Dysbiosis in inflammatory bowel disease: a role for bacteriophages?


To cite this version:

Dysbiosis in inflammatory bowel disease: a role for bacteriophages?

P Lepage, J Colombet, P Marteau, T Sime-Ngando, J Doré and M Leclerc

Gut 2008;57:424-425
doi:10.1136/gut.2007.134668

Updated information and services can be found at:
http://gut.bmj.com/cgi/content/full/57/3/424

These include:

References
This article cites 3 articles, 1 of which can be accessed free at:
http://gut.bmj.com/cgi/content/full/57/3/424#BIBL

Email alerting service
Receive free email alerts when new articles cite this article - sign up in the box at the top right corner of the article

Notes

To order reprints of this article go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to Gut go to:
http://journals.bmj.com/subscriptions/
Dysbiosis in inflammatory bowel disease: a role for bacteriophages?

Intestinal bacteria have been implicated in the initiation and amplification of inflammatory bowel disease (IBD). The dysbiosis theory, reviewed by Tamboli et al (Gut 2004;53:1), is that an imbalance between putative “harmful” versus “protective” bacterial species may promote chronic intestinal inflammation. Although several studies published so far support this hypothesis, the most vexing question posed by Tamboli et al remains “what is the origin of dysbiosis?”. Bacteriophages outnumber bacteria by a factor of 10 in many natural ecosystems, exert a strong influence on bacterial diversity and population structure, and are probably involved in dysbiosis by destabilising bacterial communities. They could be involved indirectly through gene transfer and genome reorganisation within the bacterial population or directly as immunomodulating agents or by steric competition for microbe-associated molecular patterns on bacterial surfaces. However, bacteriophages are a neglected component of the gut microbiota. The first viral metagenomic study demonstrated a wide diversity (1200 genotypes) of uncultured bacteriophage species. The present study aimed at measuring the total viral community associated with the gut mucosa and comparing viral abundance between healthy individuals and patients with Crohn’s disease (CD), and also between the ulcerated and non-ulcerated mucosa of these patients.

Fourteen healthy individuals and 19 CD patients were recruited and gave their informed consent. Biopsies were obtained during colonoscopy and, for CD patients, from non-ulcerated and ulcerated tissues. Biopsies were disrupted by ultrasonication, filtered through 0.22 μm membranes and fixed in glutaraldehyde. For epifluorescence microscopy, biopsy supernatants were filtered through 0.02 μm membranes. Virus-like particles (VLPs) were stained with SYBR Green and counted on triplicate subsamples. VLPs were detected in every sample, and no contaminating bacteria were observed (fig 1A). Strikingly high numbers of VLPs were observed (fig 1B), with an average of 1.2 × 10^9 VLPs/biopsy (4.4 × 10^7–1.7 × 10^10). Transmission electron microscopy demonstrated that viral particles corresponded to bacteriophages, with morphotypes consistent with Siphoviridae, Myoviridae and Podoviridae being the dominant families (fig 2). Each individual seemed to be colonised by one dominant phage family. This result was confirmed by pulse-field gel electrophoresis of viral genomic DNA (data not shown). At the mucosal level and with specific reference to the potential role of bacteriophages in dysbiosis, CD patients harboured significantly more VLPs than healthy individuals (2.9 × 10^9 vs 1.2 × 10^8 VLPs/biopsy; Wilcoxon test p = 0.024). Moreover, decreased amounts of VLPs were detected in CD ulcerated mucosa, with an average of 2.1 × 10^9 VLPs/biopsy compared with 4.1 × 10^9 VLPs/biopsy from non-ulcerated mucosa (fig 1). It can be hypothesised either that more viruses are produced or that they can survive longer in non-ulcerated areas. Whether bacterial composition and abundance differ when comparing non-ulcerated with ulcerated mucosa is still controversial although differing bacteriophage life cycles (lysis, lysogeny) could be responsible for some of these discrepancies.

This study shows for the first time a dense bacteriophage community specifically associated with the gut mucosa, reaching 10^10/mm^3 of tissue. Significantly more bacterio-

**Figure 1** Concentrations of virus-like particles (VLPs) assessed by epifluorescence microscopy on gut mucosal samples from healthy individuals and Crohn’s disease (CD) patients. (A) SYBR Green I-stained human mucosal viral concentrate from a healthy individual (concentrate diluted to 5 × 10^{-4}). (B) Average number of VLPs per biopsy in the different clinical groups. *p = 0.024 Wilcoxon test. NUM, non-ulcerated mucosa; UM, ulcerated mucosa.

**Figure 2** Transmission electron microscopy observation (80 kV, ×40 000 magnification) of bacteriophage morphotypes from mucosal samples. The dominant morphotypes are Myoviridae, Podoviridae and Siphoviridae. The grey bar represents 100 nm.
Disseminated tuberculosis in a Crohn’s disease patient on anti-TNFα therapy despite chemoprophylaxis

We read with interest the article by Rampton et al (Gut 2005;54:1560) concerning the prevention of tuberculosis (TB) in patients with Crohn’s disease (CD) needing treatment with anti-tumour necrosis factor α (TNFα) agents. Opportunistic infections and particularly TB are of major concern among patients who are candidates for treatment with such cytokine inhibitors. Although in rheumatoid arthritis large prospective studies reveal an increased risk for TB activation despite appropriate chemoprophylaxis,1,2 there is no similar study in CD patients. We present the case of a CD patient on therapy with infliximab who presented with symptoms of disseminated TB. To the best of our knowledge, this is the first report of disseminated TB in a patient with CD who had already received proper chemoprophylaxis prior to anti-TNFα treatment.

A 42-year-old Greek man was admitted to our department because of high fever up to 40 °C of 10 days duration, weight loss (6 kg in the last month), dry cough and headache. His medical history was significant for inflammatory ileocolonic CD with articular involvement of 13 years duration. The patient experienced clinical remission on therapy with infliximab (5 mg/kg) every 8 weeks and azathioprine (2 mg/kg) for the last 3 years. Due to a positive PPD (purified protein derivative) test, he received, with good compliance, isoniazid (INH) for 2 months prior to commencement of anti-TNFα therapy and for a total of 6.5 months. For the last 2 years he was under clinical and radiological surveillance (chest and abdominal CT) because of persistent lymphadenopathy (axillary, mediastinal and lower mesenteric) which was attributed to generalised Bartonella after a thorough work-up that included lymph node and bone marrow biopsies. He had no signs of infection on his previous infliximab infusion 1 month prior to this admission.

The physical examination revealed crackles of the medial and basal areas of the right lung. Laboratory tests were significant for leucocytosis (white blood cells (WBC) = 16,200), elevated erythrocyte sedimentation rate (ESR; 101 mm/h) and C-reactive protein (12 mg/dl; normal <0.8). CT scanning revealed diffuse micronodular lesions in both lungs with a milia pattern, and expansion of the pre-existent lymphadenopathy to the upper mesenteric. MRI of the head revealed a 4 mm lesion of the left occipitoparietal region without signs of abscess. Bronchoscopically obtained lung biopsies and sputum stains were not diagnostic and thus he underwent thoracoscopy. Empiric levofloxacin treatment (750 mg/day for 15 days) resulted in fever remission with a relapse 6 days after treatment discontinuation. Biopsies taken from the right lung confirmed the diagnosis of active tuberculosis (fig 1). Interestingly, cultures for TB in blood, bronchoalveolar lavage and sputum were negative. Therapy with rifampin, INH, pyrazinamide and ethambutol resulted in an impressive clinical and radiological improvement during the first month of treatment.

This is the first report of disseminated TB (chest, abdomen and brain) in a CD patient on anti-TNFα treatment despite chemoprophylaxis with INH. We think that it was a reactivation of latent TB rather than a new infection. Looking at the case retrospectively, it seems that the diagnosis of Bartonella (based merely on serological evidence) was actually missed TB. According to Rampton’s algorithm and current guidelines, 6–9 months of INH chemoprophylaxis has been proposed for patients with latent TB with an indication for anti-TNFα treatment. Our case indicates that in CD patients on anti-TNFα treatment appropriate INH prophylaxis does not eliminate the risk for TB activation, and a close follow-up for TB activation is required. Therapy with quinolones, a common empiric treatment in CD flares, may delay the diagnosis of TB because of their well-known anti-TB activity.3 Our case outlines the need of continuous reassessment of our practice in preventing latent TB reactivation in patients on anti-TNFα agents.

REFERENCES


Authors’ response

Bourikas et al describe a case of disseminated tuberculosis (TB) in a patient with Crohn’s disease given 6.5 months isoniazid and infliximab. Rampton’s paper4 stated that “although the incidence of infliximab-related TB may now be falling due to improved risk assessment, chemoprophylaxis, and/or reporting fatigue, complacency is clearly inappropriate” It also summarised and referred to the full text of the British Thoracic Society/British Society of Gastroenterology/British Society of Rheumatology guidelines5 published in the same year. Under section 4.6 of the full guidelines, we wrote “It should be noted...”