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With the arrival of SARS-CoV-2, it was asked whether our patients with immune-mediated inflammatory disorders, or who had an organ transplantation (IMDT patients) and/or use immunosuppressive medication (imed) are more susceptible to SARS-CoV-2 infection and/or a severe COVID-19 disease course. In the earliest reports on COVID-19, such patients were rarely described. Most reports were retrospectively collected, in various case series or cohorts without a control group.¹⁻³ The Infection and Immunomodulation Inventory Initiative cohort study was started 10 March 2020 to prospectively register self-reported periods of illness with COVID-19-like symptoms (CLS) (see questionnaire in online supplemental table 1) and compare these between IMDT patients with and without imed and controls as selected from the hospital database of the Leiden University Medical Center in March 2020. Patients were defined as being in outpatient care at the outpatient clinic for rheumatology, gastroenterology, pulmonology and/or nephrology and having an auto-inflammatory or autoimmune disease or having had a solid organ transplantation with or without imed (verified from the medical records after participant's informed consent). Controls were persons who had visited these outpatient clinics in the previous 3 years and were discharged but did not have an IMDT.

Of the 8670 individuals approached, 2110 with IMDT and 1067 controls agreed to participate (see baseline characteristics in online supplemental table 2 and differences between the non-repliers and repliers/participants in online supplemental table 3). The most prevalent diagnoses among the participants from the IMDT group were ulcerative colitis, Crohn's disease and seropositive rheumatoid arthritis (see online supplemental table 4). Between March and July 2020, 554 (33%) IMDT patients and 299 (35%) controls recorded an illness episode with at least one symptom, mostly mild with a median (IQR) duration of 4 (3–6) days in both IMDT patients and controls. Sixteen (6%) IMDT patients with imed, 8 (3%) IMDTs without imed and 5 (2%) controls were hospitalised with CLS ($p=0.8$). Logistic regression analysis showed that female gender (OR 1.45, 95% CI 1.15 to 1.82), lung disease (OR 1.50, 95% CI 1.20 to 1.88) and wearing a face mask (then not yet mandatory) (OR 1.42, 95% CI 1.13 to 1.77) were independently associated with a higher risk of experiencing CLS, whereas older age and use of imed were associated with a lower risk (see table 1).

Thus, we found a similar incidence of CLS in IMDT patients (with or without imed) and controls. However, IMDT patients on imeds with CLS had a slightly higher risk to be admitted to hospital, which may suggest worse symptom severity or an estimated greater risk of deterioration. We collected only self-reported symptoms mostly for logistical reasons. With 22% of participants reporting CLS, we may have overestimated the occurrence of COVID-19 and also included symptoms of influenza (season ended in March) and common colds, which, in turn, may have been over-reported during the anxious times of the 'first wave' of COVID-19. But since SARS-CoV-2 infection often results in mild influenza like symptoms only, we may in fact have come closer to the true infection rate than what has been reported in earlier observations based on hospitalisations and testing of worse cases.

Table 1 Univariable and multivariable analysis of variables associated with having CLS or not (OR with 95% CI)

	Data from n	Univariable	Multivariable*
Sex, female	2546	1.89 (1.58 to 2.25)	1.45 (1.15 to 1.82)
Age	2546	0.97 (0.96 to 0.97)	0.96 (0.96 to 0.97)
BMI	2391	0.99 (0.97 to 1.01)	1.00 (0.98 to 1.03)
Smoking (current)	2463	1.35 (1.02 to 1.78)	1.05 (0.74 to 1.50)
Daily alcohol use	2416	0.84 (0.71 to 1.00)	1.20 (0.96 to 1.50)
Solid organ transplantation	2546	0.74 (0.54 to 1.03)	0.79 (0.47 to 1.35)
IMDT without imed†	2546	1.00 (0.82 to 1.23)	0.94 (0.72 to 1.24)
IMDT with imed †	2546	0.79 (0.65 to 0.97)	0.68 (0.51 to 0.91)
Use of oral corticosteroids	2546	0.84 (0.66 to 1.06)	1.44 (0.95 to 2.20)
Self-reported diabetes mellitus	2381	0.69 (0.50 to 0.96)	0.89 (0.58 to 1.36)
Self-reported lung disease	2396	1.30 (1.09 to 1.54)	1.50 (1.20 to 1.88)
Self-reported heart disease	2399	0.85 (0.69 to 1.04)	1.09 (0.83 to 1.43)
Influenza vaccination‡	2415	0.71 (0.60 to 0.84)	0.96 (0.76 to 1.21)
Physical contact with family§	2220	1.47 (1.22 to 1.78)	1.22 (0.98 to 1.53)
Visiting other people (not family)	2205	1.26 (1.05 to 1.51)	0.96 (0.77 to 1.20)
Wearing a face mask	2196	1.46 (1.20 to 1.76)	1.42 (1.13 to 1.77)
Close contact (at work)	2180	1.65 (1.34 to 2.03)	1.27 (0.97 to 1.66)
Good adherence to lockdown rules	2245	1.17 (0.41 to 3.29)	2.46 (0.65 to 9.38)
Working outside the house	2435	1.39 (1.16 to 1.68)	0.92 (0.71 to 1.20)

*Number of observations: 1835.

†Control group=reference group.



‡In autumn 2019.

§Physical contact specified as 'holding/shaking hands, hugging etcetera'.

BMI, body mass index; CLS, Covid-19-like symptoms; IMDT, with immune mediated inflammatory disorders or transplant organ.

A relatively low response rate (37%) to our invitation to participate in this study means that there is a possibility of selection bias, the effect of which we cannot estimate.

In conclusion, between March and July 2020, IMDT patients, whether or not taking imeds, did not show an increased risk of reported CLS compared with controls. In our population, continuing immunosuppressant drugs as long as not ill, while following the Dutch COVID-19 rules, appears to be safe.

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