Review

Artificial intelligence and inflammatory bowel disease: practicalities and future prospects

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ABSTRACT

Artificial intelligence (AI) is an emerging technology predicted to have significant applications in healthcare. This review highlights AI applications that impact the patient journey in inflammatory bowel disease (IBD), from genomics to endoscopic applications in disease classification, stratification and selfmonitoring to risk stratification for personalised management. We discuss the practical AI applications currently in use while giving a balanced view of concerns and pitfalls and look to the future with the potential of where AI can provide significant value to the care of the patient with IBD.

INTRODUCTION

Artificial intelligence (AI) is a computational system designed by humans that, given a specific goal, takes in data, interprets it, processes it and decides on the best action to achieve the goal.

General AI systems perform nonbiological tasks that humans can do. To do this, they require the capabilities of common sense, self-awareness or the ability to define their purpose. They are not widely used in medicine. Narrow AI systems are those that we recognise in everyday use, which has one function, for example, AI optimising hospital bed flow.

AI encompasses several approaches, the most common being machine learning (ML). This applies computer algorithms to capture behaviour and patterns in systems and processes based on the input and output data. An example of this is natural language processing (NLP), in which a computer is trained to understand text and spoken word. Within ML, there is a specific subset called deep learning (DL). DL uses neural networks with multiple

Key messages

- ⇒ Artificial intelligence (AI) is a growing field which could improve patient care in inflammatory bowel disease (IBD).
- \Rightarrow Al in IBD uses two main types: machine learning and deep learning.
- ⇒ AI is currently used for IBD pathogenesis, drug discovery and endoscopic identification of disease.
- ⇒ Al could be used in IBD patient selfmonitoring, virtual management, endoscopic disease classification and patient stratification.
- ⇒ Narrow, task-focused AI methodologies can and do have a positive impact on patient care

layers, that is, networks that aim to mimic the brain through a combination of extensive data inputs, weights and biases (figure 1). An example of this is Convolutional Neural Networks (CNNs). They are used widely in image-based problems in medicine, for example, to detect and classify colonic polyps, to diagnose retinal disorders and to predict severity in chest X-rays for patients with congestive cardiac failure. Other approaches in AI DL which are less commonly used in medicine are machine reasoning and robotics. These are used in self-driving cars, but the idea of a machine diagnosing a patient with acute fulminant colitis, deciding to operate and performing a colectomy without any human intervention remains science fiction.

Inflammatory bowel disease (IBD) is a chronic debilitating condition, which has significant uncertainties along the patient journey from pre-diagnosis to the maintenance of disease control. In this review,



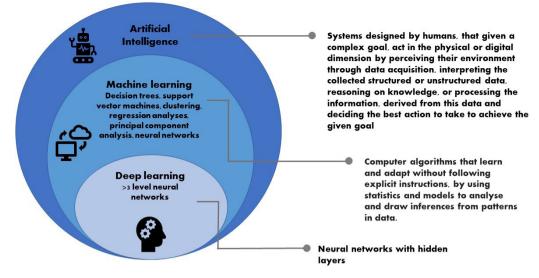


Figure 1 Artificial intelligence definition and common approaches.

we highlight the current and potential future applications of AI in the IBD journey (figure 2).

UNDERSTANDING IBD PATHOGENESIS

A significant role for AI in IBD is to aid understanding of aetiology and personalise disease management, via deeper insights into the complexity of this chronic immune condition which has both heritable and environmental components.¹

Genomics and multiomics

Interpretation of vast numbers of genetic variants in IBD and correlating these with phenotypes is highly challenging. This already uses elements of AI; in the field of genetic variant annotation, a notable example is the Combined Annotation-Dependent Depletion score.² This method uses a ML model to assign a score to any genetic variant based on the determination of whether the variant will be deleterious or not. Other examples of AI use for variant annotation occur in the paediatric field for very early onset IBD patients (aged <6 years at diagnosis) who routinely undergo genomic sequencing.³ ML models such as LEArn to Prescribe (LEAP) use big data to identify paediatric IBD patients' deleterious genetic markers with excellent accuracy (area under the curve (AUC) >0.97), providing a direct clinical impact of AI in IBD.⁴

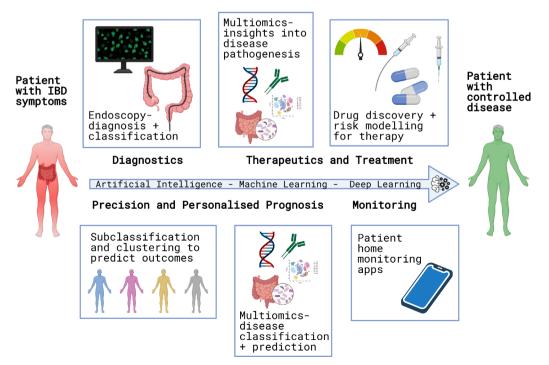


Figure 2 Application of artificial intelligence in inflammatory bowel disease (IBD).

The microbiome

The microbiome is increasingly recognised to play a role in IBD and is very well suited to AI analyses. 'Big data metagenomics' using both ML and specific DL algorithms can distinguish the microbiomes of IBD and control populations.⁵ The use of AI within host-microbiome metabolomic networks affords a predictive insight into the microbiome-host interplay in the gut covering the spectrum of synergistic effects to dysbiosis in health and disease states.⁶

A challenge of data integration lies in the heterogeneity of 'omic data inputs, from genotypes, transcript counts, bacterial species abundance, and clinical metadata. This complexity is exceptionally hard to understand, but AI systems appear successful for example, Hyams *et al* used a combination of microbial, transcriptomics and clinical data to predict 52-week steroid-free remission in paediatric ulcerative colitis (UC).⁷ Here, the authors utilised a multivariable logistic regression model and identified an AUC of 0.75 when using disease activity, week 4 remission, antimicrobial transcriptomic signature and specific bacterial abundance as input variables.

AI IN IBD DIAGNOSIS

Risk prediction, disease classification

Most of the existing literature on the use of AI in diagnosing IBD relates to developing risk prediction models using either ML or CNNs to analyse endoscopic and imaging datasets.⁸ But ML algorithms can also successfully classify disease subtypes based on genotype. Wei *et al* used ML to classify CD and UC, vs controls (AUC=0.862 and 0.826, respectively)⁹ based on single nucleotide polymorphisms (SNPs). The Crohn's disease exome challenge applied ML and DL to SNP and whole-exome sequencing to produce a classification model with an AUC of 0.72.¹⁰ Smolander *et al* have explored the use of deep belief networks, which are another type of neural network, in distinguishing UC from Crohn's disease with an accuracy of around 97%.¹¹

AI for endoscopic assessment

An area where most endoscopists value the potential for AI is endoscopic severity assessment. A recent prospective study using CNN in 875 patients with UC demonstrated the ability to detect endoscopic remission (Mayo Endoscopic Score, MES score of 0 vs 1) with 90% accuracy and histological remission with 92.9% accuracy comparable to expert reviewers.¹² In a further study, AI was able to detect MES Scores from both still and video images.¹³ Similar data is available for computer-aided diagnosis of grade 1 vs grade 3 ulcers in CD and histological inflammation by endocytoscopy. Takenaka *et al* have designed a deep neural network model for UC, trained on >40 000 endoscopic still digital images with corresponding UC Endoscopic Index of Severity (UCEIS) scores and histology. Compared with human reviewers, this system was highly accurate in assessing endoscopic and histologic remission.

A recent proof-of-concept study using an automated real-time operator-independent red density score in UC reported a good correlation between the red density score, a consensus MES and UCEIS, as well as a Robart's histological index.¹⁴ As a note of caution, translating accuracy from still images to live video analysis is a work in progress but solutions are starting to appear.

Other potential areas for application are standardised reporting of IBD endoscopy such as auto report generation through computer-aided detection or task automation such as data extraction from electronic medical records, generating reports and billing through NLP.

Clinical trials in IBD increasingly use central reading to assess endoscopic severity. Automated endoscopic scoring in high-resolution trial endoscopy videos with frame integration using recurrent neural network showed high accuracy in prediction of the correct Mayo score (70%) with a satisfactory agreement with human observers (k) of 0.84 (0.79–0.90).¹⁵ However, Yao *et al*¹³ showed that a high-quality video source is essential for optimal AI performance to be applicable in the setting of central reading. AI can also provide standardised and reproducible disease assessments and integrate images into decision support in strategy trials challenging the conventional therapeutic endpoints.

Capsule endoscopy images also benefit from AI to varying degrees. Barash *et al* developed CNN to evaluate ulceration in CD in which the overall agreement between expert consensus and an automated algorithm was 67%.¹⁶ A DL model in capsule endoscopy reading was superior to conventional reading both at the level of per-patient sensitivity (99.9% vs 74.6%, p=<0.0001) and per lesions sensitivity 99.9% vs 76.9%, p=<0.0001).¹⁷

EndoBRAIN-EYE (Cybernet Systems, Tokyo, Japan) has developed a prototype AI-based polyp detection system to detect flat dysplasia in long-standing colitis patients.

AI in radiological diagnosis

AI can also interpret IBD radiological images. Stidham *et al* reported that structural bowel damage measurements of CT enterography data in CD by semiautomated approaches are comparable to those of experienced radiologists.¹⁸ Furthermore, radiomics models can predict moderate to severe histological intestinal fibrosis more accurately than skilled radiologists.¹⁹

THERAPY AND PATIENT-FOCUSED MANAGEMENT Al and apps

AI-based healthcare apps are a growing industry, with context-aware smartphone applications for personalised outcomes for the user. Most healthcare apps use a mobile expert system model: a knowledge base containing the facts and rules relevant to the target application, which can be accessed and updated by an IBD Specialist, and a user interface, into which users place queries or results, for example, stool frequency, level of pain, mood, etc. The knowledge base and user interface are input into an inference engine that applies an IF-THEN rule (IF being the condition and THEN being the consequence), which feeds back to the user and the expert. An example of this in IBD is when a user places stool frequency at $>5 \times$ normal, with blood—the app would advise doing a faecal calprotectin and contacting the IBD service. Some apps feedback to the expert interface to alert them of a change in circumstance.

A scoping review of Digital Health apps for the Clinical care of IBD²⁰ identified 10 digital health apps for IBD, which used the app as a self-reporting/ documentation app and provided educational modules. Half of the apps had a contact through to team capability, which was user instigated, with thresholds set for investigation, for example, faecal calprotectin. One utilised ML algorithms to predict the requirement for escalation(with 95% accuracy), with treatment advice given regarding 5aminosalicy-late (5ASA) and rectal therapy.

Like other health apps, IBD apps have the potential to provide patient-centred and patient-initiated care, improve access to specialised medical services in areas that are underserved in a low-cost way. However, complexities arise with the need for encryption for patient identifiable data and the discriminant function of symptoms in identifying non -IBD pathology such as cancer in patients with IBD.

AI in IBD monitoring

Chatbots

Conversational AI methods such as chatbots may enhance timely access to IBD specialists. In using a chatbot, the input (query) must be able to be categorised effectively. In a feasibility study, an NLP algorithm was developed to classify IBD patient electronic messaging dialogue into specific queries: symptoms based, medication queries, investigation results, insurance, procedures, communications and miscellaneous. The algorithm had 90% concordance with two consultant gastroenterologists who reviewed the data and categorised it.²¹

Smart loos

With the advent of the internet of things (IoT), for example, the embedding of computing devices into everyday objects that connect via the internet, the opportunity to use everyday objects as a healthcare adjunct is arising. For IBD sufferers, the potential to use the toilet as a monitoring system for their disease is a valid and appropriate use of a combination of IoT and AI. Smart Toilets have been around since 2000, when they were initially capable of monitoring the glucose level in urine. The level of sophistication has expanded, and Smart toilets can now categorise, by DL, the stool type (Bristol Stool scale) and longitudinally log frequency and type for the user (identified by their fingerprint and anoderm). This data is accurate biometric data for the physician.²² Fisher²³ adapted the Smart Toilet to integrate an AI-based stool analysis (using CNN for stool type) and a perceptual colour quantisation for accurate real-time stool and faecal blood analysis. This has reduced recall inaccuracies and the burden of manually checking every bowel movement.

Wearables and nearables

Worn monitoring devices called wearables and nearables (neighbouring devices that interact with wearables such as smartphones) are now mainstream. These are the most likely technologies to transform future healthcare and lifestyles via self-motivation management.²⁴ Sensors to detect heart rate and step count are commonplace with integrated AI to promote healthy exercise. The value of exercise in managing fatigue and reducing the risk of flares in IBD is clear,²⁵ and in this way, AI and IoT enabled wearables and nearables will be part of the IBD journey. The prospect of wearing a sensor for early detection of a flare via sweat is an exciting wearable development for IBD, but as yet had not been AI-enabled or IoT integrated.²⁶

AI in drug design

AI is used in all drug design and discovery facets. Traditionally quantitative structure-activity relationships (QSAR) were used, which made accurate predictions of simple attributes such as solubility, which are straightforward. More complex attributes such as drug efficacy or side effect identification were not plausible using this technique.

Using Big Data and large databases, AI modelling algorithms can predict interactions between drugs and biological targets. Wen *et al*²⁷ used DL to identify drug-target interactions from DrugTarget Database. Xie *et al*²⁸ used DL-based transcriptome data classification for drug-target interaction prediction from the Library of Integrated Network-Based Cellular Signatures to identify novel targets for known drugs.

Unbiased AI-assisted methods for target identification in IBD²⁹ identified relevant gene clusters in IBD and actionable targets. The AI algorithm identified PRKAB1 as a barrier protective therapeutic target, which was then experimentally validated. The model was able to predict phase III results in clinical trials in IBD successfully.

Al to predict drug response

It is becoming possible to predict response to drugs, which is a required step for personalised medicine,^{30 31} for example, Morilla *et al* using deep neural networks found nine colonic microRNAs with five clinical features that predicted response to steroids (AUC 0.91) and infliximab (AUC 0.82).³² However, the use of integrated AI within these analyses is still in its infancy. It is essential for accurate clinical data to be used in these models and to validate findings in independent cohorts, as previous molecular biomarkers have not produced consistently reproducible results in all populations.³³

AI RISKS

The difficulty with DL modelling is while it allows modelling of biological endpoints, which reduces the overfitting, biased data and errors from small datasets, there are no universally agreed tunable parameters or criteria for the workflows, which opens the models to criticism.³⁴

A recent safety issue highlighted by IBM Watson Oncology AI algorithm, which utilised synthetic data to train the algorithm, led to incorrect and unsafe therapeutic advice, although not to an actual patient. However, this situation has cast a shadow over medical apps providing feedback to patients regarding therapeutic strategies.

CNNs are used to analyse images produced from high throughput screening and are also used to visualise 3D experimental and virtual images for ligand-protein prediction.³⁵ However, as such a wide application, it is fraught with difficulty regarding the available knowledge regarding the chemical space and the complexity of the biological context. There are at least 10 AI algorithms/ML or DL applications for structural and ligand-based virtual screening, including LS-Align, LigGrep, DrugFinder and DEEPScreen.³⁶

Despite the immense potential, the drug discovery field lags behind other successes utilising AI in terms of the amount of chemical and biological data available, the ability to represent the data, the presence of a suitable stable algorithm, or the ability to appropriately assign biological labels, for example, dose, genotype, age, etc.³⁷

AI and risks to the doctor-patient relationship

Arguably, the most important aspect of the doctorpatient relationship is trust. But AI lacks transparency because of the 'black box' nature of some algorithms. This leads to concerns about indefensible decisions and the loss of trust of the medical profession, potentially reducing patient engagement and concordance with therapy. Demystifying AI and clarity about where AI can aid the physician instead of replacing the physician will help allay such fears. Box 1 identifies current ethical and safety aspects of AI.

Training

In a recent UK-based qualitative study, perception of AI in the medical community was worryingly one of fear and concern about ethical consequences, despite

Box 1 Ethical and safety issues with artificial intelligence (AI) in medicine (see high-level expert group on AI ethics guidelines for a trustworthy AI)

A robust Al system must:

- $\Rightarrow\,$ Have respect for human autonomy.
- \Rightarrow Prevent harm have explicability.
- \Rightarrow Have human agency and oversight.
- ⇒ Have technical robustness and safety⇒ Resilience to attack and security.⇒ Have a fall-back plan.
- \Rightarrow Apply privacy and data governance laws \Rightarrow Show quality and integrity of data.
 - \Rightarrow Have accessibility to data.
- \Rightarrow Have transparency
 - \Rightarrow Be accurate.
 - \Rightarrow Show reliability and reproducibility.
 - \Rightarrow Have traceability.
 - \Rightarrow Be identifiable as an Al system (eg, chatbot).
- \Rightarrow Apply diversity, non discrimination and fairness
- \Rightarrow Have accountability
 - \Rightarrow Undertake Audit.
 - \Rightarrow Minimise and report negative impacts.
 - \Rightarrow Undertake trade-off and redress.
- \Rightarrow Apply societal and environmental well-being
 - \Rightarrow Be sustainable and environmentally friendly.
 - \Rightarrow Monitor societal impact (prevent harm).

two-thirds of the participants never having used AI applications. Nearly 90% had no working knowledge of the principles of AI.³⁸ Funding for education and engagement is going to be crucial so that clinicians understand how to use AI appropriately and appreciate its limitations. All medical AI applications must be General Data Protection Regulation (GDPR) compliant, with clear accountability and confirmation of privacy protection.

Perhaps even more important than training clinicians is to train the AI itself. Quality assurance depends on the quality of the training data sets. An AI that was not trained on gastric cancer is highly unlikely to recognise it and there may be subtle differences between patients that we are not aware of including age, sex and race differences. Transparent, standardised thresholds need to be set. The narrow nature of AI is both a blessing and a curse. for example, an algorithm designed to characterise IBD subtypes will not aid a trainee in loop resolution. But imagine having an AI that can suggest interventions to the junior endoscopist that lead to safer colonoscopy when loops form.

CONCLUSIONS

This review has highlighted areas of the IBD journey that already benefit from AI in a tangible, practical sense and suggested other areas where AI is plausible but prospective. In doing so, our aim is to familiarise, raise awareness, dispel concerns and enthuse medical colleagues as to the utility of AI within IBD. The

Colorectal

success of AI integration in the IBD journey will be measured by the value-added effect; increased understanding of the disease, better diagnostic certainty, efficacy improvements, better holistic outcomes for the patients, and improvements in the work-life balance of the IBD team. AI systems are being developed to give practitioners extra resources to provide better care.

The concept of the broad AI unit, which thinks, acts, responds and is self-aware, is rife with ethical and moral issues and is beset by the vilification ingrained within the literary and movie genres. Replacing the medical practitioner with an AI module in everyday practice remains a slightly distant concept. This review shows where narrow, task-focused AI methodologies can and do have a positive impact on patient care. Within IBD; Genomics Research, Endoscopy image classification and to a lesser extent, patient-facing home monitoring apps have led the way to confirm the place of AI in the IBD patient journey.

Overall we predict the role of AI will continue to grow in IBD and may help to address some of the clinical challenges we face, leading to improved quality in IBD care.

Glossary of terms

Algorithm: The process or rules to be followed in problem-solving operations.

Blackbox: A complex system whose internal workings are hidden or not easily understood.

Chatbot: A computer program designed to simulate conversation, usually over the internet.

CNN: Multilayered networks, specialising in image processing, designed to detect multiple features in images to correctly identify the image—for example, social media 'automatic tagging' of individuals in photos.

DL: Class of ML that uses artificial neural networks (such as CNNs) to extract progressively high-level data from raw data, for example, identification of a type of polyp in an endoscopic image.

DeepScreen: Large scale drug-target interaction prediction system using deep CNNs (can be found on github.com).

DrugFinder: Drug Target Interaction database.

Inference Engine: Processing component which makes a decision from facts/rules contained in the knowledge base of an expert system.

IoT: System of interrelated, internet connected objects, for example, a fridge which orders the weekly shopping and links to your smartphone.

LEAP: algorithm is a sequential decision making process to identify adverse drug reactions in the state of polypharmacy.

LS-Align: A tool for screening large numbers of compounds against disease relevant proteins.

ML: The use and development of computer systems that are able to learn and adapt by using algorithms

and models to analyse and draw inferences from patterns in data.

NLP: Field of linguistics, computer science and AI to train a computer to understanding language, including nuances, to be able to extract large amounts of information from documents.

Neural Networks: Series of algorithms that recognise underlying relationships in a set of data through processes that mimic the connections of neurons in the brain.

QSAR: Theoretical models that relate a quantitative measure of a chemical structure to a physical property.

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Contributors JB-W organised the authors, did the literature review wrote and edited the manuscript. JA did the literature review, wrote the manuscript and designed all the figures. AD wrote the manuscript and edited the manuscript. TT did the literature review and reviewed the manuscript for content and edited. PBA reviewed the manuscript for content and edited. SH reviewed the manuscript for content and edited. LBL wrote the manuscript and provided significant editing input. SS wrote the manuscript and provided significant editing input.

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REFERENCES

- Ashton JJ, Mossotto E, Ennis S, *et al.* Personalising medicine in inflammatory bowel disease-current and future perspectives. *Transl Pediatr* 2019;8:56–69.
- 2 Rentzsch P, Witten D, Cooper GM, *et al.* CADD: predicting the deleteriousness of variants throughout the human genome. *Nucleic Acids Res* 2019;47:D886–94.
- 3 Uhlig HH, Muise AM. Clinical genomics in inflammatory bowel disease. *Trends Genet* 2017;33:629–41.
- 4 Lai C, Zimmer AD, O'Connor R, *et al*. Leap: using machine learning to support variant classification in a clinical setting. *Hum Mutat* 2020;41:1079–90.
- 5 Iadanza E, Fabbri R, Bašić-ČiČak D, et al. Gut microbiota and artificial intelligence approaches: a scoping review. *Health Technol* 2020;10:1343–58.
- 6 Bar N, Korem T, Weissbrod O, et al. A reference map of potential determinants for the human serum metabolome. *Nature* 2020;588:135–40.
- 7 Hyams JS, Davis Thomas S, Gotman N, et al. Clinical and biological predictors of response to standardised paediatric colitis therapy (protect): a multicentre inception cohort study. Lancet 2019;393:1708–20.
- 8 Gubatan J, Levitte S, Patel A, *et al*. Artificial intelligence applications in inflammatory bowel disease: emerging technologies and future directions. *World J Gastroenterol* 2021;27:1920–35.
- 9 Wei Z, Wang W, Bradfield J, *et al*. Large sample size, wide variant spectrum, and advanced machine-learning technique boost risk prediction for inflammatory bowel disease. *Am J Hum Genet* 2013;92:1008–12.
- 10 Pal LR, Kundu K, Yin Y, et al. CAGI4 Crohn's exome challenge: marker SNP versus exome variant models for assigning risk of Crohn disease. *Hum Mutat* 2017;38:1225– 34.
- 11 Smolander J, Dehmer M, Emmert-Streib F. Comparing deep belief networks with support vector machines for classifying gene expression data from complex disorders. *FEBS Open Bio* 2019;9:1232–48.
- 12 Takenaka K, Ohtsuka K, Fujii T, *et al.* Development and validation of a deep neural network for accurate evaluation of endoscopic images from patients with ulcerative colitis. *Gastroenterology* 2020;158:2150–7.
- 13 Yao H, Najarian K, Gryak J, et al. Fully automated endoscopic disease activity assessment in ulcerative colitis. Gastrointest Endosc 2021;93:728–36.
- 14 Bossuyt P, Nakase H, Vermeire S, et al. Automatic, computeraided determination of endoscopic and histological inflammation in patients with mild to moderate ulcerative colitis based on red density. Gut 2020;69:1778–86.
- 15 Gottlieb K, Daperno M, Usiskin K, *et al.* Endoscopy and central reading in inflammatory bowel disease clinical trials: achievements, challenges and future developments. *Gut* 2021;70:418–26.
- 16 Barash Y, Azaria L, Soffer S, *et al.* Ulcer severity grading in video capsule images of patients with Crohn's disease: an ordinal neural network solution. *Gastrointest Endosc* 2021;93:187–92.
- 17 Ding Z, Shi H, Zhang H, et al. Gastroenterologist-Level Identification of Small-Bowel Diseases and Normal Variants by Capsule Endoscopy Using a Deep-Learning Model. Gastroenterology 2019;157:1044–54.
- 18 Stidham RW, Enchakalody B, Waljee AK, et al. Assessing small bowel Stricturing and morphology in Crohn's disease using semi-automated image analysis. *Inflamm Bowel Dis* 2020;26:734–42.

- 19 Li X, Liang D, Meng J, *et al*. Development and validation of a novel Computed-Tomography Enterography radiomic approach for characterization of intestinal fibrosis in Crohn's disease. *Gastroenterology* 2021;160:2303–16.
- 20 Yin AL, Hachuel D, Pollak JP, *et al.* Digital health apps in the clinical care of inflammatory bowel disease: Scoping review. *J Med Internet Res* 2019;21:e14630.
- 21 Zand A, Sharma A, Stokes Z, *et al*. An exploration into the use of a chatbot for patients with inflammatory bowel diseases: retrospective cohort study. *J Med Internet Res* 2020;22:e15589.
- 22 Park S-M, Won DD, Lee BJ, *et al.* A mountable toilet system for personalized health monitoring via the analysis of excreta. *Nat Biomed Eng* 2020;4:624–35.
- 23 Fisher D. Automated stool image analysis by artificial intelligence in A. DDW ePoster library, 2021. Available: https://eposters.ddw.org/ddw/2021/ddw-2021-virtual/320454/ deborah.fisher.automated.stool.image.analysis.by.artificial. intelligence.in.a.html [Accessed 26 Aug 2021].
- 24 Smuck M, Odonkor CA, Wilt JK, *et al.* The emerging clinical role of wearables: factors for successful implementation in healthcare. *NPJ Digit Med* 2021;4:45.
- 25 Engels M, Cross RK, Long MD. Exercise in patients with inflammatory bowel diseases: current perspectives. *Clin Exp Gastroenterol* 2018;11:1–11.
- 26 Jagannath B, Muthukumar S, Prasad S. Wearable sweat sensing device for detection of IBD biomarkers. *Inflamm Bowel Dis* 2021;27:S12.
- 27 Wen M, Zhang Z, Niu S, et al. Deep-Learning-Based drugtarget interaction prediction. J Proteome Res 2017;16:1401–9.
- 28 Xie L, He S, Song X, *et al*. Deep learning-based transcriptome data classification for drug-target interaction prediction. *BMC Genomics* 2018;19:667.
- 29 Sahoo D, Swanson L, Sayed IM, et al. Artificial intelligence guided discovery of a barrier-protective therapy in inflammatory bowel disease. Nat Commun 2021;12:4246.
- 30 Noor NM, Verstockt B, Parkes M, et al. Personalised medicine in Crohn's disease. Lancet Gastroenterol Hepatol 2020;5:80– 92.
- 31 Borg-Bartolo SP, Boyapati RK, Satsangi J. Precision medicine in inflammatory bowel disease: concept, progress and challenges. [version 1; peer review: 2 approved]. *F1000Res* 2020;9.
- 32 Morilla I, Uzzan M, Laharie D, *et al.* Colonic microRNA profiles, identified by a deep learning algorithm, that predict responses to therapy of patients with acute severe ulcerative colitis. *Clin Gastroenterol Hepatol* 2019;17:905–13.
- 33 Gasparetto M, Payne F, Nayak K, et al. Transcription and DNA Methylation Patterns of Blood-Derived CD8⁺ T Cells Are Associated With Age and Inflammatory Bowel Disease But Do Not Predict Prognosis. Gastroenterology 2021;160:232–44.
- 34 Zhou Y, Cahya S, Combs SA, *et al*. Exploring tunable Hyperparameters for deep neural networks with industrial ADME data sets. *J Chem Inf Model* 2019;59:1005–16.
- 35 Hochuli J, Helbling A, Skaist T, et al. Visualizing convolutional neural network protein-ligand scoring. J Mol Graph Model 2018;84:96–108.
- 36 Gupta R, Srivastava D, Sahu M, *et al.* Artificial intelligence to deep learning: machine intelligence approach for drug discovery. *Mol Divers* 2021;25:1315–60.
- 37 Bender A, Cortes-Ciriano I. Artificial intelligence in drug discovery: what is realistic, what are illusions? Part 2: a discussion of chemical and biological data. *Drug Discov Today* 2021;26:1040–52.
- 38 Castagno S, Khalifa M. Perceptions of artificial intelligence among healthcare staff: a qualitative survey study. *Front Artif Intell* 2020;3:578983.