



Frequency of intestinal microeukaryotes in patients undergoing screening colonoscopy for colorectal cancer

Carolina Hernández-Castro^{1,2}, Sonia del Pilar Agudelo¹, Miguel Toro¹, Jorge Botero¹, Alejandro Múnera³, Juan Camilo Correa⁴, Pamela C. Köster², David Carmena².

1. Parasitology Group, Faculty of Medicine, Academic Corporation for the Study of Tropical Pathologies, University of Antioquia. Medellín, Colombia.

2. Parasitology Reference and Research Laboratory. Spanish National Centre for Microbiology. Health Institute Carlos III. Majadahonda, Spain.

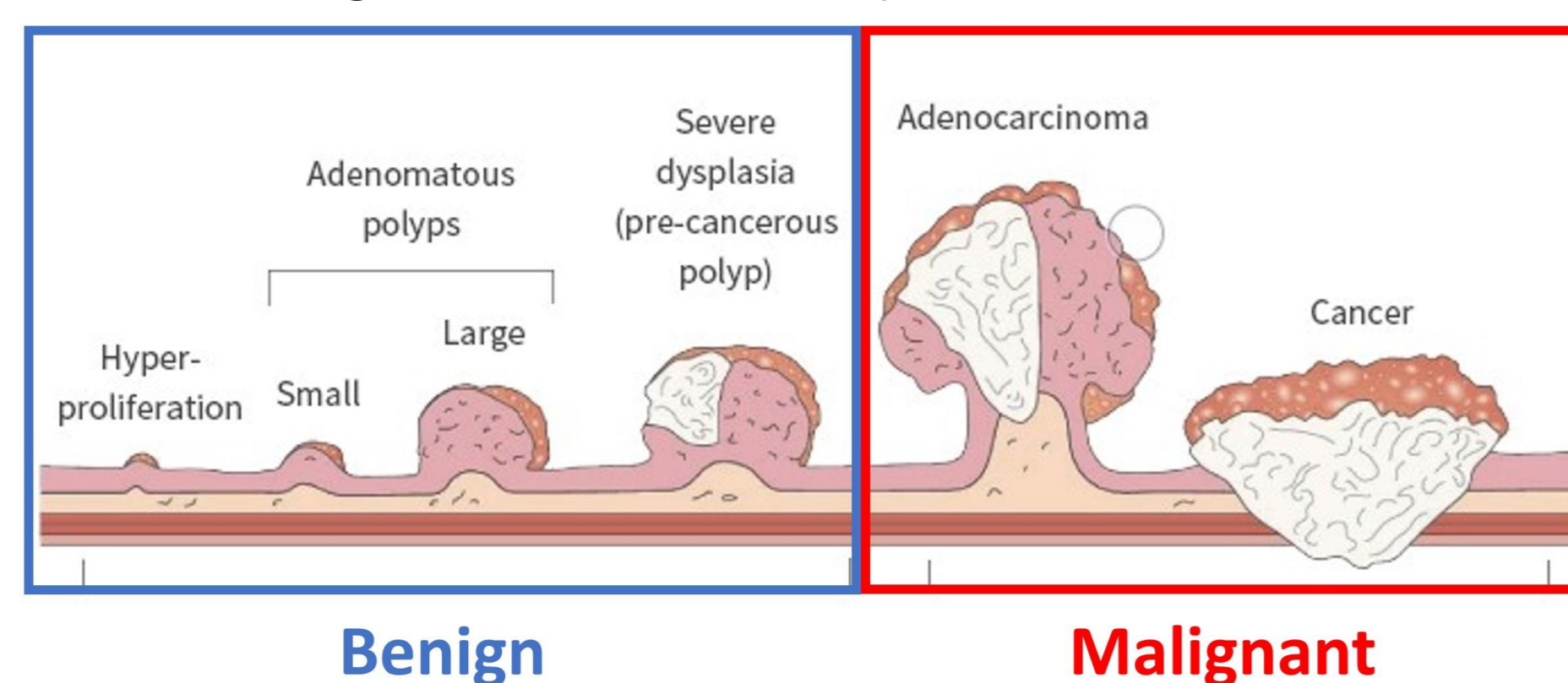
3. Institución Prestadora de Servicios de Salud "IPS Universitaria", University of Antioquia, Medellín, Colombia.

4. Clínica Medellín, Medellín, Colombia.

INTRODUCTION

The International Agency for Research on Cancer (IARC) has estimated that 16% of cancer worldwide is caused by infectious factors, including parasites¹. Colorectal cancer (CRC) is one of the most common neoplasms in humans and intestinal microeukaryotes (ME) like *Cryptosporidium* spp. or *Blastocystis* sp. has been proposed as potential cancer risk factors².

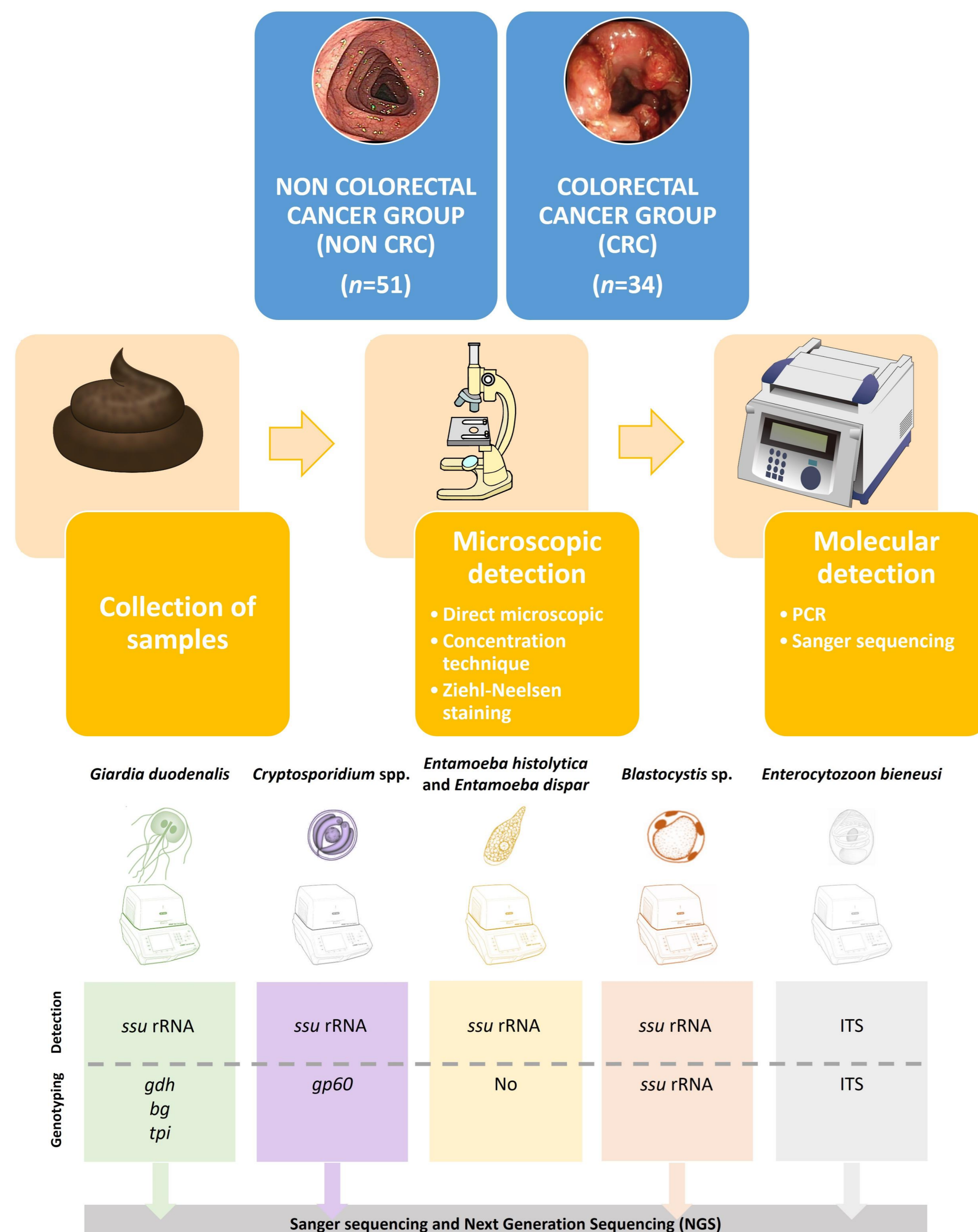
Blastocystis sp. is one of the most common ME in humans with a prevalence of up to 50% in developing countries like Colombia. Recent studies have highlighted the substantial role of *Blastocystis* and its genetic diversity in CRC³.



OBJECTIVE

This ongoing study provides preliminary data on the potential associations among intestinal microeukaryotes including *Blastocystis* sp. and an increased risk of developing CRC in Colombian patients.

MATERIALS AND METHODS



RESULTS

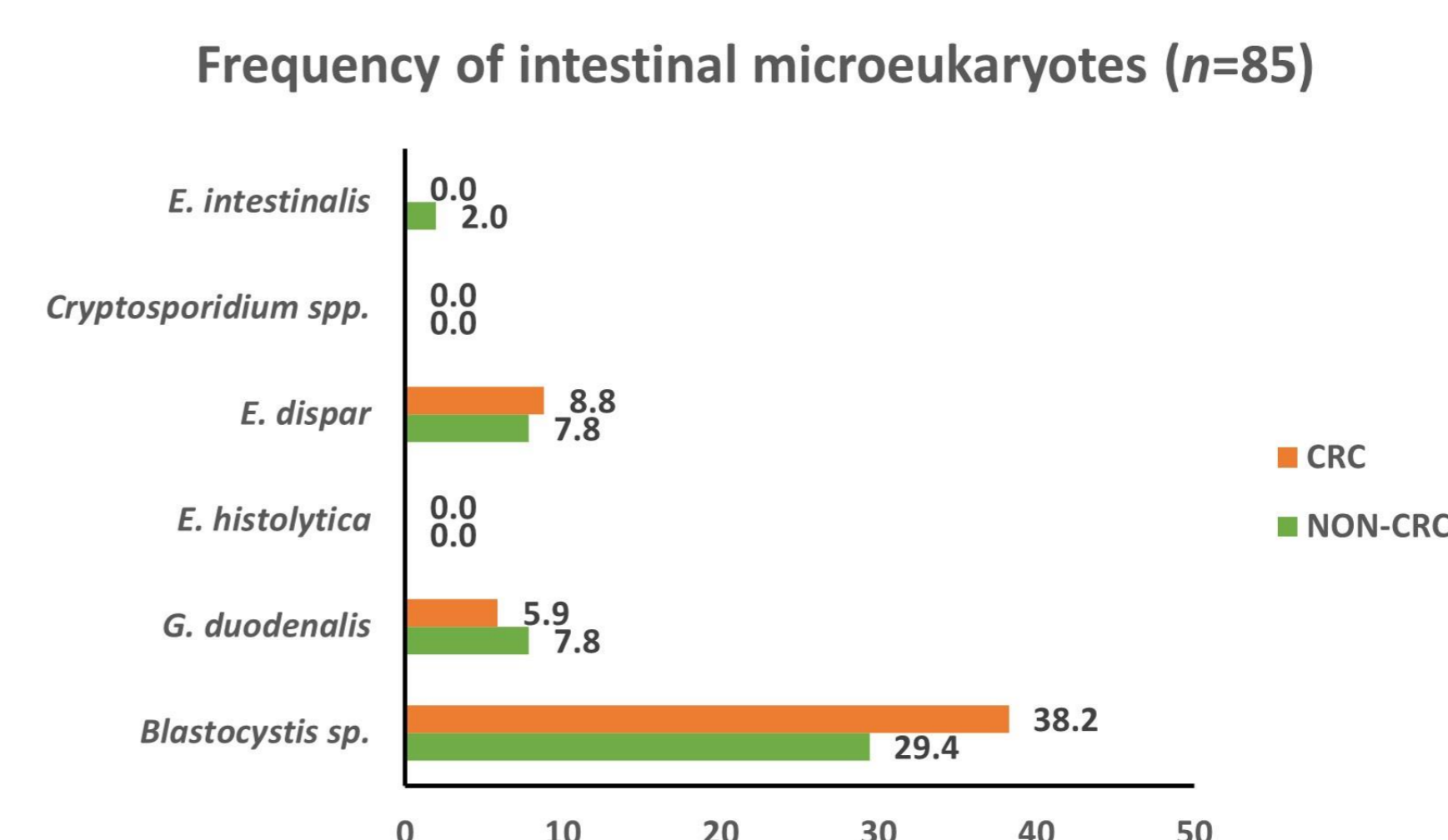


Fig. 1: Frequency of intestinal ME by microscopy and/or PCR in CRC and non-CRC patient cohorts.

	Sub-assembly/ Subtype	NO CRC	CRC
<i>Giardia duodenalis</i>	BIV	25.0% (1/4)	—
	Mixed	—	50.0% (1/2)
<i>Blastocystis</i> sp.	ST1	20.0% (3/15)	7.7% (1/13)
	ST2	13.3% (2/15)	—
	ST3	33.3% (5/15)	53.8% (7/13)
	ST6	—	7.7% (1/13)
	ST7	6.7% (1/15)	—
	ST17*	6.7% (1/15)	—
	Mixed	20.0% (3/15)	30.8% (4/13)

* First report in human population.

Table 1: Frequency of molecular diversity of *G. duodenalis* and *Blastocystis* sp. in CRC and non-CRC patient cohorts.

CONCLUSIONS

Although still preliminary, *Blastocystis* infection was higher in CRC patients (38.2%) than in non-CRC patients (29.4%).

Pathogenicity of *Blastocystis* remains disputable; this study suggest that infection by *Blastocystis* ST3 may be a risk factor for CRC development. This result is in line with those obtained in previous surveys showing the predominance of ST3 in CRC patients⁴.

Novel technologies like NGS may help unraveling the role of mixed *Blastocystis* infections involving two or more STs in chronic gastrointestinal illnesses.

REFERENCES

- Parkin DM. The global health burden of infection-associated cancers in the year 2002. *Int J Cancer*. 2006.
- Taghipour, A, et al. Are *Blastocystis hominis* and *Cryptosporidium* spp. playing a positive role in colorectal cancer risk? A systematic review and meta-analysis. *Infect Agents Cancer*. 2022.
- Ali, S.H., et al. An Association Between *Blastocystis* Subtypes and Colorectal Cancer Patients: A Significant Different Profile from Non-cancer Individuals. *Acta Parasit*. 2022.
- Sulzyc-Bielicka V, et al. Colorectal cancer and *Blastocystis* sp. infection. *Parasit Vectors*. 2021.

ACKNOWLEDGEMENTS