Autoantibodies Neutralizing Type I Interferons in 20% of COVID-19 Deaths in a French Hospital

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Abstract
Recent studies reported the presence of pre-existing autoantibodies (auto-Ab) neutralizing type I interferons (IFNs) in at least 15% of patients with critical COVID-19 pneumonia. In one study, these auto-Ab were found in almost 20% of deceased patients across all ages. We aimed to assess the prevalence and clinical impact of the auto-Ab to type I IFNs in the Seine-Saint-Denis district, which was one of the most affected areas by COVID-19 in France during the first wave. We tested for the presence of auto-Ab neutralizing type I IFNs in a cohort of patients admitted for critical COVID-19 pneumonia during the first wave in the spring of 2020 in the medicine departments at Robert Ballanger Hospital, Aulnay sous Bois. We found circulating auto-Ab that neutralized 100 pg/mL IFN-α2 and/or IFN-ω in the plasma (diluted 1/10) of 7.9% (11 of 139) of the patients hospitalized for critical COVID-19. The presence of neutralizing auto-Ab was associated with an increased risk of mortality, as these auto-Ab were detected in 21% of patients who died from COVID-19 pneumonia. Deceased patients with and without auto-Ab did not present overt clinical differences. These results confirm both the importance of type I IFN immunity in host defense against SARS-CoV-2 infection and the usefulness of detection of auto-Ab neutralizing type I IFNs in the management of patients.

Keywords Autoantibodies · type I interferons · COVID-19 · mortality

Introduction
Since the onset of the COVID-19 pandemic in December 2019, at least 220 million people have been infected, and most likely many more. Nevertheless, only about 10% of these individuals developed hypoxemic COVID-19 pneumonia (severe or critical in about 3% of cases). There have been at least 5 million deaths, and most likely closer to 8–10 million. The clinical spectrum of SARS-CoV-2 infection is therefore vast, ranging from silent infection to lethal disease. A few epidemiological risk factors have been identified. The most important one is age, with a risk of life-threatening disease doubling every 5 years. Gender as well as several other risk factors have been described (e.g., obesity) but with relatively modest effects [1, 2]. In each demographic category, however, there remains vast inter-individual clinical variability.

Recent studies showed the important role of type I Interferons (IFNs) in protective immunity against SARS-CoV-2. Inborn errors of type I IFNs immunity were described in patients with life threatening COVID-19. Rare inborn errors
of autosomal genes controlling Toll-like receptor 3 (TLR3) and interferon regulatory factor 7 (IRF7)-dependent type I IFNs immunity were initially described [3] and more recently X-linked recessive TLR7 deficiency [4]. These inborn errors impair the production or amplification of type I IFNs in response to SARS-CoV-2. Deficiency of the TLR3 pathway incriminated pulmonary epithelial cells, while that of the TLR7 pathway incriminated plasmacytoid dendritic cells.

Interestingly, several studies showed that at least 10% of patients with life threatening (critical) COVID-19 pneumonia presented autoantibodies (auto-Abs) neutralizing type I IFNs, mostly the thirteen individual IFN-α and IFN-ω [5]. These auto-Abs were present before infection by SARS-CoV-2 in the patients tested and in 0.33% of uninfected controls from before the pandemic. They were not detected in patients with asymptomatic or mild SARS-CoV-2 infection. They were also shown to block the anti viral activity of correspondent type I IFNs against SARS-CoV-2 in vitro and in vivo [5]. These findings were replicated in other cohorts in Amsterdam, Madrid, San Francisco, Lyon, and New Haven [6–13].

Recently, auto-Abs neutralizing lower, more physiological concentrations of type I IFNs (100 pg/mL, in 1/10 dilutions of plasma) were detected in 15 to 20% of patients with critical COVID-19, notably in more than 20% of critical patients over 80 years of age. When also considering auto-Abs to IFN-β, they were present in about 20% of deceased patients across all ages [14]. Surprisingly, the study of more than 34,000 individuals showed that the prevalence of these auto-Abs in the general population increases with age, notably after the age of 70 (4%), providing an explanation for the increased risk of severe COVID-19 with age [14].

In this context, we aimed to assess the prevalence and clinical impact of the auto-Abs to type I IFNs in Seine-Saint-Denis department, which was one of the most affected areas by COVID-19 in France during the first wave of the pandemic [15]. We thus tested for the presence of auto-Abs to type I IFNs neutralizing different doses in a cohort of patients admitted for critical COVID-19 pneumonia during the first wave in the spring of 2020. Moreover, a commercial ELISA kit for the determination of anti–IFN-α2 auto-Abs was also evaluated.

Materials and Methods

Study Population

A cohort of 246 patients admitted for critical COVID-19 pneumonia was constituted in Robert Ballanger Hospital, Aulnay sous Bois, France, during the first wave of the pandemic in order to assess factors associated with clinical outcomes in patients hospitalized for COVID-19 [16]. This study was approved by a research ethics committee and was registered on clinicaltrials.gov (NCT04366206).

It included patients which, after going to the emergency room, were hospitalized in medicine departments dedicated to treat COVID-19 patients. Severity criteria required a pulse oxygen saturation (SpO2) ≤ 96% despite oxygen support ≥ 6 L/min with oxygen mask, for more than 6 h. All patients therefore had “critical” COVID-19 pneumonia [14, 17]. Some of these patients required an invasive mechanical ventilation and a transfer to an intensive care unit (ICU) during their hospitalization.

From this cohort of 246 patients, 139 patients were retrospectively selected because of an available serum sample that was collected during the acute phase of disease and stored in the laboratory sample collection. The studied population included 86 men and 53 women; median age was 66 years (range: 21–92 years), and 15% of the patients was over the age of 80 years. For some of patients, serum sample was also collected about 1 year after infection.

Detection of Anti-Cytokine Auto-Abs by Gyros and ELISA

Biological serum samples were analyzed for the determination of anti-IFN-α and anti-IFN-ω auto-Abs by Gyros technology as previously described [14].

Cytokines, recombinant human (rh)IFN-α2 (Milteny Biotec, ref. number 130–108-984) or rhIFN-ω (Merck, ref. number SRP3061), were first biotinylated with EZ-Link Sulfo-NHS-LC-Biotin (Thermo Fisher Scientific, cat. number A39257), according to the manufacturer’s instructions, with a biotin-to-protein molar ratio of 1:12. The detection reagent contained a secondary antibody, Alexa Fluor 647 goat anti-human IgG (Thermo Fisher Scientific, ref. number A21445) diluted in Rextip F (Gyros Protein Technologies, ref. number P0004825; 1/500 dilution of the 2 mg/mL stock to yield a final concentration of 4 µg/mL). Buffer PBS-T 0.01% and Gyros Wash buffer (Gyros Protein Technologies, ref. number P0020087) were prepared according to the manufacturer’s instructions. Plasma or serum samples were then diluted 1/100 in PBS-T 0.01% and tested with the Bioaffy 1000 CD (Gyros Protein Technologies, ref. number P0004253) and the Gyrolab X-Pand (Gyros Protein Technologies, ref. number P0020520). Cleaning cycles were performed in 20% ethanol.

Moreover, a commercially available ELISA kit (Human Anti-IFN alpha ELISA Kit, Thermo Fisher Scientific, ref. number BMS217, hereafter referred to as “ELISA”) was also used for the quantitative detection of human anti-IFN-α2 auto-Abs. In brief, microwells that were coated with recombinant human IFN-α subtype 2c were incubated with 1:5
Research of Antinuclear Antibodies

An indirect immunofluorescence assay on HEp-2 cells (Kallestad HEp-2 kit, Biorad, ref number 30472) was used for research of antinuclear antibodies (ANA) in patients with auto-Abs neutralizing type I IFNs, with a screening dilution of 1/160.

Statistical Analyses

To compare clinic characteristics, continuous variables are shown with the median and standard deviation (SD) and dichotomous variables are presented with the number of events and percentages.

Given the small samples size, a Fisher test was used to analyze the effect of dichotomous variables and a Mann–Whitney test for continuous variables.

Results

Detection of Auto-Abs Neutralizing IFN-α2 and/or IFN-ω

We first assessed the levels of circulating auto-Abs against IFN-α2 and IFN-ω using the Gyros technology in 139 patients with critical COVID-19 pneumonia. We found that 107 (77%) patients have auto-Abs against IFN-α2 and/or IFN-ω by Gyros technology. Intermediate titer (between 30 and 100, as previously described [14]) was frequent, as found in 66.9% (93 of 139) of patients with critical COVID-19 and high levels, above 100, were found in 10.1% (14 of 139) patients with critical COVID-19.

We then tested the neutralizing activity of all these samples against IFN-α and IFN-ω in vitro at high concentrations of type I IFNs (10 ng/mL) and lower concentrations (100 pg/mL) whether they display high titer by Gyros assay or not. We found that 2.9% (4 of 139) of patients had auto-Abs neutralizing 10 ng/mL of IFN-α2 and/or IFN-ω. Three of these patients had auto-Abs neutralizing high concentrations of IFN-α2 and IFN-ω and 1 patient had only auto-Abs neutralizing high concentrations of IFN-ω. Moreover, some patients presented auto-Abs neutralizing only 100 pg/mL of IFN-α2 and/or IFN-ω but not higher concentrations. We found that 5% (7 of 139) of patients had auto-Abs neutralizing only lower concentrations of IFN-α2 and IFN-ω. Two of these patients had auto-Abs against only IFN-α2 and 5 patients had auto-Abs against IFN-ω only.

Lastly, no patients with neutralizing activity against 10 ng/mL of IFN-β were observed in our study. Overall, 7.9% of the patients displays neutralizing activity against IFN-α and/or IFN-ω.

Only 27% (3 of 11) of these patients with neutralizing auto-Abs presented high levels of auto-Abs in Gyros assay, other patients having intermediate levels in Gyros assay or even no detectable auto-Abs. Two patients with neutralizing auto-Abs against IFN-α and 3 patients with neutralizing auto-Abs against IFN-ω had no detectable auto-Abs in Gyros assay (Table 1). Moreover, unlike previous studies, where most samples with high titer of auto-Abs in Gyros assay were neutralizing in vitro [14], 79% (11 of 14) of patients...
with high levels of auto-Abs against IFN-α2 or IFN-ω in Gyros assay had no neutralizing activity in vitro. Of note, this was mostly observed for IFN-ω, where 100% of high titers was not neutralizing while 70% for IFN-α2 (Fig. 1A and B).

Detection of Auto-Abs to Type I IFNs by ELISA

All samples were tested for the presence of anti IFN-α2 auto-Abs by a commercially available ELISA kit assay. Interestingly, only 6.5% (9 of 139) of patients had anti IFN-α2 auto-Abs detectable by ELISA whereas 68% (94 of 139) patients has auto-Abs against IFN-ω by Gyros technology.

Three patients had low levels of anti IFN-α2 auto-Abs (<100 ng/mL). These patients had also intermediate titer of anti IFN-α2 auto-Abs in Gyros assays, but no neutralizing activity.

Moreover, six patients had high titer of anti IFN-α2 auto-Abs in ELISA (530.7 ng/mL or >1000 ng/mL). In Gyros assay, IFN-α2 auto-Abs were present with high levels in 3 patients, intermediate level in 1 patient and no detectable in 2 patients (Fig. 2A).

Interestingly, all the 6 patients with high titer auto-Abs in ELISA have a neutralizing activity in vitro against IFN-α2 (Fig. 2B). Three of these patients had auto-Abs neutralizing high concentrations of IFN-α2 and IFN-ω. 1 of these patients had auto-Abs neutralizing high concentrations of IFN-α2, and 2 of these patients had auto-Abs neutralizing lower concentrations of IFN-α2. High levels IFN-α2 auto-Abs were notably detected in ELISA in the 2 patients with neutralizing activity against IFN-α2 but without detectable auto-Abs in Gyros assay. Presence of high titer of auto-Abs against IFN-α2 in ELISA seems to be better correlated with neutralizing activity of auto-Abs than Gyros technology.

Among the 6 patients with high titer IFN-α2 antibodies detected by ELISA, 3 patients survived. A sample was collected 10 months after COVID-19 pneumonia for 2 of these surviving patients and analyzed by ELISA. Interestingly, high levels of IFN-α2 antibodies remained in these patients. We indeed observed a level of anti IFN-α2 auto-Abs that remained >1000 ng/mL in one patient and a level that decreased from 530.7 ng/mL to 239.3 ng/mL in another patient. These results indicate a persistence of auto-Abs at least 10 months after COVID-19 infection. These patients continue to be followed in hospital, especially in order to see if they develop again a severe COVID-19 pneumonia or other viral infections.

### Increased Mortality of Patients with Neutralizing Auto-Abs to Type I IFNs

We compared the clinical characteristics of patients having neutralizing auto-Abs against type I IFNs with patients without neutralizing auto-Abs (Table 2). Although differences were not significant with the patients without neutralizing auto-Abs, most of patients with neutralizing auto-Abs were men (82%). Eighty-two percent of patients with neutralizing auto-Abs was over the age of 65 years, and neutralizing auto-Abs was present in 12.5% of patients with critical COVID-19 over the age of 80 years.

In our study, 67% (93 of 139) of the patients was labeled as having a full engagement status, meaning with no limitation of care. The full engagement status was significantly less frequent in patient with auto-Abs (36%) than in patients without auto-Abs (70%). Nevertheless, no differences were
seen between patients with or without auto-Abs for comorbidities and biological characteristics at hospital admission. Moreover, there was no difference in the proportion of patients that were transferred to an ICU between patients with or without auto-Abs.

Finally, it was recently shown that at least 18% of patients who died of COVID-19 pneumonia had auto-Abs capable of neutralizing 100 pg/mL type I IFNs in plasma 1/10 [14]. In our study, the mortality was significantly more frequent in patients with neutralizing auto-Abs as 55% (6 of 11) of patients with auto-Abs died versus 18% (23 of 128) of patients without auto-Abs. Twenty-one percent of critical patients of our cohort who died from COVID-19 pneumonia had auto-Abs capable of neutralizing 100 pg/mL type I IFNs in plasma 1/10.

We compared clinic characteristics of deceased patients having neutralizing auto-Abs against type I IFNs with deceased patients without neutralizing auto-Abs. Although the deceased patients were usually elderly men,
no significant differences were observed between the two groups (Table 3).

**Characteristics of Patients with Non-Neutralizing Auto-Abs to Type I IFNs**

Many patients of the cohort had frequently auto-Abs, sometimes with high titer, without detected neutralizing activity. Indeed, 79% (11 of 14) of patients with high titer of auto-Abs had no apparent neutralizing activity in vitro. We compared characteristics of patients with neutralizing auto-Abs with patients with high titer auto-Abs without neutralizing activity and patients with intermediate titer auto-Abs without neutralizing activity (Table 4). Men and old patients were more frequent in patients with neutralizing Auto-Abs (82% of male patients, 9 of 11 and 82% of patients over 65 years old, 9 of 11) than in patients with auto-Abs without neutralizing activity (58% of male patients, 57 of 98 and 52% of patients over 65 years old, 51 of 98) but the difference was not significant (\(p = 0.19\) and \(p = 0.11\) respectively). Deaths were significantly more frequent in patients with neutralizing antibodies than in patients with intermediate titer auto-Abs without neutralizing activity, without the difference being significant (\(p = 0.01\)). Moreover, there were more deaths in patients with neutralizing antibodies than in patients with high titer auto-Abs without neutralizing activity, without the difference being significant (\(p = 0.18\)). These auto-Abs might be falsely positive or might be able to neutralize even lower amounts of type I IFNs.
We assessed the prevalence of auto-Abs against type I IFNs in 139 patients hospitalized for critical COVID-19 pneumonia in medicine departments dedicated to treat COVID-19 patients.

As many as 77% of patients with critical COVID-19 have auto-Abs against IFN-α and/or IFN-ω by Gyros technology, mainly with intermediate titer and less frequently with high titer. These results are much higher than those previously described as Bastard et al. described high or intermediate levels of IgG auto-Abs against IFN-α2 and/or IFN-ω in about 20% of patients with critical COVID-19 [14]. Our high proportion of intermediate titers could be explained by calibration issues of the Gyros. This nevertheless does not seem to affect the high titers, which remain the most relevant.

However, 79% of our patients with high levels of auto-Abs had no neutralizing activity. We found circulating auto-Abs that neutralized 100 pg/mL IFN-α2 and/or IFN-ω in plasma 1/10 in only 7.9% of our patients.

These results may reflect a lack of specificity of Gyros technology, especially for auto-Abs against IFN-ω and for intermediate titer of auto-Abs. Alternatively, there may be insufficient sensitivity of neutralization assays. A third possible explanation is that these auto-Abs may be specific for other type I IFN epitopes, the recognition of which would not interfere with type I IFN binding to its receptor.

We also evaluated a commercially available ELISA kit in our laboratory for detection of IFN-α2 auto-Abs. We did not search anti IFN-α2 auto-Abs by ELISA in general population. Nevertheless, according to manufacturer’s results, in a panel of 57 serum samples from randomly selected apparently healthy donors and patients suffering from various diseases, anti-IFN α2 levels ranged between 0 and 120.5 ng/mL (mean:17.4 ng/mL; SD: 26.7 ng/mL). Moreover, this commercial ELISA was recently evaluated [9]; a cut off of

### Table 2 Clinical characteristics of patients

| Patients with neutralizing auto-Abs at 100 pg/ml type I IFNs (1/10 dilution) | Patients with auto-Abs without neutralizing activity or without antibodies | p values |
|---|---|---|
| **Demographics** | | |
| n | 11 | 128 |
| Age, mean ± Standard deviation (SD), years | 68.7 ± 14.9 | 64 ± 15.8 | p=0.29 |
| ≥ 65 years old | 9 (82%) | 65 (51%) | p=0.06 |
| Sex (male) | 9 (82%) | 77 (60%) | p=0.2 |
| **Clinical features** | | |
| Full engagement (patients with no limitation of care) | 4 (36%) | 89 (70%) | p=0.04 |
| Diabetes | 3 (27%) | 51 (40%) | p=0.5 |
| Obesity | 3 (27%) | 40 (31%) | p=1 |
| Hypertension | 4 (36%) | 67 (52%) | p=0.4 |
| History of cardiovascular disease, stroke, peripheral artery disease, heart failure | 3 (27%) | 27 (21%) | p=0.7 |
| History of chronic obstructive pulmonary disease, asthma, emphysema, fibrosis | 1 (9%) | 22 (17%) | p=0.69 |
| Solid organ transplantation | 1 (9%) | 0 (0%) | p=0.08 |
| HIV | 0 (0%) | 2 (1.6%) | p=1 |
| Immunosuppressant drugs and/or long-term oral corticosteroids | 2 (18%) | 10 (9%) | p=0.24 |
| Malignancy (active) | 2 (18%) | 8 (6%) | p=0.18 |
| **Biological characteristics at hospital admission** | | |
| C protein reactive, mean ± SD, mg/L | 155 ± 89 | 124 ± 90 | p=0.2 |
| Lymphocyte count, mean ± SD, /µL | 923 ± 269 | 1196 ± 1044 | p=0.3 |
| Creatinine mean ± SD, µmol/L | 273 ± 434 | 84 ± 34 | p=0.21 |
| **Clinical outcomes** | | |
| Intubed | 1 (9%) | 21 (16%) | p=1 |
| Death | 6 (55%) | 23 (18%) | p=0.01 |

Data are presented as a number (percentage), unless otherwise noted. A Fisher test was used to analyze the effect of dichotomous variables and a Mann–Whitney test for continuous variables. Significant values are noted in italics.

### Discussion

We assessed the prevalence of auto-Abs against type I IFNs in 139 patients hospitalized for critical COVID-19 pneumonia in medicine departments dedicated to treat COVID-19 patients.

As many as 77% of patients with critical COVID-19 have auto-Abs against IFN-α and/or IFN-ω by Gyros technology, mainly with intermediate titer and less frequently with high titer. These results are much higher than those previously described as Bastard et al. described high or intermediate levels of IgG auto-Abs against IFN-α2 and/or IFN-ω in about 20% of patients with critical COVID-19 [14]. Our high proportion of intermediate titers could be explained by calibration issues of the Gyros. This nevertheless does not seem to affect the high titers, which remain the most relevant.

However, 79% of our patients with high levels of auto-Abs had no neutralizing activity. We found circulating auto-Abs that neutralized 100 pg/mL IFN-α2 and/or IFN-ω in plasma 1/10 in only 7.9% of our patients.
34 ng/mL was established and results > 100 ng/mL were considered as high. Three of our patients had anti IFN-α2 auto-Abs levels <100 ng/mL, which were considered as low, and also presented no neutralizing activity in vitro against IFN-α. Moreover, in our study, 6 patients had high titer of anti IFN-α2 auto-Abs in ELISA and presented all a neutralizing activity in vitro against IFN-α2. Goncalves et al. [9] showed similar results and determined a cut-off of 1000 ng/mL correlated with neutralization assays. In their study, ability of auto-Abs to neutralize only high concentrations of IFN-α2 (10 ng/mL) was investigated. In our study, we tested the neutralizing activity of all these samples against IFNs in vitro at high concentrations and lower concentrations. In one of our patient, a concentration of 530.7 ng/mL was shown to be correlated with ability of auto-Abs to neutralize 100 pg/mL IFN-α2 in plasma 1/10. These results have to be confirmed in more samples, and it is also necessary to determine cut off correlated with neutralizing activity against low concentration of IFN-α2. High levels IFN-α2 auto-Abs were notably detected in ELISA in 2 patients with neutralizing activity against IFN-α2 but without detectable auto-Abs in Gyros assay. The difference between Gyros technology
and Invitrogen ELISA could be explained by the nature of the conjugate, secondary antibody, goat anti-human IgG, and HRP–conjugated human IFN-α2 protein respectively, probably leading to a better detection of auto antibodies with ELISA. Overall, high titer auto-Abs against IFN-α2 in ELISA seems to be better correlated with neutralizing activity of auto-Abs than Gyros.

The proportion of patients with type I IFNs neutralizing auto-Abs is lower than those obtained in previous studies as it was described auto-Abs that neutralized 100 pg/mL IFN-α2 and/or IFN-α in plasma 1/10 in 13.6% of the critical patients [14]. However, our cohort included only patients that were at the first day of hospitalization in medicine departments dedicated to treat COVID-19 patients, with an oxygen support ≥ 6 L/min with oxygen mask, for more than 6 h. It was indeed initially constituted in order to assess associations between treatments and outcomes like need of ventilation mechanic and mortality [16]. Overall, severity of patients studied here was potentially lower than that of patients hospitalized in intensive care units [5, 6, 9–11, 14].

Previous studies showed that patients with neutralizing auto-Abs against type I IFNs were mostly men (94%) and that half were older than 65 years [5]. Moreover, it was recently shown that proportion of patients with critical COVID-19 having neutralizing auto-Abs increased with age [14]. In our study, although differences were not significant with regard to the patients without neutralizing auto-Abs, most of patients (82%) with neutralizing auto-Abs were men and over the age of 65 years. The full engagement status was significantly less frequent in patient with auto-Abs than in patients without auto-Abs but like other studies [7, 9], there was no difference for comorbidities. Unlike to study by Troya et al. [7], where a significant correlation between the presence of auto-Abs neutralizing type I IFNs, raised levels of C-protein reactive, and low lymphocytes counts was observed, in our study, no difference was seen for these biological characteristics. It is nevertheless important to note that we compare biological characteristics obtained at hospital admission, and not maximum levels of C-protein reactive and minimum levels of lymphocytes count.

Neutralizing antibodies against type I IFNs was described in almost all of patients with auto immune polyendocrinopathy syndrome type I [18–20] but also in patients with thymoma, myasthenia gravis [21] and systemic lupus erythematosus [22–24]. In our cohort, these autoimmune diseases were not described in patients with neutralizing auto-Abs, although 2 of 11 (18%) patients with neutralizing Auto-Abs against type I IFNs had a history of psoriasis. Research of antinuclear antibodies (ANA) did not reveal high titer of ANA in the 6 patients that were analyzed.

Like in a previous study [14], the presence of neutralizing auto-Abs is associated with a more frequent mortality. Up to 21% of patients who died of COVID-19 pneumonia in our study had auto-Abs capable of neutralizing 100 pg/mL type I IFNs in plasma 1/10. Deceased patients with auto-Abs did not present overt clinical differences with deceased patients without auto-Abs. These results confirm importance of the IFN-I pathway in the defense against SARS-CoV-2 infection.

Overall, the detection of neutralizing Auto-Abs against type I IFNs is therefore important due to clinical applications, notably therapeutics. All COVID-19 patients should be screened for these autoantibodies in order to predict which patients infected by SARS-CoV-2 are at higher risk of life-threatening COVID-19. Moreover, early identification of COVID-19 patients with auto-Abs should prompt early treatment and preventive management.

As high titers auto-Abs against IFN-2 in ELISA are well correlated with neutralizing activity of auto-Abs, this commercial ELISA should be used to detect easily, in routine clinical practice, neutralizing anti IFN-2 auto-Abs. Nevertheless, to our knowledge, no commercial ELISA exists to detect anti IFN- auto-Abs. It would be necessary to complete detection of anti IFN-2 auto-Abs with a functional evaluation of neutralizing anti IFN- auto-Abs, notably if low IFN-2 auto-Abs levels or no IFN-2 auto-Abs are detected.

Lastly, our study describes persistence of neutralizing Auto-Abs almost 1 year after COVID-19 infection in two patients. Risk associated with this persistence is not known to date but must be explored, especially given the potential impact of these auto-Abs on the severity of other viral diseases, as described for adverse reactions following yellow-fever vaccination [25].

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Data Availability Clinical data files are stored at Robert Ballanger Hospital. It may be shared if needed.

Declarations

Ethics Approval

Consent for Publication

Consent to Participate

Competing Interests

References

1. Levin AT, Hanage WP, Owusu-Boateng N, Cochran KB, Walsh SP, Meyerowitz-Katz G. Assessing the age specificity of infection fatality rates for COVID-19: systematic review, meta-analysis, and public policy implications. Eur J Epidemiol. 2020;35(12):1123–38. https://doi.org/10.1007/s10654-020-00698-1.

2. O’Driscoll M, Ribeiro Dos Santos G, Wang L, Cummings DAT, Azman AS, Faireux J, Fontanet A, Cauchemez S, Salje H. Age-specific mortality and immunity patterns of SARS-CoV-2. Nature. 2021;590(7844):140–5. https://doi.org/10.1038/s41586-020-2918-0.

3. Zhang Q, Bastard P, Liu Z, Le Pen J, Moncada-Velez M, Chen J, Ogishi M, Sabli IKD, Hodeib S, Korol C, Rosain J, Bilguvar K, Ye J, Bolze A, Bigio B, Yang R, Arias AA, Zhou Q, Zhang Y, Onodi F, Korniotsi S, Karpf L, Philippot Q, Cibibi M, Bonnet-Madin L, Dorgam K, Smith N, Schneider WM, Razisky BS, Hoffmann H-H, Michailidis E, Moens L, Han JE, Lorenzo L, Bizien L, Meade P, Neehus A-L, Ugurbil AC, Corneau A, Kerner G, Zhang P, Rapaport F, Seeleuthner Y, Manry J, Masson C, Schmitt Y, Schlüter A, Le Voye T, Khan T, Li J, Fellay J, Roussel L, Shahrooei M, Alosaifei MF, Mansouri D, Al-Saud H, Al-Mulla F, Amourfi F, Al-Muhsen, S. Z.; Alsohime, F.; Al Turki, S.; Hasanato, R.; van de Beek, D.; Biondi, A.; Bettini, L. R.; D’Angio’, M.; Bonfanti, P.; Imberti, L.; Sottini, A.; Paghera, S.; Quiros-Roldan, E.; Rossi, C.; Oler, A. J.; Tompkins, M. F.; Alba, C.; Vandernoot, I.; Goffard, J.-C.; Smits, G.; Migeotte, I.; Haerynck, F.; Soler-Palacin, P.; Martin-Nalda, A.; Colobran, R.; Morange, P.-E.; Keles, S.; Çolkesen, F.; Ozelik, Y.; Yasar, K. S.; Senoglu, S.; Karabela, Š. N.; Rodríguez-Gallego, C.; Novelli, G.; Hraiech, S.; Tandjaojui-Lambiotte, Y.; Duval, X.; Lauoenuan, C.; COVID-STORM Clinicians†; COVID Clinicians†; Imagine COVID Group†; French COVID Cohort Study Group†; CoV-Contact Cohort†; Amsterdam UMC Covid-19 Biobank†; COVID Human Genetic Effort†; NIAID-USUHS/TAGC COVID Immunity Group†; Snow, A. L.; Dalgard, C. L.; Milner, J. D.; Vinh, D. C.; Mogensen, T. H.; Marr, N.; Spaan, A. N.; Boisson, B.; Boisson-Dupuis, S.; Bustamante, J.; Puel, A.; Ciancanelli, M. J.; Meys, I.; Maniatis, T.; Soumelis, V.; Amara, A.; Nussenzweig, M.; Garcia-Sastre, A.; Kramer, F.; Pujol, A.; Duffy, D.; Lifton, R. P.; Zhang, S.-Y.; Gorochov, G.; Bézia, V.; Jouanguy, E.; Sancho-Shimizu, V.; Rice, C. M.; Abel, L.; Notarangelo, L. D.; Cobat, A.; Su, H. C.; Casanova, J.-L.; Inborn errors of type I IFN immunity in patients with life-threatening COVID-19. Sci Adv. 2020;37(655):eab4570. https://doi.org/10.1126/science. abd4570.

4. Asano, T.; Boisson, B.; Onodi, F.; Matouzzo, D.; Moncada-Velez, M.; Maglorius Renkilaraj, M. R. L.; Zhang, P.; Meertens, L.; Bolze, A.; Materna, M.; Korniotsi, S.; Gervais, A.; Talouarn, E.; Bigio, B.; Seeleuthner, Y.; Bilguvar, K.; Zhang, Y.; Neehus, A.-L.; Ogishi, M.; Pelham, S. J.; Le Voye, T.; Rosain, J.; Philippot, Q.; Soler-Palacin, P.; Colobran, R.; Martin-Nalda, A.; Rivière, J. G.; Tandjaojui-Lambiotte, Y.; Chaib, K.; Shahrooei, M.; Darazam, I. A.; Olayei, N. A.; Mansouri, D.; Hatipoğlu, N.; Palabiyik, F.; Ozelik, T.; Novelli, G.; Novelli, A.; Casari, G.; Aiuti, A.; Carrera, P.; Bonedan, S.; Barzaghi, F.; Rovere-Querini, P.; Tresoldi, C.; Franco, J. L.; Rojas, J.; Reyes, L. F.; Bustos, I. G.; Arias, A. A.; Morelle, G.; Christèle, K.; Troya, J.; Planas-Serra, L.; Schlüter, A.; Gut, M.; Pujol, A.; Allende, L. M.; Rodriguez-Gallego, C.; Flores, C.; Cabrera-Marante, O.; Pleguenuelo, D. E.; de Diego, R. P.; Keles, S.; Ayetkin, G.; Akan, O. M.; Bryceyn, Y. T.; Bergman, P.; Brodin, P.; Smole, D.; Smith, C. I. E.; Norlin, A.-C.; Campbell, T. M.; Covill, L. E.; Hammarsström, L.; Pan-Hammarsström, Q.; Abolhassani, H.; Mane, S.; Marr, N.; Ata, M.; Al Ali, F.; Khan, T.; Spaan, A. N.; Dalgard, C. L.; Bonfanti, P.; Biondi, A.; Tubiana, S.; Burdet, C.; Nussbaum, R.; Kahn-Kirby, A.; Snow, A. L.; COVID Human Genetic Effort†; COVID-STORM Clinicians†; COVID Clinicians†; Imagine COVID Group†; French COVID Cohort Study Group†; CoV-Contact Cohort†; Amsterdam UMC Covid-19 Biobank†; NIAID-USUHS COVID Study Group†; Bustamante, J.; Puel, A.; Boisson-Dupuis, S.; Zhang, S.-Y.; Bézia, V.; Liton, R. P.; Bastard, P.; Notarangelo, L. D.; Abel, L.; Su, H. C.; Jouanguy, E.; Amara, A.; Soumelis, V.; Cobat, A.; Zhang, Q.; Casanova, J.-L.; X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. Sci. Immunol. 2021, 6 (62), eabl4348. https://doi.org/10.1126/sciimmunol.abl4348.

5. Bastard, P.; Rosen, L. B.; Zhang, Q.; Michailidis, E.; Hoffmann, H.-H.; Zhang, Y.; Dorgham, K.; Philippot, Q.; Rosain, J.; Bézia, V.; Manry, J.; Shaw, E.; Haljasmägi, L.; Peterson, P.; Lorenzo, L.; Bizien, L.; Trouillet-Assant, S.; Dobbs, K.; de Jesus, A. A.; Belot, A.; Kallaste, A.; Catherinot, E.; Tandjaoui-Lambiotte, Y.; Le Pen, J.; Kerner, G.; Bigio, B.; Seeleuthner, Y.; Yang, R.; Bolze, A.;
to COVID Group§; NH-COVAIR Study Group§; Danish CHGE§; Danish Blood Donor Study§; St. James’s Hospital; SARS CoV2 Interest group§; French COVID Cohort Study Group§; Imagine-COVID Group§; The Milieu Intérieur Consortium§; CoV-Contact Cohort§; Amsterdam UMC Covid-19; Biobank Investigators§; COVID Human Genetic Effort§; CONSTATANCES cohort§; 3C-Dijon Study§; Cerba Health-Care§; Etablissement du Sang study group§; Anderson, M. S.; Boisson, B.; Bézat, V.; Zhang, S.-Y.; Vandrekos, E.; Hermine, O.; Pujol, A.; Peterson, P.; Mogensen, T. H.; Rowen, L.; Mond, J.; Debette, S.; de Lamballerie, X.; Duval, X.; Mentré, F.; Zins, M.; Soler-Palacin, P.; Colobran, R.; Gorochov, G.; Solanich, X.; Susen, S.; Martinez-Picado, J.; Raout, D.; Vasse, M.; Gregersen, P. K.; Piemonti, L.; Rodríguez-Gallego, C.; Notarangelo, L. D.; Su, H. C.; Ksidis, K.; Okada, S.; Puel, A.; Jouanguy, E.; Rice, C. M.; Tiberghien, P.; Zhang, Q.; Cohat, A.; Abel, L.; Casanova, J.-L.; Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. Sci. Immunol. 2021, 6 (62), eab43430. https://doi.org/10.1126/sciimmunol.ab14340.

15. Region, L. P. La surmortalité durant l’épidémie de Covid-19 dans les départements franciliens https://www.ors-idf.org/nos-travaux/publications/la-surmortalite-durant-epidemie-de-covid-19-dans-les-departements-franciliens/ (accessed 2021-09-09).

16. Rossi B, Nguyen LS, Zimmermann P, Boucenna F, Dubret L, Baucher L, Guillot H, Boudouyre M-A, Allenbach Y, Salem J-E, Barsoum P, Oufella A, Gros H. Effect of tocilizumab in hospitalized patients with severe COVID-19 pneumonia: a case-control cohort study. Pharmaceuticals. 2020;13(10):317. https://doi.org/10.3390/ph13100317.

17. Clinical Spectrum https://www.covid19treatmentguidelines.nih.gov/overview/clinical-spectrum/ (accessed 2021-09-14).

18. Meager A, Visvalingam K, Peterson P, Möll K, Murumägi A, Krohn K, Eskelin P, Perheentupa J, Husebye E, Kadota Y, Willcox N. Anti-interferon autoantibodies in autoimmune polyendocrinopathy syndrome type 1. PLoS Med. 2006;3(7): e289. https://doi.org/10.1371/journal.pmed.0030299.

19. Kärner J, Meager A, Laan M, Maslovskaja J, Pihlap M, Remm A, Juronen E, Wolff ASB, Husebye ES, Podkrajšek KT, Bratanic N, Battelino T, Willcox N, Peterson P, Kisand K. Anti-cytokine autoantibodies suggest pathogenetic links with autoimmunity polyendocrinopathy syndrome type 1. PLoS Med. 2006;3(7): e289. https://doi.org/10.1371/journal.pmed.0030289.

20. Bastard P, Orlova E, Sozaeva L, Lévy R, James A, Schmitt MM, Ochoa S, Kareva M, Rodina Y, Gervais A, Le Voyer T, Rosair J, Philippon Q, Neehus A-L, Shaw E, Migaud M, Bizien L, Ekwall O, Berg S, Beccuti G, Ghizzoni L, Thiriez G, Pavot A, Goujard C, Frémond M-L, Carter E, Rothenbuhler A, Linglart A, Mignot B, Comte A, Cheikh N, Hermine O, Breivik L, Husebye ES, Humbert S, Rohrllich P, Coaquette A, Vuoto F, Faure K, Mahtlaoui N, Kotnik P, Battelino T, Trebušak Podkrajšek K, Kisand K, Ferré EMM, DiMaggio T, Rosen LB, Burbelo PD, McIntyre M, Kann NY, Scherbina A, Pavlova M, Kolodkina A, Holland SM, Zhang S-Y, Crow YJ, Notarangelo LD, Su HC, Abel L, Anderson MS, Jouanguy E, Neven B, Puel A, Casanova J-L, Lionakis MS. Pre-existing autoantibodies to type I IFNs underlie critical COVID-19 pneumonia in patients with APS-1. J Exp Med. 2021:218(7): e20210554. https://doi.org/10.1084/jem.20210554.

21. Meager A, Wadhwa M, Dilger P, Bird C, Thorpe R, Newsom-Davis J, Willcox N. Anti-cytokine autoantibodies in autoimmunity: preponderance of neutralizing autoantibodies against interferon-alpha, interferon-omega and interleukin-12 in patients with thymoma and/or myasthenia gravis: cytokine autoantibodies, thymoma and myasthenia gravis. Clin Exp Immunol. 2003;132(1):128–36. https://doi.org/10.1046/j.1365-2249.2003.02113.x.

22. Howe HS, Leung BPL. Anti-cytokine autoantibodies in systemic lupus erythematosus. Cells. 2019;9(1):72. https://doi.org/10.3390/cells9010072.

23. Morimoto AM, Flesher DT, Yang J, Wolslegel K, Wang X, Brady A, Abbas AR, Quarmby V, Wakshull E, Richardson B, Townsend MJ, Behrens TW. Association of endogenous anti-interferon-α autoantibodies with decreased interferon-pathway and disease activity in patients with systemic lupus erythematosus. Arthritis Rheum. 2011;63(8):2407–15. https://doi.org/10.1002/art.30399.

24. Gupta S, Tatoulis IP, Rosen LB, Hasni S, Alevizos I, Manza ZG, Rivera J, Jiang C, Siegel RM, Holland SM, Moutsopoulos HM, Browne SK. Distinct functions of autoantibodies against interferon in systemic lupus erythematosus: a comprehensive analysis of anticytokine autoantibodies in common rheumatic diseases: anti-interferon functions in SLE. Arthritis Rheumatol. 2016;68(7):1677–87. https://doi.org/10.1002/art.36067.

25. Bastard P, Michailidis, E.; Hoffmann, H.-H.; Chbihi, M.; Le Voyer, T.; Rosair, J.; Philippon, Q.; Seeleuthner, Y.; Gervais, A.; Materna, M.; de Oliveira, P. M. N.; Maia, M. de L. S.; Dinis Ano Bom, A. P.; Azamor, T.; Araújo da Conceição, D.; Gourdouris, E.; Homma, A.; Slesak, G.; Schäfer, J.; Pulendran, B.; Miller, J. D.; Huits, R.; Yang, R.; Rosen, L. B.; Bizien, L.; Lorenzo, L.; Chrabieh, M.; Erazo, L. V.; Rozenberg, F.; Jeljeli, M. M.; Bézat, V.; Holland, S. M.; Cohat, A.; Notarangelo, L. D.; Su, H. C.; Ahmed, R.; Puel, A.; Zhang, S.-Y.; Abel, L.; Seligman, S. J.; Zhang, Q.; MacDonald, M. R.; Jouanguy, E.; Rice, C. M.; Casanova, J.-L. Auto-antibodies to type I IFNs can underlie adverse reactions to yellow fever live attenuated vaccine. J. Exp. Med. 2021; 218 (4), 20202486. https://doi.org/10.1084/jem.20202486.

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