

# NHS Lothian Research and Development Conference 2025



# Reduced Risk of Colorectal Cancer in Inflammatory Bowel Disease with Biologic Therapies or 5-Aminosalicylates but not Immunomodulators: A Systematic Review and Meta-analysis

Amirhosein Kefayat<sup>1,2</sup>, Ross J. Porter<sup>3,4</sup>, Antonia M.D. Churchhouse<sup>1,2</sup>, Jonathan Blackwell<sup>2</sup>, Eleanor F. Watson<sup>2</sup>, A. John Morris<sup>4</sup>, Morris Gordon<sup>5</sup>, Gaurav B. Nigam<sup>6</sup>, James E. East<sup>6</sup>, Matthew D. Rutter<sup>7,8</sup>, Christopher A. Lamb<sup>9,10</sup>, Tim Raine<sup>11</sup>, Alexander C. Ford<sup>12,13</sup>, Shahida Din<sup>1,2</sup>

<sup>1</sup>Institute of Genetics and Cancer, University of Edinburgh, Edinburgh, United Kingdom., <sup>2</sup>Edinburgh Inflammatory Bowel Diseases Unit, Western General Hospital, Edinburgh, United Kingdom., <sup>3</sup>School of Cancer Sciences, College of Medical, Veterinary and Life Sciences, University of Glasgow, United Kingdom., <sup>4</sup>Department of Gastroenterology, Queen Elizabeth University Hospital, Glasgow UK., <sup>5</sup>School of Medicine, University of Central Lancashire, Preston, Lancashire, UK., <sup>6</sup>Translational Gastroenterology Unit, Nuffield Department of Medicine, John Radcliffe Hospital, University of Oxford, Oxford, UK., <sup>7</sup>Department of Gastroenterology, North Tees and Hartlepool NHS Foundation Trust, Stockton-on-Tees, UK, <sup>8</sup>Faculty of Medical Sciences, Newcastle University, Newcastle-upon-Tyne, UK, <sup>9</sup>Translational & Clinical Research, Institute, Newcastle University, Newcastle upon Tyne, UK. <sup>10</sup>Gastroenterology, Newcastle Upon Tyne Hospitals NHS Foundation Trust, Newcastle Upon Tyne, UK. <sup>11</sup>Department of Gastroenterology, Addenbrooke's Hospital, Cambridge University Hospitals NHS Trust, Cambridge, UK. <sup>12</sup>Leeds Gastroenterology Institute, St. James's University Hospital, Leeds, United Kingdom.

### **Introduction:**

The incidence and mortality of colorectal cancer (CRC) remains elevated in the inflammatory bowel disease (IBD) population. We aimed to examine the association between biologics, 5-aminosalicylates (5-ASAs), and immunomodulators, with the risk of CRC and / or dysplasia in patients with IBD in a contemporaneous systematic review and meta-analysis.

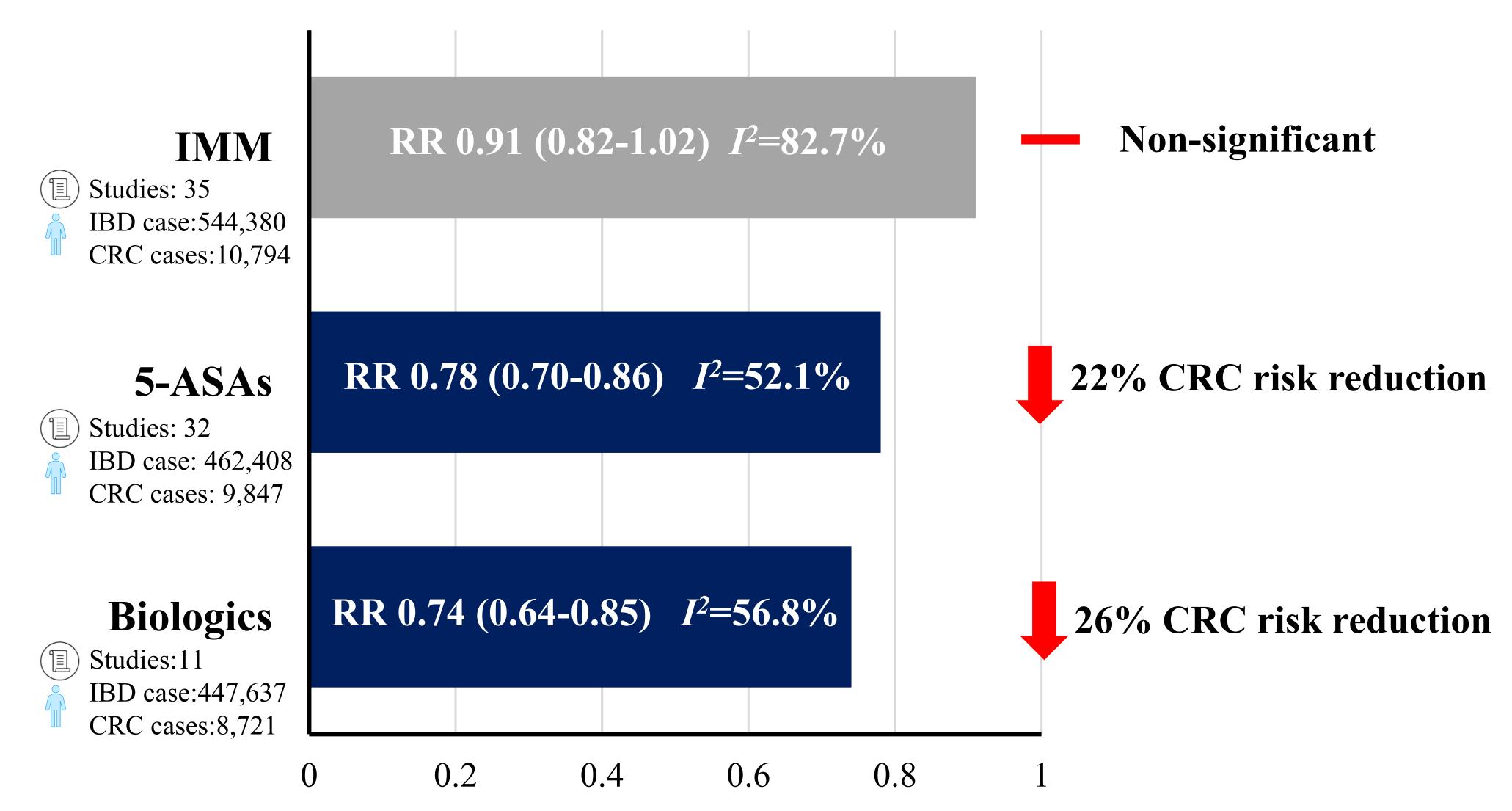
## **Method:**





**Embase**°

We searched Web of Science, PubMed, MEDLINE, and EMBASE from inception to 15th June 2024 for all studies assessing the impact of biologics, 5-ASAs and immunomodulators on the occurrence of CRC or dysplasia in adults (≥16 years) with IBD. Data were pooled using a random effects model generating relative risk (RR) estimates with 95% confidence intervals (CIs).



#### **Results:**

- If the RR was less than 1 and the 95% CI did not cross 1, there was a significantly reduced likelihood of the AE with active drug.
- We assessed inter-study heterogeneity, using the  $I^2$  statistic. The  $I^2$  ranges between 0% and 100%, with values of 25% to 49% considered low, 50% to 74% moderate, and  $\geq$ 75% high heterogeneity between the studies.
- Biologic therapies and 5-ASAs were associated with a reduced risk of CRC and /or dysplasia in patients with IBD.
- Immunomodulators were not associated with a reduced risk.

### **Conclusions:**

The use of biologics and 5-ASAs, but not immunomodulators, are associated with a reduced risk of CRC and / or dysplasia. Further studies are need to reveal the reason but some studies suggest it might be inflammation control by these medications.

: @ShahidaDin1

sdin@ed.ac.uk
akefayat@ed.ac.uk