RESEARCH PAPER

Are presymptomatic SARS-CoV-2 infections in nursing home residents unrecognised symptomatic infections? Sequence and metadata from weekly testing in an extensive nursing home outbreak

Judith H. van den Besselaar[1], Reina S. Sikkema[2], Fleur M. H. P. A. Koene[3,4], Laura W. van Buul[5], Bas B. Oude Munnink[2], Ine Frénay[6], René te Witt[7], Marion P. G. Koopmans[2], Cees M. P. M. Hertogh[5], Bianca M. Buurman[1]

[1]Department of Internal Medicine, Section of Geriatric Medicine, Amsterdam Public Health Research Institute, Amsterdam University Medical Center, 1105 AZ Amsterdam, the Netherlands
[2]Department of Viroscience, Erasmus Medical Center, 3015 CN Rotterdam, the Netherlands
[3]Department of Medical Microbiology, Amsterdam University Medical Center, 1105 AZ Amsterdam, the Netherlands
[4]Department of Infectious Diseases, Public Health Laboratory, Public Health Service of Amsterdam, 1018WT Amsterdam, The Netherlands
[5]Department of Medicine for Older People, Amsterdam Public Health Research Institute, Amsterdam University Medical Center, 1081 BT Amsterdam, The Netherlands
[6]Regional Laboratory for Medical microbiology (RLM) Dordrecht- Gorinchem, 3318 AT Dordrecht, The Netherlands
[7]Eurofins|NMDL-LCPL, 2280 CA Rijswijk, The Netherlands

Address correspondence to: Judith H. van den Besselaar, Department of Internal Medicine, Section of Geriatric Medicine, Amsterdam Public Health Research Institute, Amsterdam University Medical Center, Meibergdreef 9, 1105 AZ Amsterdam, the Netherlands. Tel: 0031 6 831 65 744. Email: j.h.vandenbesselaar@amsterdamumc.nl

Abstract

Background: Sars-CoV-2 outbreaks resulted in a high case fatality rate in nursing homes (NH) worldwide. It is unknown to which extent presymptomatic residents and staff contribute to the spread of the virus.
Aims: To assess the contribution of asymptomatic and presymptomatic residents and staff in SARS-CoV-2 transmission during a large outbreak in a Dutch NH.
Methods: Observational study in a 185-bed NH with two consecutive testing strategies: testing of symptomatic cases only, followed by weekly facility-wide testing of staff and residents regardless of symptoms. Nasopharyngeal and oropharyngeal testing with RT-PCR for SARs-CoV-2, including sequencing of positive samples, was conducted with a standardised symptom assessment.
Results: 185 residents and 244 staff participated. Sequencing identified one cluster. In the symptom-based test strategy period, 3/39 residents were presymptomatic versus 38/74 residents in the period of weekly facility-wide testing (*P*-value < 0.001). In total, 51/59 (91.1%) of SARS-CoV-2 positive staff was symptomatic, with no difference between both testing strategies (*P*-value 0.763). Loss of smell and taste, sore throat, headache or myalga was hardly reported in residents compared to staff (*P*-value <0.001). Median Ct-value of presymptomatic residents was 21.3, which did not differ from symptomatic (20.8) or asymptomatic (20.5) residents (*P*-value 0.624).
Conclusions: Symptoms in residents and staff are insufficiently recognised, reported or attributed to a possible SARS-CoV-2 infection. However, residents without (recognised) symptoms showed the same potential for viral shedding as residents with

symptoms. Weekly testing was an effective strategy for early identification of SARS-Cov-2 cases, resulting in fast mitigation of the outbreak.

Keywords: nursing home, COVID-19, presymptomatic, residents, staff, older people

Key Points

- Subjective symptoms such as sore throat or loss of smell/taste are hardly reported in SARS-CoV-2 positive nursing home (NH) residents.
- Facility wide testing of staff revealed limited presymptomatic or asymptomatic SARS-CoV-2 cases.
- Staff did not recognise (self) report or attribute symptoms to SARS-CoV-2 in NH residents and themselves.
- Weekly facility wide testing is an effective strategy for early identification of SARS-CoV-2 cases.

Introduction

Worldwide, nursing homes (NHs) are facing outbreaks of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) with high case fatality rates [1, 2]. The current ECDC-guideline recommends expanded viral testing of asymptomatic residents in NHs if a single new case of a SARS-CoV-2 infection is detected, based on data of previous NH outbreaks which suggest an important role for presymptomatic spread of SARS-COV-2 among residents [3–9]. However, it remains unknown to which extent asymptomatic and presymptomatic cases contribute to the spread of SARS-CoV-2. Also, specifically in the NH setting, it remains unclear to what extent asymptomatic cases are truly without symptoms. Sole reliance on symptoms for testing in NHs could be insufficient because self-reporting of complaints is often compromised in residents due to limited ability to communicate (e.g. in residents with dementia) [10]. The Dutch guideline for COVID-19 in NHs states that only residents with possible symptoms of SARS-CoV-2 should be tested [11] and no policy for testing of asymptomatic residents or staff is facilitated in the Netherlands.

Multiple reports have been published about the prevalence of asymptomatic and presymptomatic residents and staff in NHs after the implementation of a facility-wide testing strategy during an outbreak [4, 6, 9, 12, 13]. The prevalence of asymptomatic staff and residents differed from single cases to up to half of the infected cases. Low cycle threshold (Ct) values were found in asymptomatic and presymptomatic cases, suggesting potential of viral shedding [6, 9]. A large registry of 857 Dutch residents with confirmed SARS-CoV-2 showed that 93% of cases expressed any of the symptoms cough, shortness of breath, or fever. A large range of other symptoms were also reported such as fatigue, diminished intake, gastro-intestinal symptoms, malaise or rhinorrhea [14]. However, the presentation of SARS-CoV-2 can be difficult to recognise in NH residents, which can cause delay in testing, isolation and treatment [14, 15]. In addition, during a community-wide outbreak it can be difficult to distinguish residential outbreaks from multiple introductions without sequencing of viruses from cases [16].

Viral spread by presymptomatic or unrecognised symptomatic cases has important implications for Personal Protective Equipment (PPE) use, facility-wide testing and isolation measures in NHs for the prevention of outbreaks. The aim of this study is to analyse the contribution of presymptomatic spread of SARS-CoV-2 in all staff and residents of a NH in the Netherlands by serial weekly point prevalence surveys, PCR and sequencing.

Methods

Setting and study population

The study took place in a 185-bed NH in the province South Holland which provides long-term care and is specialised in dementia care. All residents and staff working during the outbreak were invited to participate in the study. Data were collected retrospectively before May 18th and prospectively from May 18th onwards. NH details are presented in Appendix 1, Supplementary data are available in *Age and Ageing* online.

SARS-CoV-2 testing and analysis

Two phases in the NH test strategy can be distinguished: First, until May 11th, a symptom-based testing strategy was followed, according to national guidelines: cases were tested when they experienced any symptoms. The only exception of this strategy was at May 6th: at the ward where the outbreak started, all negative residents were tested regardless of symptoms. Second, from May 12th the NH implemented a policy of facility-wide weekly testing in addition to the symptom-based testing strategy, implying SARS-CoV-2 testing of all residents without a previous positive test and regardless of the presence of any symptoms. Staff was tested regardless of symptoms in the week of May 18th and June 1st.

Samples were transported to collaborating laboratories at the end of each test day, where they were tested for SARS-CoV-2 polymerase chain reaction (PCR) targets. Three

different laboratories collaborated because of the large number of tests which were conducted: As a result, different PCR platforms were used, however, the used targets were similar (RdRp- gene, E-gene, N- gene; see Appendix 2, Supplementary data are available in *Age and Ageing* online).

Sequencing, phylogenetic analysis and cluster definition

PCR-positive samples with Ct-value below 32 were selected for sequencing using a SARS-CoV-2 specific amplicon based Nanopore sequencing approach, as previously described [17]. The consensus genome was generated only including positions with a coverage >30 as described previously [18]. Additional details on sequencing methods are provided in Appendix 3, Supplementary data are available in *Age and Ageing* online.

Data collection

A standardised symptom-assessment form of 16 symptoms was completed by the research team for each assenting resident, using electronic health record review. Staff was invited to complete a first questionnaire electronically (via email) in the week of May 18th (Appendices 4 and 5, Supplementary data are available in *Age and Ageing* online).

A participant was classified symptomatic if he had at least one new or worsened symptom in the 14 days prior to a positive test result. A participant was classified asymptomatic if no new or worsened symptoms were present and no symptoms would develop in the 14 days following the positive test. Participants were classified pre-symptomatic if they had no symptoms at moment of testing, but developed symptoms in the 2 weeks following a positive test [19].

Analyses

Data are reported as mean/median with range and standard deviations (SD) and counts with percentages as appropriate. Differences between groups were assessed with student's T-test and Mann–Whitney U for continuous variables and Chi-square test for categorical data. Differences were considered statistically significant at $P < 0.05$ (two-tailed). All analyses were done using SPSS, version 26 (IBM, Armonk, NY) and Excel.

Ethics

Written information about the study was sent out to residents and their legal representatives at May 18th, with the possibility to opt-out. Health care professionals were asked informed consent for participating in the study prior to digital questionnaire completion. The Medical Ethics Committee of the VU University Medical Centre in Amsterdam reviewed the study protocol and confirmed that the study does not fall under the scope of the Medical Research Involving Human Subjects Act.

Results

At April 29th, when the first resident tested positive for SARS-CoV-2, 185 residents lived and 384 staff worked in the NH. Four legal representatives of residents and 34 staff members declined participation. Baseline characteristics are described in Table 1. Residents who tested positive for SARS-CoV-2 were older and more likely to have cognitive impairment. Staff positive for SARS-CoV-2 consisted mostly of (registered) health care assistants and health-care aids. Appendix 6 (Supplementary data are available in *Age and Ageing* online) shows the STROBE diagram of participating residents and staff.

Introduction of the virus and outbreak

The first positive resident (29 April) had been admitted from 17 to 23 April at the geriatric department of the local hospital with a urosepsis. She had a negative PCR for SARS-CoV-2 and her chest X-ray was classified as CORADS-1, suggesting a very low probability of COVID-19 [20]. On April 29th, she developed a fever and was readmitted to the hospital, where retrospectively an outbreak had occurred, and tested positive for SARS-CoV-2. Previously, three NH staff members tested positive for SARS-CoV-2 in April, but none of them worked in the period they were contagious. Figure 1 shows the timeline with date of onset of COVID-19 for participating residents and staff and the NH policy.

Sequencing

In total, 53 sequences of residents and NH staff were available. In addition, 7 sequences of the hospital outbreak were generated. All sequences cluster together, also sequences detected at the geriatric department of the hospital outbreak were near identical. Two subclusters appear to be present in sequences of residents and staff, without differences when considering wards where residents lived and staff worked (Figure 2).

Symptomatic, presymptomatic and asymptomatic cases during symptomatic and weekly testing strategy

Results of the standardised symptom-assessment are presented in Table 2. Except for the symptoms fever and nausea, residents and staff showed different prevalence for all symptoms. Because of retrospective and prospective data collection in staff, we performed sensitivity analyses which compared symptoms in prospective and retrospective questionnaires. Also we repeated comparisons between positive staff and residents only using prospective questionnaires of staff.

Staff was tested twice regardless of symptoms, while residents were tested four times regardless of symptoms: We performed a third sensitivity analysis where we compared residents/staff who were tested on the dates of the point prevalence surveys regardless of symptoms to residents/staff
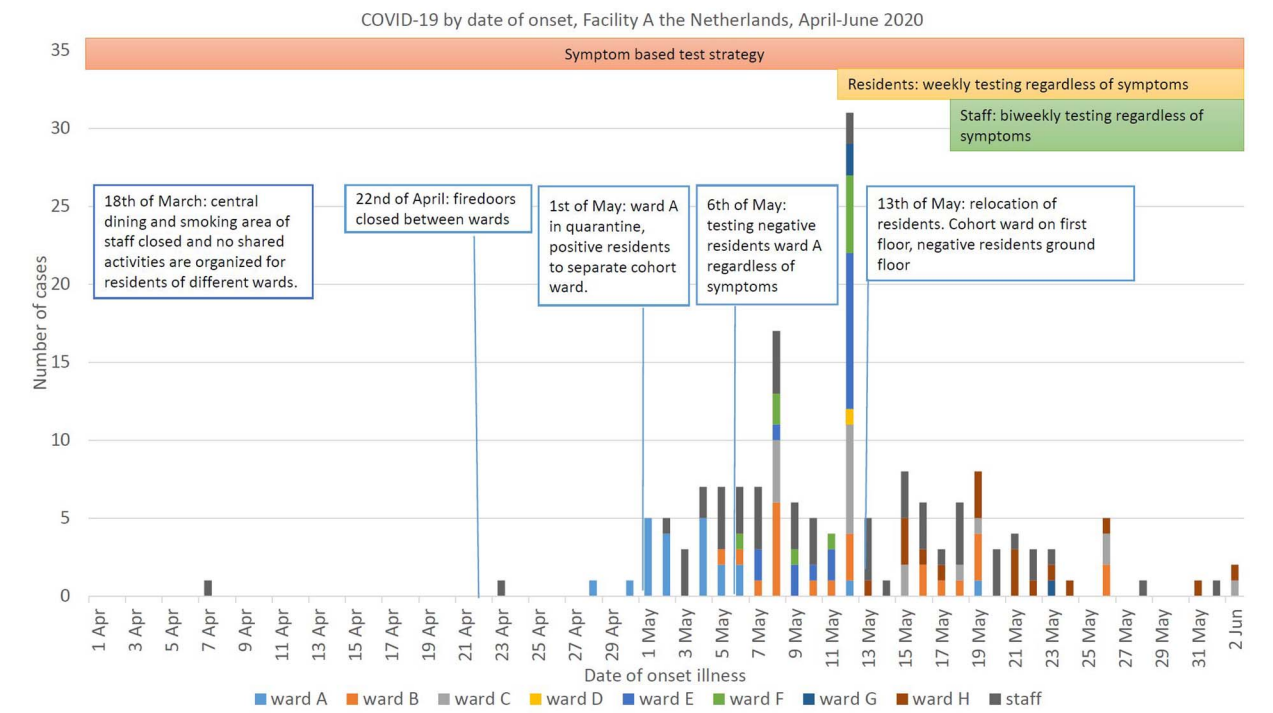
Figure 1. COVID-19 by date of onset and NH policy, shows the date of onset of COVID-19 for participating residents of the different wards and participating staff from the 15th of April until the 2nd of June. Key changes in NH policy for infection prevention and testing are indicated. On May 13th, facility management decided to move all positive tested residents to the first floor of the building, while residents who tested negative were moved to the ground floor of the building. PPE used on the first floor included isolation gown, gloves over the wrists, goggles and a surgical mask; on the ground floor surgical masks and gloves were used.

Table 1. Baseline characteristics residents and staff

	SARS-CoV-2 test results		P-value (95% CI)
Residents	Positive N = 113	Negative N = 68	
Age (median/range)	85.0 (44–99)	81.5 (48–100)	0.001 (−9.009; −2.237)
Female n (%)	82 (72.6)	50 (73.5)	0.888
Coexisting conditions			
Pulmonary disease n (%)	12 (10.6)	3 (4.4)	0.142
Cardiovascular disease n (%)	40 (35.4)	15 (22.1)	0.059
Cerebrovascular disease n (%)	23 (20.4)	9 (13.2)	0.224
Diabetes n (%)	18 (15.9)	18 (26.5)	0.085
Cognitive impairment n (%)	104 (92.0)	53 (77.9)	0.007
Reduced kidney function n (%)	7 (6.2)	2 (2.9)	0.329
Obesity n (%)	5 (4.4)	8 (11.8)	0.064
Staff[a]	Positive N = 56	Negative N = 188	
Age (median/range)	43.0 (18–74)	46.5 (18–74)	0.853 (−5.764; −3.942)
Female n (%)	47 (83.9)	175 (93.1)	0.036
Profession, n (%)			0.027
Health care assistants and aids	39 (69.6)	88 (46.8)	
Nurse	3 (5.4)	11 (5.9)	
Physical therapist	0	7 (3.7)	
Physician	0	6 (3.2)	
Other[b]	14 (24.6)	76 (40.4)	
Reporting contact with Covid-19 suspected or confirmed residents, n (%)			0.296
Yes	43 (76.8)	159 (84.6)	
No	5 (8.8)	8 (4.3)	
Unknown	8 (14.0)	21 (11.2)	

[a]34 Staff members declined participation, 106 staff did not complete the questionnaire. [b]Staff working in kitchen, logistics, occupational therapists, psychologists, management.

Figure 2. Zoom-in of Dutch phylogenetic tree, with sequences of NH A in red (clients) and orange (employees). Sequences in blue originate from the related hospital outbreak. Sequences in black originate from a Dutch national reference database.

who were tested at all other dates in the study period. This did not alter results (Appendix 7, Supplementary data are available in *Age and Ageing* online).

A significant difference in presymptomatic residents was found between the two testing strategies (*P*-value < 0.001). Before the start of facility-wide weekly testing, 39 residents tested positive: 36 (92.3%) were symptomatic and 3 (7.7%) residents were presymptomatic. The three presymptomatic residents were tested at May 6th when all residents of the ward where the outbreak started were tested regardless of symptoms. In the period of weekly testing, 74 residents tested positive, of which 29 (39.2%) were symptomatic at the time of testing, 38 (51.4%) were presymptomatic and 7(9.5%) were asymptomatic.

A total of 56 staff tested positive and completed the questionnaire: 51 (91.1%) were symptomatic at the moment of testing, 2 (3.9%) were pre-symptomatic and 3 (5.9%) staff members were asymptomatic. No difference in symptomatic,

presymptomatic and asymptomatic staff members was found between symptom based or additional weekly testing strategy (*P*-value 0.763).

Symptom onset and presentation with symptomatic and weekly testing strategy

Until May 11th, 39 residents tested positive and all developed symptoms. Symptoms developed between 6 days before testing and 3 days after testing, with a median of development of symptoms the day before the test (interquartile range 2 days to 1 day before test) (Figure 3A). After the addition of weekly testing regardless of symptoms, 74 residents tested positive of which 67 residents developed symptoms between 11 days before testing and 8 days after testing, with a median of development of symptoms the day of the test (interquartile range 2 days before the test to 3 days after the test) (Figure 3B). The time between onset of symptoms and test

Table 2. Symptom assessment of residents and staff with a positive SARS-CoV-2 PCR-test

Symptom assessment n (%)	Residents N = 113			Staff N = 56			P-value
Symptomatic	65 (57.5)			51 (91.1)			<0.001
Presymptomatic	41 (36.3)			2			
Asymptomatic	7 (5.2)			3[a]			
Cough	31 (27.4)			26 (46.4)			0.014
Dyspnea	13 (11.5)			20 (35.7)			<0.001
Fever	30 (26.5)			15 (26.8)			0.974
Saturation	27 (23.9)			NA			
Delirium	16 (14.2)			NA			
Chills	4 (3.5)			22 (39.3)			<0.001
Malaise	25 (22.1)			24 (42.9)			0.005
Fatigue	19 (16.8)			42 (75.0)			<0.001
Myalgia	2 (1.8)			26 (46.6)			<0.001
Headache	5 (4.4)			36 (64.3)			<0.001
Sore throat	2 (1.8)			21 (37.5)			<0.001
Nasal congestion	15 (13.3)			34 (60.7)			<0.001
Diarrhoea	10 (8.8)			14 (25.0)			0.005
Nausea	9 (8.0)			7 (12.5)			0.343
Diminished intake	17 (15.0)			23 (41.1)			<0.001
Loss of smell or taste	0			27 (48.2)			<0.001
Testing strategy	Symptom based N = 39	Weekly N = 74	P-value	Symptom based N = 26	Weekly N = 30	P-value	
Symptomatic, n (%)	36 (92.3)	29 (39.2)	<0.001	23 (94.6)	28 (93.3)	0.763	
Presymptomatic, n (%)	3 (7.7)	38 (51.4)		1 (3.8)	1 (3.3)		
Asymptomatic, n (%)	0	7 (9.5)		2[a] (7.6)	1 (3.3)		

[a]Two staff members did not complete follow-up questionnaire.

date differed significantly between the two testing strategies (P-value = 0.000). With both test strategies, symptomatic residents had symptoms for multiple days without testing.

Ct-values

Ct values were available for 97/113 positive residents; the median Ct-value was 21.3 (range 14.5–40). Symptomatic residents (N = 59) had a median Ct-value of 20.8 (range 14.5–38.1), presymptomatic resident (N = 33) had a median Ct-value of 21.3 (range 16.1–40) and asymptomatic resident (N = 5) had a median Ct-value of 20.5 (range 17.3–39.7). There was no difference in Ct-value between these groups (P = 0.624).

Ct-values were available for 38/56 staff members; with a median of 24.6 (range 13.7–38.1). Of one asymptomatic staff Ct-value was 34.6. The two presymptomatic staff members Ct-values were 29.8 and 32.3. Symptomatic staff members (N = 35) had a median Ct-value of 23.7 (range 13.7–38.1).

Discussion

We describe a large SARS-CoV-2-outbreak in a NH which most likely started by an infected resident discharged from a local hospital where SARS-CoV-2 prevailed. The addition of weekly facility-wide testing regardless of symptoms identified 38 (52.7%) presymptomatic residents and 7 (8.1%) asymptomatic residents. These cases were found up to 8 days before symptoms occurred. In staff limited, presymptomatic and asymptomatic cases were identified. The absence of subjective symptoms (such as loss of smell or taste) in residents compared to staff who are infected by the same SARS-CoV-2 strain suggests the under-reporting of symptoms in residents. As such, it is not possible to make a distinction between a/presymptomatic and unrecognised symptomatic residents in this study. However, a/presymptomatic residents have the same high viral load as symptomatic residents, which suggests the same potential for viral shedding. These results support the guidelines of the ECDC and CDC to test asymptomatic residents and staff to identify pre- and asymptomatic cases of SARS-CoV-2.

The high prevalence of presymptomatic cases in residents and the limited registration of subjective symptoms is comparable to other studies [3, 5, 6, 9]. Studies performing facility wide testing regardless of symptoms of staff in NHs with a confirmed COVID-19 case found the same limited number of asymptomatic staff as in our study [3, 4, 13]. To our knowledge, we are the first study reporting on symptoms from residents and staff in a large outbreak of the same virus strain. The large difference between presymptomatic staff and residents found in this study has three possible explanations: First, a large number of residents in this NH are cognitive impaired, which makes it difficult for them to express their symptoms. Second, staff reporting on residents' symptoms were not aware of all the symptoms related to COVID-19. During the outbreak symptoms of residents
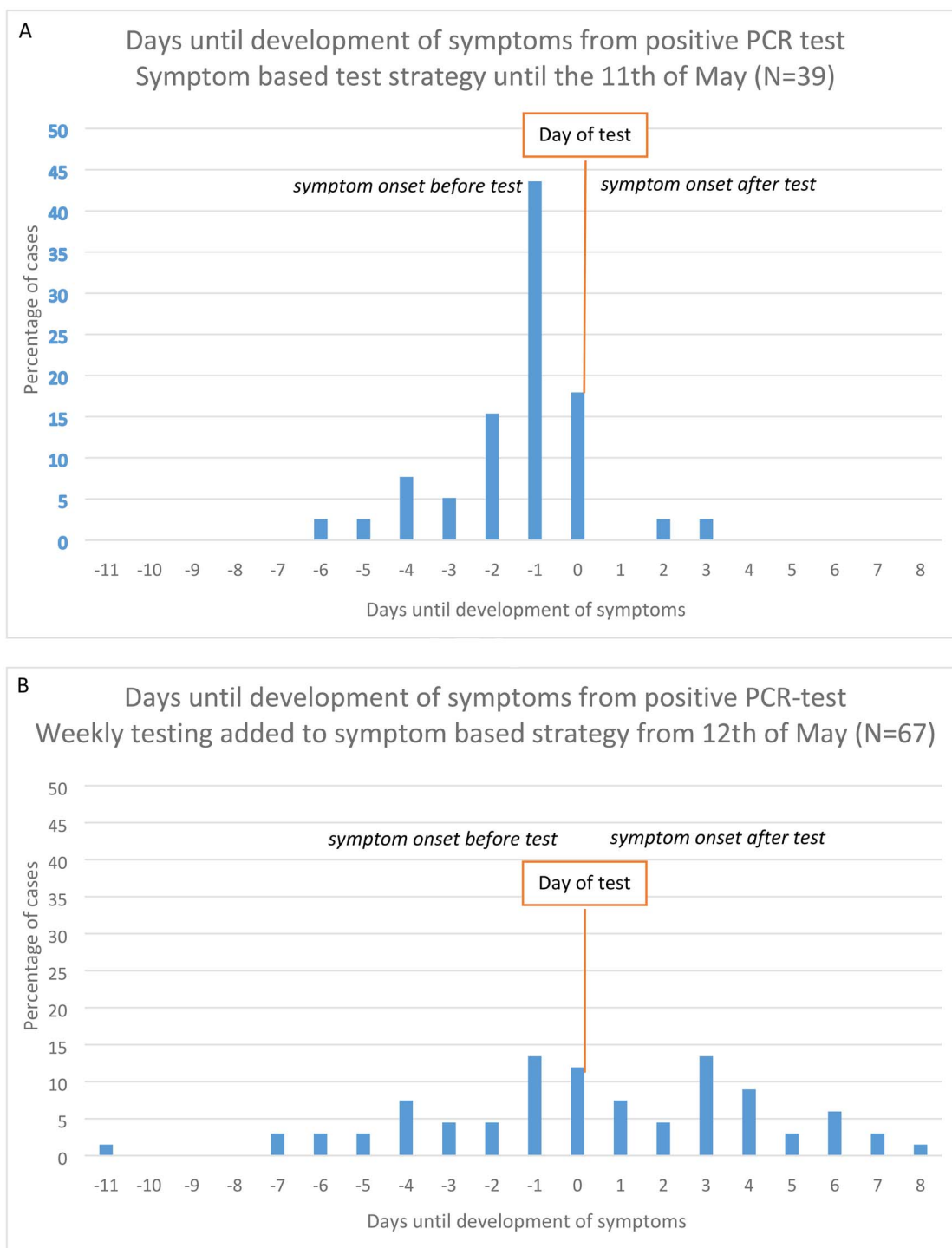
Figure 3. Frequency plot of days until development of symptoms from positive PCR-test of residents. Negative values represent symptomatic residents, while positive values represent presymptomatic residents. The value 0 means that residents developed symptoms at the day of PCR-test: whether the symptoms developed before or after testing determines if they were presymptomatic or symptomatic. (A) symptomatic testing strategy until the 11th of May. (B) Addition of facility-wide weekly testing strategy regardless of symptoms from the 12th of May.

were sometimes documented for multiple days, but they were nevertheless not tested. Third, understaffing because of the outbreak could have led to suboptimal symptom registration: mild or subjective symptoms were missed, because staff had to take care of residents they were not familiar to work with, or because of limited time to register symptoms. Understaffing as a risk for under-recognition of new cases is supported by data of Gorges and Li which shows NHs with

at least one case, higher nurse aide [21] and total nursing hours [21, 22] are associated with a lower probability of experiencing an outbreak and with fewer deaths.

Our study showed no difference between Ct-values of symptomatic, presymptomatic, or asymptomatic cases in residents, similar to previous studies [6, 9]. All these studies have the same risk of underreporting of (mild) symptoms of SARS-CoV-2 and incorrect classifying residents as pre- or asymptomatic. This suggest that these residents should be treated the same: as possibly infectious. Timely isolation of these residents and PPE could be important interventions to prevent further spread of the virus.

The new approach of mass repeated testing, irrespective of symptoms, in skilled nursing facilities has been advocated since May [23]. After this, studies have been published describing this approach, often resulting in reduced SARS-CoV-2 transmission after the implementation of this testing strategy [24, 25]. However, limited additional cases were found after a weekly testing strategy was implemented in three Dutch NHs after their first cases of SARS-CoV-2 [26]. Possibly, the testing was early in the outbreak and led to rapid isolation, combined with the increased availability of PPE or because cases per capita in the community were very low [26]. Cases per capita in the community have been identified as an important predictor for outbreaks in NHs [21, 27].

The testing of staff regardless of symptoms could be important because previous research showed that health care workers have difficulty in recognising possible COVID-19 symptoms for themselves: 65% reported working while exhibiting symptoms [28]. This is reflected in our results, as we found that almost none of the staff was asymptomatic at the moment of testing, even after the implementation of a testing strategy regardless of symptoms. The WHO advises that syndromic surveillance of health workers for COVID-19 symptoms should be performed before entering the workplace. If human resources and logistics permit it, active syndromic surveillance is recommended. Symptoms that should be monitored at minimum are fever, dry cough, myalgia, arthralgia, fatigue, headache, shortness of breath, anosmia and ageusia [29].

Implementing sequencing, combined with epidemiological information, is important to understand the extent of intramural transmission versus introductions from the community. In addition, transmission clusters and risk factors for transmission can be identified, which can be used to implement infection prevention measures to prevent further spread. Previous research has shown that whole genome sequencing can generate evidence for transmission routes that would not have been identified with traditional epidemiological investigations [16, 17].

Limitations

Not all staff members who tested positive participated in the study. In addition, some staff members had to answer the questionnaire retrospectively, which gives the risk of recall bias. Sensitivity analyses did not alter results.

Further, the difference between symptomatic staff and residents could perhaps be explained by the fact that staff was tested less frequent than residents. This may have contributed partly to the higher proportion of symptomatic staff. In our sensitivity where we compared staff who were tested on the dates of the point prevalence surveys to staff who were tested on other dates during the study period because of symptoms (Appendix 7, Supplementary data are available in *Age and Ageing* online). Except for loss of smell and taste no difference in symptoms between staff tested regardless of symptoms or outside the point prevalence surveys was observed.

Also, the low rate of reported symptoms in residents could be explained by the high proportion of residents with cognitive impairment. The NH in this study was specialised in psychogeriatric care; in a representative sample of Dutch NH, 59% of residents were diagnosed with cognitive impairment [30].

Last, not all SARS-CoV-2 positive samples were sequenced. However, a lot of time points could be analysed and they show all the same cluster which makes it unlikely that multiple clusters were circulating in the NH.

Conclusion

Our study suggests that a proportion of the presymptomatic cases in NHs are possibly unrecognised symptomatic cases and supports the guideline of the CDC and ECDC that facility-wide testing of residents and staff needs to be undertaken after the first confirmed SARS-CoV-2 case in the facility [7, 8]. If there is limited viral testing capacity, initial testing of (asymptomatic) close contacts is advised [8]. This will identify possible asymptomatic, presymptomatic cases and unrecognised symptomatic cases and prevent further spread of the virus. Sequencing should be performed to discriminate ongoing intramural transmission and multiple introductions. *Box 1* summarises the lessons learned during this study.

Box 1. Lessons learned of SARS-CoV-2 outbreak

Lessons learned

1. Preparing for an outbreak

Educate staff about all the possible symptoms of COVID-19: Take routine temperature and saturation of residents for reference values. Also sufficient staffing and staff dedicated to a few patients is necessary for early recognition of symptoms. NHs should make protocols with a local laboratory so when an outbreak occurs, rapid testing is possible.

Increasing cases per capita in the population

When cases per capita in the general population of the area are increasing, staff and visitors should wear at least surgical face mask to prevent introduction of the virus. In this outbreak, a resident transferring from another health care facility with a negative SARS-CoV-2 PCR was the index case and presymptomatic at the moment of transfer. Consider quarantine of residents who are admitted, regardless of a recent, negative PCR-test.

During an outbreak

Recognition of start of possible COVID-19 symptoms is very difficult, especially in residents with dementia. Weekly testing during an outbreak identifies presymptomatic or unrecognised symptomatic residents and makes timely isolation and use of PPE possible. We support international recommendations to consider routine testing of staff as soon a positive case of COVID-19 is identified in either staff or residents [29].

References

1. Comas-Herrera A, Zalakain J, Litwin C *et al.* Mortality associated with COVID-19 outbreaks in care homes: early international evidence. Article in LTCcovid. org, International Long-Term Care Policy Network, CPEC-LSE, 26. 2020.

2. Yourish L, Ivory S. One-Third of all U.S. Coronavirus Deaths Are Nursing Home Residents or Workers. New York Times, 2020.

3. Roxby AC, Greninger AL, Hatfield KM *et al.* Outbreak investigation of COVID-19 among residents and staff of an independent and assisted living community for older adults in Seattle, Washington. JAMA Intern Med 2020; 180: 1101–5.

4. Guery R, Delaye C, Brule N *et al.* Limited effectiveness of systematic screening by nasopharyngeal RT-PCR of medicalized nursing home staff after a first case of COVID-19 in a resident. Med Mal Infect 2020; 50: 748–50.

5. Dora AV WA, Jatt LP. Universal and serial laboratory testing for SARS-CoV-2 at a long-term care skilled nursing facility for veterans — Los Angeles, California, 2020. Morb Mortal Wkly Rep 2020; 21: 651–5.

6. Arons MM, Hatfield KM, Reddy SC, Kimball A, James A, Jacobs JR *et al.* Presymptomatic SARS-CoV-2 infections and transmission in a skilled nursing facility. New Engl J Med 2020; 382: 2081–90.

7. European Centre for Disease Prevention and Control. Surveillance of COVID-19 in long-term care facilities in the EU/EEA. Stockholm: ECDC, 2020.

8. CDC. Testing Guidelines for Nursing Homes. Interim SARS-CoV-2 Testing Guidelines for Nursing Home Residents and Healthcare Personnel. In: Centre for Disease Control and Prevention, 2020.

9. Kimball A, Hatfield KM, Arons M. Asymptomatic and Presymptomatic SARS-CoV-2 Infections in Residents of a Long-Term Care Skilled Nursing Facility-King County, Washington, 2020. MMWR Morb Mortal Wkly Rep. 2020: 377–81.

10. Huffman JC, Kunik ME. Assessment and understanding of pain in patients with dementia. Gerontologist 2000; 40: 574–81.

11. Behandeladvies COVID-19 Acute fase en nazorg. Verenso; 2020. Available from: https://www.verenso.nl/_asset/_public/Thema-en-projecten/Infectieziekten/Covid-19/200825-15-00-COVID-19-behandel-advies-DEFINITIEF.pdf.

12. Goldberg SA, Pu CT, Thompson RW, Mark E, Sequist TD, Grabowski DC. Asymptomatic spread of COVID-19 in 97 patients at a skilled nursing facility. J Am Med Dir Assoc 2020; 21: 980–1.

13. Graham NS, Junghans C, Downes R *et al.* SARS-CoV-2 infection, clinical features and outcome of COVID-19 in United Kingdom nursing homes. J Infect 2020; 8: 411–9.

14. Rutten JJS, van Loon SM, Joling KJ, Smalbrugge MS, van Buul LW, Hertogh CMPM. Covid-19 in verpleeghuizen. Ned Tijdschr Geneesk 2021; 164: D5173.

15. D'Adamo H, Yoshikawa T, Ouslander JG. Coronavirus disease 2019 in geriatrics and long-term care: the ABCDs of COVID-19. J Am Geriatr Soc 2020; 68: 912–7.

16. Voeten HA, Sikkema RS, Damen M *et al.* Unravelling the modes of transmission of severe acute respiratory syndrome Coronavirus 2 (SARS-CoV-2) during a nursing home outbreak: looking beyond the church super-spread event. Clin Infect Dis 2020: ciaa1664. doi: 10.1093/cid/ciaa1664.

17. Oude Munnink BB, Nieuwenhuijse DF, Stein M *et al.* Rapid SARS-CoV-2 whole-genome sequencing and analysis for informed public health decision-making in the Netherlands. Nat Med 2020; 26: 1405–10.

18. Oude Munnink BB, Nieuwenhuijse DF, Sikkema RS, Koopmans M. Validating whole genome nanopore sequencing, using Usutu virus as an example. J Vis Exp 2020. doi: 10.3791/60906.

19. World Health Organization. Clinical management of COVID-19: interim guidance, 27 May 2020. (No. WHO/2019-nCoV/clinical/2020.5). World Health Organization 2020. https://apps.who.int/iris/handle/10665/332196.

20. Prokop M, Everdingen Wv, Vellinga TvR et al. CO-RADS: a categorical CT assessment scheme for patients suspected of having COVID-19—definition and evaluation. Radiology 2020; 296: E97–104.

21. Gorges RJ, Konetzka RT. Staffing levels and COVID-19 Cases and outbreaks in US nursing homes. J Am Geriatr Soc 2020; 68: 2462–6.

22. Li Y, Temkin-Greener H, Shan G, Cai X. COVID-19 infections and deaths among Connecticut nursing home residents: facility correlates. J Am Geriatr Soc 2020; 68: 1899–1906.

23. Gandhi M, Yokoe DS, Havlir DV. Asymptomatic transmission, the Achilles' heel of current strategies to control Covid-19. N Engl J Med 2020; 382: 2158–60.

24. Sanchez GV, Biedron C, Fink LR, Hatfield KM, Polistico JMF, Meyer MP et al. Initial and repeated point prevalence surveys to inform SARS-CoV-2 infection prevention in 26 skilled nursing facilities-Detroit, Michigan, March-May 2020. MMWR Morb Mortal Wkly Rep 2020; 69: 882–6.

25. Escobar DJ, Lanzi M, Saberi P et al. Mitigation of a COVID-19 outbreak in a nursing home through serial testing of residents and staff. Clin Infect Dis 2021; 72: e394–6.

26. van Buul LW, van den Besselaar JH, Koene F et al. Asymptomatic Cases and Limited Transmission of SARS-CoV-2 in Residents and Healthcare Workers in Three Dutch Nursing Homes. Gerontol Geriatr Med 2020; 6:2333721420982800. doi: 10.1177/2333721420982800.

27. Shi SM, Bakaev I, Chen H, Travison TG, Berry SD. Risk factors, presentation, and course of coronavirus disease 2019 in a large, academic long-term care facility. J Am Med Dir Assoc 2020; 21: 1378–83.e1

28. Chow EJ, Schwartz NG, Tobolowsky FA, Zacks RLT, Huntington-Frazier M, Reddy SC et al. Symptom screening at illness onset of health care personnel with SARS-CoV-2 infection in King County, Washington. JAMA 2020; 323: 2087–9.

29. World Health Organization. Prevention, identification and management of health worker infection in the context of COVID-19. World Health Organization. 2020. Report No.: WHO/2019-nCoV/HW_infection/2020.1.

30. Rutten JJS, van Loon AM, van Kooten J, van Buul LW, Joling KJ, Smalbrugge M et al. Clinical suspicion of COVID-19 in nursing home residents: symptoms and mortality risk factors. J Am Med Dir Assoc 2020; 21: 1791–7.e1.