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Impaired immunogenicity to COVID-19 vaccines in autoimmune systemic diseases. High prevalence of non-response in different patients' subgroups

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Abbreviations: Autoimmune systemic diseases, ASD; Rheumatoid factor, RF; Anti-citrullinated protein antibodies, ACPA; Rheumatoid arthritis, RA; Systemic lupus erythematosus, SLE; Systemic sclerosis, SSC; Cryoglobulinemic vasculitis, CV; Adverse events, AEs; Neutralizing antibody, NAb; World Health Organization, WHO; Binding Antibody Units, BAU.

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ABSTRACT

Autoimmune systemic diseases (ASD) may show impaired immunogenicity to COVID-19 vaccines. Our prospective observational multicenter study aimed to evaluate the seroconversion after the vaccination cycle and at 6-12-month follow-up, as well the safety and efficacy of vaccines in preventing COVID-19.

The study included 478 unselected ASD patients (mean age 59 ± 15 years), namely 101 rheumatoid arthritis (RA), 38 systemic lupus erythematosus (SLE), 265 systemic sclerosis (SSc), 61 cryoglobulinemic vasculitis (CV), and a miscellanea of 13 systemic vasculitis. The control group included 502 individuals from the general population (mean age 59 ± 14 SD years). The immunogenicity of mRNA COVID-19 vaccines (BNT162b2 and mRNA-1273) was evaluated by measuring serum IgG-neutralizing antibody (NAb) (SARS-CoV-2 IgG II Quant antibody test kit; Abbott Laboratories, Chicago, IL) on samples obtained within 3 weeks after vaccination cycle.

The short-term results of our prospective study revealed significantly lower NAb levels in ASD series compared to controls [286 (53–1203) vs 825 (451–1542) BAU/mL, $p < 0.0001$], as well as between single ASD subgroups and controls. More interestingly, higher percentage of non-responders to vaccine was recorded in ASD patients compared to controls [13.2% (63/478), vs 2.8% (14/502); $p < 0.0001$]. Increased prevalence of non-response to vaccine was also observed in different ASD subgroups, in patients with ASD-related interstitial lung disease ($p = 0.009$), and in those treated with glucocorticoids ($p = 0.002$), mycophenolate-mofetil ($p < 0.0001$), or rituximab ($p < 0.0001$). Comparable percentages of vaccine-related adverse effects were recorded among responder and non-responder ASD patients.

Patients with weak/absent seroconversion, believed to be immune to SARS-CoV-2 infection, are at high risk to develop COVID-19. Early determination of serum NAb after vaccination cycle may allow to identify three main groups of ASD patients: responders, subjects with suboptimal response, non-responders. Patients with suboptimal response should be prioritized for a booster-dose of vaccine, while a different type of vaccine could be administered to non-responder individuals.

1. Introduction

Observational cohorts demonstrated that patients with autoimmune systemic diseases (ASD) have a higher prevalence of COVID-19 compared to the general population [1–5]. Moreover, worse clinical outcomes, including higher rates of hospitalization and death, have been reported in older ASD patients and/or in those with pre-existing ASD-related lung fibrosis [2,6–8]. Therefore, ASD are now considered among patients' populations highly vulnerable to SARS-CoV-2 infection and morbidity possibly due to their deep immune-system alterations with the possible contribution of ongoing immunosuppressive treatments [2,9–11]. In the early stages of the COVID-19 pandemic, the risk of developing COVID-19 in these patients has been minimized by the stringent lockdown measures adopted by national health authorities together with the broad use of telemedicine allowing the essential contact between patients and referral centers. Subsequently, with the introduction of the COVID-19 mass vaccination campaign, ASD patients were included among population subgroups prioritized for vaccination in several countries, including Italy (<https://www.trovanorme.salute.gov.it/norme/renderPdf.spring?seriegu=SG&datagu=24/03/2021&redaz=21A01802&artp=1&art=1&subart=1&subart1=10&vrs=1&prog=002>) [12]. Data from clinical trials and real-life studies confirmed a good immunogenicity and safety of available vaccines in ASD patients, in particular those mRNA-based [13–16]. However, a number of factors including immune-system disorders, underlying chronic inflammatory diseases, disease activity at the time of vaccination, concurrent immune-modifier therapies (high glucocorticoid dosages, rituximab, etc.), and older age have been suggested to negatively affect the immune response to vaccines in individual patients and/or particular disease subsets [14–17]. Starting from the last February 2021 the COVID-19 & Italian ASD Study Group organized a multicenter prospective observational study with the main purpose of evaluating the immunogenicity and safety of COVID-19 vaccination in ASD patients, the possible effects of ongoing treatments, as well as its ability to prevent symptomatic SARS-CoV-2 infection over long-term clinical follow-up. Here we report the data regarding the COVID-19 vaccine immunogenicity and safety evaluated in a large ASD patients' series after the completion of vaccination cycle.

2. Patients and methods

Our prospective, long-term, observational, multicenter survey, involved 21 centers of the COVID-19 & ASD Italian Study Group equally distributed among the Italian regions. The study included some well-defined ASD in order to deal with sufficiently homogeneous and representative disease subgroups. In particular, we consecutively recruited patients with rheumatoid factor (RF) and/or anti-citrullinated protein antibodies (ACPA) positive rheumatoid arthritis (RA), systemic lupus erythematosus (SLE), systemic sclerosis (SSc), cryoglobulinemic vasculitis (CV), and a miscellanea of other systemic vasculitis; disease definition and patients' clinical assessment were carried out following updated classification criteria and current methodologies [18]. The primary aim of the study was the evaluation of the immunogenicity of COVID-19 vaccines at the end of vaccination cycle; secondary aims were to investigate the safety of COVID-19 vaccines and the impact of ongoing immune-modifier medications. The serum samples were collected between March and July 2021 in order to detect the early antibody response (1–3 weeks after the administration of the second dose of vaccine).

Demographic characteristics, specific vaccine administered and vaccine-related adverse events (AEs), measures of disease activity, ASD-related complications, and ongoing treatment at the time of vaccination were recorded. Regarding the latter, the peri-vaccination management of immune-modifier medications followed the recommendations of the Italian Society of Rheumatology (<https://www.reumatologia.it/vaccinazioni>), taking in account both individual patient's clinical characteristics (e.g. major comorbidities) and disease activity. Individuals treated with rituximab or cyclophosphamide during the last 6 months before vaccination ($n = 7$) were excluded from the study. Additional exclusion criteria were pregnancy, history of allergic reactions after previous vaccination, and/or recent SARS-CoV-2 infection.

The monitoring of safety in all participants was carried out by telephone interview 1–3 weeks before the first dose, then 2 weeks after the first and second dose in order to detect possible vaccine-related AEs using a standardized symptoms assessment questionnaire. The immunogenicity of COVID-19 vaccines is evaluated by measuring the titer of IgG neutralizing antibody (NAb) against SARS-CoV-2 trimeric spike S1/S2 glycoproteins on serum samples obtained within 1–3 weeks after completion of the vaccination cycle using the SARS-CoV-2 IgG II Quant

antibody test kit (Abbott Laboratories, Chicago, IL). As recommended by World Health Organization (WHO), antibody titers are expressed as Binding Antibody Units (BAU)/ml, with a cut-off for positive testing of 7 BAU/ml.

The study protocol was approved by local ethic committee (RETRO-CoV2 study code #17886_bio); informed consent was obtained from all patients before participation.

A sample of 502 individuals (73% females, mean age 59 ± 14 years)

Table 1

Comparison between SARS-CoV-2 IgG neutralizing antibodies (NAb) negative and positive individuals.

	ASD Whole series (n = 478)	SARS-CoV-2 IgG NAb negative (n = 63)	SARS-CoV-2 IgG NAb positive (n = 415)	P value Odds Ratio C.I.
Age, years	59 ± 15	61 ± 15	59 ± 16	0.284
Female gender, n (%)	401 (83.9)	50 (79.4)	351 (84.6)	0.294
Past COVID-19, n (%)	15 (3.1)	0 (0)	15 (3.6)	0.125
ASD-related complications				
Interstitial lung disease, n (%)	112 (23.4)	23 (36.5)	89 (21.4)	0.009 2.106 (1.198–3.702) 0.889 1.050 (0.534–2.064)
Skin ulcers, n (%)	88 (18.4)	12 (19.0)	76 (18.3)	0.248 1.942 (0.618–6.098)
Renal involvement, n (%)	18 (3.8)	4 (6.3)	14 (3.4)	0.084 2.469 (0.858–7.110)
Pulmonary arterial hypertension, n (%)	19 (4.0)	5 (7.9)	14 (3.4)	0.727 0.768 (0.173–3.405)
Treatment				
Glucocorticoids, n (%)	204 (42.7)	38 (60.3)	166 (40.0)	0.002 2.280 (1.327–3.919)
Methotrexate, n (%)	78 (16.3)	9 (14.3)	69 (16.6)	0.639 0.836 (0.394–1.772)
Sulfasalazine, n (%)	4 (0.8)	1 (1.6)	3 (0.7)	0.483 2.215 (0.227–21.631)
Hydroxychloroquine, n (%)	112 (23.4)	9 (14.3)	103 (24.8)	0.066 0.505 (0.241–1.058)
Leflunomide, n (%)	4 (0.8)	0 (0)	4 (1.0)	0.434
Mycophenolate, n (%)	79 (16.5)	21 (33.3)	58 (14.0)	<0.0001 3.078 (1.701–5.567)
Azathioprine, n (%)	22 (4.6)	1 (1.6)	21 (5.1)	0.220 0.303 (0.040–2.290)
TNF inhibitors, n (%)	23 (4.8)	1 (1.6)	22 (5.3)	0.199 0.288 (0.038–2.176)
Abatacept, n (%)	11 (2.3)	1 (1.6)	10 (2.4)	0.685 0.653 (0.082–5.192)
Rituximab, n (%)	26 (5.4)	12 (19.0)	14 (3.4)	<0.0001 6.739 (2.956–15.367)
IL-6 inhibitors, n (%)	21 (4.4)	1 (1.6)	20 (4.8)	0.243 0.319 (0.042–2.416)
JAK inhibitors, n (%)	14 (2.9)	3 (4.8)	11 (2.7)	0.354 1.836 (0.498–6.773)
Belimumab, n (%)	14 (2.9)	2 (3.2)	12 (2.9)	0.901 1.101 (0.241–5.039)
Vaccines administered				
BNT162b2, n (%)	448 (93.7)	60 (95.2)	388 (93.5)	0.595 1.392 (0.409–4.730)
mRNA-1273, n (%)	30 (6.3)	3 (4.8)	27 (6.5)	0.595 0.719 (0.211–2.442)
Vaccine-related adverse events				
No adverse events, n (%)	221 (46.2)	29 (46.0)	192 (46.3)	0.972 0.991 (0.582–1.686)
Pain at the injection site, n (%)	182 (38.1)	22 (34.9)	160 (38.6)	0.580 0.855 (0.491–1.489)
Fever, n (%)	42 (8.8)	3 (4.8)	39 (9.4)	0.226 0.482 (0.144–1.609)
Headache, n (%)	30 (6.3)	5 (7.9)	25 (6.0)	0.560 1.345 (0.495–3.652)
Fatigue, n (%)	36 (7.5)	8 (12.7)	28 (6.7)	0.095 2.010 (0.872–4.633)
ASD flares, n (%)	10 (2.1)	2 (3.2)	8 (1.9)	0.519 1.668 (0.346–8.039)
SARS-CoV-2 IgG NAb titre, BAU/mL	825 (451–1542)	0.6 (0.2–3.3)	446.5 (114.5–1360.1)	<0.0001

Data are presented as mean ± standard deviation, number (percentage), or median [25th–75th percentile], as appropriate. P values represent comparison between SARS-CoV-2 IgG NAb negative and positive subgroups and have been calculated using the Student's T test for continuous variables or the Fisher's exact test for binary variables.

Legend: ASD, autoimmune systemic diseases; COVID-19, coronavirus disease 2019; IL-6, interleukin-6; JAK, Janus kinase; NAb, neutralizing antibodies; TNF, tumor necrosis factor.

from the general population was used as control group. The same exclusion criteria were applied to the control group; moreover, subjects with a past diagnosis of any ASD and/or a past exposure for any reason to immunosuppressive treatments were excluded ($n = 6$).

Vaccine was administered to patients and controls by intramuscular injection in the deltoid muscle according to the manufacturer indications and Italian national guidelines; the BNT162b2 and mRNA-1273 vaccines were administered in 448 (93.7%) and 30 (6.3%) ASD patients, respectively, as well as in comparable percentages of control individuals.

2.1. Statistical analysis

Data are expressed as mean \pm standard deviation (SD), median (25th–75th percentile) or number (percentage) as appropriate. The Student's T test was used for comparing means of continuous variables between two groups; highly skewed variables were ln-transformed before the analysis. Fisher's exact test was used to compare categorical variables. The one-way analysis of variance (ANOVA) and the Kruskal–Wallis H test were used for comparing three or more groups, as appropriate. Univariate and multivariate logistic regression models were built to assess the predictivity of continuous or categorical variables for a dichotomic dependent variable, expressed as odds ratio (OR) and 95% confidence interval (95% CI). All tests were two tailed. Analyses were performed using the Statistic Package for Social Sciences (SPSS)[®] software, Version 23 (IBM).

3. Results

A total of 478 patients (84% females, mean age 59 ± 15 years) were consecutively recruited by the participating centers, including patients with RF and/or ACPA positive RA ($n = 101$), SLE ($n = 38$), SSc ($n = 265$), CV ($n = 61$), and a miscellanea of other systemic vasculitis ($n = 13$).

General features of the study population are reported in Table 1. The serum titer of anti-SARS-CoV-2 IgG NAb, evaluated at the end of the vaccination cycle, was significantly lower in the whole ASD patients series compared to control group [286 (53–1203) vs 825 (451–1542) BAU/mL, $p < 0.0001$]; comparable differences were also detected between individual ASD patients' subgroups and controls (Fig. 1A). In addition, the percentage of vaccine non-responders (defined as anti-SARS-CoV-2 IgG NAb below threshold) was significantly higher in the

whole ASD patients' series than controls [13.2% (63/478), vs 2.8% (14/502); $p < 0.0001$; Fig. 1B]. Consistently, all ASD patient subgroups showed significantly higher percentages of non-responders compared to controls (Fig. 1B); more evident for both CV and other systemic vasculitis when compared to other subgroups, namely RA, SLE, and SSc patients. (Fig. 1B).

The comparison between anti-SARS-CoV-2 IgG NAb negative ($n = 63$) and positive ASD ($n = 415$) revealed no differences in gender and age distribution; on the other hand, the number of non-responders was significantly higher in patients with concurrent ASD-related interstitial lung disease (Table 1). Moreover, anti-SARS-CoV-2 IgG NAb negative ASD patients were more frequently treated with glucocorticoids, mycophenolate mofetil, and/or rituximab. Comparable percentages of vaccine-related AEs were recorded among responder and non-responder ASD patients (Table 1).

The comparison between ASD subgroups (Table 2) demonstrated a number of differences at least in part correlated to demographic and clinical features specific for each disorder; namely, an increased prevalence of interstitial lung disease and skin ulcers in SSc patients, as well as of renal involvement in SLE and CV patients. Similarly, differences were also observed as regards the therapeutic regimens; namely, methotrexate and TNF inhibitors were more frequently employed in RA patients, steroids, hydroxychloroquine, and belimumab in SLE, mycophenolate mofetil in SSc, and rituximab in CV and other systemic vasculitis (Table 2). Of note, none of the patients discontinued immunomodifier medications during the vaccination cycle.

With regard to the safety of vaccine, transitory, mild side effects following the first or second dose of vaccine were recorded in about half of the ASD patients and controls, more frequently headache in SLE and CV and pain at injection site in systemic vasculitis patients (Table 2).

In addition, disease flares after completion of the vaccination cycle were reported by 10/478 (2.1%) ASD patients. Flares included the recurrence of purpura in 6 CV patients, and of skin ulcers in 1 SSc subject, the reactivation of arthritis in a woman with RA, and de novo onset paresthesia in 2 SLE patients.

The possible correlation between impaired vaccine immunogenicity and ongoing immune-modifier treatments was also investigated. A total of 38/63 SARS-CoV-2 IgG NAb negative patients (60.3%) were receiving glucocorticoids compared to 166/415 (40.0%) in SARS-CoV-2 IgG NAb positive ($p = 0.002$). Of note, only 4 (6.3%) and 10 (2.4%) patients were receiving a dose of glucocorticoids greater than 5 mg prednisone equivalent, respectively. Moreover, the percentage of patients treated

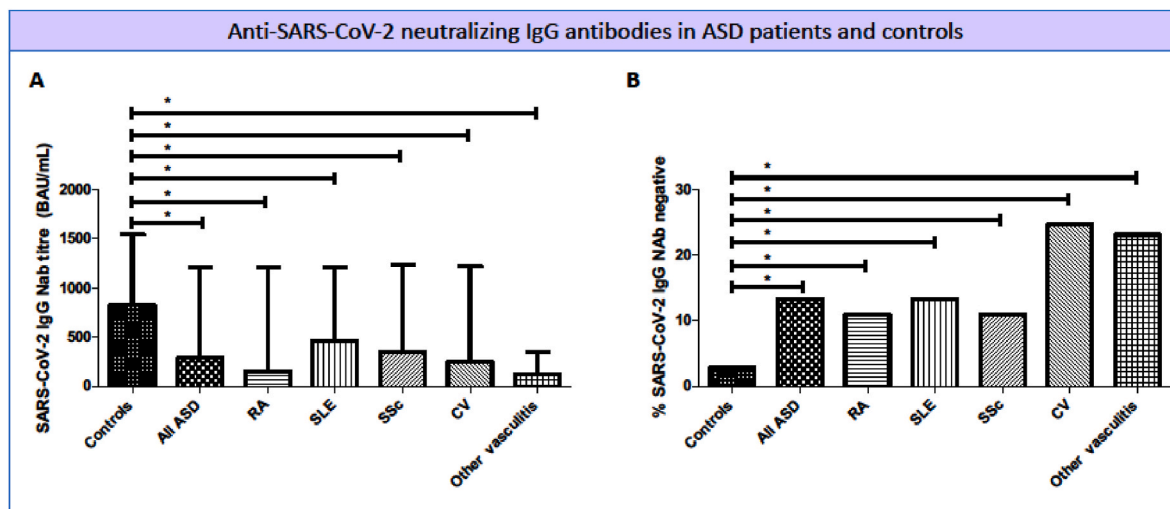


Fig. 1. Anti-SARS-CoV-2 IgG neutralizing antibodies (NAb) titer (A) and percentage of non-responders (B) in autoimmune systemic diseases (ASD) and controls. Anti-SARS-CoV-2 IgG NAb were measured 1–3 weeks after the second dose of vaccine in ASD patients and controls. A statistically significant difference (* indicates $p < 0.05$) was between each individual ASD and controls for both NAb titer (panel A) and percentage of non-responders (panel B). Legend: BAU, binding antibody units; CV, cryoglobulinemic vasculitis; RA, rheumatoid arthritis; SLE, systemic lupus erythematosus; SSc, systemic sclerosis.

Table 2
Analysis of the cohort after stratification for specific autoimmune systemic disease.

	RA (n = 101) (a)	SLE (n = 38) (b)	SSc (n = 265) (c)	CV (n = 61) (d)	Other vasculitis (n = 13) (e)	P value
Age, years	61 ± 12 _{b, d}	46 ± 17 _{a, c, d}	57 ± 15 _{b, d}	74 ± 11 _{a, b, c, e}	51 ± 18 _d	<0.0001
Female gender, n (%)	71 (70.3)	36 (94.7) _{a, e}	236(89.1) _{a, e}	51(83.6)	7(53.8)	<0.0001
Disease duration, years	8 ± 7 _b	14 ± 12 _{a, d}	10 ± 8	9 ± 6 _b	4 ± 5	0.008
Past COVID-19, n (%)	5 (5.0)	3 (7.9)	7 (2.6)	0 (0)	0 (0)	0.163
ASD-related complications						
Interstitial lung disease, n (%)	2 (2.0)	1 (2.6)	107 (40.4) _{a, b}	0 (0)	2 (15.4)	<0.0001
Skin ulcers, n (%)	0 (0)	4 (10.5)	78 (29.4) _d	5 (8.2)	1 (7.7)	<0.0001
Renal involvement, n (%)	0 (0)	6 (15.8) _c	4 (1.5)	8 (13.1) _c	0 (0)	<0.0001
Pulmonary art. hypert., n (%)	0 (0)	1 (2.6)	18 (6.8)	0 (0)	0 (0)	0.012
Gastrointestinal inv., n (%)	0 (0)	0 (0)	19 (7.2)	0 (0)	0 (0)	0.003
Treatment						
Glucocorticoids, n (%)	44(43.6)	31 (81.6) _a	105 (39.8)	16 (26.2) _a	8 (61.5)	<0.0001
Methotrexate, n (%)	57 (56.4) _{b, c}	3 (7.9)	15 (5.7)	0 (0)	3 (23.1)	<0.0001
Sulfasalazine, n (%)	4 (4.0)	0 (0)	0 (0)	0 (0)	0 (0)	0.005
Hydroxychloroquine, n (%)	10 (9.9)	25 (65.8) _{a, c, d, e}	75 (28.3) _{a, d}	1 (1.6)	1 (7.7)	<0.0001
Leflunomide, n (%)	2 (2.0)	1 (2.6)	1 (0.4)	0 (0)	0 (0)	0.359
Mycophenolate, n (%)	0 (0)	4 (10.5)	73 (27.5) _a	0 (0)	2 (15.4)	<0.0001
Azathioprine, n (%)	0 (0)	4 (10.5)	15 (5.7)	0 (0)	3 (23.1) _c	<0.0001
TNF inhibitors, n (%)	21 (20.8) _c	0 (0)	1 (0.4)	0 (0.0)	1 (7.7) _c	<0.0001
Abatacept, n (%)	6 (5.9) _c	0 (0)	5 (1.9)	0 (0)	0 (0)	0.067
Rituximab, n (%)	2 (2.0)	0 (0)	14 (5.3)	8 (13.1) _a	2 (15.4)	0.007
IL-6 inhibitors, n (%)	6 (5.9)	0 (0)	13 (4.9)	0 (0)	2 (15.4)	0.060
JAK inhibitors, n (%)	14 (13.9)	0 (0)	0 (0)	0 (0)	0 (0)	<0.0001
Belimumab, n (%)	0 (0)	14 (36.8)	0 (0)	0 (0)	0 (0)	<0.0001
Vaccines administered						
BNT162b2, n (%)	93 (92.1)	34 (89.5)	253 (95.5)	55 (90.2)	13 (100)	0.269
mRNA-1273, n (%)	8 (7.9)	4 (10.5)	12 (4.5)	6 (9.8)	0 (0)	0.268
Vaccine-related adverse events						
No adverse events, n (%)	56 (55.4) _d	21 (55.3) _d	125 (47.2) _d	15 (24.6)	4 (30.8)	0.002
Pain at the injection site, n (%)	33 (32.7)	15 (39.5)	96 (36.2)	28 (45.9)	10 (76.9) _{a, c}	0.021
Fever, n (%)	8 (7.9)	5 (13.2)	22 (8.3)	4 (6.6)	3 (23.1)	0.312
Headache, n (%)	9 (8.9)	6 (15.8) _c	8 (3.0)	7 (11.5) _c	0 (0)	0.004
Fatigue, n (%)	5 (5.0)	1 (2.6)	28 (10.6)	2 (3.3)	0 (0)	0.077
SARS-CoV-2 IgG NAb titre, BAU/ mL	153.05 [52.72–1140.23]	455.87 [129.16–1028.31]	349.48 [63.21–1224.2]	245.08 [7.33–1072.00]	118.83 [46.92–317.90]	0.184
Vaccine non-response, n (%)	11 (10.9)	5 (13.2)	29 (10.9)	15 (24.6) _{a, b, c}	3 (23.1) _{a, b, c}	0.047

Data are presented as mean ± standard deviation, number (percentage), or median [25th–75th percentile], as appropriate. P values have been calculated using one-way analysis of variance (ANOVA) for continuous variables or Kruskal–Wallis H test for binary variables. A letter has been assigned to each disease group as reported in first row; subscript letters have been used to highlight post-hoc contrasts between pairs; when a subscript letter is associated to an individual value it represents that the value differs from that observed in the group associated with the letter.

Legends: ASD, autoimmune systemic diseases; COVID-19, coronavirus disease 2019; CV, cryoglobulinemic vasculitis; IL-6, interleukin-6; JAK, Janus kinase; mRSS, modified Rodnan skin score; NAb, neutralizing antibodies; RA, rheumatoid arthritis; SLE, systemic lupus erythematosus; SSc, systemic sclerosis; TNF, tumor necrosis factor.

with mycophenolate mofetil ($p < 0.0001$) or rituximab ($p < 0.0001$) was significantly higher in non-responders compared to the responder subgroup.

Univariate logistic regression analysis was carried out to investigate possible predictors of vaccine non-response in ASD patients. Pre-existent ASD-related interstitial lung disease (OR: 2.106, 95% CI: 1.198–3.702, $p = 0.010$), treatment with glucocorticoids (OR: 2.27, 95% CI: 1.321–3.903, $p = 0.003$), mycophenolate mofetil (OR: 3.078, 95% CI: 1.701–5.567, $p < 0.0001$) or rituximab (OR: 6.739, 95% CI: 2.956–15.367, $p < 0.0001$) were associated with higher odds of vaccine non-response (Table 3). In a multivariate logistic regression model including only variables that yielded significant association with vaccine non-response in univariate analysis to avoid model overfitting, only treatment with rituximab maintained its predictivity in impairing the immunogenicity of anti-SARS-CoV-2 vaccine (OR: 4.031, 95% CI: 1.464–11.105, $p = 0.007$).

Finally, the ASD responders and non-responders' subgroups did not show significant correlations with the type of vaccine administered or with the prevalence of vaccine-related side effects (Table 1).

4. Discussion

The primary end-point of the present study was to investigate the immunogenicity of mRNA-based COVID-19 vaccines, evaluated by measurement of serum anti-SARS-CoV-2 IgG NAb, in a large series of ASD patients. According to our data, a significantly higher percentage of patients failed to mount a response within 1–3 weeks after completion of the vaccination cycle when compared to controls, particularly for CV or systemic vasculitis. Consistently, serum levels of anti-SARS-CoV-2 IgG NAb were significantly lower in the whole ASD series and in different patients' subgroups. Interestingly, failure to mount an antibody response to COVID-19 vaccine correlated with coexistent ASD-related interstitial lung disease and some ongoing treatments, namely glucocorticoids, mycophenolate mofetil, and/or rituximab. With regard to the overall safety of both mRNA vaccines, no severe AEs or disease flares were observed in our ASD series; only mild and transitory AEs were recorded in comparable percentages among vaccine responder and non-responder patients.

During the last few months, other observational studies on the same

Table 3

Univariate logistic regression analysis using vaccine non-response as dependent variable.

	OR	P value
Age	0.990 (0.973–1.008)	0.283
Female gender	1.426 (0.733–2.775)	0.296
ASD-related complications		
Interstitial lung disease	2.106 (1.198–3.702)	0.010
Skin ulcers	1.050 (0.534–2.064)	0.889
Renal involvement	1.942 (0.618–6.098)	0.256
Pulmonary arterial hypertension	2.469 (0.858–7.110)	0.094
Gastrointestinal involvement	0.768 (0.173–3.405)	0.728
Treatment		
Glucocorticoids	2.27 (1.321–3.903)	0.003
Methotrexate	0.836 (0.394–1.772)	0.640
Sulfasalazine	2.215 (0.227–21.631)	0.494
Hydroxychloroquine	0.505 (0.241–1.058)	0.070
Mycophenolate	3.078 (1.701–5.567)	<0.0001
Azathioprine	0.303 (0.040–2.290)	0.247
TNF inhibitors	0.288 (0.038–2.176)	0.228
Abatacept	0.653 (0.082–5.192)	0.687
Rituximab	6.739 (2.956–15.367)	<0.0001
IL-6 inhibitors	0.319 (0.042–2.416)	0.268
JAK inhibitors	1.836 (0.498–6.773)	0.361
Belimumab	1.101 (0.241–5.039)	0.901

Legend: ASD, autoimmune systemic diseases; IL-6, interleukin-6; JAK, Janus kinase; TNF, tumor necrosis factor.

topic have been published [13–16,19]. Only few of these studies investigated large series of patients with variegated diagnoses [14–16]; on the other hand, the remaining evidence is based on small case series focusing on single disorders such as SLE or RA [13,20–22].

The largest, controlled analysis was reported by Furer and coworkers that assessed the immunogenicity and safety of two-dose regimen BNT162b2 mRNA vaccine in adult with ASD [16]. The authors compared 686 patients with a smaller number of healthy controls (n = 121) and found a significantly reduced immunogenicity in the ASD cohort (86% vs 100%) [16].

Similar results (86% seroconversion) have been reported in a smaller, not controlled, cohort of ASD patients [15] while a higher immunogenic response (94% seroconversion) was obtained in RA and SLE [14]. However, in the latter study the most likely explanation was a young age of the enrolled subjects [14]. Besides the evidences of lower seroconversion rate in patients with ASD compared to the general population, the titer of anti-SARS-CoV-2 IgG NAb was also reported to be lower in ASD patients compared to controls [13]. This observation was confirmed by studies on specific diseases such as SLE [23] and RA [20]. Finally, a markedly impaired serological response to the COVID-19 vaccine was reported in a cohort including 73 patients with RA and 61 patients with SLE [22]; the overall immunogenicity rate was 77%, a lower figure compared to previous studies, probably due to the timing of the assessment which was performed only one week after the second dose of vaccine [22]. Fewer patients with RA (49, 67%) than those with SLE (54, 89%) had measurable antibodies against SARS-CoV-2 [22].

The results of the present study, including homogeneous subgroups of ASD are in keeping with the main findings of the few available reports [22]. Although the possibility of a direct comparison is limited by the different study protocols and cohort composition, our findings further support the role of some risk factors for impaired vaccine immunogenicity, mainly treatment with mycophenolate mofetil [14–16] or rituximab [13–16,24]; the latter representing the main cause of non-response to anti-SARS-CoV-2 vaccine (39% seropositivity) [14–16,24–26], as also suggested by the results of univariate analysis focusing on the factors associated with vaccine non-response. A recent review article

underlined the role of anti-CD20 therapies in blunting the humoral immune response to vaccines, which could be counterbalanced by preserved T-cell response, as well as the need to develop personalized peri-vaccination management strategy [5,24,25,27].

On the other hand, the prejudicial role of low-dose glucocorticoids [14, 16], methotrexate [14–16,21,22], and abatacept [16] on COVID-19 vaccine immunogenicity still remains controversial, while other molecules, such as belimumab, has been shown to don't interfere with antibody responses [22]. The role of increasing age in blunting immunogenicity of vaccines is suggested by only one monocentric cohort study [16]. Previous studies consistently report that, after the completion of the vaccination cycle, overall disease activity remained mostly stable [13,15,16], while only a few cases of flares were reported [15,16]. De novo onset of arthritis was seen in a patient with interstitial lung disease and positive RF [15] and a fulminant hemorrhagic cutaneous vasculitis with subsequent fatal sepsis caused the death of a patient with ANCA-associated vasculitis [16]. The highest percentage (6.5%) of post-vaccination flare was described by an Italian study focusing on RA cases [28]. Pre-vaccine discontinuation of the anti-rheumatic therapies in this population of patients may account for the discrepancy with other reports. In our series, disease reactivation was rare and overall, not alarming. Of note, the majority of patients experiencing flares had cryoglobulinemic vasculitis (60%), a condition not adequately evaluated by previous studies [15] or included, as anecdotal cases, in a miscellanea of ASD patients [13,16].

Although the blunted humoral response to COVID-19 vaccination is a sufficient reason of concern, the complete lack of NAb response represents the most crucial consequence of the immunological derangement characterizing ASD patients, along with few immune-modifier drugs such as rituximab. Despite some well-identified iatrogenic risk factors, the non-response observed in over 13% of ASD patients (25% in CV) remains largely unpredictable; therefore, it becomes crucial to early identify NAB negative individuals. Another compelling open question is the management of patients who show inadequate or absent humoral response. These seronegative patients, presumed to be immunized against SARS-CoV-2, may be at high risk of symptomatic infection. Although a percentage of seronegatives may develop a delayed seroconversion, around 23% as suggested by a recent observational study [29], this does not affect the significant difference between ASD patients and controls in the prevalence of non-responders. Consequently, in our opinion, the early measurement of serum NAB titer after completion of the vaccination cycle should be included in the routine assessment of all immunocompromised patients; this information may guide the subsequent management in terms of both timing and type of vaccine to be administered as a booster-dose, as well as the concurrent immune-modifier treatments. Indeed, in an early report, a booster-dose of vaccine in patients with ASD seems able to induce an amplified antibody response in the majority of subjects [30,31].

A proper management strategy for ASD patients during pre- and post-vaccination cycle should take in account the possible immunosuppressive effect of different treatments; the latter include rituximab and mycophenolate mofetil, mainly responsible for impaired immunogenicity, and some commonly used drugs such as corticosteroids. Therefore, the production of adequate therapeutic guidelines on the basis of recently available real-life experiences is eagerly awaited. The determination of NAB serum levels within the first 3 weeks from the second dose of vaccine would allow the identification of essentially three groups of patients: full responders, patients with suboptimal response, non-responders. In our opinion, patients showing a suboptimal response should be prioritized for a booster-dose of vaccine, while the administration of a different type of vaccine (for example from mRNA to vector-based vaccine) could be attempted in non-responder individuals. In any case, a careful evaluation of individual patients can model a personalized strategy, such as the tapering/discontinuation or delayed

therapeutic sessions of immunosuppressive treatments, mainly for rituximab, before a booster-dose of vaccine.

5. Conclusion

The ongoing pandemic still represents a complex challenge for the management of frail patients. A continuous adjustment of management strategies is still required, considering the evolution of pandemic, the clinic-pathogenetic peculiarities of different patients' settings, and their unpredictable response to vaccination. Hopefully, a widespread availability of serological testing may allow to recognize patients previously infected by SARS-CoV-2; rather it may be decisive to early identify patients with impaired/absent seroconversion needing an individualized therapeutic strategy.

Authors' statements

Clodoveo Ferri: conceptualisation, study design, supervision, writing, investigation, methodology, project administration, literature search; Francesco Ursini: formal analysis, figures, data interpretation, writing; Laura Gragnani: literature search, writing, methodology; Vincenzo Raimondo: data collection; Dilia Giuggioli: data collection; Rosario Foti: data collection; Maurizio Caminiti: data collection; Domenico Olivo: data collection; Giovanna Cuomo: data collection; Marcella Visentini: data collection; Fabio Cacciapaglia: data collection; Roberta Pellegrini: data collection; Erika Pigatto: data collection; Teresa Urraro: data collection; Caterina Naclerio: data collection; Antonio Tavoni: data collection; Lorenzo Puccetti: data collection; Giuseppe Varcasia: data collection; Ilaria Cavazzana: data collection; Massimo L'Andolina: data collection; Piero Ruscitti: data collection; Marta Vadacca: data collection; Pietro Gigliotti: data collection; Francesca La Gualana: data collection; Franco Cozzi: data collection; Amelia Spinella: data collection; Elisa Visalli: data collection; Ylenia Dal Bosco: data collection; Giorgio Amato: data collection; Francesco Masini: data collection; Giuseppa Pagano Mariano: data collection; Raffaele Brittell: data collection; Vincenzo Aiello: data collection; Rodolfo Caminiti: data collection; Daniela Scorpiniti: data collection; Giovanni Rechichi: data collection; Tommaso Ferrari: data collection; Monica Monti: data collection; Giusy Elia: data collection; Franco Franceschini: data collection; Riccardo Meliconi: data collection; Milvia Casato: data collection; Florenzo Iannone: data collection; Roberto Giacomelli: data collection; Poupak Fallahi: data collection; Stefano Angelo Santini: methodology; Anna Linda Zignego: data collection; Alessandro Antonelli: conceptualisation, writing. All authors critically revised the paper and approved the submitted version.

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Data availability

Data will be made available on request.

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