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Rapid measurement of SARS-CoV-2 spike T cells in whole blood from vaccinated and naturally infected individuals

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Defining the correlates of protection necessary to manage the COVID-19 pandemic requires the analysis of both antibody and T cell parameters, but the complexity of traditional tests limits virus-specific T cell measurements. We tested the sensitivity and performance of a simple and rapid SARS-CoV-2 spike protein–specific T cell test based on the stimulation of whole blood with peptides covering the SARS-CoV-2 spike protein, followed by cytokine (IFN-γ, IL-2) measurement in different cohorts including BNT162b2-vaccinated individuals (n = 112), convalescent asymptomatic and symptomatic COVID-19 patients (n = 130), and SARS-CoV-1–convalescent individuals (n = 12). The sensitivity of this rapid test is comparable to that of traditional methods of T cell analysis (ELISPOT, activation-induced marker). Using this test, we observed a similar mean magnitude of T cell responses between the vaccinees and SARS-CoV-2 convalescents 3 months after vaccination or virus priming. However, a wide heterogeneity of the magnitude of spike-specific T cell responses characterized the individual responses, irrespective of the time of analysis. The magnitude of these spike-specific T cell responses cannot be predicted from the neutralizing antibody levels. Hence, both humoral and cellular spike–specific immunity should be tested after vaccination to define the correlates of protection necessary to evaluate current vaccine strategies.

Introduction

SARS-CoV-2, the etiological agent of COVID-19, has spread worldwide, resulting in a global health and economic crisis that mass vaccinations are trying to resolve. The host’s capacity to be protected from viral infection or from the development of severe diseases requires a coordinated activation of different components of the immune system that ultimately leads to the production of neutralizing and antigen-binding antibodies and antiviral T cells. Evidence that antibodies and T cells are required for protection has been found in monkeys challenged with SARS-CoV-2 (1). Similarly, antibodies and T cells are present in the majority of SARS-CoV-2–infected individuals who have controlled infection without severe symptoms (2–6), and a robust CD8+ T cell response is associated with mild disease in oncological patients with humoral defects (7).

Recently developed SARS-CoV-2 vaccines that protect more than 90% of the vaccinated individuals from severe COVID-19 can induce spike-specific antibodies and T cells (8, 9, 10). However, it is not entirely clear what level of antibodies and/or T cells is necessary to confer such protection or whether differences in antibody and T cell levels in fact exist in vaccinated persons. Efforts to define the protective threshold of antibodies through mathematical modeling (11) have shed some light on this issue, but such work on T cell responses has so far been absent. Although experimental data have shown that high levels of neutralizing antibodies can be sufficient to protect against experimental infection, lower levels require the presence of T cells (1). Neutralizing antibody titers are, however, extremely heterogeneous after natural infection (12), and while most of the new SARS-CoV-2 vaccines induce high neutralizing antibody levels (10, 13), their persistence over time needs to be evaluated. Instead, virus-specific T cells appear to persist for a long time after viral clearance (i.e., 17 years after SARS-CoV-1 infection), and detection of SARS-CoV-2–specific T cells in patients with COVID-19 with waning antibody titers has been reported by different groups (5, 4, 14, 15). Furthermore, the protective role of spike-specific T cells in vaccinated individuals...
has also been highlighted by a recent analysis of the early profile of spike-specific immunity (16).

We think that the correlates of protection induced by vaccinations should therefore be derived from large prospective studies in which the levels of both antibodies and T cells are measured. However, although tests for antibodies are routinely performed, the technical complexity of SARS-CoV-2 T cell measurements has so far limited this analysis, with some exceptions (17), to a small number of individuals characterized in a few specialized laboratories. This is because T cells specific for a defined pathogen constitute a minuscule fraction of total T cells (often less than 1%-3%) present in the blood and can be distinguished mainly by complex functional assays that preserve the viability of the T cells during the assay. In addition, methods that are technically simple and do not require complex laboratory equipment, like the ELISPOT assay, need to be performed in cells that have been purified from whole blood. This introduces into the assay the lengthy and technically demanding processes of PBMC separation. Other assays that can directly measure the frequency and function of virus-specific T cells through expression of activation markers or cytokine production necessitate more complex equipment (i.e., a flow cytometer) and highly specialized personnel that might not be available in every routine diagnostic laboratory.

A possible rapid and simple alternative to these methods is the direct addition of stimulatory antigens or peptides to whole blood that induce the secretion of cytokines (usually IFN-γ) in plasma, which is subsequently quantified. This assay is routinely applied in the diagnosis of active tuberculosis (18), and it has also been shown to measure the presence of SARS-CoV-2–specific T cells in asymptomatic (5) and symptomatic SARS-CoV-2–infected (19, 20) patients. However, to our knowledge, its accuracy and validation have not been adequately analyzed over time in individuals who have been vaccinated against SARS-CoV-2, and only responses immediately after vaccination have been tested (16). Therefore, in this study, we applied a range of cellular methods to measure SARS-CoV-2 T cell responses in individuals vaccinated with the pre-fusion-stabilized, full-length SARS-CoV-2 spike protein (BNT162b2) or in those naturally infected with SARS-CoV-2. We demonstrated that the detection and relative quantification of spike-specific T cells in vaccinated individuals can be easily and rapidly achieved through the simple addition of spike peptide pools to whole blood. Utilization of different peptide pools to stimulate whole blood provides the flexibility to derive rapid information about the kinetics and magnitude of spike-specific T cell responses induced by vaccination and compare them with those present in convalescent individuals.

Results
Rapid quantification of SARS-CoV-2 spike–specific T cells by direct peptide stimulation of whole peripheral blood. We characterized the initial kinetics of spike-specific T cells induced by 2 doses of the mRNA vaccine BNT162b2 over a 51-day period using different methods of antigen-specific T cell analysis in fresh blood as well as in cryopreserved PBMCs. Whole blood from 6 healthy individuals was collected before (day 0) and 7, 10, and 20 days after the prime dose and then 7, 10, 20, and 30 days after the boost dose. Whole fresh blood (2–6 hours after collection) was either directly stimulated with peptides for a cytokine release assay (CRA) or processed by Ficoll density gradient centrifugation to obtain isolated PBMCs (Figure 1A). PBMCs were either used fresh in the IFN-γ ELISPOT assay (Figure 1A) or cryopreserved for further analysis. Fresh blood and fresh PBMCs were stimulated with the SpG peptide pool containing fifty-five 15 mer peptides (Supplemental Table 1; supplemental material available online with this article; https://doi.org/10.1172/JCI152379DS1) covering spike-specific T cell epitopes that are immunogenic in 95% of SARS-CoV-2–infected individuals (5). Two negative controls consisting of the vehicle control with an identical DMSO concentration present in the SpG peptide pool and a peptide pool covering SARS-CoV-2 nucleoprotein (NP) (41 peptides covering the C-terminal half of the NP; Supplemental Table 1) were used. We measured IFN-γ and IL-2 levels in the whole blood after 14–18 hours of incubation and enumerated the spots by ELISPOT assay following overnight incubation.

Figure 1B shows that the 2 different assays detected a predominant spike-specific response in all the individuals and defined a matching profile of spike-specific T cell responses after the prime and boost vaccinations. The number of IFN-γ spots detected after boost vaccination matched that observed in the phase I/II trial involving individuals vaccinated with BNT162b2 (10) and with a similar preparation consisting of the trimerized secreted version of the spike receptor–binding domain (BNT162b1; ref. 13), including a trial conducted in Chinese individuals vaccinated with BNT162b1 (21). In our study, although stimulation with the NP–specific peptide pool remained largely negative, the levels of IFN-γ in the blood and the number of IFN-γ spots showed identical peak responses that occurred 7–10 days after the first dose in individuals V4 and V5, and 7–10 days after the second dose in individuals V1, V3, and V6. There were, however, some minor discrepancies. The CRA did not detect boosting of spike-specific T cells induced by the second vaccine dose in subjects V4 and V5, perhaps in relation to the transient lymphopenia induced by the mRNA vaccination (13). IL-2 cytokine measurement (Figure 1C) revealed a pattern of spike-specific T cell responses equivalent to that achieved through IFN-γ release. However, IL-2 levels exceeded those of IFN-γ in all individuals 21 days after the first and second vaccine doses. Overall, we found a very strong correlation between IL-2 and IFN-γ secretion and the number of IFN-γ spots (Figure 1D), which allowed a precise estimation of the quantity of IFN-γ–producing cells related to the quantity of cytokines detected in whole blood (Table 1).

Assessment of the spike-specific T cell response directly from fresh whole blood yields results comparable to those obtained with classical T cell assays. Since T cell analysis is often performed in a single centralized laboratory using cryopreserved samples collected at different sites, we also analyzed the spike-specific T cell response after vaccination by performing ELISPOT activation-induced cellular marker (AIM) assays using cryopreserved samples stimulated with an SpG peptide pool. We then compared the results with those from the ELISPOT and CRA performed using the corresponding fresh whole blood. As already shown (22), the quantity of spike-specific spots detected by ELISPOT in cryopreserved PBMCs was reduced in comparison with the quantity detected in freshly isolated PBMCs (Supplemental Figure 1A), but the dynamics of the spike-specific response remained consistent with fresh
PBMCs (Supplemental Figure 1A) as also evidenced by the high correlation between the ELISPOT results from the differently processed samples (Figure 2A and Supplemental Figure 1B). The AIM assay, in our case, was less precise at detecting the dynamic expanded samples (Figure 2A and Supplemental Figure 1B). The AIM correlation between the ELISPOT results from the differently processed PBMCs (Supplemental Figure 1A) as also evidenced by the high correlation of IL-2 concentrations in SpG peptide pool–stimulated whole blood, with the results from the whole-blood CRA. We found that cytokines in whole blood remained well correlated with ELISPOT secreted in whole blood pulsed with different peptides constitutes a reliable method to gauge the presence and magnitude of functional T cells specific for epitopes covered by the peptides used.

A total spike-specific T cell response is accurately represented by the T cells specific for the SpG peptide pool. Although the whole-blood CRA using the 7 overlapping peptide pools of the spike protein could provide us information on the immunogenicity of the different regions of spike and the total spike-specific T cell response, assessment of the response in larger numbers of individuals requires a more streamlined approach. Thus, we analyzed the relation between the total spike-specific T cell response and the response to our selected SpG peptide pool. A schematic representation of the localization of peptide pools 1–7 in relation to the S1 (N-terminal), RBD, and S2 (C-terminal) regions of the spike protein is shown in Figure 3A. These different spike peptide pools were used in the whole-blood CRA and ELISPOT with freshly isolated PBMCs. The results of these different assays performed at the indicated time points are first shown in 2 representative vaccine recipients (Figure 3B), whereas the results obtained in all 6 individuals are represented as a heatmap in Figure 3C. The 3 different measurements (IFN-γ and IL-2 CRA and IFN-γ ELISPOT) provided very similar information in relation to the T cell response induced by BNT162b2 in healthy individuals. Even though some differences can be noted, like in individual V3, in whom the dominant IFN-γ response (CRA and ELISPOT) was induced by pool 3, while pool 7 induced the dominant IL-2 response, overall, all the assays were largely equivalent. Consistent across the 3 different measurements, spike-specific T cells preferentially targeted the S2 chain of spike (covered by pools 5, 6, and 7 spanning spike 700–1273 aa), with responses in all 6 of the individuals tested at different time points. The whole-blood CRA and ELISPOT also showed that the region 501–705 aa contained the T cells specific for epitopes covered by the peptides used.

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ELISPOT), in which both the total spike protein and SpG peptide pool–specific T cell responses were determined in the same sample through stimulation with the corresponding peptide pools, we observed a strong positive linear relationship indicating that the T cell response against the SpG peptide pool was highly representative of the total spike T cell response (Figure 4B). In fact, the SpG peptide pool–specific T cell response constituted approximately 60%–80% of the total T cell response against the entire spike protein (Figure 4C). This was further supported by a linear regression analysis that also showed a good correlation between the IFN-γ and IL-2 CRA data obtained through individual spike pool stimulation (pools 1–7) and total spike protein IFN-γ ELISPOT data (Supplemental Figure 3). Hence, using the whole-blood CRA with SpG peptide pool as a stimulant, we proceeded to analyze the spike-specific T cell response in a larger cohort of BNT162b2-vaccinated individuals and of individuals who had recovered from SARS-CoV-2 and SARS-CoV-1 infection.

T cell responses to spike after vaccination or after natural infection with SARS-CoV-2 or SARS-CoV-1. A total of 112 individuals vaccinated with BNT162b2 (201 samples), 62 and 68 individuals who recovered from symptomatic (115 samples) and asymptomatic (62 samples) SARS-CoV-2 infection, respectively, and 12 individuals who recovered from SARS-CoV-1 infection 18 years ago (12 samples) were studied longitudinally using the whole-blood CRA with the SpG peptide pool and measuring both IFN-γ and IL-2.

First, we analyzed samples collected 3 months or more after the boost vaccination (3 months) or SARS-CoV-2 infection clearance (3–12 months) to understand whether the whole blood CRA remained reliable for quantification of the spike-specific T cell responses at later time points beyond that tracked in Figures 1 and 5. Linear regression analysis of IFN-γ and IL-2 secretion in a whole-blood CRA and the corresponding frequency of SpG-reactive T cells in cryopreserved PBMCs quantified by either IFN-γ ELISPOT or AIM assay (n = 6; 24 samples). Dotted lines denote the 95% CI.
Figure 3. Immunodominance of spike-specific T cells in vaccinated individuals. (A) Schematic representation of the 7 spike-specific peptide pools containing 15 mer overlapping peptides spanning the entire spike protein. Pools 1–4 contain peptides from the signal peptide and the S1 chain, whereas pools 5 and 6 encompass the S2 chain together with the transmembrane and cytoplasmic domains. (B) Plots show the longitudinal evaluation of spike-specific T cell responses (pools 1–7) by quantification of IFN-γ (left) or IL-2 (middle) in peptide-stimulated whole blood, or by IFN-γ ELISPOT (right) in 2 representative vaccinees. (C) Heatmap shows the spike-specific T cell responses quantified longitudinally in all vaccinees (n = 6) using the 3 different assays described above. “X” denotes time points that were untested.
from SARS-CoV-1 infection 17 years ago, T cells specific for SARS-CoV-2 spike protein also remained detectable (8 of 12 were IFN-γ positive), similar to the NP-specific T cell responses described previously (25), despite the low aa conservation of the SpG peptides between the 2 viruses (Figure 6A). Clearly, further analysis is needed to determine whether T cells induced by vaccines will be maintained at levels similar to those induced by natural infection beyond the 3-month period.

We also assessed whether there were differences in the magnitude of the responses between the different groups. We compared the spike-specific T cell responses detected at similar time points 2–3 months after boost vaccination or viral clearance. Unlike the reports of higher neutralizing antibody titers in vaccinees (13), we found that individuals with symptomatic SARS-CoV-2 infection and vaccinees mounted equivalent magnitudes of spike-specific T cell responses (both IFN-γ and IL-2 secretion), whereas higher levels of IFN-γ secretion were only detected in individuals who had an asymptomatic SARS-CoV-2 infection (Figure 6B). The latter observation is in line with previous analyses of asymptomatic and symptomatic SARS-CoV-2–infected individuals within 1 month of viral clearance, in which the former showed increased cytokine production with comparable frequencies of virus-specific T cells (5). Subtle qualitative differences in the spike-specific T cell response were also observed in the different groups. Upon vaccination, spike peptide–induced IFN-γ and IL-2 secretion levels were comparable, but gradually diverged over time, leading to higher levels of detectable IL-2 two to 3 months after boost vaccination (Figure 6C). Symptomatic individuals also produced significantly more IL-2 than IFN-γ more than 6 months after viral clearance, while this difference was less pronounced in asymptomatic individuals even at the latest time points tested (9–12 months after viral clearance), which did not reach statistical significance (Figure 6C). In individuals with previous SARS-CoV-1 infection, whole-blood CRA also detected higher IL-2–secreting, spike-specific T cell responses 18 years after resolution of infection (Figure 6A). Hence, quantification of IL-2 secretion provides better sensitivity than IFN-γ in identifying individuals with a long-term spike-specific memory T cell response.

Whole-blood CRA detects the wide dynamic range and heterogeneous function of spike-specific T cell responses in vaccinated individuals. In addition to evaluating the kinetics as well as quantitative and qualitative differences in the T cell responses, the whole-blood CRA also detected a wide range of spike-specific T cell responses in vaccinated individuals. Figure 7A shows the paired longitudinal samples of 27 vaccinees approximately 14 and 90 days after boost vaccination. The levels of secreted IFN-γ and IL-2 in the whole-blood CRA differed between the 2 time points and among individuals. Interestingly, the quantity of cytokines detected 2 weeks after boost-dose vaccination did not always predict the level of spike-specific T cell responses measurable by day 90. Some individuals had a greater than 20-fold reduction in IFN-γ and IL-2 levels on day 90 after boost vaccination, while in others, the levels, particularly those for IL-2, were more stable (Figure 7A). Indeed, some individuals have a more pronounced decline in IFN-γ levels than IL-2 levels, or vice versa, as indicated in Figure 7B, in which

Figure 4. Frequency of the SpG peptide pool and total spike protein–specific T cells. (A) Schematic representation of the individual 15 mer overlapping peptides contained in the SpG peptide pool. (B) Linear regression analysis of the T cell response against the SpG peptide pool and the total spike protein (pools 1–7) as evaluated by ELISPOT (left) or by the quantification of IFN-γ (middle) or IL-2 (right) in peptide-stimulated whole blood (n = 6; 42 samples). (C) The SpG peptide pool–specific T cell response quantified by each assay is expressed as a fraction of the total spike protein T cell response observed (n = 6; 42 samples). Bars indicate the mean.

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neutralizing antibody quantities cannot predict the corresponding spike-specific T cell responses in an individual, and this further stresses how T cell response information from the whole-blood CRA complements existing antibody assessments.

**Discussion**

The complexity of virus-specific T cell characterization relegates their analysis to studies performed in selected laboratories experienced in the complex methods of T cell analysis. The rapidity, simplicity, and accuracy of the CRA in whole blood can allow for routine measurement of SARS-CoV-2 T cells in large populations and thus further our understanding of the role of antiviral T cells during the current COVID-19 pandemic. It is important to note that measurement of cytokine release in stimulated whole blood not only detects the mere presence of T cells but determines their functionality as well. This is an important feature that differentiates the CRA (and other assays like ELISPOT and AIM) from the recently developed T-Detect COVID test (Adaptive Biotechnologies), which uses next-generation sequencing of T cell receptors (TCRs) to determine the presence or absence of cellular immunity to SARS-CoV-2 (27). At the same time, although a functional read-out is of benefit, the whole-blood CRA, like ELISPOT, does not differentiate between functional CD4⁺ and CD8⁺ T cells. Hence, such an approach is not suitable for determining the quantities or unique functionality of different T cell subsets. The use of peptides to stimulate and detect specific T cells could also bias the detection toward high-avidity T cell responses. However, this is not unique to the whole-blood CRA, as other assays like ELISPOT, intracellular cytokine staining (ICS), and AIM also use isolated PBMCs stimulated with similar quantities of peptide. Nonetheless, we think that the ability of the whole-blood CRA to measure the wide, dynamic range of functional spike-specific T cells and not just their presence will be an important asset that will more precisely evaluate the protective ability of T cells after infection or vaccination.

In this study, by sequentially testing vaccinated and SARS-CoV-2–convalescent individuals, we show that IL-2 and IFN-γ quantification in whole blood measured spike-specific T cell responses with an accuracy equivalent to that obtained with ELISPOT assays performed in freshly purified PBMCs. Minor discrepancies between the magnitudes of T cells were detected only at early time points when the CRA was able to detect a signal in the absence of ELISPOT results. Furthermore, analysis 2 and 3 months after vaccination showed, on average, a better sensitivity of IL-2 than IFN-γ in detecting spike-specific T cell responses in individuals. The superior ability of IL-2 to detect long-term memory spike-specific T cells was also supported by the analysis of SARS-CoV-2–convalescent individuals 12 months after infection and also 17 years after infection in the case of SARS-CoV-1–infected individuals. Interestingly, this difference in sensitivity was less pronounced in convalescent COVID-19 patients with asymptomatic infection. Though speculative at the moment, this lack of difference between IFN-γ and IL-2 secretion levels could reflect a better functionality of spike-specific T cell responses that would hence plausibly contribute to the benign disease trajectory for these individuals.

The ability of the CRA to measure the dynamic range of spike-specific T cell responses allowed us to study a wide group of
vaccinated and COVID-19–convalescent individuals. We found that, despite the homogeneous cohort of vaccinated adults (21–60 years of age, healthy and SARS-CoV-2–naive individuals), the spike-specific T cell response in vaccinated individuals (n = 112; 201 samples) and in convalescent asymptomatic (n = 62; 62 samples) and symptomatic (n = 68; 115 samples) COVID-19 patients were longitudinally quantified by measuring IFN-γ secretion in whole blood after SpG peptide pool stimulation. Cross-reactive SARS-CoV-2 spike–specific T cells were also quantified in whole blood from individuals who were infected with SARS-CoV-1 eighteen years ago (n = 12; 12 samples). The responses of individuals before receiving BNT162b2 vaccination are shown for reference. Pie chart shows the number of peptides in the SpG peptide pool that are conserved or unique between SARS-CoV-2 and SARS-CoV-1. The sampling timespan (highlighted in yellow) is shown, and the number of samples analyzed at each time point is indicated in parentheses. Dashed lines denote the detection cutoff for the measured cytokines. D–14, day ~14. (B) Quantities of secreted IFN-γ (red) and IL-2 (blue) in SpG peptide pool–stimulated whole blood from vaccinees and COVID-19 patients sampled 2–3 months after a boost vaccination dose (Vacc.) or viral clearance. The bars indicate the median value for each group, and the dashed lines indicate the detection cutoff for the measured cytokines. Significant differences were analyzed and are displayed as above. (C) Longitudinal dynamics of secreted IFN-γ (red) and IL-2 (blue) in SpG peptide pool–stimulated whole blood from vaccinees and COVID-19. Dashed lines indicate the detection cutoff for the measured cytokines. Significant differences in each group were analyzed by 1-way ANOVA, and the P value (adjusted for multiple comparisons) are shown. NS = P > 0.05, *P ≤ 0.05, **P ≤ 0.01, ***P ≤ 0.001, and ****P ≤ 0.0001.

Another important observation that further supports the concept of marked heterogeneity of the immune response induced by the BNT162b2 vaccine was the lack of correlation between the magnitudes of humoral and cellular immunity. The substantial independence of different components of the immune system after the initial induction phase has been demonstrated in COVID-19 convalescents (29, 30) and can be explained by recent data showing that neutralizing antibodies can be produced without follicular T cell help (31). The CRA also revealed similar profiles in both COVID-19–convalescent and vaccinated individuals, as no correlation was detected between the neutralizing antibody titers and the magnitude of cytokine secretion in these individuals. COVID-19 convalescents were studied more than 1 year after infection and had different levels of neutralizing antibodies, whereas vaccinated individuals mostly had consistently high neutralizing antibody titers with little variation among those tested 3 months after vaccination. In other studies, antibody persistence was still observed up to 6 months after mRNA vaccination at the time of the report (32). We do not know whether antibodies induced by vaccination will show a rate of decline similar to that observed after natural infection beyond the 6-month follow-up date, which makes the analysis of cellular immunity even more important. Our data at the moment show that 3 months after vaccination, the levels of neutralizing antibodies cannot be used as a surrogate for the spike-specific cellular immunity induced by vaccination.

This quantitative heterogeneity was, however, not mirrored by the regions of spike protein targeted by T cells induced by BNT162b2 vaccination. We observed a substantial similarity of immunodominance among the different vaccinated individuals, with a large part of the T cell response directed toward the spike 2 chain and with an almost complete lack of T cell determinants within the N-terminal region of S1 (region 501–705 aa). A reduced presence of T cell epitopes in this region was already observed in SARS-CoV-2 convalescents (23). It will be interesting to test whether this documented profile of spike–T cell specificity will also occur in individuals vaccinated with different products, where subtle differences in codon usage, signal peptides, and aa modifications (pre-fusion conformation stabilization and furin cleavage site mutations) were introduced (33–36). Of note, an intermediate dominance of the RBD-containing spike pool 3 was also observed, kines further increased the heterogeneity of the vaccine-induced immunity in different individuals, as IFN-γ and IL-2 quantities did not decrease in parallel in all the individuals. Some vaccinees displayed stable IL-2 production levels associated with a profound decrease in IFN-γ levels, whereas others showed exactly the opposite. Whether these differences can be attributed to the presence of different populations of effector/memory T cells or different ratios of CD4+ and CD8+ T cells, and whether such differences have an impact on protection will need to be analyzed in a large clinical study. In addition, it will be important to continue monitoring the spike-specific T cell response beyond the 3-month observation period to understand whether the spike-specific T cell response induced by vaccines will behave like the one observed after natural infection. Importantly, the rapid cytokine assay was able to detect spike-specific T cells in approximately 84% of individuals 1 year after infection with SARS-CoV-2 and also in 8 of 12 individuals 18 years after SARS-CoV-1 infection.
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Figure 7. Heterogeneity of spike-specific T cell responses in vaccinated individuals. (A) SARS-CoV-2 spike–specific T cell responses were evaluated by SpG peptide pool stimulation of whole blood from vaccinated individuals (n = 27) 2 weeks (green circle) and 3 months (red circle) after the boost vaccination dose. The secreted IFN-γ and IL-2 concentrations are shown. ****P ≤ 0.0001, by Wilcoxon matched-pairs signed-rank test. (B) Bivariate dot plots of secreted IFN-γ and IL-2 concentrations. Arrows connect paired individuals analyzed on day −14 and day 90. Dashed lines indicate the detection cutoff for the measured cytokines.

with possible implications for T cell recognition of the region and T cell immune escape by emerging viral variants of concern.

In conclusion, we show that the rapid measurement of cytokine production in whole blood after peptide-specific activation is a quick and simple assay that can reliably detect the wide, dynamic range of functionally heterogeneous spike-specific T cell responses induced after vaccination or infection in different individuals. Even though T cells cannot prevent infection in the absence of antibodies, their pivotal role in the protection from severe disease has been shown in natural infection of healthy individuals (2, 3) and oncological patients (7), as well as in vaccinated individuals (16). As such, since the quantity of spike-specific T cells cannot be predicted by the simple measurement of antibodies, this high-throughput, simple assay can be feasibly performed as part of routine testing to complement existing antibody measurements and thus help to define the correlates of protection necessary for the design of current vaccine strategies.

Methods

Study participants. Vaccinated individuals were between 21 and 60 years of age, healthy, and had no history of SARS-CoV-2 infection. Whole-blood and serum samples were collected at the indicated intervals for serological and T cell response analysis. RBD-binding antibodies in the serum of vaccinees V1–V6, before receiving the BNT162b2 vaccine, were quantified using cPASS (GenScript), and all were negative.

Peptides. Peptides of 15 mer that overlapped by 10 aa spanning the entire SARS-CoV-2 spike protein (GISAID EPI_ISL_410713) were synthesized (GenScript) and pooled into 7 pools of approximately 40 peptides in each pool (Supplemental Table 1). Fifty-five spike peptides covering the immunogenic regions of the SARS-CoV-2 spike protein that represent 40.5% of the whole spike protein formed the SpG peptide pool as described previously (5).

CRA of whole peripheral blood stimulated with SARS-CoV-2 spike peptide pools. Freshly drawn whole blood (320 μL; drawn within 6 hours of venipuncture) was mixed with 80 μL RPMI and stimulated with the indicated SARS-CoV-2 spike peptide pools (Supplemental Table 1) at 2 μg/mL or with DMSO as a control. After 16 hours of culturing, the culture supernatant (plasma) was collected and stored at −80°C. Cytokine concentrations in the plasma were quantified using an Ella machine with microfluidic multiplex cartridges that measured IFN-γ and IL-2 according to the manufacturer’s instructions (ProteinSimple). The levels of cytokines present in the plasma of DMSO controls were subtracted from the corresponding peptide pool–stimulated samples. The positivity threshold was set at 10 times the lower limit of quantification of each cytokine (IFN-γ = 1.7 pg/mL; IL-2 = 5.4 pg/mL) after DMSO background subtraction.

PBMC isolation. Peripheral blood was collected from all individuals in heparin-containing tubes, and PBMCs from all collected blood samples were isolated by Ficoll-Paque density gradient centrifugation.

SARS-CoV-2 spike–specific T cell quantification. The frequency of SARS-CoV-2 spike–specific T cells was quantified as described previously (5). Briefly, freshly isolated or cryopreserved PBMCs (as indicated) were stimulated with SpG peptide pool in an IFN-γ ELISPOT assay. ELISPOT plates (MilliporeSigma) were coated with human IFN-γ antibody overnight at 4°C. A total of 400,000 PBMCs were seeded per well and stimulated for 18 hours with the SpG peptide pool at 2 μg/mL. The plates were then incubated with a human biotinylated IFN-γ detection antibody, followed by streptavidin–alkaline phosphatase (streptavidin-AP) and developed using the KPL BCIP/NBT phosphatase substrate (Seracare Life Sciences). To quantify positive peptide-specific responses, 2× mean spots of the unstimulated wells were subtracted from the corresponding peptide pool–stimulated samples. The positivity threshold was set at 10 times the lower limit of quantification for each cytokine (IFN-γ = 1.7 pg/mL; IL-2 = 5.4 pg/mL) after DMSO background subtraction.

RBD-hACE2 binding inhibition assay. Antibodies inhibiting the binding of virus to the host cell were measured using a commercial RBD-human angiotensin-converting enzyme 2 (hACE2) binding inhibition assay called cPASS (GenScript). Following the manufacturer’s instructions, serum was diluted 1:10 in the kit sample buffer and mixed 1:1 with HRP-conjugated RBD and incubated for 30 minutes at 37°C. RBD-antibody mixtures were
then transferred onto ELISA plates coated with recombinant hACE2 receptor and incubated for 15 minutes at 37°C. Following incubation, the plates were washed with the kit wash solution followed by incubation with 3,3′,5,5′-tetramethylbenzidine (TMB) substrate for 15 minutes, and the reaction was stopped with stop solution. Absorbance was measured at OD$_{450}$ nm. The percentage of inhibition of RBD-hACE2 binding was computed using the following equation: percentage of inhibition $= (1 – (OD \text{ of serum} + RBD)/(OD \text{ of negative control} + RBD)) \times 100$. As recommended in the cPASS kit instructions, a cutoff of 20% and above was used to determine positive RBD-hACE2 inhibition.

AIM assay. Cryopreserved PBMCs were thawed and stimulated for 24 hours at 37°C with the SpG peptide pool (2 μg/mL) in AIM-V media supplemented with 2% pooled human AB serum. Cells were then stained with the Fixable Yellow Dead Cell Stain Kit (Invitrogen, Thermo Fisher Scientific) followed by staining with the following surface markers as previously described (16): anti-CD3, anti-CD4, anti-CD8, anti-CD69, anti-CD134 (OX40), and anti-CD137 (4-1BB). All samples were acquired on a BD-LSR II Analyzer (BD) and analyzed with Flowjo software (BD). The gating strategy is shown in Supplemental Figure 2.

Antibodies. The following antibodies were used in the ELISPOT assay: anti-human IFN-γ-coating antibody (Mabtech, catalog 3420-3-1000) and anti-human IFN-γ biotin (Mabtech, catalog 3420-6-1000). The following antibodies were used in the AIM assay: anti-human CD3 BV605 (BioLegend, catalog 317321); anti-human CD8 V500 (BD, catalog 560774); anti-human CD4 V650 (BD, catalog 563875); anti-human CD69 AF700 (BioLegend, catalog 310921); anti-human CD134 (OX40) PE (BioLegend, catalog 350004); and anti-human CD137 (4-1BB) APC (BD, catalog 550890).

Statistics. All statistical analyses were performed using GraphPad Prism, version 9 (GraphPad Software). Significant differences in each group were analyzed by 1-way ANOVA with adjustment for multiple comparisons or by the Wilcoxon matched-pairs signed-rank test. Where applicable, the statistical tests used and the definition of center are indicated in the figure legends. Statistical significance was set at a P value of less than 0.05. In all instances, n refers to the number of patients analyzed.

Study approval. Individuals who had recovered from SARS-CoV-2 infection (asymptomatic: n = 62; symptomatic: n = 68), were vaccinated with BNT162b2 (n = 112), or had SARS-CoV-1 infection 17 years ago (n = 12) were enrolled in this study as part of the PROTECT study (National Healthcare Group Domain Specific Review Board [NHG DSRB], ref. 2012/00917); the healthcare Worker Vaccination study (SingHealth Centralized Institutional Review Board [CIRB], CIRB ref. 2021/2014); the Novel Pathogens study (CIRB ref. 2018/3045); and the SARS Recall study (NHG DSRB ref. 2020/00091). All participants provided written informed consent.

Author contributions ATT, NLB, and AB designed the experiments. WNC, RDA, EEO, and LFW performed and analyzed the antibody experiments. JMEL, KK, AC, MDCQ, and NT performed all other experiments and analyzed the data. ATT, NLB, and AB interpreted all the data. ATT, NLB, and AB prepared the figures and wrote the manuscript. DY, JXYS, MICC, BEY, LYH, JGHL, and DCL recruited the COVID-19 patients, the SARS-Recall patients, and the vaccinees and provided all clinical samples and data. AB designed and coordinated the study.

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