INTRODUCTION

Food and eating can be a source of pleasure, means of social interaction and peer acceptance. Having Inflammatory Bowel Disease (IBD) may alter these psychosocial factors because of painful or embarrassing symptoms and/or undernutrition resulting in activity limitation. However little is currently known about the impact of IBD on the psychosocial factors of food and quality of life.

AIMS

This study aimed to determine patients’ experiences of the social and psychological impacts of food on people with IBD.

METHODS

Semi-structured interviews were carried out with 28 patients regarding their experiences of food and eating in relation to their IBD. Interviews were recorded and transcribed verbatim. Concepts were labelled through line by line coding using a constant comparative approach based on grounded theory.

RESULTS

<table>
<thead>
<tr>
<th>Disease Type</th>
<th>Male (%)</th>
<th>Age (Mean, SD)</th>
<th>Years since diagnosis (Mean, SD)</th>
<th>Surgery (%)</th>
<th>Active disease (%)</th>
<th>BMI (Mean, SD)</th>
<th>MUST score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crohn’s</td>
<td>43.75%</td>
<td>37.59 (12.06)</td>
<td>10.34 (10.86)</td>
<td>43.75%</td>
<td>56.25%</td>
<td>23.33 (5.58)</td>
<td>Low (63.63%)</td>
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<tr>
<td>Ulcerative</td>
<td>50%</td>
<td>37.26 (12.48)</td>
<td>10.41 (10.80)</td>
<td>8.30%</td>
<td>25%</td>
<td>23.22 (5.67)</td>
<td>Medium (50%)</td>
</tr>
</tbody>
</table>

CONCLUSION

Eating and drinking is a significant challenge for many IBD patients, particularly during flares; however CD patients appear to have more sustained eating issues during remission.

Cognitive and emotional factors such planning and preparation, eating an extra consideration and frustration around missing out and unpredictability impact on quality of life and enjoyment of eating, especially away from home.

Acknowledging difficulties around eating and drinking could improve quality of life and adjustment to IBD.

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