort**

The institutional review board approved this retrospective study cohort and waived patient informed consent. Approval was granted for analysis of the CT images, medical records, and laboratory data. Patient diagnosis and treatment have not been modified for the purpose of this study. This study includes patients that underwent a chest CT in our department between March 13 and November 30, 2020. Thus, this study covers the period of the entire first wave and the “lockdown light” phase of the second wave (rise of the second wave) in Germany in 2020. Patients were admitted through the emergency department of the hospital or via different hospital units for other elective reasons for admission.

The inclusion criteria were an initially negative qPCR test and suspected SARS-CoV-2-infection based on respiratory symptoms, which included cough, shortness of breath, and/or need for oxygen supply, and a chest CT examination after an initial qPCR test. The exclusion criteria were follow-up examinations and CT examinations following a positive qPCR test result, see Fig. 1. Patients with an initially negative, but subsequently positive qPCR test, without any CT examination during their treatment were not included in this study.

**qPCR**

At our clinic, patients are tested for suspected COVID-19 at the time of referral using a standardized qPCR method according to a standardized protocol of our virology department [15]. Initial qPCR tests were performed in the emergency room on respiratory tract specimens (nasopharyngeal and oropharyngeal) at the virology department. Further tests were taken at the different hospital units. In our hospital, even in the early pandemic phase, the qPCR tests were usually available after 6–8 hours. Results of the qPCR...
Antibody tests were performed in patients in poor clinical condition with suspected infection despite multiple negative qPCR tests. At the beginning of the pandemic, it was not yet possible to make a statement about the test accuracy of the COVID-19 qPCR test, since no evaluated comparative method was available. Here, in addition to CT examination, antibody tests were also carried out occasionally in order to detect a false-negative qPCR test in the case of antibody detection. The presence of IgA and IgG antibodies directed against corona epitopes was tested by the virology department.

CT acquisition

Due to the radiation exposure during CT examinations and the risk of stochastic and deterministic radiation damage including risk of radiation-induced cancer, CT was not performed as the primary diagnostic method for the detection of infection in our hospital in every patient. Only if patients had clinical indications (e.g., worsening of symptoms resulting in unstable patients, oxygen supply, intubation, and life-threatening diagnoses like pulmonary embolism), CT was performed. Even though we occasionally had many patients with respiratory symptoms in the emergency department, the indication for CT was restrictive.

All included patients were examined on one of three state-of-the-art CT scanners (Somatom Definition Edge – scanner A, during day period – or Somatom Definition Flash (scanner B) and Somatom Definition AS (scanner C), during the night and on weekends (all Siemens Healthineers, Forchheim, Germany). Patients were imaged in supine position with elevated arms and in breath-hold technique following maximal inspiration. The scan range was defined from the lung apex to the base. The applied CT protocol parameters were 100–120 kVp, with automatic exposure control. Images were reconstructed iteratively using ADMIRE (scanner A) and SAFIRE (scanner B, C) (both Siemens Healthineers, Forchheim, Germany). According to the national recommendations for the diagnosis of COVID-19, a non-contrast-enhanced chest CT examination was performed. Only in cases with an additional diagnostic question (n = 3, e.g. lung embolism), contrast agents were applied [16].

CT image evaluation

CT images were reviewed in consensus by one resident and two board-certified radiologists. A structured reporting system is used in the institution to evaluate each CT scan according to the same criteria. The following CT criteria were considered: a) lesion characteristics: ground-glass opacities, consolidation, crazy paving pattern, interlobular septal thickening, air bronchogram, bronchiectasis, cavens, pleural thickening and pneumothorax, b) lesion location: left, right, or bilateral lung parenchyma, peripheral or central accentuation. All additional pathologies were described in a “further findings” section.

Based on the literature, DRG, and the RSNA recommendations, the suspicion of an infection was raised in the presence of the following CT patterns [3, 5–8, 10, 11, 17]: ground-glass opacities, consolidation, crazy paving pattern, and bilateral patterns with emphasis on the lung periphery. Consolidations and crazy paving patterns were optional and not necessarily suspicious as long as bilateral, peripherally accentuated ground-glass opacities with a patchy appearance were present. In the presence of non-typical changes, e.g., evidence of bacterial infection such as consolidations in one lobe, tree-in-bud phenomenon, peripheral avoidance, ubiquitous mosaic ground-glass patterns with septal thickening and enlarged heart and/or unilateral occurrence, patients were classified as non-infectious. Especially in an early state of infection, unilateral occurrence may also be shown by SARS-CoV-2 in-

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fection, as published after this study, but we internally classified unilateral occurrence as unlikely at the time of study [18].

Data collection

CT image data, volumetric CT dose index (CTDIvol), dose length product (DLP), and scan length were collected in the local picture archive and communication system (SECTRA Medical, Sweden). Patient weight and height were documented.

The effective dose was calculated using the tube potential-specific conversion factors published by Deak et al. \(k_{100kVp} = 0.0144 \text{ mSv/(mGy cm)}\), \(k_{120kVp} = 0.0145 \text{ mSv/(mGy cm)}\), using the tissue weighting factors in ICRP publication 103 [19].

Data analysis

Statistical analysis was performed using SPSS version 27 (SPSS Inc. Chicago, IL) and Microsoft Excel 2016 (Redmond, WA, USA). For continuous values, mean and standard deviation with the corresponding ranges (minimum-maximum) are provided. The sensitivity, specificity, positive and negative predictive values (PPV/NPV), and disease prevalence including 95% confidence intervals (CI) were calculated by applying the contingency table with the Fisher’s exact and Wilson-Brown test using qPCR results as the reference with GraphPad Prism 8.0 (GraphPad Software, Inc., San Diego, California, USA). A contingency table with Fisher’s exact test as well as Chi-square test was performed to assess differences in CT findings between COVID-19 true-positive, false-positive, and true-negative patients according to the structured report. The level of significance was \(p < 0.05\). Tests were adjusted for all pairwise comparisons using Bonferroni correction.

Results

Patient population

A total of 437 examinations were included in the final study cohort (see Fig. 1). Patient characteristics and CT exposure parameters can be found in Table 1.

qPCR results and antibody test results

All included patients were initially negative according to the primary qPCR test. Despite the initially negative qPCR test, subsequent qPCR tests were positive in seven patients. Therefore, the initial qPCR was false-negative in 7/437 (1.6%) patients.

In 9/437 (2%) patients, antibody tests were carried out during hospitalization. Two of the tests detected IgA or IgG antibodies.

Diagnostic performance of CT

Of all included patients, 60/437 (14%) patients had CT findings in accordance with COVID-19. CT detected all of the abovementioned 7/60 (12%) patients with positive subsequent qPCR tests correctly (CT true positives) (see Fig. 2). In 53/60 (88%) patients, the qPCR remained negative (CT false positives). In 377/437 (86%) patients, CT findings were not in accordance with COVID-19 (CT true negatives). There were no false-negative results on CT evaluation.

For the detection of COVID-19 with chest CT, the sensitivity was 100% [95% CI: 65–100%], the specificity was 88% [95% CI: 84–90%], the PPV was 12% [95% CI: 6–22%], and the NPV was 100% [95% CI: 99–100%]. The prevalence of COVID-19 in the cohort was 2% [95% CI: 1–3%]. The PLR equaled 8.11 [95% CI: 6.31–10.44], whereas the NLR was 0 (see Table 2).

CT findings

CT findings for the total cohort and separated into CT true-positive, true-negative and false-positive patients are presented in

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After categorization of the patients according to the typical manifestations of COVID-19, some pulmonary findings were significantly more frequent in CT true-positive (CTP) patients compared to CT true-negative (CTN) patients, see Table 4. These were: ground-glass opacities (100%:51% (CTP:CTN), \( p \leq 0.01 \)), consolidation (100%:46%, \( p \leq 0.01 \)), and crazy paving pattern (57%:6%, \( p \leq 0.01 \)). Differences regarding spatial distribution were also notable. Bilateral manifestation (100%:36%, \( p \leq 0.01 \)) as well as emphasis of the peripheral (100%:20%, \( p \leq 0.01 \)) lung lobes were significantly more common in patients with true-positive CT findings. Pleural thickening, bronchiectasis, caverns, pleural and pericardial effusion were seen equally in both groups.

In 206/437 (47%) patients, CT findings were consistent with pneumonia other than COVID-19. In 53/437 (12%) patients, no findings on CT were found despite their symptoms.

CT findings of false-positive patients
In total, 53 patients were classified as false positives by CT. In these patients ground-glass opacities were visible in 96%, consolidations in 87%, and thickened interlobular septa in 68% (Fig. 3). In some patients, other viral pathogens, such as cytomegalovirus or parenchymal changes due to an underlying disease, were found to be the cause of the parenchymal changes.

CT findings of true-positive patients
The most common manifestations were ground-glass opacities and consolidation (each 100%), crazy paving pattern, thickened interlobular septa, and air bronchogram (all 57%). A bilateral manifestation (100%) was visible with emphasis in the periphery.
In this special study setting, CT was able to reliably detect the presence of COVID-19 in patients with initially negative qPCR. Patients with initially false-negative qPCR but active SARS-CoV-2 infection showed typical and in some cases already pronounced pulmonary patterns of COVID-19. The study was performed in a tertiary care hospital in a pandemic situation, where fast differentiation between SARS-CoV-2 infected and non-infected patients was necessary to prevent disease transmission within the hospital. Despite a rush of patients, the hospital was able to handle the patients and treat those in need. At the time of the study, no data on the sensitivity and specificity of the qPCR test was available due to the lack of a comparable method. Nevertheless, there were patients in whom infection was ruled out by qPCR but who were still strongly suspected of having an infection (e.g., first-degree contacts, steadily worsening respiratory symptoms with oxygen demand, additional loss of taste and smell, and diarrhea). In these patients, there was a clinically justified indication to perform a CT examination. Therefore, in these cases, we wanted to investigate how likely an infection is to be visible on CT despite a negative qPCR test and whether CT can identify infected patients.

Our data indicate a high sensitivity (100 %) with a moderate specificity of 88 % with respect to diagnosing COVID-19 using CT in patients with initially false-negative qPCR. In a study by Long et al., all patients with initially negative qPCR but subsequently detected the patients as infected after the second or third test. (100 %) and lower lobes (57 %) (Fig. 4, 5). In all cases, qPCR detected the patients as infected after the second or third test.

**CT findings of CT true-positive patients compared to CT false-positive patients**

No statistically significant differences in lung manifestations between CT true-positive and false-positive patients were determined, see Table 4.

**Discussion**

In this special study setting, CT was able to reliably detect the presence of COVID-19 in patients with initially negative qPCR.
positive qPCR already had positive CT findings for COVID-19 pneumonia at their initial presentation, resulting in an identical sensitivity of 100 % in these six patients [20]. Unfortunately, they do not present any data on the specificity.
Published sensitivities reported by Li et al. (80%), Bai et al. (70–93%), Kim et al. (94%), Fang et al. (98%), Long et al. (97%), and Caruso et al. (97%) [20–25] and specificities reported by Kim et al. (37%), Caruso et al. (56%), Li et al. (82%), and Bai et al. (93–100%) [21, 22, 25, 26] cannot be compared directly to the cohort presented here. A limitation is the pre-selection of our cohort in which we included only patients with suspected disease who initially had a negative qPCR test result. Patients with an initially positive qPCR result were excluded in contrast to the last-mentioned studies. This reduces the total number of patients as well as the number of SARS-CoV-2-infected patients in our cohort, thereby limiting a comparison with other published studies in terms of infestation pattern and onset of symptoms.

In this study, the suspicion of an infection was assessed based on the literature and the RSNA recommendations [5–8, 17]. Compared to the abovementioned studies [20–25], the presence of the same CT patterns was evaluated (e.g., peripheral distribution, ground-glass opacities, crazy paving pattern, vascular thickening, consolidations). Although there are now different categorization strategies (CO-RADS, RSNA, DRG recommendations), they all assess the same manifestations more or less. We are aware that consensus evaluation with 3 radiologists does not correspond to everyday clinical practice. Since experience was limited at the beginning of the pandemic, it was important for us to make a consensus decision in order to have as few false-negative patients as possible. In this way, we aimed to interrupt a potential chain of infection. One reason for our high sensitivity but moderate specificity could be the internal procedure of strict evaluation of typical pulmonary changes. Nevertheless, our results show no significant difference in the presence of the patterns between true-positive and false-positive patients but significant differences between false-positive and true-negative patients. Therefore, we conclude that the configuration of the parenchymal changes is decisive for the evaluation of the presence of acute infection. For example, ubiquitous ground-glass opacities are less likely to be present at the onset of infection. In contrast, round-shaped ground-glass opacities, which are predominantly found in the periphery and the basal lobes, are more likely to be associated with acute SARS-CoV-2 infection [27, 28]. Nevertheless, other respiratory viral infections may be present and can cause similar parenchymal patterns in comparison to COVID-19. These similar viral parenchymal patterns will probably make it more difficult to diagnose SARS-CoV-2 infection by CT in the future after the pandemic situation when different viral infections are present in the patient population. In addition, preexisting conditions of the lung parenchyma that can lead to a false diagnosis may be present [29]. Especially in cases with mild parenchymal changes, a misinterpretation is possible. Hence, in the case of uncertain CT findings, patients were classified and treated as infected and potentially contagious to ensure that no positive cases were missed. Thus, the sensitivity is increased at the expense of specificity. One source of error could be the swabbing procedure for the qPCR test. Insufficient execution might result in false-negative results. Nevertheless, the number of correct negative swabs as well as the presence of a trained and permanent team in the emergency room show that the swabs were taken correctly and in a qualified manner. One explanation for false-negative qPCR tests discussed in the literature could be the viral load of the sample and the amount of sputum at the time of the test [30]. Especially in the first days of infection, these false-negative results can occur, as studies have already shown [31].

In addition, the pretest probability is a relevant factor. The relatively low number of false-negative findings is related to the low prevalence at our hospital at the time of the study. However,
a high local prevalence also means a higher probability of false-negative tests [31].

**Clinical Relevance**

In conclusion, CT can identify infected patients before qPCR in this particular study setting in which patients with an initially negative qPCR test underwent CT for additional diagnosis in a pandemic situation. Yet, CT cannot perfectly distinguish between COVID-19 and other respiratory infections in a cohort with initially negative qPCR. Especially patients with initially false-negative qPCR who tested positive only after repeated qPCR tests illustrate the necessity for CT examinations in patients with COVID-19-compatible symptoms but negative initial qPCR test. Nevertheless, due to the radiation exposure during CT examinations and the risk of stochastic and deterministic radiation damage including the risk of radiation-induced cancer, CT cannot replace qPCR tests as a screening method. Yet, CT can help to interrupt infection pathways and identify initially false-negative patients to ensure adequate treatment.

**Conflict of Interest**

The authors declare that they have no conflict of interest.

**References**

[1] Lippi G, Simundic A, Plebani M. Potenzial preanalytical and analytical vul-
nerabilities in the laboratory diagnosis of coronavirus disease 2019 (COV-

[2] Rubin GD, Ryerson CJ, Haramati LB et al. The Role of Chest Imaging in Pa-

tient Management During the COVID-19 Pandemic. Chest 2020; 158:

106–116


the diagnosis of COVID-19 – a systematic, prospective comparison with

PCR. Dtsch Aerzteblatt Online 2020; 117: 389–395


215: 338–343


J Roentgenol 2020; 215: 87–93


tients with suspected corona virus disease 2019 (COVID-19). Br J Radiol

2020; 93: 20200243


E45


masquerades. BJR:case Reports 2020; 6: 20200667

[13] Li X, Fang X, Bian Y et al. Comparison of chest CT findings between

COVID-19 pneumonia and other types of viral pneumonia: a two-center

retrospective study. Eur Radiol 2020; 30: 5470–5478

[14] Koo HJ, Lim S, Choe J et al. Radiographic and CT features of viral pneu-

monia. Radiographics 2018; 38: 719–739


virus (2019-nCoV) by real-time RT-PCR. Eurosurveillance 2020; 25: 1–8


of the Thoracic Imaging Section of the German Radiological Society for

clinical application of chest imaging and structured CT reporting in the

COVID-19 pandemic. RöFo – Fortschrritte Auf Dem Gebiet Der Röntgen-

strahlen Und Der Bildgebung Verfahren 2020; 192: 633–640


Expert Consensus Document on Reporting Chest CT Findings Related to

COVID-19: Endorsed by the Society of Thoracic Radiology, the American

College of Radiology, and RSNA. Radiol Cardiothorac Imaging 2020; 2:

e200152

[18] El Homsi M, Chung M, Bernheim A et al. Review of chest CT manifesta-

tions of COVID-19 infection. Eur J Radiol Open 2020; 7: 100239

[19] Deak PD, Smal Y, Kalender WA. Multisection CT Protocols: Sex- and

Age-specific Conversion Factors Used to Determine Effective Dose from


(COVID-19): rRT-PCR or CT? Eur J Radiol 2020; 126: 108961


331

[22] Bai HX, Hsieh B, Xiong Z et al. Performance of Radiologists in Differenti-

ating COVID-19 from Non-COVID-19 Viral Pneumonia at Chest CT.

Radiology 2020; 296: E46–E54

[23] Kim H, Hong H, Yoon SH. Diagnostic Performance of CT and Reverse

Transcriptase Polymerase Chain Reaction for Coronavirus Disease 2019:


Comparison to RT-PCR. Radiology 2020; 296: E115–E117


Rome, Italy. Radiology 2020; 296: E79–E85

[26] Kim JY, Choe PG, Oh Y et al. The First Case of 2019 Novel Coronavirus

Pneumonia Imported into Korea from Wuhan, China: Implication for


Radiol Cardiothorac Imaging 2020; 2: e200208

[28] Schmitt W, Marchiori E. COVID-19: Round and oval areas of ground-


[29] Iino M. Interstitial pneumonitis associated with the immunomodulatory


