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Increasing incidence of microscopic colitis in a population-based cohort study in Switzerland

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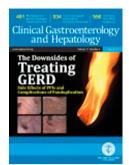
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3	Incre	asing incidence of microscopic colitis in a population-based cohort						
4	study	v in Switzerland						
5 6	Hugo	Maye ¹ , Ekaterina Safroneeva ² , Sébastien Godat ¹ , Christine Sempoux ³ , Pu						
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44 **INTRODUCTION**

Microscopic colitis (MC) is a chronic inflammatory disease of the colon that 45 presents with chronic, non-bloody watery diarrhea and only few or no endoscopic 46 abnormalities. Histologic examination discriminates lymphocytic colitis (LyC; 47 presence of ≥20 intraepithelial lymphocytes per 100 surface epithelial cells) and 48 collagenous colitis (CC: colonic subepithelial collagen band >10 micrometers in 49 50 diameter).[1,2] MC not otherwise specified (NOS) describes a subgroup of 51 patients who do not fulfill the diagnostic criteria for either CC or LyC.[1,2] 52 Population-based epidemiologic data regarding MC are scarce. We aimed to 53 evaluate the clinical presentation at diagnosis, incidence and prevalence of MC 54 in Cantons of Vaud and Fribourg, Switzerland.

55

56 **METHODS**

Cantons of Vaud and Fribourg lie in the French speaking, Western part of 57 58 Switzerland. As of 12/2017, both cantons together had a population of 1,109,230 inhabitants. After having identified MC patients through databases of all 59 60 Pathology institutes (n=6) serving both cantons and a histology slide review to assure correctness of diagnosis, we performed a chart review in practices of all 61 gastroenterologists covering both cantons (n=42). The study was approved by 62 the ethics committee of Cantons of Vaud and Fribourg (CER-VD 306/15). Two 63 64 hundred and fifty-two patients with MC, diagnosed between January 1994 and 65 December 2017, were identified. Of these, 34 were excluded for having NOS. We calculated incidence rates using data provided by the Institutes of population
 statistics of Canton of Vaud and Fribourg.

68

69 **RESULTS**

Of the 218 patients with MC, 123 (56.4%) had LyC and 95 (43.6%) had CC. 70 Seventy-four percent (162/218) of MC patients were female, mean age at first 71 72 symptoms was 62±15.4 years (range 24-89), mean age at MC diagnosis was 73 63.2±14.3 years (range 29-89). All MC patients suffered from diarrhea, followed by abdominal pain (31.7%), weight loss (31.2%), bloating (20.6%), fatigue 74 75 (9.6%), nausea / vomiting (3.2%). Exposure to risk factors for MC were frequently found and included HMG-CoA reductase inhibitors (27.1%), non-steroidal anti-76 inflammatory drugs (14.2%), proton-pump inhibitors (22.5%), serotonine reuptake 77 78 inhibitors (22.5%), and smoking (20.2%). Infectious agents were searched and excluded as cause of chronic diarrhea in all included patients. 79

A colonoscopy was performed as diagnostic tool in all of the 218 patients. In 74.3% of patients the colonoscopy was normal. Polyps were found in 16.5% of MC patients, followed by edema (9.2%), erythema (4.6%), and an erosion (0.5%). Median thickness of the subepithelial collagen band in patients with CC was 25µm, whereas patients with LyC had a median of 35 intra-epithelial lymphocytes per 100 epithelial cells.

Oral budesonide was most frequently used as first therapy (72.9%), followed
by loperamide (66.1%), aminosalicylates (16.1%), and cholestyramine (12.8%).

Incidence rates were calculated and are shown together with the cumulative prevalence in **Table 1**. No patient was diagnosed with MC prior to 1994. Incidence of MC significantly increased from 0.36/100,000 person-years in 1994-1997 to 6.85/100,000 person-years in 2017 (p=0.025, trend test). The cumulative prevalence of MC, LyC, and CC in 2017 was 19.65/100,000, 11.09/100,000, and 8.56/100,000, respectively. As such, the current prevalences for MC, LyC, and CC are 1/5,088 persons, 1/9018 persons, and 1/11,676 persons, respectively.

96

97 **DISCUSSION**

Our population-based study from Western Switzerland found a steady increase 98 in incidence of MC during the last two decades. Findings of our study are in 99 100 accordance with the results of a systematic review and meta-analysis that 101 reported pooled incidence rates for CC of 4.14 (95% CI 2.89-5.40) per 100,000 102 person-years and 4.85 (95% CI 3.45-6.25) for LyC. Bergman et al. assessed the incidence of MC in Sweden from 1995-2015 in a nationwide cohort. Among 103 13,844 patients, incidence of MC was 10.5/100,000 as from 2006 which is 104 105 roughly 6 times higher when compared to our findings.[4] Our data are with the results of Fernandez-Banares et al. who found lower 106 comparable incidences (2.2/100,000 for LyC and 2.6/100,000 for CC) among 290,000 107 108 inhabitants in Spain.[5] These results reinforce the existence of a north-south 109 gradient of MC which has been described by several groups.[3]

110 Strengths of our study are that all gastroenterologists and pathologists 111 working in Cantons of Vaud and Fribourg collaborated in this project which is

121	Medical writing: none
120	Funding sources: none
119	
118	countries, MC incidences are low in the population we studied.
117	incidence was steadily increasing over the last two decades. Compared to other
116	In conclusion, in the first Swiss population-based study we found that MC
115	response to different drugs.
114	to evaluate questions regarding the natural history of MC such as therapeutic
113	related to its retrospective design that impairs the generation of high-quality data
112	crucial for the generation of population-based data. Limitations of our study are

- Conflict of interest relevant to this study: none for all authors 122 ourno
- 123

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TABLES

Table 1: Incidence (plus 95% confidence intervals) and prevalence of MC, LyC, and CC from 1994 to 2017. The incidence is shown per 100,000 inhabitants, stratified according to 4 year intervals and per year. The prevalence was calculated per 100,000 inhabitants at the end of the respective 4 year interval.

Interval	1994-	1998-	2002-	2006-	2010-	2014-
	1997	2001	2005	2009	2013	2017
Population Vaud +	839,965	856,091	901,168	967,802	1,027,985	1,109,230
Fribourg					8	
MC new cases	3	8	22	45	64	76
MC incidence per	0.36,	0.93,	2.44,	4.7,	6.23,	6.85,
4 year interval	0-0.48	0.48-1.4	2.2-2.68	3.32-5.8	5.44-7	6.12-7.56
MC incidence per	0.09	0.23	0.61	1.18	1.56	1.71
year				×		
MC prevalence	0.36	1.29	3.66	8.06	13.81	19.65
LyC new cases	1	5	16	27	35	39
LyC incidence per	0.12,	0.58,	1.78,	2.79,	3.41,	3.52,
4 year interval	0-0.48	0.48-	1.32-2.2	2.48-	2.72-3.88	2.88-3.96
		0.92		3.32		
LyC incidence per	0.03	1.5	0.45	0.7	0.85	0.88
year						
LyC prevalence	0.12	0.7	2.44	5.06	8.17	11.09
CC new cases	2	3	6	18	29	37
CC incidence per	0.24,	0.35,	0.67,	1.86,	2.92,	3.34,
4 year interval	0-0.48	0-0.48	0.48-	1.24-	2.72-3.12	2.88-3.6
			0.89	2.48		
CC incidence per	0.06	0.09	0.17	0.47	0.73	0.84
year						
CC prevalence	0.24	0.58	1.22	2.99	5.64	8.56

TABLES

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CC incidence per	0.06	0.09	0.17	0.47	0.73	0.84
year						
CC prevalence	0.24	0.58	1.22	2.99	5.64	8.56

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