

## **Psoriasis is Candidate for Intestinal Microbiota Transplantation?**

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#### **Abstract**

More frequently we observe articles that deal with the role of intestinal microbiota in other conditions that are not strictly the impact of *Clostridium difficile*.

We see good results in intestinal inflammatory processes such as Nonspecific Chronic Ulcerative Colitis, and although to a lesser extent in Crohn's Disease and other problems.

But What's going on in other conditions such Psoriasis?

We presented a patient with this pathology and observed improvement, although not remission.

What happened with comorbidities presented by the patient in question?

Keywords: Psoriasis and Microbiota; Fecal Microbiota Transplant (FMT); Intestinal Microbiota Transplant (IMT)

#### Introduction

Psoriasis is a chronic disease, in which immunity intervenes. People may have cutaneous, joint and systemic manifestations that generate significant morbidity and increased risk of mortality.

#### **Presentation of the Case**

A 34-year-old male, who has been suffering from Psoriasis in plaques for 25 years, characterized by small rashes on the scalp and on the right side of the auricular pavilion.

## **Explanation for figure 1**

The lesions appeared later in both the auricular pavilions, as well as in the frontal region. Erythematous plaques and whitish scales extend to the nose and eyebrows, accompanied by itching, without Koebner's phenomenon. He has used a spray, lotion and creams, based on cortisone, with little improvement.

Edema in the hands when carrying something heavy. It happened 7 times and now no longer; It was 2 - 3 years ago. Anxiety, 24 points on Hamilton's scale (Hamilton M. The assessment of anxiety states by rating. Br J Med Psychol. 1959; 32: 50-55).

Abdominal bloating when eat spicy or exaggerates his social responses. Dry eyes. It moistens them when blinking and there is no need for drops. Insomnia.

Ultrasound before FMT with grade I Non-alcoholic hepatic steatosis. Glutamic pyruvic transaminase 57. Alkaline phosphatase 38. Total bilirubins 1.52; Direct 0.59 and indirect 1.23. Blood count with mean corpuscular hemoglobin in 26. Normal coagulation tests. Weight 180.78 pounds. Size 5'9". Pulse 90 X'; blood pressure 120 - 180; Temperature 97.7°F; Abdominal perimeter 39.76 inches.

#### It is diagnosed:

- 1. Plaque psoriasis (scalp and face).
- 2. Anxiety degree 24 Hamilton scale.
- 3. Non-alcoholic hepatic steatosis Grade 1.
- 4. Abdominal perimeter 39.76 inches. For investigating cause.
- 5. Spastic colon.
- 6. Overweight grade 1.
- 7. Mycosis in small fingers of both feet.

IMT is performed, finding only spastic colon. They are deposited: 250 ml of Microbiota in ascending colon, 250 ml in transverse colon and 200 ml in descending colon. No incidents photographs of scalp are taken, including small toe nail, right, where there is fungal infection (black color) in external third of it. (There is in both feet, only that in the left, it is smaller). It is prescribed Probiolog (*Lactobacillus acidophilus* and *Bifidobacterium lactis*). 1 x 3; one month.

Two weeks after the IMT, reported a 10% reduction in the manifestation of Psoriasis. The one located on the forehead. Pruritus has reduced to 4 on a scale of 1 to 10, in all injuries. He is more hungry. The dream is refreshing. There was a feeling of evacuation (6 times a day), after IMT, evacuating semiliquid, on 3 occasions; giving without medication.



Figure 1: Frontal eruption.

## **Explanation for figure 2**

Abdominal perimeter 40.55 inches (Increased). The current Hamilton survey is 9 points. Reduced 15 points. Probiolog  $1 \times 3$  is prescribed; one more month.

Ultrasound of control liver without steatosis. A month later, he reports improvement. Dermatological lesions only increase in color when angry. Now he gets angry less.

The current color of Psoriasis located in the right temporal region is smaller in size and in coloration. It tends to match the color of the skin. Eat twice a day Masseter contracture occurs, although sparingly.

We advise you to exercise and eat 3 times a day. We prescribe: Rebiot I, ingestible solution. One a day (*Lactobacillus paracasei*/Vitamin B-6), *Bifidobacterium lactis* (BB-12), Vitamin B-12); BB-12; FOS fiber, Inulin).

The nail lesions and both small toes of the feet remain the same.



Figure 2: Decrease in frontal rash.

## **Discussion**

Psoriasis is a chronic disease, in which immunity intervenes. People may have cutaneous, joint and systemic manifestations that generate significant morbidity and increased risk of mortality. In addition, it affects the quality of life and is a risk factor for cardiovascular diseases, metabolic syndrome, peripheral vascular disease, inflammatory bowel disease, malignant tumors and nephropathies [1-3].

The skin is usually the most affected organ, but the joints, nails, eyes and the cardiovascular system can be compromised [4-6]. All of the above has stimulated us to look for treatment schemes that help solve this significant problem. That is why we have carried out an Intestinal Microbiota Transplant in the case in question, in order to offer other therapeutic alternatives, since there are few publications that treat Psoriasis with IMT, since they usually refer to the pathology. In children, nail involvement or complications such as arthritis [7-12] or something more complicated, such as cardiovascular [13,14] ocular [15-17] or the implementation of brilliant standards or meta-analysis studies [18-20]. Also, we see interesting reports, where Psoriasis is linked to digestive disorders, such as intestinal inflammatory processes, affecting the possibility that there is a link between these two diseases Psoriasis and Intestinal Inflammatory Disease [21-23].

Di Cesare., et al. they approach the disease from the immunological point of view, providing diverse tools for development in this sense [24].

Also, processes such as tobacco or some medications have been referenced as causes of Psoriasis and its complications [25]. Although we began to see other considerations, such as alterations of the microbiota, intrinsically related to Psoriasis, such as the study by Scher, et al. [26] those who observe coincidence with psoriatic arthritis, dysbiosis and psoriatic manifestations. They conclude that patients with PsA (psoriatic arthritis), as well as patients with cutaneous Psoriasis had relatively lower abundance of intestinal bacteria multiples.

Although some genera decreased concomitantly in both conditions, the PsA samples had lower abundance of taxa that reportedly are beneficial. This intestinal microbiota profile in PsA was similar to that described previously in patients with Inflammatory Bowel Disease and was associated with changes in specific inflammatory proteins exclusive of this group and different from those of patients with cutaneous Psoriasis and healthy controls. Therefore, the role of the gut microbiome in the pathogenesis of Psoriasis-PsA and the associated immune response deserves further study.

Numerous studies address the coincidence of Inflammatory Bowel Disease and dermatological processes [21,27-29].

There are few observations that indicate the importance of specific bacteria in the psoriatic processes, which highlights the great interaction that exists between these two major conditions [30-32].

With all the above, we cannot underestimate the management with Intestinal Microbiota Transplantation, and thus, have a new tool that contributes to the integral management of the psoriatic processes [34-37].

It has been found that serum concentrations of interleukin-23 (IL-23) are elevated and polymorphisms in the IL-23 receptor are associated with ankylosing spondylitis, which translates the importance of the inflammatory process [38].

#### **Conclusions**

We consider that the intrinsic relationship between Intestinal Microbiota and dermatological alterations exists, as is the case of Psoriasis, and intervenes in the development process of this skin disease.

The dysbiosis, generated by imbalance in the concentration of bacteria, with poor development of the microbiota, contributes much to some dermatological alterations. Often being accompanying phenomena.

For all of the above, the Intestinal Microbiota Transplant should not be underestimated in these patients. Even when, as in our case, it improved a bit, there was a clear improvement in anxiety, digestive, hepatic and psychological alterations, which undoubtedly are part of the dysbiosis, and which generates a good number of added medical alterations.

## Conflicts of Interest

The authors declare that they do NOT have affiliation or participation in organizations with financial interests.

## **Ethical Approval**

The present report does not contain any study with human or animal subjects made by the authors.

## **Informed Consent**

The authors obtained written informed consent from the patient in order to develop the present case.

#### **Bibliography**

- 1. Greb JE., et al. "Psoriasis". Nature Reviews Disease Primers 2 (2016): 16082.
- 2. Carrascosa JM., et al. "Obesity and psoriasis: inflammatory nature of obesity, relationship between psoriasis and obesity, and therapeutic implications". Actas Dermo-Sifiliograficas 105.1 (2014): 31-44.
- 3. Michalek IM., et al. "A systematic review of worldwide epidemiology of psoriasis". *Journal of the European Academy of Dermatology and Venereology* 31.2 (2017): 205-212.

- 4. Rachakonda TD., et al. "Psoriasis prevalence among adults in the United States". *Journal of the American Academy of Dermatology* 70.3 (2014): 512-516.
- 5. Takeshita J., et al. "Psoriasis and comorbid diseases: Epidemiology". *Journal of the American Academy of Dermatology* 76.3 (2017): 377-390.
- 6. Ritchlin CT., et al. "Psoriatic Arthritis". The New England Journal of Medicine 376.10 (2017): 957-970.
- 7. Tollefson MM., et al. "Incidence of psoriasis in children: a population-based study". *Journal of the American Academy of Dermatology* 62.6 (2010): 979-987.
- 8. Crawford GM. "Psoriasis of the nails". Archives of Dermatology and Syphilology 38.4 (1938): 583-594.
- 9. Langenbruch A., *et al.* "Nail involvement as a predictor of concomitant psoriatic arthritis in patients with psoriasis". *The British Journal of Dermatology* 171.5 (2014): 1123-1128.
- 10. Armstrong AW., et al. "Psoriasis comorbidities: results from the National Psoriasis Foundation surveys 2003 to 2011". *Dermatology* (Basel, Switzerland) 225.2 (2012): 121-126.
- 11. Klaassen KM., et al. "Nail psoriasis: a questionnaire-based survey". The British Journal of Dermatology 169.2 (2013): 314-319.
- 12. Tan ES., et al. "Nail psoriasis: a review". American Journal of Clinical Dermatology 13.6 (2012): 375-388.
- 13. Ahlehoff O., et al. "Psoriasis is associated with clinically significant cardiovascular risk: a Danish nationwide cohort study". Journal of Internal Medicine 270.2 (2011): 147-157.
- 14. Gelfand JM., et al. "Risk of myocardial infarction in patients with psoriasis". *Journal of the American Medical Association* 296.14 (2006): 1735-1741.
- 15. Rehal B., et al. "Ocular psoriasis". Journal of the American Academy of Dermatology 65.6 (2011): 1202-1212.
- 16. Chandran NS., *et al.* "Psoriasis and the eye: Prevalence of eye disease in Singaporean Asian patients with psoriasis". *The Journal of Dermatology* 34.12 (2007): 805-810.
- 17. Tsippora Shainhouse. "Ocular Manifestations of Psoriasis". EC Ophthalmology 5.5 (2017): 172-176.
- 18. Gottlieb A., et al. "Guidelines of care for the management of psoriasis and psoriatic arthritis: Section 2. Psoriatic arthritis: overview and guidelines of care for treatment with an emphasis on the biologics". *Journal of the American Academy of Dermatology* 58.5 (2008): 851-864.
- 19. Armstrong EJ., *et al.* "Psoriasis and Major Adverse Cardiovascular Events: A Systematic Review and Meta-Analysis of Observational Studies". *Journal of the American Heart Association* 2.2 (2013): e000062.
- 20. Gisondi P., *et al.* "Prevalence of metabolic syndrome in patients with psoriasis: a hospital-based case-control study". *The British Journal of Dermatology* 157.1 (2007): 68-73.
- 21. Li WQ., et al. "Psoriasis, psoriatic arthritis and increased risk of incident Crohn's disease in US women". *Annals of the Rheumatic Diseases* 72.7 (2013): 1200-1205.

- 22. Cohen AD., et al. "Psoriasis associated with Ulcerative colitis and Crohn's disease". *Journal of the European Academy of Dermatology and Venereology* 23.5 (2009): 561-565.
- 23. Toussirot É and Aubin F. "Paradoxical reactions under TNF-α blocking agents and other biological agents given for chronic immune-mediated diseases: an analytical and comprehensive overview". *RMD Open* 2.2 (2016): e000239.
- 24. Di Cesare A., *et al.* "The IL-23/Th17 Axis in the Immunopathogenesis of Psoriasis". *Journal of Investigative Dermatology* 129.6 (2009): 1339-1350.
- 25. Armstrong AW., et al. "Psoriasis and smoking: a systematic review and meta-analysis". The British Journal of Dermatology 170.2 (2014): 304-314.
- 26. Scher JU., *et al.* "Decreased bacterial diversity characterizes the altered gut microbiota In patients with psoriatic arthritis, resembling dysbiosis in inflammatory bowel disease". *Arthritis and Rheumatology* 67.1 (2015): 128-139.
- 27. Manichanh C., *et al.* "Reduced diversity of faecal microbiota in Crohn's disease revealed by a metagenomic approach". *Gut* 55.2 (2006): 205-211.
- 28. Png CW., et al. "Mucolytic bacteria with increased prevalence in IBD mucosa augment in vitro utilization of mucin by other bacteria". American Journal of Gastroenterology 105.11 (2010): 2420-2428.
- 29. Vigsnaes LK., *et al.* "Gram-negative bacteria account for main differences between faecal microbiota from patients with ulcerative colitis and healthy controls". *Beneficial Microbes* 3.4 (2012): 287-297.
- 30. Rath HC., *et al.* "Normal luminal bacteria, especially Bacteroides species, mediate chronic colitis, gastritis, and arthritis in HLA-B27/human beta2 microglobulin transgenic rats". *Journal of Clinical Investigation* 98.4 (1996): 945-953.
- 31. Scher JU., et al. "Expansion of intestinal Prevotella copri correlates with enhanced susceptibility to arthritis". Elife 2 (2013): e01202.
- 32. Stebbings S., *et al.* "Comparison of the faecal microflora of patients with ankylosing spondylitis and controls using molecular methods of analysis". *Rheumatology (Oxford)* 41.12 (2002): 1395-1401.
- 33. Arumugam M., et al. "Enterotypes of the human gut microbiome". Nature 473.7346 (2011): 174-180.
- 34. Scarpa R., *et al.* "Microscopic inflammatory changes in colon of patients with both active psoriasis and psoriatic arthritis without bowel symptoms". *Journal of Rheumatology* 27.5 (2000): 1241-1246.
- 35. Ciccia F., et al. "Interleukin-22 and interleukin-22-producing NKp44+ natural killer cells in subclinical gut inflammation in ankylosing spondylitis". Arthritis and Rheumatology 64.6 (2012): 1869-1878.
- 36. Nielsen OH., et al. "Rectal dialysate and fecal concentrations of neutrophil gelatinase-associated lipocalin, interleukin-8, and tumor necrosis factor-alpha in ulcerative colitis". American Journal of Gastroenterology 94.10 (1999): 2923-2928.
- 37. Willing BP., *et al.* "A pyrosequencing study in twins shows that gastrointestinal microbial profiles vary with inflammatory bowel disease phenotypes". *Gastroenterology* 139.6 (2010): 1844-1854.
- 38. Sherlock JP, *et al.* "IL-23 induces spondyloarthropathy by acting on ROR-gammat+CD3+CD4-CD8- entheseal resident T cells". *Nature Medicine* 18.7 (2012): 1069-1076.

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