





ORIGINAL RESEARCH

Intestinal microbiota changes induced
by TNF-inhibitors in IBD-related
spondyloarthritis

Maria Chiara Ditto ¹, Simone Parisi ¹, Gianpiero Landolfi,² Richard Borrelli,¹ Cristina Realmuto,¹ Annacarla Finucci,¹ Gian Paolo Caviglia,³ Davide Giuseppe Ribaldone,⁴ Marco Astegiano,⁴ Anna Zanetti,² Greta Carrara,² Carlo Alberto Scirè ², Marco Antivalle ⁵, Piercarlo Sarzi-Puttini,⁵ Enrico Fusaro¹

To cite: Ditto MC, Parisi S, Landolfi G, *et al*. Intestinal microbiota changes induced by TNF-inhibitors in IBD-related spondyloarthritis. *RMD Open* 2021;**7**:e001755. doi:10.1136/rmdopen-2021-001755

► Additional supplemental material is published online only. To view, please visit the journal online (<http://dx.doi.org/10.1136/rmdopen-2021-001755>).

Manuscripts previously presented at EULAR 2021 and published as a conference abstract. Ditto MC, Parisi S, Landolfi G, *et al*. POS0230 Intestinal Microbiota changes TNF-Inhibitors induced in IBD-related spondyloarthritis. *Ann Rheum Dis* 2021;80:335.

Received 28 May 2021
Accepted 30 July 2021



© Author(s) (or their employer(s)) 2021. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Dr Simone Parisi;
simone.parsi@hotmail.it

ABSTRACT

Background The close relationship between joints and gut inflammation has long been known and several data suggest that dysbiosis could link spondyloarthritis (SpA) to inflammatory bowel diseases (IBD). The introduction of biological drugs, in particular tumour necrosis factor inhibitors (TNFi), revolutionised the management of both these diseases. While the impact of conventional drugs on gut microbiota is well known, poor data are available about TNFi.

Aim To investigate the impact of TNFi on gut microbiota.

Methods We evaluated 20 patients affected by enteropathic arthritis, naïve for biological drugs, treated with TNFi at baseline and after 6 months of therapy. All patients followed a Mediterranean diet. Patients performed self-sampling of a faecal sample at baseline and after 6 months of therapy. NGS-based ITS and 16S rRNA gene sequencing was performed, followed by the taxonomic bioinformatics analysis.

Results After 6 months of therapy, we detected a remarkable increase in Lachnospiraceae family ($\Delta +10.3$, $p=0.04$) and Coprococcus genus ($\Delta +2.8$, $p=0.003$). We also noted a decreasing trend in Proteobacteria ($\Delta -8.0$, $p=0.095$) and Gammaproteobacteria ($\Delta -9$, $p=0.093$) and an increasing trend in Clostridia ($\Delta +8.2$, $p=0.083$). We did not find differences between TNFi responders (SpA improvement or IBD remission achieved) and non-responders in terms of alpha and beta diversity.

Conclusions Our findings are consistent with the hypothesis that TNFi therapy tends to restore the intestinal eubiosis.

INTRODUCTION

Enteropathic arthritis is included in a cluster of inflammatory joint diseases,¹ according to the European Spondyloarthropathy Study Group classification criteria.² Patients with enteropathic spondyloarthritis (SpA) show clinical features of axial³ and peripheral⁴ SpA, associated with either Crohn's disease (CD) or ulcerative colitis (UC). The exact pathogenetic mechanisms linking gut inflammation with joint inflammation are still unclear,

Key messages**What is already known about this subject?**

- To date, the manipulation of the intestinal microbiota is considered pivotal for the cure or control of several pathologies, which are sustained and fuelled by dysbiosis.
- Scarce data are currently available about dysbiosis and inflammatory arthritis.
- The impact of microbiota on rheumatic diseases and the effect of tumour necrosis factor (TNF) treatment on microbiota is inadequately investigated and not still completely understood.

What does this study add?

- To our knowledge, this is the first study that evaluates the impact of anti-TNFs on the microbiota in patients with enteropathic spondyloarthritis.

How might this impact on clinical practice or further developments?

- Our study contributes to reinforce the hypothesis that the therapy with anti-TNF alpha can affect microbiota by restoring intestinal eubiosis.
- Based on this, the microbiota could play a key role in future targeted therapies.

although shared genes and environmental factors might be involved.⁵

The role of microbiota in host health is well established, mainly through the maintenance of the mucosal integrity and anti-inflammatory immune response⁶⁻⁸; therefore, it is not surprising that dysbiosis, defined as the loss of microbiome diversity and composition in an individual, can promote autoimmune diseases.

Pieces of evidence regarding dysbiosis and SpA have been published⁹⁻¹⁹ and data would suggest a link between SpA and inflammatory bowel diseases (IBD).^{9 20 21}

The introduction of biological drugs, in particular tumour necrosis factor inhibitors (TNFi), revolutionised the handling of enteropathic SpA.^{22–26} Infliximab and adalimumab are approved for the treatment of CD and UC, and the 2019 recommendations encourage the use of these two TNFi monoclonal antibodies in patients with seronegative SpA and coexisting IBD.²⁷

While the impact of TNFi on restoring gut permeability is well known,^{28–30} unsatisfactory data are available on microbiota and TNFi. Data acquired on animals would suggest that TNFi can restore gut permeability and microbiota.³¹ Few studies about the impact on TNFi on microbiota in humans affected by IBD or SpA would indicate a role of TNFi in restoring eubiosis^{32–34}; yet, data on enteropathic SpA are still not available.^{32 33}

Our study aimed to investigate the impact of TNFi on gut microbiota as well as the association of microbiome characteristics with inflammatory parameters in a population of enteropathic arthritis patients, who were treated with either adalimumab or infliximab.

MATERIALS AND METHODS

Patients recruitment and study design

Twenty patients were recruited at the Rheumatologic-Gastroenterologic Diseases clinic in Città della Salute e della Scienza di Torino, Italy, from June 2017 to December 2018.

Inclusion criteria comprised: patients of age greater than 18 years old and capable of providing informed consent before study participation; patients on a typical Mediterranean diet; CD or UC diagnosis in patients fulfilling criteria for axial or peripheral SpA (ASAS 2009^{3 35}); patients able to perform the sample's test after appropriate training as indicated in the BMR Genomics protocol; patients naïve to TNFi, and treated for arthritis, IBD or both, according to current guidelines for rheumatic disease³⁶ and IBD^{37 38}; stable concomitant treatment in the previous 3 months and for all the duration of the study.

Exclusion criteria were: history of major bowel resection; history of minor bowel resection or digestive stoma in the previous 5 years; any contra-indication to TNFi therapy; refusal to sign the informed consent; linguistic or cognitive difficulties that did not allow a full understanding of the consent form process; pregnancy or breastfeeding.

Clinical history, physical examination, instrumental examinations, biochemical examination including C reactive protein (CRP), erythrocyte sedimentation rate (ESR), HLA-B27 and faecal calprotectin at the baseline and after 6 months were performed.

All patients were screened for Hepatitis B Virus (HBV), Hepatitis C Virus (HCV) and *Mycobacterium tuberculosis* (HBsAg, HBcAb, HBsAb; HCV-Ab; quantiferon-TB gold assay, chest X-ray) before starting biologicals, as per normal clinical practice. Faecal samples were collected for every participant: the first within 24 hours before starting TNFi, while the second after 6 months of therapy. After

the discussion at the interdisciplinary rheumatic/gastroenteric board, treatment decisions were made based on clinical requirements.

All participants filled out a food daily diary to verify the Mediterranean diet, the absence of food intolerance with the consequent bias due to food restrictions, and the intake of drugs different from standard therapy.

The design of the prospective study did not affect the treatment or other clinical actions. Patients were naïve to TNFi treatment or other biologicals. TNFi included infliximab and adalimumab.

The study was approved by the ethics committee and subjects provided prior written informed consent.

Samples collection, processing and analysis

Faecal samples were collected in 2 mL faeces collection tubes with screw caps, filled with DNA-stabilisation buffer provided by BMR Genomics, and stored at -80°C until further use. All patients were trained for faecal sample collection, to reduce as much as possible the sample contamination; all samples complied with quality control.

The processing performed through metagenomic NGS (next-generation sequencing or sequencing in parallel) by BMR Genomics laboratories included the amplification of the V3 and V4 regions of the 16S-gene using universal primers according to Takahashi *et al*³⁹ followed by sequencing on the Illumina MiSeq platform (San Diego, California, USA) in the Paired-End 2×300 bp format and the standard bioinformatic taxonomic analysis of the 16S NGS.

Patient assessment

All patients were scheduled for biological therapy and received the treatment at least until week 24. Clinical disease indices were evaluated at baseline and the end of the study using: (i) Visual Analogue Scale to measure pain and disease activity in all patients; (ii) Bath Ankylosing Spondylitis Disease Activity Index (BASDAI) for axial involvement; (iii) Clinical Disease Activity Index (CDAI); (iv) Health Assessment Questionnaire-Disability Index (HAQ-DI) for peripheral involvement; (v) Harvey-Bradshaw Index for CD (HBI)⁴⁰ or partial Mayo (pMAYO) score for UC. The BASDAI clinical response was defined as a BASDAI <4 .⁴¹ The improvement in HAQ-DI was defined as a minimum clinically important difference of 0.22.⁴² Responders in the HBI for CD were defined by a decrease in the HBI score greater than or equal to 2 (or HBI ≤ 4 at 6 months), corticosteroid-free therapy. For patients with UC remission was defined as the reduction of the pMAYO⁴³ by at least two points, or final score ≤ 1 , in the absence of corticosteroids.

Statistical analysis

Continuous characteristics were presented as median and IQR or mean \pm SD. For proportions, absolute and relative frequencies were reported.

Mc-Nemar test was performed to analyse qualitative differences, while Wilcoxon test was applied to analyse mean differences (between paired samples).

Univariate regression models were implemented to test the microbiota differences between baseline and after 6 months (linear model for clinimetric variables and logistic model for IBD success or SpA improvement).

A *p* value <0.05 was considered to indicate statistical significance. The analysis was performed using R Statistical Software (Foundation for Statistical Computing, Vienna, Austria).⁴⁴

Alpha diversity⁴⁵ is a measure of the diversity of species within an ecosystem or sample; it is expressed by the number of species or Operational Taxonomic Units (OTU) observed in the sample, which is a measure of species richness and evenness in that sample. Alpha diversity was measured through the Simpson's Diversity Index (equation 1) and Shannon-Wiener Diversity Index (equation 2) from the OTU occurrence matrix.⁴⁶

The Simpson's Diversity Index (*D*) was calculated using the following equation:

$$D = 1 - \left[\frac{\sum n_i \cdot (n_i - 1)}{N \cdot (N - 1)} \right] \quad (1)$$

in which *n_i* is the total number of OTUs in a particular species, while *N* is the total number of OTUs in all species.

The Shannon-Wiener Diversity Index (*H*) was calculated using the following equation:

$$H = \sum [p_i \cdot \ln(i)] \quad (2)$$

in which *p_i* is *n_i* divided by *N*.

Beta diversity⁴⁵ is a measure of the diversity of species between two or more ecosystems or samples; it is expressed as the total number of unique species among the samples which are under comparison. Beta diversity therefore provides the variability of observed taxa among samples. The beta diversity Index was calculated using the PAST software by inputting the OTU occurrence matrix.⁴⁶ Principal Coordinates Analysis (PCoA) plots of Beta diversity were performed using the Bray-Curtis similarity index, with the transformation exponent equal to 2. To test the significance between groups, one-way analysis of similarity (ANOSIM) was carried out using Bray-Curtis distances, with the number of permutation set to 9999.

RESULTS

Twenty patients met inclusion criteria and were accepted to participate in the study. Informed consent was obtained from study participants.

The overview of patient characteristics is summarised in [table 1](#).

Surgery (ileocecal resection) had been performed in 9 (45%) patients before the last 5 years.

At the baseline 90% of patients received mesalazine, 60% of patients took systemic corticosteroids and 20% took an immunosuppressant (azathioprine). All patients followed a Mediterranean diet.

Table 1 Demographic and biometrics characteristics

Characteristics	Value
Median age (IQR)	53 (43.8–61.5)
Sex, n (%)—male	12 (60)
Smokers, n (%)	4 (20)
Family history of RD/IBD, n (%)	3 (15)
Years of illness (years)	14.5 (1–38)
Arthritis, n (%)—PsA	1 (5)
Arthritis, n (%)—AS	8 (40)
Arthritis, n (%)—SpA	11 (55)
Axial, n (%)	10 (50)
Peripheral, n (%)	12 (60)
IBD, n (%)—Crohn's	17 (85)
IBD, n (%)—UC	3 (15)
Skin involvement, n (%)	4 (20)
Eye involvement, n (%)	6 (30)
HLA-B27, n (%)	3 (15)
Endoscopic score, n (%)—1	2 (10)
Endoscopic score, n (%)—2	13 (65)
Endoscopic score, n (%)—3	5 (25)
Gut localisation, n (%)—E3	2 (10)
Gut localisation, n (%)—L1	8 (40)
Gut localisation, n (%)—L2	3 (15)
Gut localisation, n (%)—L3	6 (30)
Gut localisation n (%)—L4	1 (5)

AS, Ankylosing Spondylitis; IBD, inflammatory bowel diseases; PsA, Psoriatic Arthritis; RD, Rheumatic Disease; SpA, spondyloarthritis; UC, ulcerative colitis.

Clinical outcomes

After 6 months of therapy, no patient discontinued TNFi due to adverse events; 100% of patients achieved clinical IBD remission, while the success of the therapy (corticosteroids free) was only achieved in 65% of patients (13 out of 20). The improvement of rheumatic manifestations was reached in 70% of patients.

CRP decreased from a median value of 8.2 mg/L (4.8–20.8 mg/L) at T0 to a median value of 2.9 mg/L (1–4 mg/L) at T6 (*p*=0.001). Similarly, the ESR decreased from a median value of 21.5 mm/hour (10.8–34 mm/hour) at T0 to 11 mm/hour (7.8–21 mm/hour) at T6 (*p*=0.003). Calprotectin decreased from a median value of 207.5 µg/g (125.5–446.2 µg/g) at T0 to a median value of 81 µg/g (50–197.2 µg/g) at T6 (*p*=0.004).

As regards clinimetry, we noticed a BASDAI decrease from 5.2 (4.1–5.6) to 2.8 (2.5–4.3, *p*=0.013), CDAI decrease from 13 (10.5–16) to 7 (5.2–11, *p*=0.004) (see [table 2](#)).

Description of the microbiota composition and effect of TNFi on microbiota

The faecal microbiota at baseline was characterised by a large proportion of Firmicutes (43.6% (29.5–53.8)) followed by Bacteroidetes (35.5% (19.1–50.3)),

Table 2 Comparison of clinical variables between baseline and T6 (6 months)

Clinical variables	T0	T6	P value
Faecal calprotectin ($\mu\text{g/g}$)—median (IQR)	207.5 (125.5–446.2)	81 (50–197.2)	0.004
CRP (mg/L)—median (IQR)	8.2 (4.8–20.8)	2.9 (1–4)	0.001
ESR (mm/hour)—median (IQR)	21.5 (10.8–34)	11 (7.8–21)	0.003
VAS_pain—median (IQR)	50 (38.8–60)	35 (10–42.5)	0.001
VAS_disease—median (IQR)	50 (38.8–50)	37.5 (25–42.5)	0.006
HAQ—mediana (IQR)	0.6 (0.1–0.8)	0.2 (0.1–0.6)	0.004
BASDAI_score—median (IQR)	5.2 (4.1–5.6)	2.8 (2.5–4.3)	0.013
CDAI_activity—median (IQR)	13 (10.5–16)	7 (5.2–11)	0.004
IBD activity, n (%)—0	11 (55)	20 (100)	0.174
IBD activity, n (%)—1	6 (30)	0 (0)	
IBD activity, n (%)—2	2 (10)	0 (0)	
IBD activity, n (%)—3	1 (5)	0 (0)	

BASDAI, Bath Ankylosing Spondylitis Disease Activity Index; CDAI, Clinical Disease Activity Index; CRP, C reactive protein; ESR, erythrocyte sedimentation rate; HAQ, Health Assessment Questionnaire; IBD, inflammatory bowel diseases; VAS, Visual Analogue Scale.

Proteobacteria (12.6% (3.4–23.3)), Actinobacteria (0.8% (0.1–4.5)) (see [figure 1](#)).

We compared the global composition of the faecal microbiota between T0 and T6. After 6 months of therapy, we observed noteworthy differences among the family of Lachnospiraceae ($\Delta +10.3$, $p=0.04$) and the genus of Coprococcus ($\Delta +2.8$, $p=0.003$), as shown in [figure 2](#). We also observed an increase in the proportion of unclassified phyla ([figure 1](#), $\Delta +10.3$, $p=0.015$).

Despite the absence of statistically significant differences, we observed a trend in Proteobacteria ($\Delta -9.0$, $p=0.087$), Clostridia ($\Delta +8.2$, $p=0.083$) and Gammaproteobacteria ($\Delta -9$, $p=0.093$), Lachnospiraceae ($\Delta +9.4$, $p=0.044$) and Dorea ($\Delta +1.2$, $p=0.059$).

The univariate analysis for the comparison of relevant microbiota variables (outcome) between clinimetric variables (covariates) and the delta (T6-T0) exhibited a significant result only for CRP ($p=0.033$) ([table 3](#)).

The univariate analysis for the comparison between clinimetric variables (covariates) and the delta (T6-T0) of relevant microbiota variables (covariates) and the IBD success/SpA improvement outcome did not show any significant correlation ([table 4](#)).

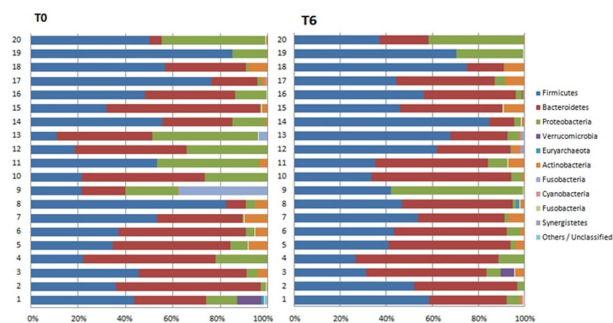


Figure 1 Distribution of phylum at the baseline (T0) and after 6 months (T6).

Finally, we compared alpha (intra-sample) and beta (inter-sample) diversity among patients at baseline and after 6 months of therapy (online supplemental files 1–6). We found a decrease of alpha diversity assessed by the Shannon-Weiner index in the class subgroup after therapy ([figure 3](#)). Comparing the responders group versus the non-responders one, we noted a meaningful reduction of alpha diversity determined by the Shannon-Weiner index only in the non-responder class group ([figure 4](#)) (online supplemental file 7); however, these results have not been confirmed when alpha diversity was assessed through the Simpson index ([figure 5](#)) (online supplemental file 8). When beta diversity was measured among population groups we did not identify differences either between T0 and T6 or between responders and non-responders.

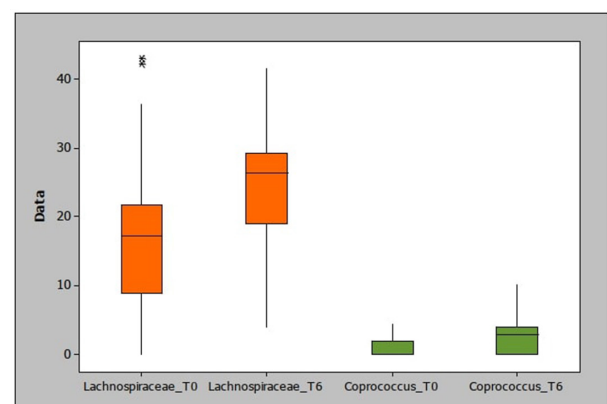


Figure 2 Comparison between Lachnospiraceae and Coprococcus at the baseline (T0) and after 6 months (T6) of therapy.

Table 3 Univariate analysis to evaluate the association between clinimetric variables (covariates) and the delta (T6-T0) of relevant microbiota variables (outcome)

Variable	Lachnospiraceae		Coprococcus	
	Coeff	P value	Coeff	P value
Faecal calprotectin	-0.003	0.408	-0.000	0.413
CRP	-0.157	0.540	-0.099	0.033
ESR	0.062	0.652	-0.022	0.409
VAS pain	-0.044	0.783	0.002	0.946
VAS disease	-0.028	0.868	0.013	0.679
HAQ	1.717	0.774	1.208	0.287
BASDAI	2.669	0.507	-0.512	0.560
CDAI	-1.366	0.062	0.017	0.867

BASDAI, Bath Ankylosing Spondylitis Disease Activity Index; CDAI, Clinical Disease Activity Index; CRP, C reactive protein; ESR, erythrocyte sedimentation rate; HAQ, Health Assessment Questionnaire; VAS, Visual Analogue Scale.

DISCUSSION

The human gut microbiota is composed of approximately 100 trillion bacteria, belonging to several hundreds of different species.⁴⁷ More than 90% of all colonic bacteria belongs to four phyla in the Bacteria domain: primarily in Firmicutes and Bacteroidetes phyla, followed by Actinobacteria and Proteobacteria phyla. Other less represented phyla include Verrucomicrobia and Fusobacteria.⁴⁸ The distribution of these phyla varies along the gastrointestinal tract, depending on the different microenvironments and the nutrient availability.⁴⁹

A healthy microbiota contributes to a healthy host. Indeed, microbiota not only plays a fundamental role in the metabolism of nutrients and drugs but it also contributes to the development and maintenance of the intestinal mucosal barrier integrity; additionally, it participates in the immune system regulation and defence against pathogens.^{6,7}

The relationship between dysbiosis and several diseases was robustly established; for example, in periodontal disease,⁵⁰ cancer,⁵¹ obesity,^{52,53} diabetes^{54,55} and chronic fatigue syndrome.⁵⁶

Pieces of evidence on dysbiosis both in SpA and in IBD were also found, which would suggest a link between these two inflammatory diseases. For instance, germ-free rats HLA-B27 + not only developed gut inflammation but also joints inflammation after recolonisation.⁵⁷

Table 4 Univariate analysis to evaluate the association between the delta (T6-T0) of relevant microbiota variables (covariates) and the IBD success/SpA improvement (outcome)

Variable	IBD success		SpA improvement	
	Exp (Coeff)	P value	Exp (Coeff)	P value
Delta Lachnospiraceae	1.053	0.201	0.925	0.100
Delta Coprococcus	1.412	0.234	1.251	0.394

IBD, inflammatory bowel diseases; SpA, spondyloarthritis.

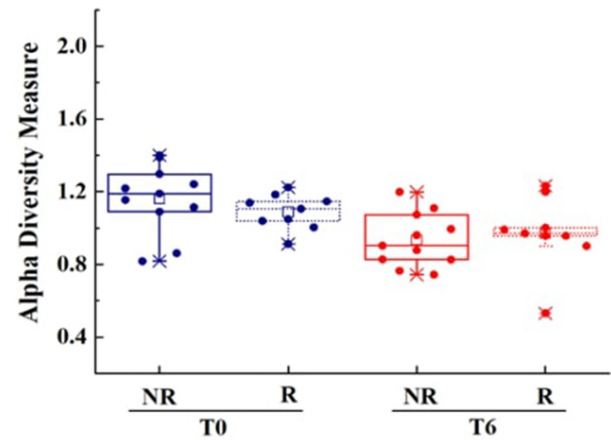


Figure 3 Determination of alpha diversity at the class level assessed by the Shannon-Weiner index. Plot of alpha diversity against patients at basal (T0, blue) and after 6 months of treatment (T6, red); a comparison between non-responder patients (NR, line) and responder patients (R, dots) is also shown both at T0 and T6. Alpha diversity was calculated using the Shannon-Wiener Diversity Index (Eq. 2); the index for patients at T0 is significantly different from those at T6 (two-tailed t-test with unequal variance, p value=0.0013).

While the effect of some drugs on microbiota, such as metphormine, Proton-pump inhibitor (PPI) and antibiotics are well known, poor data are available on TNFi.

Since the efficacy of these drugs on SpA and IBD has been widely demonstrated, a dramatic improvement in inflammatory parameters both in the acute phase index (CRP, ESR) and in faecal calprotectin is not surprising.

The aim of our study was to check microbiota modification after TNFi in a population affected by enteropathic arthritis. While some data are available on IBD or SpA alone, no literature data exist on this cluster of patients. TNFi can have an impact on disease improvements not only by modifying the intestinal permeability but also with a direct action on the microbiota. For instance, animal models of PG-induced arthritis showed an impaired abundance on three of the most representative phyla and TNFi treated animals did not; TNFi was able to restore the relative abundance of such species, including *Lactobacillus* and *Clostridium*.³⁰ Additionally, a decrease in the major phyla of Firmicutes, in particular *Faecalibacterium prausnitzii* and *Clostridium leptum* species in both SpA and IBD,²⁰ as well as an increase of Firmicutes to Gammaproteobacteria has

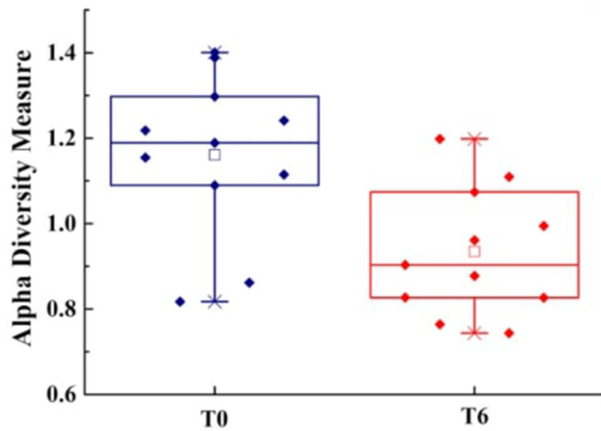


Figure 4 Determination of the alpha diversity between non-responders at the class level assessed by the Shannon-Weiner index. Plot of the alpha diversity against non-responder patients at basal (T0, blue) and after 6 months of treatment (T6, red). Alpha diversity was calculated using the Shannon-Wiener Diversity Index (Eq. 2); the index for patients at T0 is significantly different from those at T6 (two-tailed t-test with unequal variance, p value=0.0053).

been observed in CD.⁵⁸ Therefore, we first investigated the microbiota composition before and after treatment at the taxonomic levels (phylum, order, family, genus and species), by comparing the faecal samples before and after 6 months of TNFi (adalimumab or infliximab) therapy. Focusing on microbiota at the phylum level, we observed a reduction of Proteobacteria from 12.6 (3.4–23.3) to 4.6 (2–6.9), which is consistent with literature despite the absence of statistical significance ($p=0.09$). Similarly, we ascertained the same trend for the Gammaproteobacteria class (Proteobacteria

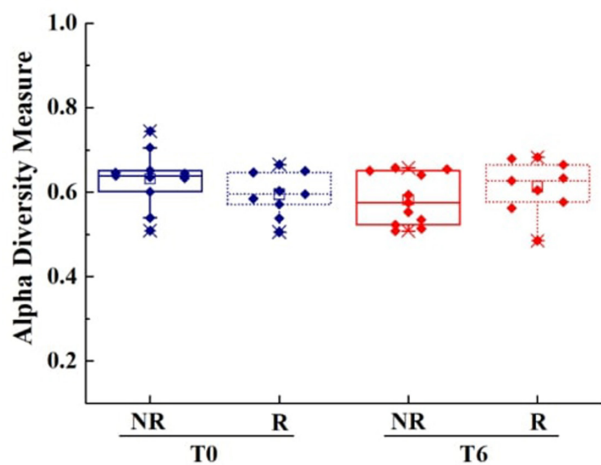


Figure 5 Determination of alpha diversity at the class level assessed by Simpson index. Plot of alpha diversity against patients at the basal (T0, blue) and after 6 months of treatment (T6, red); a comparison between non-responders (NR, line) and responders (R, dots) is also shown both at T0 and T6. Alpha diversity was calculated using the Simpson's Diversity Index (equation 1); the index for patients at T0 is not significantly different from those at T6 (two-tailed t-test with unequal variance, p value=0.3288).

phylum) with a reduction from 10.8 (0–22.8) to 1.8 (0–6, $p=0.09$); contrarily, we spot an increase in the Clostridia class (Firmicutes phylum), which raised from 34 (18.9–47.2) to 42.2 (34.3–52.1, $p=0.08$). These results are consistent with the increase of Firmicutes/Proteobacteria ratio in IBD patients and seem to indicate a shift versus eubiosis after TNFi.

In the overall Lachnospiraceae family, the gut microbiota showed a discordant trend between SpA and IBD, as it increased in the former and reduced in the latter. However, two genera of the Lachnospiraceae family shared a similar trend between SpA and IBD; indeed, both in SpA and IBD, the *Ruminococcus gnavus* genus increased and the *Roseburia* genus decreased.⁵⁹ From the analysis at the family level, we observed a significant increase of the members of the Lachnospiraceae family, which switched from 17.1 (9.2–21.1) to 26.5 (20–29.1, $p=0.04$); of particular interest was the *Coprococcus* genus, which switched from 0 (0–1.6) to 2.8 (0–4, $p=0.003$). Our data again would suggest an impact of TNFi on gut microbiota (figure 2).

Literatures reports on the microbiotic population trends of SpA and IBD are often conflicting. In some cases, the population trend decreases in SpA and grows in IBD; in other cases, the population trend increases in SpA and decreases in IBD; in still other cases, the population trend in SpA is very similar to that in IBD.

In our study, the correct interpretation of results can be further complicated as, in some cases, samples were taken from patients affected both by SpA and IBD. Although, a typical dysbiosis in our cohort should be more similar to the IBD dysbiosis—which might indicate a greater impact of IBD than SpA on the microbiota of these patients—detected trends in SpA and IBD did not demonstrate a full statistical significance; we also did not identify any known bacterial alteration in IBD.

We discovered a significative decrease of the alpha diversity determined using the Shannon-Weiner index in the class subgroup between T0 and T6, but these findings have not been confirmed by the Simpson index (the best index to use with a small sample size). As to the evaluation of beta diversity, we did not spot differences between patients at T0 and T6 and between responders and non-responders.

Our data are consistent with findings from the Aden and colleagues study, in which two groups of patients affected by IBD or rheumatic diseases were treated with TNFi. They discovered that the treatment restored the microbiota with an increase in the alpha and beta diversity only for IBD patients; no differences were identified between responders and not responders.³²

Another study concluded that the response to TNFi was affected by high disease activity and unstable microbiota composition, and patients with reduced biodiversity at baseline were more likely to fail the TNFi treatment. TNFi abolished the differences among patient groups, suggesting that the treatment can

restore the microbiota, which happens independently to the clinical response.³³ As a consequence of this, we performed further analysis to verify the impact of the inflammatory status on the change of the microbiota and check the relationship between the movement of the microbiota and the course of the disease. The univariate analysis showed that the CRP trend—a parameter of inflammation both for SpA and IBD—is consistent with the Coprococcus trend; this would confirm the hypothesis that the TNFi therapy tends to restore the intestinal eubiosis, by decreasing inflammation. We did not spot differences between responders and not responders, suggesting that TNFi can restore faecal microbiota and independently to clinical response.³²

The sample size of our population was small, which heavily affected the power of our statistical analysis.

Our study did not compare the intestinal microbiota of patients with that of healthy people and, therefore, it does not establish the nature of the dysbiosis, which might play a key role in the pathogenesis of SpA. Furthermore, we did not compare groups of patients treated by TNFi or other concomitant treatment with those drugs regularly used by patients as maintenance therapy (such as mesalazine, azathioprine, steroids and antibiotics)⁶⁰; thus, we cannot strongly affirm that modifications of microbiota are related to TNFi. To partially solve this bias, patients maintained the same therapy before and during the study period; moreover, all patients were treated with infliximab or adalimumab (etanercept was excluded). These drugs in vitro similarly mediate complement-dependent cytotoxicity and antibody-dependent cell-mediated cytotoxicity; they also affect in the same manner the proportion of cells undergoing apoptosis and the level of granulocyte degranulation.⁶¹

We analysed microbiota in faecal samples, which were used as a surrogate for the entire gut microflora. Although it is well known that the composition and function of the faecal microflora differ from that of the mucosal microflora⁴⁸ and it also includes microbiota and virota, the evaluation of stool samples is currently the easiest way to assess intestinal microbiota and it allows the evaluation of the microflora from the entire gut tract.

Our study is based on 16S rDNA sequencing, which allowed us to investigate only bacteria and exclude the virobiota, the mycobiota and eucaryota inhabiting intestinal tract and playing a consistent role in the pathophysiology of several diseases.⁶²

We included into the study only patients with a typical Mediterranean diet, but we did not perform any specific evaluation on the diet; for instance, we did not determine the percentages of carbohydrates, fats and proteins intake, which might have affected the results.⁶³

Although enteropathic arthritis is a well-known nosological entity, there are not clinimetric systems to evaluate the progress of this disease; therefore, the trend of both IBD and arthritis have been ascertained separately.

Furthermore, literature concerning dysbiosis in these two pathologies often differs, leading to a distortion of our results.

Despite these limits and independently of the achievement of clinical success, our results unequivocally reveal a modification of the microbiota towards eubiosis related to TNFi. Based on this, the microbiota could play a key role in targeted therapies in the future.

CONCLUSIONS

Our study is a proof of concept study, which has explored for the first time the modification of microbiota composition towards eubiosis in relationship with the TNFi treatment. Our study would indicate an association of TNFi with the decrease of Proteobacteria and with the increase of Lachnospiraceae and Coprococcus. The study data would support the hypothesis the TNFi therapy, by decreasing inflammation, tends to restore the intestinal eubiosis. Our study is characterised by some limitations, including the small number of patients under analysis and the absence of a control, which prevents from a definitive conclusion; yet, it can be considered a starting point for future examinations on this topic.

Author affiliations

¹Rheumatology Unit, Azienda Ospedaliero Universitaria Città della Salute e della Scienza di Torino, Turin, Italy

²Epidemiology Unit, Italian Society of Rheumatology, Milan, Italy

³Gastroenterology Unit, University of Turin Department of Medical Sciences, Turin, Italy

⁴Gastroenterology Unit, Azienda Ospedaliero Universitaria Città della Salute e della Scienza di Torino, Turin, Italy

⁵Rheumatology Unit, Ospedale Luigi Sacco, Milan, Italy

Twitter Simone Parisi @Dott.Simone Parisi

Contributors MCD and SP contributed in the same way to the realisation of the study and to the writing of the paper. GL contributed to the development of the specific part of statistics on biodiversity and to review of the paper. GPC contributed to the analysis of the samples, CR contributed to the data collection, AZ, GC, CAS contributed to the statistical analysis of the study. RB, AF, DGR, MAstegiano, MAntivalle, PS-P and EF contributed to the review of the study and the paper.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests None declared.

Patient consent for publication Not required.

Ethics approval Comitato Etico Interaziendale of AOU Città Della Salute e della Scienza di Torino (n of approval 0056924).

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request. All data relevant to the study are included in the article or uploaded as supplementary information. The study dataset is available upon reasonable request and is archived by the corresponding author. The protocol deposited with the Ethics Committee is also available.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iDs

Maria Chiara Ditto <http://orcid.org/0000-0001-5025-287X>
 Simone Parisi <http://orcid.org/0000-0003-4496-8315>
 Carlo Alberto Scirè <http://orcid.org/0000-0001-7451-0271>
 Marco Antivalle <http://orcid.org/0000-0003-2215-1777>

REFERENCES

- Zochling J, Smith EUR. Seronegative spondyloarthritis. *Best Pract Res Clin Rheumatol* 2010;24:747–56.
- Dougados M, van der Linden S, Juhlin R, et al. The European Spondylarthropathy Study Group preliminary criteria for the classification of spondylarthropathy. *Arthritis Rheum* 1991;34:1218–27.
- Rudwaleit M, van der Heijde D, Landewé R, et al. The development of assessment of spondyloarthritis International Society classification criteria for axial spondyloarthritis (Part II): validation and final selection. *Ann Rheum Dis* 2009;68:777–83.
- Rudwaleit M, van der Heijde D, Landewé R, et al. The assessment of spondyloarthritis International Society classification criteria for peripheral spondyloarthritis and for spondyloarthritis in general. *Ann Rheum Dis* 2011;70:25–31.
- Rudwaleit M, Baeten D, Ankylosing spondylitis and bowel disease. *Best Pract Res Clin Rheumatol* 2006;20:451–71.
- Hooper LV, Midtvedt T, Gordon JI. How host-microbial interactions shape the nutrient environment of the mammalian intestine. *Annu Rev Nutr* 2002;22:283–307.
- Jandhyala SM, Talukdar R, Subramanyam C, et al. Role of the normal gut microbiota. *World J Gastroenterol* 2015;21:8787–803.
- Vael C, Desager K. The importance of the development of the intestinal microbiota in infancy. *Curr Opin Pediatr* 2009;21:794–800.
- Breban M, Tap J, Leboime A, et al. Faecal microbiota study reveals specific dysbiosis in spondyloarthritis. *Ann Rheum Dis* 2017;76:1614–22.
- Beaud D, Tailliez P, Anba-Mondoloni J. Genetic characterization of the beta-glucuronidase enzyme from a human intestinal bacterium, *Ruminococcus gnavus*. *Microbiology* 2005;151:2323–30.
- Prindiville T, Cantrell M, Wilson KH. Ribosomal DNA sequence analysis of mucosa-associated bacteria in Crohn's disease. *Inflam Bowel Dis* 2004;10:824–33.
- Joossens M, Huys G, Cnockaert M, et al. Dysbiosis of the faecal microbiota in patients with Crohn's disease and their unaffected relatives. *Gut* 2011;60:631–7.
- Stebbing S, Munro K, Simon MA, et al. Comparison of the faecal microflora of patients with ankylosing spondylitis and controls using molecular methods of analysis. *Rheumatology* 2002;41:1395–401.
- Taurog JD, Maika SD, Satumtira N, et al. Inflammatory disease in HLA-B27 transgenic rats. *Immunol Rev* 1999;169:209–23.
- Ciccia F, Guggino G, Rizzo A, et al. Dysbiosis and zonulin upregulation alter gut epithelial and vascular barriers in patients with ankylosing spondylitis. *Ann Rheum Dis* 2017;76:1123–32.
- Morgan XC, Tickle TL, Sokol H, et al. Dysfunction of the intestinal microbiome in inflammatory bowel disease and treatment. *Genome Biol* 2012;13:R79.
- Scher JU, Ubeda C, Artacho A, et al. Decreased bacterial diversity characterizes the altered gut microbiota in patients with psoriatic arthritis, resembling dysbiosis in inflammatory bowel disease. *Arthritis Rheumatol* 2015;67:128–39.
- Sokol H, Seksik P. The intestinal microbiota in inflammatory bowel diseases: time to connect with the host. *Curr Opin Gastroenterol* 2010;26:327–31.
- Cao Y, Shen J, Ran ZH. Association between Faecalibacterium prausnitzii reduction and inflammatory bowel disease: a meta-analysis and systematic review of the literature. *Gastroenterol Res Pract* 2014;2014:872725.
- Gill T, Asquith M, Rosenbaum JT, et al. The intestinal microbiome in spondyloarthritis. *Curr Opin Rheumatol* 2015;27:319–25.
- Nagalingam NA, Kao JY, Young VB. Microbial ecology of the murine gut associated with the development of dextran sodium sulfate-induced colitis. *Inflam Bowel Dis* 2011;17:917–26.
- Van den Bosch F, Kruijthof E, De Vos M, et al. Crohn's disease associated with spondyloarthritis: effect of TNF-alpha blockade with infliximab on articular symptoms. *Lancet* 2000;356:1821–2.
- Ellman MH, Hanauer S, Sitrin M, et al. Crohn's disease arthritis treated with infliximab: an open trial in four patients. *J Clin Rheumatol* 2001;7:67–71.
- Herfarth H, Obermeier F, Andus T, et al. Improvement of arthritis and arthralgia after treatment with infliximab (remicade) in a German prospective, open-label, multicenter trial in refractory Crohn's disease. *Am J Gastroenterol* 2002;97:2688–90.
- Generini S, Giacomelli R, Fedi R, et al. Infliximab in spondyloarthritis associated with Crohn's disease: an open study on the efficacy of inducing and maintaining remission of musculoskeletal and gut manifestations. *Ann Rheum Dis* 2004;63:1664–9.
- Kaufman I, Caspi D, Yeshurun D, et al. The effect of infliximab on extraintestinal manifestations of Crohn's disease. *Rheumatol Int* 2005;25:406–10.
- Ward MM, Deodhar A, Gensler LS, et al. 2019 update of the American College of Rheumatology/Spondylitis association of America/Spondyloarthritis research and treatment network recommendations for the treatment of ankylosing spondylitis and Nonradiographic axial spondyloarthritis. *Arthritis Rheumatol* 2019;71:1599–613.
- Xu P, Elamin E, Elizalde M. Modulation of Intestinal Epithelial Permeability by Plasma from Patients with Crohn's Disease in a Three-dimensional Cell Culture Model. *Sci Rep* 2019;9:1–11.
- Al-Sadi R, Guo S, Ye D, et al. TNF- α modulation of intestinal epithelial tight junction barrier is regulated by ERK1/2 activation of Elk-1. *Am J Pathol* 2013;183:1871–84.
- Suenaert P, Bulteel V, Lemmens L, et al. Anti-tumor necrosis factor treatment restores the gut barrier in Crohn's disease. *Am J Gastroenterol* 2002;97:2000–4.
- Liu B, Yang L, Cui Z, et al. Anti-TNF- α therapy alters the gut microbiota in proteoglycan-induced ankylosing spondylitis in mice. *Microbiologyopen* 2019;8:e927.
- Aden K, Rehman A, Waschina S, et al. Metabolic Functions of Gut Microbes Associate With Efficacy of Tumor Necrosis Factor Antagonists in Patients With Inflammatory Bowel Diseases. *Gastroenterology* 2019;157:1279–92.
- Bazin T, Hooks KB, Barnette T, et al. Microbiota composition may predict anti-TNF alpha response in spondyloarthritis patients: an exploratory study. *Sci Rep* 2018;8:5446.
- Ribaldone DG, Caviglia GP, Abdulle A, et al. Adalimumab Therapy Improves Intestinal Dysbiosis in Crohn's Disease. *J Clin Med* 2019;8:1646.
- Rudwaleit M, Landewé R, van der Heijde D, et al. The development of assessment of spondyloarthritis International Society classification criteria for axial spondyloarthritis (Part I): classification of paper patients by expert opinion including uncertainty appraisal. *Ann Rheum Dis* 2009;68:770–6.
- van der Heijde D, Sieper J, Maksymowych WP, et al. 2010 update of the International ASAS recommendations for the use of anti-TNF agents in patients with axial spondyloarthritis. *Ann Rheum Dis* 2011;70:905–8.
- Reinglas J, Gonczi L, Kurt Z, et al. Positioning of old and new biologicals and small molecules in the treatment of inflammatory bowel diseases. *World J Gastroenterol* 2018;24:3567–82.
- Orlando A, Armuzzi A, Papi C, et al. The Italian Society of gastroenterology (sigE) and the Italian group for the study of inflammatory bowel disease (IG-IBD) clinical practice guidelines: the use of tumor necrosis factor-alpha antagonist therapy in inflammatory bowel disease. *Dig Liver Dis* 2011;43:1–20.
- Takahashi S, Tomita J, Nishioka K, et al. Development of a prokaryotic universal primer for simultaneous analysis of bacteria and archaea using next-generation sequencing. *PLoS One* 2014;9:e105592.
- Bennebroek Evertsz' F, Hoeks CCMQ, Nieuwkerk PT, et al. Development of the patient Harvey Bradshaw index and a comparison with a clinician-based Harvey Bradshaw index assessment of Crohn's disease activity. *J Clin Gastroenterol* 2013;47:850–6.
- Calin A, Garrett S, Whitelock H, et al. A new approach to defining functional ability in ankylosing spondylitis: the development of the Bath ankylosing spondylitis functional index. *J Rheumatol* 1994;21:2281–5.
- Behrens F, Koehm M, Schwaneck EC, et al. Use of a "critical difference" statistical criterion improves the predictive utility of the Health Assessment Questionnaire-Disability Index score in patients with rheumatoid arthritis. *BMC Rheumatol* 2019;3:51.
- Travis SPL, Schnell D, Krzeski P, et al. Reliability and initial validation of the ulcerative colitis endoscopic index of severity. *Gastroenterology* 2013;145:987–95.

- 44 Team R. A language and environment for statistical computing. *Computing* 2006;1.
- 45 Walters KE, Martiny JBH. Alpha-, beta-, and gamma-diversity of bacteria varies across habitats. *PLoS One* 2020;15:e0233872.
- 46 Hammer O, Harper DAT, Ryan PD. Past: paleontological statistics software package for education and data analysis.
- 47 Sender R, Fuchs S, Milo R. Are we really Vastly Outnumbered? revisiting the ratio of bacterial to host cells in humans. *Cell* 2016;164:337–40.
- 48 Eckburg PB, Bik EM, Bernstein CN, *et al.* Diversity of the human intestinal microbial flora. *Science* 2005;308:1635–8.
- 49 Becattini S, Taur Y, Pamer EG. Antibiotic-Induced changes in the intestinal microbiota and disease. *Trends Mol Med* 2016;22:458–78.
- 50 Nath SG, Raveendran R. Microbial dysbiosis in periodontitis. *J Indian Soc Periodontol* 2013;17:543–5.
- 51 Sheflin AM, Whitney AK, Weir TL. Cancer-Promoting effects of microbial dysbiosis. *Curr Oncol Rep* 2014;16:406.
- 52 Turnbaugh PJ, Ley RE, Mahowald MA, *et al.* An obesity-associated gut microbiome with increased capacity for energy harvest. *Nature* 2006;444:1027–31.
- 53 Turnbaugh PJ, Hamady M, Yatsunenkov T, *et al.* A core gut microbiome in obese and lean twins. *Nature* 2009;457:480–4.
- 54 Abdellatif AM, Sarvetnick NE. Current understanding of the role of gut dysbiosis in type 1 diabetes. *J Diabetes* 2019;11:632–44.
- 55 Sircana A, Framarin L, Leone N, *et al.* Altered gut microbiota in type 2 diabetes: just a coincidence? *Curr Diab Rep* 2018;18:98.
- 56 Lakhan SE, Kirchgessner A. Gut inflammation in chronic fatigue syndrome. *Nutr Metab* 2010;7:79.
- 57 Hill DA, Artis D. Intestinal bacteria and the regulation of immune cell homeostasis. *Annu Rev Immunol* 2010;28:623–67.
- 58 Willing BP, Dicksved J, Halfvarson J, *et al.* A pyrosequencing study in twins shows that gastrointestinal microbial profiles vary with inflammatory bowel disease phenotypes. *Gastroenterology* 2010;139:1844–54.
- 59 Mondot S, Barreau F, Al Nabhani Z, *et al.* Altered gut microbiota composition in immune-impaired Nod2(-/-) mice. *Gut* 2012;61:634–5.
- 60 Manichanh C, Borruel N, Casellas F, *et al.* The gut microbiota in IBD. *Nat Rev Gastroenterol Hepatol* 2012;9:599–608.
- 61 Nesbitt A, Fossati G, Bergin M, *et al.* Mechanism of action of certolizumab pegol (CDP870): in vitro comparison with other anti-tumor necrosis factor alpha agents. *Inflamm Bowel Dis* 2007;13:1323–32.
- 62 Clemente JC, Ursell LK, Parfrey LW, *et al.* The impact of the gut microbiota on human health: an integrative view. *Cell* 2012;148:1258–70.
- 63 Bibbò S, Ianiro G, Giorgio V, *et al.* The role of diet on gut microbiota composition. *Eur Rev Med Pharmacol Sci* 2016;20:4742–9.