


BMJ Open Use of external control arms in immune-mediated inflammatory diseases: a systematic review

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ABSTRACT

Objectives External control arms (ECAs) provide useful comparisons in clinical trials when randomised control arms are limited or not feasible. We conducted a systematic review to summarise applications of ECAs in trials of immune-mediated inflammatory diseases (IMIDs).

Design Systematic review with an appraisal of ECA source quality rated across five domains (data collection, study populations, outcome definitions, reliability and comprehensiveness of the dataset, and other potential limitations) as high, low or unclear quality.

Data sources Embase, Medline and Cochrane Central Register of Controlled Trial were searched through to 12 September 2023.

Eligibility criteria Eligible studies were single-arm or randomised controlled trials (RCTs) of inflammatory bowel disease, pouchitis, rheumatoid arthritis, juvenile idiopathic arthritis, ankylosing spondylitis, psoriatic arthritis, psoriasis and atopic dermatitis in which an ECA was used as the comparator.

Data extraction and synthesis Two authors independently screened the search results in duplicate. The characteristics of included studies, external data source(s), outcomes and statistical methods were recorded, and the quality of the ECA data source was assessed by two independent authors.

Results Forty-three studies met the inclusion criteria (inflammatory bowel disease: 16, pouchitis: 1, rheumatoid arthritis: 12, juvenile idiopathic arthritis: 1, ankylosing spondylitis: 5, psoriasis: 3, multiple indications: 4). The majority of these trials were single-arm (33/43) and enrolled adult patients (34/43). All included studies used a historical control rather than a contemporaneous ECA. In RCTs, ECAs were most often derived from the placebo arm of another RCT (6/10). In single-arm trials, historical case series were the most common ECA source (19/33). Most studies (31/43) did not employ a statistical approach to generate the ECA from historical data.

Conclusions Standardised ECA methodology and reporting conventions are lacking for IMIDs trials. The establishment of ECA reporting guidelines may enhance the rigour and transparency of future research.

BACKGROUND

Randomised controlled trials (RCTs) are the gold standard for evaluating the efficacy

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ This systematic review used a comprehensive predefined approach to identify published, interventional immune-mediated inflammatory disease studies in which external data have been used as the comparator.
- ⇒ Although we devised a sensitive search strategy comprising keywords and controlled vocabulary, the language used to describe external control arm (ECA) methodology lacks standardisation and ECA-specific MeSH terms do not exist.
- ⇒ Our assessment of ECA data source quality was conducted with an instrument that has not been validated to appraise ECA methodology, as no such instrument has been developed for use in the context of systematic reviews.

and safety of medical interventions.¹ The strengths of randomisation include minimisation of differences of known and unknown confounders among experimental groups, reduced patient selection bias and facilitation of robust statistical analyses. The randomised controlled design also permits blinding, which mitigates potential bias introduced by study personnel, participants and outcome assessors. Despite these advantages, RCTs can present practical, logistical and ethical challenges.¹ For example, it can be difficult to enrol a sufficient number of participants when the disease is rare; there are multiple competing drug development programmes, or trial avoidance where patients are concerned about being allocated to placebo. Furthermore, placebo-controlled RCTs may be unethical if there is a lack of clinical equipoise, or an existing well-established standard of care therapy available.

As noted by the US Food and Drug Administration (FDA)² and the European Medicines Agency (EMA),³ there are instances where it may be acceptable to use external data to form a control arm, instead of a concurrently



randomised internal control group. External control arms (ECAs) comprise participants who are not part of the same study as the group receiving the investigational agent.⁴ Both regulatory agencies have indicated that a well-characterised disease course, large anticipated treatment effect, sufficiently similar treatment and external populations, and the use of objective outcome measures are important factors in externally controlled designs.^{2,3} Patient data used to form the ECA can be collected during the same or similar time period (contemporaneous control) or derived from a previously treated patient population where retrospective or retrospectively analysed data are used as a comparator (historical control).²⁻⁷ These data are typically sourced from prior clinical trials, disease registries, cohort studies, electronic health records and claims databases. External data can also be used to augment the sample size of an internal control arm. The use of an ECA can reduce sample size requirements, avoid ethical pitfalls of using placebo and provides an opportunity to use historical control arm data which may be both valid and large.^{2,4-6} Yet, potential bias introduced by differences in study populations, trial design and outcome assessment along with the inability to control for unknown confounders and lack of blinding necessitates careful consideration of data source selection, analytical methods and missing data handling.

To date, ECAs have most frequently been used in clinical trials of oncology and rare diseases.⁸ Both the FDA and EMA have published guidance documents outlining recommendations for using real-world data and real-world evidence in regulatory decision-making,⁹⁻¹³ and key recommendations for the design and conduct of externally controlled clinical trials.² Study designs involving comparisons with ECAs are increasingly being submitted to regulatory agencies, and explored in novel therapeutic areas.^{14,15} Between 2005 and 2017, 43 product applications for indications in haematology and rare diseases were submitted for regulatory approval, of which 98% and 79% were approved by the FDA and EMA, respectively.¹⁴ In an analysis of 45 FDA approvals of non-oncology products that relied on external controls, 20% of approved products were for the treatment of a common disease (ie, affecting more than 200 000 patients in the USA) and 91% were recognised as addressing an unmet medical need.¹⁵ Given the willingness of regulators to consider ECA data in clinical development programmes and practical issues inherent to RCT design, there is growing interest in exploring how to best adopt this methodology within novel contexts.

Conducting RCTs for immune-mediated inflammatory diseases (IMIDs), such as inflammatory bowel disease, can be challenging due to slow and competitive patient recruitment. Contributing factors include a large clinical development pipeline, the availability of several approved treatment options in routine clinical care, and increasingly stringent eligibility criteria, particularly with respect to the use of prior and concomitant therapies.¹⁶ In addition, the chance of exposure to placebo can be

a deterrent for participation in clinical trials for both patients and investigators. The incorporation of ECA data into IMID trial design may provide an efficient alternative or supplement to a traditional randomised control arm. Currently, it is unclear how often and in what setting ECAs have been incorporated into IMID trials, and what data sources and statistical techniques have been used.

This systematic review aimed to identify and describe trials of IMIDs (ie, Crohn's disease, ulcerative colitis, chronic refractory pouchitis, rheumatoid arthritis, juvenile idiopathic arthritis, ankylosing spondylitis, psoriatic arthritis, psoriasis and atopic dermatitis) that have used an ECA, and assess the methodological quality of these studies.

METHODS

This systematic review is reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Statement¹⁷ and was conducted following an a priori developed protocol (available on request).

Search strategy

Embase, Medline and the Cochrane Central Register of Controlled Trials were searched without language restriction from database inception to 12 September 2023, using predefined strategies (online supplemental appendix 1). A semiautomated recursive search of the bibliographies of relevant publications, including review articles and meta-analyses, was also performed using the Systematic Review Accelerator SpiderCite tool (Bond University, Gold Coast, Qld, Australia).

Eligibility criteria

Eligible studies: (1) compared data from a randomised or single-arm clinical trial with a control arm from an external source (ie, previous clinical trials, disease registries, commercial insurance or national health insurance claims databases, or electronic health records); (2) enrolled adults or children with Crohn's disease, ulcerative colitis, chronic refractory pouchitis, rheumatoid arthritis/juvenile idiopathic arthritis, ankylosing spondylitis, psoriatic arthritis, psoriasis, or atopic dermatitis; and (3) assessed a medical intervention. The ECA comprised patients from placebo, standard of care, or active comparator groups. Trials were eligible for inclusion regardless of whether the ECA was prospectively or retrospectively integrated into the study design. Protocols for upcoming trials were considered, in addition to studies published in full text or abstract form. Studies were excluded if they were non-interventional or if the active arm data were not collected as part of a clinical trial. Additionally, we excluded studies that did not report the ECA dataset source.

Screening and data extraction

Search results were independently screened in duplicate by two authors (AZ and JC) and data extraction was

completed by two independent authors (AZ and RE) using a standardised spreadsheet. Data collected from each study included: the publication year, study design, trial phase, indication, study arms, number of participants, inclusion/exclusion criteria, interventions, source of ECA data, statistical methods, outcomes assessed, outcome type (dichotomous vs continuous) and rationale for using an ECA. Disagreements encountered during screening and data extraction were resolved by consensus or, if consensus could not be reached, through arbitration by a third author.

Quality assessment

The methodological quality of the ECA data source was assessed by two authors (AZ and RE) using a checklist adapted from Thorlund *et al.*¹⁸ The checklist comprised 5 key domains: (1) quality of data source; (2) similarity of the study populations; (3) similarities of the outcome definitions; (4) the reliability and comprehensiveness of the dataset; and (5) other potential limitations. Key judgement criteria are reported in online supplemental table 1. Each domain was classified as being of high (all criteria are met), low (no criteria are met) or unclear quality (some criteria are met and/or insufficient information to ascertain whether all criteria are met).

Patient involvement

No patients were involved in the study concept and design, acquisition and interpretation of data, or drafting of the manuscript.

RESULTS

Search results

A total of 2192 records were identified from database and recursive searching, from which 610 duplicates were removed. Based on the information provided in the titles and abstracts of the remaining 1582 records, 1442 were excluded as non-applicable. Full-text review was performed for 140 records, of which 97 were excluded with reasons (online supplemental table 2). Forty-three studies met the eligibility criteria and were included (figure 1).

Randomised controlled trials

Characteristics of included studies

Ten of the 43 included studies (23%) compared the active arm(s) of an RCT to an ECA (table 1).^{19–28} Five of the included RCTs (5/10; 50%) prospectively leveraged external control data as part of the study design.^{19–23} Four of these studies (4/5; 80%) randomised participants to 1 of 2 active arms and used an ECA to completely replace

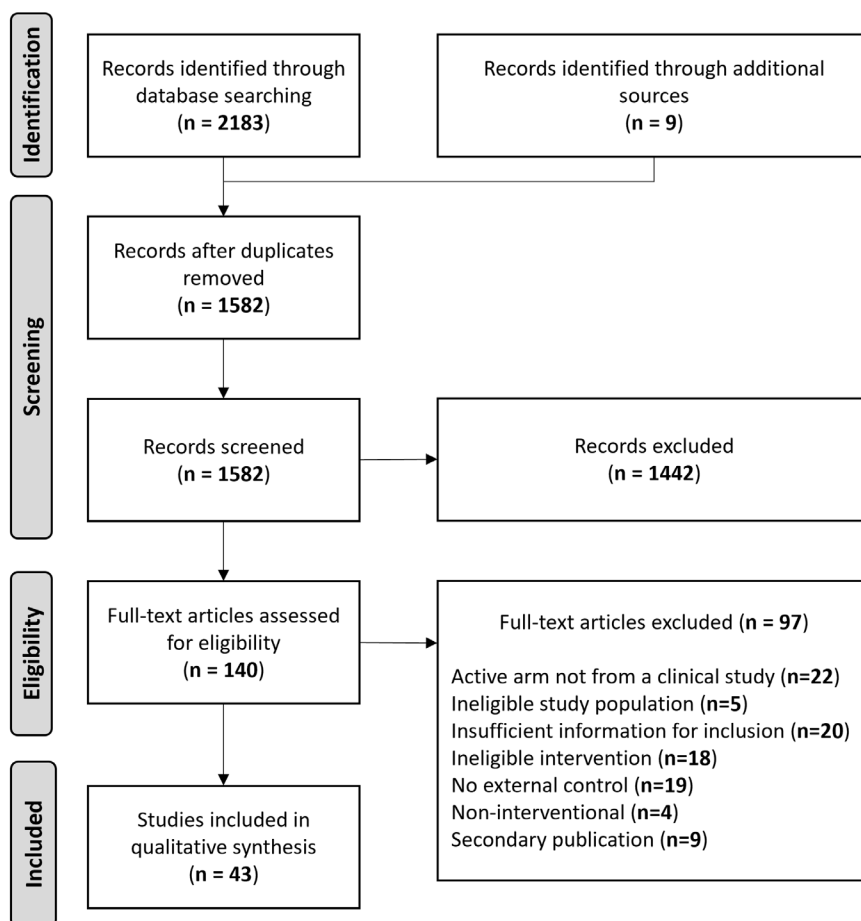


Figure 1 Study flow diagram of study inclusion. Adapted from Page *et al.*¹⁷

Table 1 Characteristics of included randomised controlled trials

| Study ID | Indication | Active arm | | | ECA | | | Control |
|--|------------|------------|-----|-------|-------------------------------|-----|---|-------------|
| | | Population | N | Phase | Intervention | N | Source | |
| Hueber <i>et al.</i> , 2012 ¹⁹ | CD | Adult | 39 | 2 | Secukinumab | 671 | 6 RCTs (phase 2–3; published) | Placebo |
| Hyams <i>et al.</i> , 2012 ²⁰ | UC | Paediatric | 60 | 3 | Infliximab | 244 | RCT (phase 3; published) | Placebo |
| Hodkinson <i>et al.</i> , 2013 ²¹ | RA | Adult | 87 | NR | DMARDs+tight control strategy | 145 | Historical case series* (unpublished) | DMARDs |
| Croft <i>et al.</i> , 2021 ²² | UC | Paediatric | 93 | 3 | Adalimumab | NR | 4 RCTs (phase 3; published) | Placebo |
| Hupf <i>et al.</i> , 2021† ²⁵ | AS | Adult | 23 | 2 | Secukinumab | 513 | 8 RCTs (phase 2–3; published) | Placebo |
| Reed <i>et al.</i> , 2021† ²⁴ | RA | Adult | NR | 1–3 | Tofacitinib | NR | Disease registry (published) | DMARDs |
| Rudrapatna <i>et al.</i> , 2021† ²⁶ | CD | Adult | 73 | 2b | Tesnatilimab | 56 | EHR data (unpublished) | Ustekinumab |
| Wang <i>et al.</i> , 2021† ²⁷ | RA | Adult | 54 | 2 | Upadacitinib | 159 | RCT (phase 3; published) | Placebo |
| Magnolo <i>et al.</i> , 2022 ²³ | PsO | Paediatric | 84 | 3 | Secukinumab | 812 | 5 RCTs (phase 3; published) | Placebo |
| Norvang <i>et al.</i> , 2022† ²⁸ | RA | Adult | 188 | NR | DMARDs+stringent T2T strategy | 328 | Prospective observational study (published) | DMARDs |

*Historical control comprised of patients who have previously received treatment at the same site(s) as active arm study participants.

†Primary study objective was to examine methodological approaches for complementing prior or concurrent RCT data with external control data.

AS, ankylosing spondylitis; CD, Crohn's disease; DMARD, disease-modifying antirheumatic drug; ECA, external control arm; EHR, electronic health record; NR, not reported; PsO, psoriasis; RA, rheumatoid arthritis; RCT, randomised controlled trial; T2T, treat-to-target; UC, ulcerative colitis.

the function of a placebo arm, while 1 (1/5; 20%) was a placebo-controlled RCT that incorporated external data to augment the sample size of the placebo group. Among the five RCTs that prospectively used external data were three phase 3^{20 22 23} (60%) and one phase 2¹⁹ (20%) trials. The remaining five RCTs (5/10; 50%) retrospectively compared data from the active arm(s) of prior (4/5; 80%) or ongoing (1/5; 20%) phase 1–3 trials with an ECA.^{24–28}

Most of the included RCTs enrolled adults (7/10; 70%), with 3 of the 10 trials (30%) conducted in paediatric populations. Study populations comprised patients with inflammatory bowel disease (4/10; 40% (ulcerative colitis: 2/10; 20%, Crohn's disease: 2/10; 20%)), rheumatoid arthritis (4/10; 40%), psoriasis (1/10; 10%), and ankylosing spondylitis (1/10; 10%). Among these studies, biologic therapies (6/10; 60%) were the most common intervention under investigation, including secukinumab (3/10; 30%), infliximab (1/10; 10%), adalimumab (1/10; 10%) and tesnatilimab (1/10; 10%). Two out of 10 studies (20%) evaluated small molecules (tofacitinib (1/10; 10%) and upadacitinib (1/10; 10%)). Treatment strategies were assessed in two RCTs (tight control strategy (1/10; 10%) and stringent treat-to-target strategy (1/10; 10%)).

The majority of studies (6/10; 60%) made comparisons with placebo data derived from 1 or more previously conducted RCTs. An external active control group was used by 2 out of 10 studies (20%), with data sourced from a disease registry (1/10; 10%) or electronic health records (1/10; 10%). Lastly, 2 studies (2/10; 20%) formed a standard of care control group using data from a historical case series (1/10; 10%) or a prior prospective cohort study (1/10; 10%).

External control methodology

All 10 studies used a historical ECA; we did not identify any studies that employed contemporaneously collected external data. Various methods were used to generate ECAs among the 10 studies that compared the active arm(s) of an RCT to external control data (table 2). Statistical methods used to adjust the external dataset included propensity score matching (2/10; 20%), propensity score weighting (1/10; 10%) and Bayesian dynamic borrowing (2/10; 20%). One study did not specify the weighting method employed (1/10; 10%). Four studies (4/10; 40%) did not make any statistical adjustments to ECA data to balance characteristics between active and external arms. Nine out of 10 studies (90%) made comparisons between the active arm and ECA for at least one efficacy outcome. In all cases, efficacy outcomes were defined by an indication-specific disease activity index and were assessed as dichotomous (9/9; 100%) or continuous (1/9; 11.1%) endpoints. Two studies (2/10; 20%) used external data to contextualise safety outcomes (ie, adverse events and serious adverse events).

Table 2 External control arm methodology in randomised controlled trials

| Study ID | Indication | Statistical methods for generating ECA | Outcome(s) compared with ECA | Outcome timepoint (weeks) | Evidence of treatment benefit compared with ECA* | Rationale for using an ECA |
|---|------------|--|---|---------------------------|--|--|
| Hueber <i>et al</i> , 2012 ¹⁹ | CD | Weighting | Efficacy (CDAI response rate) | 6 | No | Small study sample size |
| Hyams <i>et al</i> , 2012 ²⁰ | UC | Unadjusted | Efficacy (MCS response rate) | 8 | Yes | Unethical to use internal placebo arm |
| Hodkinson <i>et al</i> , 2013 ²¹ | RA | Unadjusted | Efficacy (DAS 28 response and remission rates) | 26 | Yes | NR |
| Croft <i>et al</i> , 2021 ²² | UC | Unadjusted | Efficacy (MCS remission rate) | 8, 52 | Yes | Unethical to use internal placebo arm advised by regulatory organisations; low recruitment |
| Hupf <i>et al</i> , 2021† ²⁵ | AS | Bayesian dynamic borrowing | Efficacy (ASAS 20 response rate) | 6 | NA | ECA methodology study |
| Reed <i>et al</i> , 2021† ²⁴ | RA | Propensity score matching | Safety (SAEs) | NR | NA | ECA methodology study |
| Rudrapatna <i>et al</i> , 2021† ²⁶ | CD | Unadjusted | Efficacy (mean change in CDAI, steroid-free CDAI remission rate) | 12, 24 | NA | ECA methodology study |
| Wang <i>et al</i> , 2021† ²⁷ | RA | Propensity score matching | Efficacy (ACR 20 response rate) | 12 | NA | ECA methodology study |
| Magnolo <i>et al</i> , 2022 ²³ | PsO | Bayesian dynamic borrowing | Efficacy (PASI 75, PASI 90 & IGA 0/1 response rates) Safety (AEs) | 12 | Yes | Unethical to use internal placebo arm; advised by regulatory organisations |
| Norvang <i>et al</i> , 2022† ²⁸ | RA | Propensity score weighting | Efficacy (DAS 28 remission rate) | 24, 52, 104 | NA | ECA methodology study |

*As defined by the included study.

†Primary study objective was to examine methodological approaches for complementing prior or concurrent RCT data with external control data.

ACR, American College of Rheumatology; AE, adverse event; AS, ankylosing spondylitis; ASAS, Assessment of SpondyloArthritis International Society; CD, Crohn's disease; CDAI, Crohn's Disease Activity Index; DAS, Disease Activity Score; ECA, external control arm; IGA, Investigator's Global Assessment; MCS, Mayo Clinical Score; NA, not applicable; NR, not reported; PASI, Psoriasis Area Severity Index; PsO, psoriasis; RA, rheumatoid arthritis; SAE, serious adverse event; UC, ulcerative colitis.

Indications for use of an ECA

Three of the included RCTs (3/10; 30%) reported that comparisons to external data were made due to ethical issues associated with randomising participants to a placebo arm (table 2).^{20 22 23} All were phase 3 trials in paediatric populations, with 2 of these studies (2/3; 66.7%) adopting a study design that included an ECA following consultations with regulatory agencies.^{22 23} One study (1/10; 10%) additionally cited poor study recruitment due to non-acceptance of an internal control in the paediatric setting as a reason for incorporating external data.^{22 23} Augmenting the small sample size of the internal placebo group was provided as the rationale for introducing external placebo data into the primary analysis in 1 of the included RCTs (1/10; 10%).¹⁹

Single-arm studies

Characteristics of included studies

The majority of the included studies (33/43; 77%) were single-arm trials that relied on external data to provide context to treatment outcomes (table 3).^{29–61} Among these were one phase 2a trial (1/33; 3%), one phase 2b trial (1/33; 3%), one phase 2–3 trial (1/33; 3%), four phase 3 long-term RCT extensions (4/33; 12%) and 4 post-marketing surveillance studies (4/33; 12%).

Most of the included single-arm trials enrolled adult participants (27/33; 81%), with some studies conducted in paediatric (4/33; 12%) and mixed-age (2/33; 6%) populations. Therapies for the treatment of inflammatory bowel disease (13/33; 40% (ulcerative colitis: 5/33; 15%, Crohn's disease: 1/33; 3%)), rheumatoid arthritis

Table 3 Characteristics of included single-arm studies

| Study ID | Indication | | Population | | Active arm | | ECA | | Source | Control |
|--|------------|----------------------|------------|-------------------------------------|------------|--|--|--|--------|---------|
| | | | | | N | Intervention | N | | | |
| Baker and Jewell, 1989 ²⁹ | IBD | Adult and paediatric | 24 | SOC+ciclosporin | 406 | Multiple* (published) | Multiple* (published) | SOC | | |
| Rosenberg <i>et al</i> , 1990 ³⁰ | UC | Adult | 20 | Methylprednisolone (high dose) | 175 | Multiple† (published) | Multiple† (published) | Prednisolone or methylprednisolone (standard dose) | | |
| Stenger <i>et al</i> , 1998 ³¹ | RA | Adult | 139 | Sulfasalazine+methotrexate | 89 | Historical case series† (unpublished) | Historical case series† (unpublished) | NSAIDs | | |
| Marotte <i>et al</i> , 2007 ³² | RA | Adult | 90 | Infliximab | 99 | Historical case series† (unpublished) | Historical case series† (unpublished) | Methotrexate | | |
| Shen <i>et al</i> , 2007 ³⁴ | Pouchitis | Adult | 16 | Ciprofloxacin+tindazole | 10 | two historical case series (published) | two historical case series (published) | Mesalamine | | |
| Moul <i>et al</i> , 2007 ³³ | PsO | Adult | 16 | Alefacept (high dose) | 166 | RCT (phase 3; published) | RCT (phase 3; published) | Alefacept (standard dose) | | |
| van der Heijde <i>et al</i> , 2008 ³⁶ | AS | Adult | 201 | Infliximab | 192 | Disease registry (unpublished) | Disease registry (unpublished) | SOC (anti-TNF naïve) | | |
| Boyle <i>et al</i> , 2008 ³⁵ | IBD | Adult | 14 | Laparoscopic resection | 16 | Historical case series† (unpublished) | Historical case series† (unpublished) | Open colon resection | | |
| van der Heijde <i>et al</i> , 2009 ³⁷ | AS | Adult | 307 | Adalimumab | 169 | Disease registry (unpublished) | Disease registry (unpublished) | SOC (anti-TNF naïve) | | |
| Garg <i>et al</i> , 2011 ³⁸ | IBD | Adult | 100 | Rapid iron polymaltose infusion | 486 | two historical case series (published) | two historical case series (published) | Slow iron polymaltose infusion | | |
| Yong <i>et al</i> , 2011† ³⁹ | PsO | Adult | NR | Etanercept | NR | Administrative claims database (unpublished) | Administrative claims database (unpublished) | SOC (non-biologics) | | |
| Carter <i>et al</i> , 2012*** ⁴⁰ | RA | Adult | 50 | Rituximab+prednisone | NR | RCTs (package insert; published) | RCTs (package insert; published) | Rituximab+methylprednisolone | | |
| Kwok and Leung, 2012 ⁴¹ | RA | Adult | 20 | SOC+T2T strategy | 20 | Historical case series† (unpublished) | Historical case series† (unpublished) | SOC | | |
| Baraliakos <i>et al</i> , 2014 ⁴³ | AS | Adult | 22 | Infliximab | 34 | Historical case series† (unpublished) | Historical case series† (unpublished) | SOC (anti-TNF naïve) | | |
| Horneff <i>et al</i> , 2014†† ⁴⁴ | Multiple†† | Paediatric | 127 | Etanercept | 323 | 7 RCTs (published) | 7 RCTs (published) | Placebo | | |
| Lichtenstein <i>et al</i> , 2014§ ⁴⁵ | UC | Adult | 388 | Mesalamine | 83 | RCTs (phase 3+package insert; published) | RCTs (phase 3+package insert; published) | Placebo | | |
| Gardenbroek <i>et al</i> , 2015 ⁴² | UC | Adult | 15 | Early surgical closure | 29 | Historical case series† (unpublished) | Historical case series† (unpublished) | SOC | | |
| Mahajan <i>et al</i> , 2018 ⁴⁸ | UC | Adult | 41 | Faecal microbiota transplantation | 38 | Historical case series† (unpublished) | Historical case series† (unpublished) | Azathioprine | | |
| Rozette <i>et al</i> , 2018 ⁴⁹ | Multiple§§ | Adult and paediatric | 66 | Infliximab (rapid infusion) | 50 | Historical case series† (unpublished) | Historical case series† (unpublished) | Infliximab (standard infusion) | | |
| Tweehuysen <i>et al</i> , 2018 ⁵⁰ | Multiple¶¶ | Adult | 625 | Etanercept biosimilar (SB4) | 600 | Historical case series† (unpublished) | Historical case series† (unpublished) | Etanercept | | |
| Horneff <i>et al</i> , 2018**** ⁴⁶ | JIA | Paediatric | 18 | Golimumab | 3620 | Historical case series (unpublished) | Historical case series (unpublished) | Anti-TNF or methotrexate | | |
| Inui <i>et al</i> , 2018 ⁴⁷ | RA | Adult | 13 | Etanercept+on-demand dosing regimen | 16 | Historical case series† (unpublished) | Historical case series† (unpublished) | Etanercept | | |

Continued

Table 3 Continued

| Study ID | Indication | | Population | | Active arm | | ECA | | Source | Control |
|---|-------------|--------------|------------|--|------------|--|--------|--|---|---------|
| | N | Intervention | N | | N | | N | | | |
| Braun <i>et al</i> , 2019 ^{§51} | AS | Adult | 168 | Secukinumab | 69 | RCT (phase 4; published) | 69 | RCT (phase 4; published) | NSAIDs | |
| Kawashiri <i>et al</i> , 2020 ^{***†††52} | RA | Adult | NR | Infliximab biosimilar (CT-P13) | NR | 2 cohort studies (published) | NR | 2 cohort studies (published) | SOC (anti-TNF) | |
| Muskens <i>et al</i> , 2020 ^{§3} | Multiple††† | Adult | 84 | Etanercept biosimilar | 89 | Historical case series† | 89 | Historical case series† (unpublished) | Etanercept | |
| Sanchez-Hernandez <i>et al</i> , 2020 ⁵⁴ | IBD | Adult | 81 | Infliximab+proactive TDM | 76 | Historical case series† | 76 | Historical case series† (unpublished) | Infliximab | |
| Sagami <i>et al</i> , 2021 ^{***57} | IBD | Adult | 700 | CT-P13 | 44 274 | Administrative claims database (unpublished) | 44 274 | Administrative claims database (unpublished) | Infliximab | |
| Gasparetto <i>et al</i> , 2021 ⁵⁶ | IBD | Paediatric | 30 | Infliximab+proactive TDM | 37 | Historical case series† | 37 | Historical case series† (unpublished) | Infliximab | |
| Danese <i>et al</i> , 2021 ⁵⁵ | UC | Adult | 50 | RVT-3101 | 234 | 2 RCTs (phase 3; published) | 234 | 2 RCTs (phase 3; published) | Placebo | |
| Holland <i>et al</i> , 2022 ⁵⁹ | CD | Paediatric | 37 | Infliximab+proactive TDM | 37 | Historical case series† | 37 | Historical case series† (unpublished) | Infliximab | |
| Huang <i>et al</i> , 2022 ⁶⁰ | RA | Adult | 111 | Multiple†††+tight control T2T strategy | 389 | Historical case series† | 389 | Historical case series† (unpublished) | Multiple†††+T2T strategy | |
| Truong <i>et al</i> , 2022 ⁵⁸ | IBD | Adult | 65 | Transanal approach restorative proctocolectomy | 100 | Historical case series† | 100 | Historical case series† (unpublished) | Open approach restorative proctocolectomy | |
| Takei <i>et al</i> , 2022 ⁶¹ | RA | Adult | 60 | Hydroxychloroquine+DMARDs | 276 | Cohort study (published) | 276 | Cohort study (published) | SOC (DMARDs) | |

*Control data sourced from two cohort studies, one historical case series and one RCT.
†Control data sourced from two historical case series and one RCT.
‡Historical control comprised patients who have previously received treatment at the same site(s) as active arm study participants.
§Phase 3 RCT extension.
¶Primary study objective was to examine methodological approaches for complementing prior or concurrent RCT data with external control data.
**Phase 2/3 trial.
††Phase 3b trial.
‡‡JIA, enthesitis-related arthritis, and psoriatic arthritis.
§§CD, UC, and JIA.
¶¶RA, psoriatic arthritis, and AS.
***Post-marketing surveillance study.
†††Methotrexate, leflunomide, or a combination of DMARD and corticosteroid therapy.
‡‡‡Protocol for upcoming single-arm study.
AS, ankylosing spondylitis; CD, Crohn's disease; DMARD, disease-modifying antirheumatic drug; ECA, external control arm; IBD, inflammatory bowel disease; JIA, juvenile idiopathic arthritis; NR, not reported; NSAID, non-steroidal anti-inflammatory drug; Pso, psoriasis; RA, rheumatoid arthritis; RCT, randomised controlled trial; SOC, standard of care; T2T, treat-to-target; TDM, therapeutic drug monitoring; TNF, tumour necrosis factor; UC, ulcerative colitis.

and juvenile idiopathic arthritis (9/33; 27%), ankylosing spondylitis (4/33; 12%), psoriasis (2/33; 6%), and chronic refractory pouchitis (1/33; 3%) were evaluated. Four of the 33 single-arm trials (12%) administered treatment to a study population comprising participants with differing IMIDs. Pharmacological interventions were most frequently assessed (21/33; 64%). These included biologics (9/33; 31%), biosimilars (4/33; 12%), corticosteroids (2/33; 6%), immunosuppressives (3/33; 9%), antibiotics (1/33; 3%) and 5-aminosalicylic acid (1/33; 3%). The remaining 12 single-arm trials investigated a dosing strategy (4/33; 12%), treatment algorithm (2/33; 6%), surgical procedure (3/33; 9%), rapid infusion protocol (2/33; 6%), or faecal microbiota transplantation (1/33; 3%).

Unpublished historical case series of patients previously treated at the same site(s) as active arm participants was the most common ECA source (17/33; 52%) (table 3). External data were also derived from 1 or more prior phase 3–4 RCTs (6/33; 18%), product package inserts (2/33; 6%), published case series (2/33; 6%), disease registries (2/33; 6%), administrative claims databases (2/33; 6%), and previously conducted cohort studies (2/33; 6%). Two studies (2/33; 6%) pooled participant data from several different types of sources. The ECAs formed in single-arm trials comprised participants who had received standard of care (20/33; 61%), an active control therapy (10/33; 30%), or placebo (3/33; 9%).

External control methodology

All 33 included single-arm trials used a historical control, rather than a contemporaneous ECA. In most studies, external datasets were not statistically adjusted for known covariates to improve the comparability of external and target study populations (26/33; 79%) (table 4). Five of the 33 studies applied a matching methodology (15%), including propensity score matching (2/33; 6%), and 1 study employed a propensity score weighting (3%) approach to improve the comparability of active and external groups. Twenty-four of the included single-arm trials (73%) evaluated one or more efficacy outcomes. These studies assessed dichotomous (20/33; 60%) and continuous (7/33; 21%) efficacy endpoints based on an indication-specific disease activity index (18/33; 54.5%). Safety outcomes were compared with external data in 15 studies (45.4%). These included the incidence of adverse events (10/33; 30%), adverse infusion reactions (3/33; 9%), postsurgical complications (1/33; 3%) and serious adverse events (1/33; 3%).

Indication for use of an ECA

In 9 out of 33 single-arm trials (27%),^{32 36 37 43–46 51 52} it was considered unethical to randomise participants to an internal control arm due to the availability of approved therapies with established efficacy and safety (7/33; 21%), long-term assessment of treatment outcomes (6/33; 18%), or paediatric study population (2/33; 6%) (table 4). Six trials (6/33; 18%)^{33 35 40 42 55 57} cited

practical challenges as the rationale for using an ECA. In five of these cases (5/33; 15%), an adequately powered RCT would reportedly not have been feasible due to the relatively small patient population available for study participation.

Quality assessment

The results of the methodological quality assessment are reported in online supplemental file 1. Of the 210 (43 studies × 5 domains) ratings, 99 (46%) were high quality, 113 (52%) were unclear quality and 3 (1%) were low quality with respect to the source used to generate the ECA. The domains that were rated for each study included the quality of the data source (40% high quality, 60% unclear quality), similarity of study populations (43% high quality, 57% unclear quality), similarity of outcome definitions (72% high quality, 21% unclear quality, 7% low quality), and reliability and comprehensiveness of the external dataset (21% high quality, 79% unclear quality).

DISCUSSION

In recent years, there has been increased interest among researchers, regulators and industry partners in using external control data to address the practical limitations associated with traditional RCT designs. This systematic review aimed to summarise how ECAs are currently being integrated into studies of Crohn's disease, ulcerative colitis, chronic refractory pouchitis, rheumatoid arthritis, juvenile idiopathic arthritis, ankylosing spondylitis, psoriatic arthritis, psoriasis and atopic dermatitis, and examine the methodological quality of this research. We identified 43 trials in which an ECA was used to contextualise the efficacy or safety of a medical therapy. No trials of atopic dermatitis were identified. Most of the included clinical trials were single-arm studies, and all used historical control data to form an ECA. Among included RCTs, external data were most often sourced data from multiple previously conducted RCTs. Conversely, external controls comprised participants who previously received treatment at the same site as the active arm population was the most frequent source of the ECA when the clinical trial was a single-arm design. In most cases, no adjustment methods were used to balance baseline characteristics between the treatment arm and historical data used to form the ECA. We did not identify any specific characteristics of data sources, outcome measure selection, analysis method or ECA type that were associated with a positive or negative study result.

All primary phase 3 trials of IMIDs identified in this systematic review that were submitted to or relied on by regulatory agencies for decision-making were in the paediatric setting (4/43; 9%), where an internal placebo group may have been deemed unethical by stakeholders. In a phase 3 trial that supported FDA and EMA approval of infliximab in paediatric ulcerative colitis, Hyams *et al*²⁰ created a historical control group using pooled placebo data from prior infliximab RCTs in adults. A positive

Table 4 External control arm methodology in single-arm studies

| Study ID | Indication | Statistical methods for generating ECA | Outcome(s) compared with ECA | Outcome timepoint (weeks) | Evidence of treatment benefit compared with ECA* | Rationale for using an ECA |
|--|------------|--|--|---------------------------|--|---------------------------------------|
| Baker and Jewell, 1989 ²⁹ | IBD | Unadjusted | Efficacy (severe attack rate) | 6 | No | Well-characterised disease course |
| Rosenberg <i>et al</i> , 1990 ³⁰ | UC | Unadjusted | Efficacy (response and surgical rates) | >1 | No | NR |
| Stenger <i>et al</i> , 1998 ³¹ | RA | Unadjusted | Efficacy (cumulative CRP-AUC, median change in Sharp-score) Safety (AEs) | 104 | Yes | NR |
| Marotte <i>et al</i> , 2007 ³² | RA | Matching | Efficacy (DAS28 response rate) | 52 | Yes | Unethical to use internal placebo arm |
| Shen <i>et al</i> , 2007 ³⁴ | Pouchitis | Unadjusted | Efficacy (PDAI clinical remission and response rates) Safety (AEs) | 4 | Yes | NR |
| Moul <i>et al</i> , 2007 ³³ | PsO | Unadjusted | Efficacy (PASI 50 and PASI 75 response rates) | 12 | No | Small study sample size |
| van der Heijde <i>et al</i> , 2008 ³⁶ | AS | Matching | Efficacy (median change in mSASSS) | 104 | No | Unethical to use internal placebo arm |
| Boyle <i>et al</i> , 2008 ³⁵ | IBD | Matching | Safety (LOS, postoperative hospitalizations) | LOS | Yes | Impractical to use internal placebo |
| van der Heijde <i>et al</i> , 2009 ³⁷ | AS | Unadjusted | Efficacy (median change in mSASSS) | 104 | No | Unethical to use internal placebo arm |
| Garg <i>et al</i> , 2011 ³⁸ | IBD | Unadjusted | Safety (AEs) | >1 | NA | NR |
| Yong <i>et al</i> , 2011† ³⁹ | PsO | NR | Safety (SAEs) | NR | NA | ECA methodology study |
| Carter <i>et al</i> , 2012 ⁴⁰ | RA | Unadjusted | Safety (acute infusion reactions) | >1 | Yes | Small study sample size |
| Kwok and Leung, 2012 ⁴¹ | RA | Unadjusted | Efficacy (DAS20 and ACR response rate) | 52 | Yes | NR |
| Baraliakos <i>et al</i> , 2014 ⁴³ | AS | Unadjusted | Efficacy (mean change in mSASSS) | 8 years | No | Unethical to use internal placebo arm |
| Horneff <i>et al</i> , 2014 ⁴⁴ | Multiple‡ | Unadjusted | Efficacy (JIA ACR 30 response rate) | 12 | Yes | Unethical to use internal placebo arm |
| Lichtenstein <i>et al</i> , 2014 ⁴⁵ | UC | Unadjusted | Safety (AEs) | 104 | Yes | Unethical to use internal placebo arm |
| Gardenbroek <i>et al</i> , 2015 ⁴² | UC | Unadjusted | Efficacy (anastomotic healing, closure of anastomotic defect and function pouch rates) | 24 | Yes | Impractical to use internal placebo |
| Mahajan <i>et al</i> , 2018 ⁴⁸ | UC | Unadjusted | Efficacy (clinical, endoscopic, and steroid-free clinical remission rates) Safety (AEs) | 22 | Yes | NR |
| Rozette <i>et al</i> , 2018 ⁴⁹ | Multiple § | Unadjusted | Safety (adverse infusion reactions) | > 1 | NA | NR |
| Tweehuysen <i>et al</i> , 2018 ⁵⁰ | Multiple¶ | Unadjusted | Efficacy (CRP and DAS28-CRP response rates) Safety (AEs) | 24 | No | NR |

Continued



Table 4 Continued

| Study ID | Indication | Statistical methods for generating ECA | Outcome(s) compared with ECA | Outcome timepoint (weeks) | Evidence of treatment benefit compared with ECA* | Rationale for using an ECA |
|---|------------|--|--|---------------------------|--|--|
| Horneff <i>et al</i> , 2018 ⁴⁶ | JIA | Unadjusted | Efficacy (JADAS10 response rate) Safety (AEs) | NR | NA | Unethical to use internal placebo arm |
| Inui <i>et al</i> , 2018 ⁴⁷ | RA | Unadjusted | Efficacy (DAS28-ESR and Sharp score response rates) | 52, 104 | No | NR |
| Braun <i>et al</i> , 2019 ⁵¹ | AS | Unadjusted | Efficacy (mean change in mSASSS) | 104 | No | Unethical to use internal placebo arm |
| Kawashiri <i>et al</i> , 2020 ^{**52} | RA | Unadjusted | Efficacy (DAS28-ESR clinical relapse rate) | 24 | NA | HESDE previously established |
| Muskens <i>et al</i> , 2020 ⁵³ | Multiple†‡ | Unadjusted | Safety (AEs) | 52 | NA | NR |
| Sanchez-Hernandez <i>et al</i> , 2020 ⁵⁴ | IBD | Unadjusted | Efficacy (treatment failure and IBD-related surgery and hospitalisation rates) Safety (AEs, acute infusion reactions) | 156 | Yes | NR |
| Sagami <i>et al</i> , 2021 ⁵⁷ | IBD | Unadjusted | Safety (AEs) | 104 | NA | Impractical to use internal placebo arm |
| Gasparetto <i>et al</i> , 2021 ⁵⁶ | IBD | Unadjusted | Efficacy (dose escalation rate) | 52 | Yes | NR |
| Danese <i>et al</i> , 2021 ⁵⁵ | UC | Propensity score weighting | Efficacy (Mayo endoscopic score) | 14 | Yes | Impractical to use internal placebo arm; low recruitment |
| Holland <i>et al</i> , 2022 ⁵⁹ | CD | Matching | Efficacy (mPCDAI response rate) | 26, 52, 78 | No | NR |
| Huang <i>et al</i> , 2022 ⁶⁰ | RA | Propensity score matching | Efficacy (SDAI, RA-CDAI and DAS28 response rates) | 12, 26, 52, 104 | Yes | NR |
| Truong <i>et al</i> , 2022 ⁵⁸ | IBD | Unadjusted | Safety (postoperative complications) | 4 | No | Lack of large single-institution studies |
| Takei <i>et al</i> , 2022 ⁶¹ | RA | Propensity score matching | Efficacy (ACR20, DAS28-ESR, SDAI response rates) | 24 | Yes | NR |

*As defined by the included study.

†Primary study objective was to examine methodological approaches for complementing prior or concurrent RCT data with external control data.

‡Juvenile idiopathic arthritis, enthesitis-related arthritis and psoriatic arthritis.

§Crohn's disease, ulcerative colitis and juvenile idiopathic arthritis

¶Rheumatoid arthritis, psoriatic arthritis and ankylosing spondylitis.

**Protocol for an upcoming single-arm study with an ECA.

ACR, American College of Rheumatology; AE, adverse event; AS, ankylosing spondylitis; AUC, area under the curve; CD, Crohn's disease; CRP, C reactive protein; DAS, disease activity score; ECA, external control arm; ESR, erythrocyte sedimentation rate; HESDE, historical evidence of sensitivity to drug effects; IBD, inflammatory bowel disease; JADAS, Juvenile Arthritis Disease Activity Score; JIA, juvenile idiopathic arthritis; LOS, length of stay; mPCDAI, modified Paediatric Crohn's Disease Activity Index; mSASSS, modified Stoke Ankylosing Spondylitis Spinal Score; NA, not applicable; NR, not reported; PASI, Psoriasis Area Severity Index; PDAI, Pouchitis Disease Activity Index; PsO, psoriasis; RA, rheumatoid arthritis; RA-CDAI, Rheumatoid Arthritis Clinical Disease Activity Index; SAE, serious adverse event; SDAI, Simplified Disease Activity Index; UC, ulcerative colitis.

result was determined by using the upper limit of the 95% CI for the primary endpoint in the adult trials to define the lower 95% CI threshold in the paediatric trial. Croft *et al*²² assessed the efficacy and safety of adalimumab in a

phase 3 study of paediatric ulcerative colitis. This study encountered recruitment delays due to poor acceptance of the placebo group and, following consultation with the FDA and EMA, a meta-analysis of historical adult placebo

groups was used to replace the internal control. Adalimumab subsequently received regulatory approval in this patient population.²² In indications where historical paediatric data may be limited, leveraging prior adult placebo data may be acceptable and advantageous to shorten the time between regulatory approval of advanced therapies in adult and paediatric patients.

In a phase 3 study that supported FDA and EMA approval of secukinumab in paediatric psoriasis, Magnolo *et al*²³ adopted a Bayesian dynamic borrowing approach using historical adult and paediatric placebo data from previous RCTs. Low-dose and high-dose secukinumab were found to be superior to historical placebo with respect to the coprimary and key secondary outcomes, with an estimated probability of a positive treatment effect of 100% compared with external placebo for both dosing regimens. Finally, Horneff *et al*⁴⁴ conducted a single-arm phase 3b study to expand EMA approval to three under-studied categories of juvenile idiopathic arthritis (extended oligoarticular juvenile idiopathic arthritis, enthesitis-related arthritis, and psoriatic arthritis). Given the paediatric population, and well-established efficacy and safety of etanercept in polyarticular-course juvenile idiopathic arthritis, it was deemed unethical to randomise participants to receive placebo. Although etanercept response rates were statistically superior compared with the ECA, external placebo data were primarily sourced from a meta-analysis of paediatric RCTs that had been published nearly a decade prior.

If the historical ECA and target study sample substantially differ due to population drift, changes in eligibility criteria, or evolving standards of care, treatment effects may be misestimated. The primary method for mitigating estimation bias is careful selection of the studies from which the historical ECA is drawn. Statistical methods can also be used to adjust for imbalances between historical and target study samples. Propensity score matching involves performing regression on a set of prespecified baseline covariates to determine a propensity score for each participant. This enables the creation of matched sets of historical and target study participants with similar propensity score values.^{55 62} However, propensity score matching and other methods of balancing that use prespecified covariates have important limitations. First, confounders must be correctly identified and accurately measured. When matching is based on *observed* covariates, the historical ECA may not be representative of the parent population and therefore not suitable for comparison with the target study.⁶² Second, propensity score matching may result in 'pruning', where individuals with extreme covariate values have no suitable matches and are systematically excluded from the ECA, which can increase imbalance and bias.⁶³ With Bayesian dynamic borrowing, data are weighted based on the level of discrepancy between the historical control and target study.^{23 64} When correctly performed, dynamic borrowing can provide additional information and augment sample size with a sufficient degree of scepticism to reduce bias.

As with propensity score matching, dynamic borrowing requires correct specification of potential confounders to determine whether the ECA and target study data are closely matched. Evidence suggests that dynamic borrowing may carry an inflated chance of type I error; however, the benefits of additional power and precision may outweigh this risk.⁶⁵

When evaluating the data collection process, similarity of the populations, similarity of the outcome definitions, the reliability and comprehensiveness of the datasets, and other limitations, only 3 of the 215 items (1%) assessed were rated as low quality. However, half of the items (113/215; 52.5%) could not be assigned a definitive low-quality or high-quality rating due to insufficient detail reported in the study publication. The included studies frequently failed to describe in detail the eligibility criteria and baseline characteristics of the active arm and ECA, outcome definitions, covariates, whether there were missing data, and the analytical methods employed to balance the cohort.

Comprehensive reporting is therefore necessary to critically appraise the quality of ECA studies, particularly when real-world data are used. This is underscored by the external control quality checklist proposed by Thorlund *et al*¹⁸ and in draft guidance by the FDA and EMA which state that adequate documentation during data curation and transformation is essential to increase confidence in the resultant data.^{11 13} The FDA has additionally issued draft guidance outlining key considerations for designing externally-controlled clinical trials, selecting fit-for-use ECA data sources, analysing comparisons and accounting for potential bias.² While this document does not recommend specific statistical methodologies for making comparisons to external data, it is noted that underlying assumptions should be identified and further examined using sensitivity analyses and model diagnostics. The need to assess the impact of missing and misclassified data in the ECA, which may be more common when using real-world data sources, is emphasised.⁶⁶

Moreover, objective and reliable measurements for the data of interest are recommended to mitigate bias due to misclassified data and lack of blinding. Within the context of IMIDs, this may involve restricting outcome measures to the most objective item(s) included in composite disease active indices. For example, centrally read endoscopic improvement, as defined by the Mayo Endoscopic subscore, was selected as the primary outcome in a statistically significant study by Danese *et al*⁵⁵ evaluating a novel biologic in a phase 2 ulcerative colitis trial. Trial patients were matched with historical placebo patients from prior RCTs using propensity score analysis, and placebo endoscopic response rates were used to determine the null hypothesis.

Regulatory submissions are evaluated on a case-by-case basis, and approvals are often conditional on additional evidence from postapproval studies.¹⁴ Our results suggest that comprehensive ECA guidelines may be particularly beneficial for IMID research. The patient populations,

standards of care, outcome definitions and study time-points have substantially evolved over time in IMIDs, and all studies identified in the current systematic review included historical controls. Ensuring that appropriate source data and optimal statistical methodologies are selected may therefore be of particular importance within this context.

Our study has some important strengths. We examined how external controls have been implemented in IMID trials using a predefined systematic approach. To our knowledge, this is the first systematic review on the use of ECAs in these populations. This qualitative summary may help inform ECA data source selection in future externally controlled trials of IMIDs and help in trial design where a treatment arm might be useful in the future as an ECA. However, the limitations of this study should also be acknowledged. First, while we aimed to devise a highly sensitive search strategy using text words and controlled vocabulary, the language used to describe ECA methodology is varied, and ECA-specific MeSH terms do not exist. In an effort to ensure that all eligible studies were identified, bibliographies of relevant review articles were also searched. Nine of the 2192 records screened were identified using this approach. Furthermore, only published studies were searched and included in this systematic review. Second, the quality of included studies was rated using an instrument that was not developed and validated to appraise ECA methodology, as no such instrument currently exists. Our quality assessment exclusively focused on the source of the external control data. We did not evaluate the methods used to generate external controls given that trials were eligible for inclusion regardless of whether they used trial-level or patient-level data.

In conclusion, we found that external control data have been applied in a variety of IMID settings to contextualise efficacy and safety outcomes. The reporting of the methodology used to generate and analyse ECAs has been incomplete and heterogeneous. The establishment of authoritative reporting guidelines may serve as a catalyst for transparent reporting and rigorous study design for ECA studies in IMIDs.

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Supplemental Appendix 1. Search strategies.**Embase (Ovid, inception-12 September 2023)**

- 1 ("external control arm*" or "external control group*" or "external control*" or "external comparator*").mp.
- 2 ("synthetic control arm*" or "synthetic control group*" or "synthetic control*" or "synthetic comparator*").mp.
- 3 ("externally controlled" or "externally-controlled" or "synthetically controlled" or "synthetically-controlled").mp.
- 4 "external data".mp.
- 5 ("baseline control*" or "baseline-control*" or "registry control*" or "registry-control*" or "sequential control*" or "sequential-control*" or "historical control*" or "historical-control*").mp.
- 6 ("real world" and "control").mp.
- 7 "external placebo".mp.
- 8 "historical placebo".mp.
- 9 "historic cohort".mp.
- 10 exp Crohn disease/
- 11 Crohn*.mp.
- 12 exp ulcerative colitis/
- 13 colitis.mp.
- 14 exp arthritis, rheumatoid/
- 15 "rheumatoid arthritis".mp.
- 16 exp Ankylosing spondylitis/
- 17 Ankylosing spondylitis.mp.
- 18 "inflammatory bowel disease".mp.
- 19 "IBD".mp.
- 20 exp psoriatic arthritis/
- 21 "psoriatic arthritis".mp.
- 22 exp psoriasis/
- 23 "psoriasis".mp.
- 24 exp dermatitis, atopic/
- 25 "atopic dermatitis".mp.
- 26 or/1-9
- 27 or/10-25
- 28 26 and 27

MEDLINE (Ovid, inception-12 September 2023)

- 1 ("external control arm*" or "external control group*" or "external control*" or "external comparator*").mp.
- 2 ("synthetic control arm*" or "synthetic control group*" or "synthetic control*" or "synthetic comparator*").mp.
- 3 ("externally controlled" or "externally-controlled" or "synthetically controlled" or "synthetically-controlled").mp.
- 4 "external data".mp.

- 5 ("baseline control*" or "baseline-control*" or "registry control*" or "registry-control*" or "sequential control*" or "sequential-control*" or "historical control*" or "historical-control*").mp.
- 6 ("real world" and "control").mp.
- 7 "external placebo".mp.
- 8 "historical placebo".mp.
- 9 "historic cohort".mp.
- 10 exp Crohn disease/
11 Crohn*.mp.
- 12 exp ulcerative colitis/
13 colitis.mp.
- 14 exp arthritis, rheumatoid/
15 "rheumatoid arthritis".mp.
- 16 exp Ankylosing spondylitis/
17 Ankylosing spondylitis.mp.
- 18 "inflammatory bowel disease".mp.
- 19 "IBD".mp.
- 20 exp psoriatic arthritis/
21 "psoriatic arthritis".mp.
- 22 exp psoriasis/
23 "psoriasis".mp.
- 24 exp dermatitis, atopic/
25 "atopic dermatitis".mp.
- 26 or/1-9
27 or/10-25
28 26 and 27

Cochrane Central Register of Controlled Trials (CENTRAL) (inception-12 September 2023)

- #1 "external control arm*" or "external control group*" or "external control*" or "external comparator*"
- #2 "synthetic control arm*" or "synthetic control group*" or "synthetic control*" or "synthetic comparator*"
- #3 "externally controlled" or "externally-controlled" or "synthetically controlled" or "synthetically-controlled"
- #4 "external data"
- #5 "baseline control*" or "baseline-control*" or "registry control*" or "registry-control*" or "sequential control*" or "sequential-control*" or "historical control*" or "historical-control*"
- #6 "real world" and "control"
- #7 "external placebo"
- #8 "historical placebo"
- #9 "historic cohort"
- #10 MeSH descriptor: [Crohn Disease] explode all trees
#11 Crohn*
- #12 MeSH descriptor: [Colitis, Ulcerative] explode all trees
#13 colitis
- #14 MeSH descriptor: [Arthritis, Rheumatoid] explode all trees
#15 "rheumatoid arthritis"

- #16 MeSH descriptor: [Spondylitis] explode all trees
- #17 Ankylosing spondylitis
- #18 "inflammatory bowel disease"
- #19 "IBD"
- #20 MeSH descriptor: [Arthritis, Psoriatic] explode all trees
- #21 "psoriatic arthritis"
- #22 MeSH descriptor: [Psoriasis] explode all trees
- #23 "psoriasis"
- #24 MeSH descriptor: [Dermatitis, Atopic] explode all trees
- #25 "atopic dermatitis"
- #26 #1 or #2 or #3 or #4 or #5 or #6 or #7 or #8 or #9
- #27 #10 or #11 or #12 or #13 or #14 or #15 or #16 or #17 or #18 or #19 or #20 or #21 or #22 or #23 or #24 or #25
- #28 #26 and #27

Supplemental Table 1. External control data source quality checklist.^a

| Key domain | Criteria for judgement |
|-----------------------------------|--|
| Quality of data source | <ul style="list-style-type: none">• Patient data sourced from RCTs or prospective cohort studies• Baseline characteristics data relevant to the target population were collected |
| Similarity of study populations | <ul style="list-style-type: none">• Sufficiently similar disease definitions and inclusion/exclusion criteria• Sufficiently similar patient demographics• Appropriate use of statistical adjustments, subgroup analyses, or sensitivity analyses if patient characteristics differ |
| Similarity of outcome definitions | <ul style="list-style-type: none">• Sufficiently similar outcome definitions and timepoints of assessment• Appropriate use of sensitivity analyses if timepoints differ |
| Reliability and comprehensiveness | <ul style="list-style-type: none">• Use of multiple sources of data• Sufficient sample size and covariates• Low proportion of missing data or appropriate use of sensitivity analyses to assess the effect of missing data |
| Other limitations | <ul style="list-style-type: none">• No other dataset limitations that may limit the reliability and validity of comparisons apparent |

Abbreviations: RCT, randomised controlled trial.

^aAdapted from Thorlund et al.¹

Supplemental Table 2. Excluded studies.

| # | Study ID | Reason for exclusion |
|-----|---------------------------------|--|
| 1. | Backhouse 2012 ² | Non-interventional |
| 2. | Banerjee 2016 ³ | Active arm not from a clinical study |
| 3. | Baraliakos 2011 ⁴ | Secondary publication of included study (Baraliakos 2014) |
| 4. | Braun 2018 ⁵ | Secondary publication of included study (Braun 2019) |
| 5. | Braun 2018 ⁶ | Secondary publication of included study (Braun 2019) |
| 6. | Carter 2009 ⁷ | Secondary publication of included study (Carter 2012) |
| 7. | Carter 2011 ⁸ | Secondary publication of included study (Carter 2012) |
| 8. | Cavallaro 2019 ⁹ | Active arm not from a clinical study |
| 9. | Cozzi 2022 ¹⁰ | No external control |
| 10. | Cruz 2010 ¹¹ | Insufficient information for inclusion; study population not specified |
| 11. | Curtis 2015 ¹² | Secondary publication of included study (Curtis 2015) |
| 12. | Deery 2020 ¹³ | Ineligible study population |
| 13. | Descalzo 2012 ¹⁴ | Non-interventional |
| 14. | Eslamy 2014 ¹⁵ | Intervention for the treatment of immune-mediated disease not assessed |
| 15. | Favalli 2018 ¹⁶ | No external control |
| 16. | Flor-de-Lima 2013 ¹⁷ | Active arm not from a clinical study |
| 17. | Garg 2010 ¹⁸ | Secondary publication of included study (Garg 2011) |
| 18. | Girleanu 2019 ¹⁹ | Insufficient information for inclusion; control group source not specified |
| 19. | Glintborg 2019 ²⁰ | Active arm not from a clinical study |
| 20. | Gorgun 2004 ²¹ | Active arm not from a clinical study |
| 21. | Grass 2019 ²² | Active arm not from a clinical study |
| 22. | Haifer 2019 ²³ | No external control |
| 23. | Hallgren 1990 ²⁴ | Insufficient information for inclusion; control group source not specified |
| 24. | Harold 2018 ²⁵ | Intervention for the treatment of immune-mediated disease not assessed |
| 25. | Harris 2019 ²⁶ | Insufficient information for inclusion; control group source not specified |
| 26. | Haugeberg 2019 ²⁷ | Insufficient information for inclusion; control group source not specified |
| 27. | Hjuler 2018 ²⁸ | Intervention for the treatment of immune-mediated disease not assessed |
| 28. | Holroyd 2018 ²⁹ | No external control |
| 29. | Hunter 2014 ³⁰ | Insufficient information for inclusion; control group source not specified |
| 30. | Hunter 2014 ³¹ | Insufficient information for inclusion; control group source not specified |
| 31. | Iannone 2016 ³² | No external control |
| 32. | Ito 2002 ³³ | Ineligible study population |
| 33. | Jamal 2019 ³⁴ | No external control |
| 34. | Jani 2020 ³⁵ | Active arm not from a clinical study |
| 35. | Jani 2022 ³⁶ | Active arm not from a clinical study |
| 36. | Karakan 2013 ³⁷ | Insufficient information for inclusion; control group source not specified |
| 37. | Kaur 2018 ³⁸ | Non-interventional; historical control not used to assess the efficacy or safety of a medical intervention |
| 38. | Kaur 2020 ³⁹ | Non-interventional; historical control not used to assess the efficacy or safety of a medical intervention |
| 39. | Klein 2019 ⁴⁰ | Active arm not from a clinical study |
| 40. | Lamplot 2015 ⁴¹ | Intervention for the treatment of immune-mediated disease not assessed |
| 41. | Lebwohl 2015 ⁴² | No external control |
| 42. | Lebwohl 2016 ⁴³ | No external control |
| 43. | Lee 2019 ⁴⁴ | No external control |
| 44. | Lippuner 1996 ⁴⁵ | Ineligible study population |
| 45. | Loftus 2009 ⁴⁶ | No external control |
| 46. | Loftus 2009 ⁴⁷ | No external control |
| 47. | Luglio 2016 ⁴⁸ | No external control |
| 48. | Maas 2016 ⁴⁹ | Insufficient information for inclusion; control group source not specified |
| 49. | Mahil 2021 ⁵⁰ | Intervention for the treatment of immune-mediated disease not assessed |
| 50. | McDaniel 2020 ⁵¹ | Intervention for the treatment of immune-mediated disease not assessed |
| 51. | Medel 2011 ⁵² | Intervention for the treatment of immune-mediated disease not assessed |
| 52. | Murphy 2013 ⁵³ | Intervention for the treatment of immune-mediated disease not assessed |
| 53. | Neva 2003 ⁵⁴ | Intervention for the treatment of immune-mediated disease not assessed |
| 54. | Okazaki 2016 ⁵⁵ | Ineligible study population |
| 55. | Pal 2018 ⁵⁶ | Insufficient information for inclusion; control group source not specified |
| 56. | Pal 2019 ⁵⁷ | Insufficient information for inclusion; control group source not specified |
| 57. | Papa 2012 ⁵⁸ | Insufficient information for inclusion; control group source not specified |

| # | Study ID | Reason for exclusion |
|-----|-----------------------------------|--|
| 58. | Papa 2012 ⁵⁹ | Insufficient information for inclusion; control group source not specified |
| 59. | Papa 2012 ⁶⁰ | Insufficient information for inclusion; control group source not specified |
| 60. | Papadopoulou 1995 ⁶¹ | Insufficient information for inclusion; control group source not specified |
| 61. | Pena-Sagredo 2009 ⁶² | Active arm not from a clinical study |
| 62. | Petit 2020 ⁶³ | Intervention for the treatment of immune-mediated disease not assessed |
| 63. | Petit 2021 ⁶⁴ | Intervention for the treatment of immune-mediated disease not assessed |
| 64. | Prats-Urbe 2020 ⁶⁵ | No external control |
| 65. | Reddy 2006 ⁶⁶ | Intervention for the treatment of immune-mediated disease not assessed |
| 66. | Reich 2001 ⁶⁷ | No external control |
| 67. | Rodewald 2001 ⁶⁸ | Active arm not from a clinical study |
| 68. | Schulert 2018 ⁶⁹ | Active arm not from a clinical study |
| 69. | Sebastian 2021 ⁷⁰ | Intervention for the treatment of immune-mediated disease not assessed |
| 70. | Shashi 2020 ⁷¹ | Active arm not from a clinical study |
| 71. | Sigurdardottir 2018 ⁷² | Active arm not from a clinical study |
| 72. | Sigurdardottir 2019 ⁷³ | Active arm not from a clinical study |
| 73. | Simpson 2010 ⁷⁴ | Insufficient information for inclusion; control group source not specified |
| 74. | Spinelli 2013 ⁷⁵ | No external control |
| 75. | Spinelli 2015 ⁷⁶ | No external control |
| 76. | Spinelli 2019 ⁷⁷ | Intervention for the treatment of immune-mediated disease not assessed |
| 77. | Stein 2016 ⁷⁸ | Intervention for the treatment of immune-mediated disease not assessed |
| 78. | Strand 2021 ⁷⁹ | No external control |
| 79. | To 2012 ⁸⁰ | Insufficient information for inclusion; control group source not specified |
| 80. | Tweehuysen 2017 ⁸¹ | Secondary publication of included study (Tweehuysen 2018) |
| 81. | Tweehuysen 2018 ⁸² | Secondary publication of included study (Tweehuysen 2018) |
| 82. | Vadlamudi 2010 ⁸³ | Active arm not from a clinical study |
| 83. | Varkas 2016 ⁸⁴ | Active arm not from a clinical study |
| 84. | Weidner 2000 ⁸⁵ | Intervention for the treatment of immune-mediated disease not assessed |
| 85. | Weinstein 2021 ⁸⁶ | Intervention for the treatment of immune-mediated disease not assessed |
| 86. | Wolf 2015 ⁸⁷ | Active arm not from a clinical study |
| 87. | Wolf 2015 ⁸⁸ | Active arm not from a clinical study |
| 88. | Wolf 2015 ⁸⁹ | Active arm not from a clinical study |
| 89. | Wolf 2016 ⁹⁰ | Active arm not from a clinical study |
| 90. | Yoshii 2016 ⁹¹ | Ineligible study population |
| 91. | Zhou 2004 ⁹² | Intervention for the treatment of immune-mediated disease not assessed |
| 92. | Feagan 2023 ⁹³ | Insufficient information for inclusion; control group source not specified |
| 93. | Chou 2022 ⁹⁴ | Active arm not from a clinical study |
| 94. | So 2023 ⁹⁵ | Insufficient information for inclusion; control group source not specified |
| 95. | Hagino 2023 ⁹⁶ | No external control |
| 96. | Loughlin 2022 ⁹⁷ | Insufficient information for inclusion; control group source not specified |
| 97. | Hagino 2022 ⁹⁸ | No external control |

Supplemental Table 3. External control data quality assessment.^a

| Study ID | Quality of data source | Similarity of study populations | Similarity of outcome definitions | Reliability and comprehensiveness of the datasets | Other limitations |
|---------------------------------------|------------------------|---------------------------------|-----------------------------------|---|-------------------|
| Baker 1989 ⁹⁹ | Unclear | Unclear | High | Unclear | Unclear |
| Rosenberg 1990 ¹⁰⁰ | Unclear | Unclear | High | Unclear | Unclear |
| Stenger 1998 ¹⁰¹ | Unclear | Low | Low | Unclear | Low |
| Marotte 2007 ¹⁰² | Unclear | Low | Low | Low | Low |
| Shen 2007 ¹⁰³ | Unclear | Low | Low | Unclear | Low |
| Moul 2007 ¹⁰⁴ | Low | Unclear | Low | Unclear | Low |
| van der Heijde 2008 ¹⁰⁵ | Low | Low | Low | Unclear | Unclear |
| Boyle 2008 ¹⁰⁶ | Unclear | Unclear | Low | Unclear | Unclear |
| van der Heijde 2009 ¹⁰⁷ | Low | Unclear | Low | Unclear | Low |
| Garg 2011 ¹⁰⁸ | Unclear | Unclear | Unclear | Unclear | Unclear |
| Yong 2011 ^{109 b} | Unclear | Unclear | Unclear | Unclear | Low |
| Carter 2012 ¹¹⁰ | Low | Unclear | Low | Unclear | Unclear |
| Hueber 2012 ¹¹¹ | Low | Low | Low | Low | Low |
| Hyams 2012 ¹¹² | Low | Unclear | Unclear | Low | Low |
| Hodkinson 2013 ¹¹³ | Unclear | Unclear | Low | Unclear | Unclear |
| Kwok 2012 ¹¹⁴ | Unclear | Unclear | Low | Unclear | Unclear |
| Baraliakos 2014 ¹¹⁵ | Unclear | Unclear | Low | Unclear | Unclear |
| Horneff 2014 ¹¹⁶ | Low | Unclear | High | Unclear | Unclear |
| Lichtenstein 2014 ¹¹⁷ | Low | Low | Unclear | Unclear | Unclear |
| Gardenbroek 2015 ¹¹⁸ | Unclear | Low | Low | Unclear | Unclear |
| Mahajan 2018 ¹¹⁹ | Unclear | Unclear | Low | Unclear | Low |
| Rozette 2018 ¹²⁰ | Unclear | Unclear | Low | Unclear | Unclear |
| Tweehuysen 2018 ¹²¹ | Unclear | Unclear | Low | Unclear | Unclear |
| Horneff 2018 ¹²² | Unclear | Unclear | Low | Unclear | Unclear |
| Inui 2018 ¹²³ | Unclear | Unclear | Low | Unclear | Low |
| Braun 2019 ¹²⁴ | Low | Low | Low | Unclear | Unclear |
| Kawashiri 2020 ^{125 c} | Low | Unclear | Unclear | Unclear | Unclear |
| Muskens 2020 ¹²⁶ | Unclear | Low | Low | Unclear | Low |
| Sanchez-Hernandez 2020 ¹²⁷ | Unclear | Low | Low | Low | Low |
| Croft 2021 ¹²⁸ | Low | Unclear | Low | Low | Low |
| Hupf 2021 ^{129 b} | Low | Unclear | Unclear | Low | Low |

| Study ID | Quality of data source | Similarity of study populations | Similarity of outcome definitions | Reliability and comprehensiveness of the datasets | Other limitations |
|---|------------------------|---------------------------------|-----------------------------------|---|-------------------|
| Reed 2021 ¹³⁰ ^b | Low | Unclear | Unclear | Unclear | Unclear |
| Rudrapatna 2021 ¹³¹ ^b | Unclear | Unclear | Low | Unclear | Low |
| Wang 2021 ¹³² ^b | Low | Low | Low | Low | Low |
| Sagami 2021 ¹³³ | Unclear | Low | Low | Unclear | Low |
| Gasparetto 2021 ¹³⁴ | Unclear | Unclear | Unclear | Unclear | Low |
| Danese 2021 ¹³⁵ | Low | Low | Unclear | Low | Low |
| Magnolo 2022 ¹³⁶ | Low | Low | Low | Low | Low |
| Norvang 2022 ¹³⁷ ^b | Low | Low | Low | Unclear | Low |
| Holland 2022 ¹³⁸ | Unclear | Low | Low | Unclear | Unclear |
| Huang 2022 ¹³⁹ | Unclear | Low | Low | Unclear | Low |
| Truong 2022 ¹⁴⁰ | Unclear | Low | Low | Unclear | Low |
| Takei 2022 ¹⁴¹ | Unclear | Low | Low | Unclear | Unclear |

Abbreviations: ECA, external control arm.

^a The quality assessment is based on the control quality checklist developed by Thorlund et al.¹

^b Primary study objective was to examine methodological approaches for complementing prior or concurrent RCT data with external control data.

^c Protocol for an upcoming single-arm study with an ECA.

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