

Preferences for preventive treatments for rheumatoid arthritis

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Original article

Preferences for preventive treatments for rheumatoid arthritis: discrete choice survey in the UK, Germany and Romania

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Abstract

Objective. To quantify preferences for preventive therapies for rheumatoid arthritis (RA) across three countries.

Methods. A web-based survey including a discrete choice experiment was administered to adults recruited via survey panels in the UK, Germany and Romania. Participants were asked to assume they were experiencing arthralgia and had a 60% chance of developing RA in the next 2 years and completed 15 choices between no treatment and two hypothetical preventive treatments. Treatments were defined by six attributes (effectiveness, risks and frequency/route of administration) with varying levels. Participants also completed a choice task with fixed profiles reflecting subjective estimates of candidate preventive treatments. Latent class models (LCMs) were conducted and the relative importance of attributes, benefit–risk trade-offs and predicted treatment uptake was subsequently calculated.

Results. Completed surveys from 2959 participants were included in the analysis. Most participants preferred treatment over no treatment and valued treatment effectiveness to reduce risk more than other attributes. A five-class LCM best fitted the data. Country, perceived risk of RA, health literacy and numeracy predicted class membership probability. Overall, the maximum acceptable risk for a 40% reduction in the chance of getting RA (60% to 20%) was 21.7%, 19.1% and 2.2% for mild side effects, serious infection and serious side effects, respectively. Predicted uptake of profiles reflecting candidate prevention therapies differed across classes.

Conclusion. Effective preventive pharmacological treatments for RA were acceptable to most participants. The relative importance of treatment attributes and likely uptake of fixed treatment profiles were predicted by participant characteristics.

Key words: RA, preventive treatment, patient preferences, discrete choice experiment

Rheumatology key messages

- Acceptable preventive treatments for RA could reduce pain, disability and societal costs on a huge scale.
- This study quantifies preferences for treatments to reduce the risk of RA development across three countries.
- This guides trial design and development of decision-making support for individuals at risk of RA.

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Introduction

Increased understanding of the biological mechanisms involved in the pre-clinical stages of rheumatoid arthritis (RA) [1] and the ability to identify individuals at risk of developing RA [2] have highlighted opportunities for preventive intervention [3–5]. There is emerging research interest in assessing whether a short course of therapy will prevent the onset of RA in at-risk individuals. B cell depletion, with a single infusion of 1000 mg of rituximab, has been shown to delay the onset of RA in individuals with arthralgia and autoantibody positivity [6]. The preventive efficacy of abatacept in this group [7] and HCQ in asymptomatic autoantibody-positive individuals [8] is under investigation and novel cellular therapies to prevent RA are being explored [9].

Difficulties in recruitment to RA prevention trials have been reported [10]. A trial assessing the efficacy of 40 mg atorvastatin daily for 3 years was prematurely discontinued due to a low inclusion rate resulting from unwillingness to participate [11]. Recent EULAR recommendations highlight the importance of understanding the perspectives of at-risk groups to facilitate tailored strategies to enhance engagement with prevention studies and increase understanding about what types of preventive treatments are acceptable [12]. There is growing awareness of the value of insights into patient preferences for drug development, approval and reimbursement [13–15]. Increased understanding of the relative importance of treatment attributes and acceptable benefit-risk trade-offs for preventive treatments could prevent investment in studies of unacceptable treatments, guide selection of outcomes and endpoints and inform the development of resources to support decision making around participation in prevention trials.

While preferences for treatments for established RA have been well studied [16, 17], few studies have quantified preferences for preventive treatment of at-risk groups [18]. Among those at risk for RA, there is uncertainty regarding not only treatment effectiveness and safety, but also the individual's risk of developing RA, the timeline for that risk and the severity of disease if RA does develop [19]. Qualitative studies have explored stakeholder perceptions of preventive treatment for RA [20]. A survey of individuals with arthralgia and autoantibody positivity found that 40% reported a willingness to accept preventive treatments that were 100% effective with minor side effects of immune suppression if their risk of developing RA was 70% [21]. A pilot study in Switzerland [22] and two Canadian studies [23, 24] of preventive treatment choices found that treatments with mild side effects were acceptable, although the relative importance of treatment attributes varied across studies. The aim of this study was to quantify preferences and preference heterogeneity for preventive treatments for RA in large samples of the general population across three European countries.

Methods

This study is part of a case study for the Innovative Medicines Initiative (IMI) project 'Patient Preferences in

Benefit-Risk Assessments during the Medical Product Lifecycle' (PREFER), which aims to develop evidence-based recommendations on how and when preference studies can inform decision making during drug development [13]. The study was approved by the London-Hampstead Research Ethics Committee (19/LO/0407) and the Ethics Committee of the Friedrich-Alexander-Universität Erlangen-Nürnberg (92_17 B). The protocol has been published [25].

Participant identification and recruitment

Participants were recruited in the UK, Germany and Romania via online survey panels (surveyengine.com) using age- and gender-matched samples informed by a prior study of individuals at risk of RA [26]. These countries were chosen to reflect a geographically and culturally diverse range across Europe to facilitate assessment of preference consistency across populations with distinct healthcare systems. Members of online survey panels in each country received an e-mail invitation ([Supplementary Data S1](#), available at *Rheumatology* online) to take part in and link to the online survey. The initial questions of the survey confirmed respondent's eligibility to participate (age ≥ 18 years and without a diagnosis of RA). Eligible respondents were provided with participant information and asked to provide anonymous electronic informed consent. Participants provided written informed consent. After completing the survey, panel members were credited with panel points (equivalent to $\sim \text{€}2\text{--}3$). Translation of the survey from English to German and Romanian was undertaken in accordance with ISO17100 standards [27]. Recruitment continued until 1000 surveys were completed in each country. For sample size assessment details see [Supplementary Data S1](#), available at *Rheumatology* online.

Survey development

A discrete choice experiment (DCE) within a web-based survey was developed following best practices [28, 29]. This method has been widely applied to elicit preferences for healthcare interventions [30, 31]. The DCE consists of a series of choice tasks. For each task, participants were asked to imagine they were experiencing joint pain that was impacting on their daily activities and had received test results indicating their risk of developing RA in the next 2 years was 60%; their doctor has suggested they consider taking a treatment to reduce their risk; and treatment would be for 1 year. Participants were asked to choose between two treatment options or to opt out (no treatment), where treatments could lower the chance of getting RA in exchange for a risk of side effects and different modes and frequencies of administration. Each treatment option was described by their level of six attributes. See [Supplementary Data S1](#), available at *Rheumatology* online for an example choice task.

DCE attribute and level selection and experimental design

The selection of treatment attributes included in the DCE was informed by a previous literature review [18],

qualitative study and attribute ranking survey [25]. Final attributes (Table 1) were agreed upon by an international team of clinical researchers, consultant rheumatologists, preference elicitation experts and patient partners. Attribute levels were estimated with input from clinical experts, including researchers leading the development and/or clinical trials of preventive treatments for RA. The text used to describe attributes and levels to participants is available in [Supplementary Data S1](#), available at *Rheumatology* online. The combinations of attribute levels that define each treatment option and the set of treatment options in each choice question of a DCE constitute the experimental design. This must have statistical properties that allow estimation of the preference weights of interest obtained from responses to a number of choice tasks that is feasible for participants to complete. Ngene (ChoiceMetrics, Sydney, NSW, Australia) was used to construct a Bayesian D-efficient fractional factorial design [32]. Prior information on attribute importance was initially based on previous literature and expert best guesses, then the outcomes of conditional logit analysis of a pilot survey ($N = 100$) were used as priors for the final experimental design. Sixty unique choice tasks were generated, divided over four blocks. Each participant answered one block (15 choice tasks) only. Participants were randomized to blocks upon entering the survey using the minimization method to ensure equal proportions of participants across blocks. The design was restricted to exclude the following

combinations of attribute levels: taking a pill monthly or every 6 months, having an injection daily or having a drip daily or weekly. Interactions between effectiveness and chance of a serious infection and between effectiveness and the chance of serious side effects were included.

Survey content

The survey was co-developed with patient partners and pretested in a convenience sample ($N = 15$) using 'think-aloud' interviews. It included background information about RA (see [Supplementary Data S1](#), available at *Rheumatology* online), questions to assess comprehension of the background material, introduction to the choice tasks and treatment attributes and levels, a guided 'walk-through' choice task example, warm-up choice tasks and 15 DCE choice tasks. During completion of each task, participants could choose to view the explanation of each attribute and its levels. Participants then completed an additional choice task including four treatment options with fixed profiles, which reflected drug characteristics and known risks of agents under investigation as preventive treatments for RA (HCQ, atorvastatin, abatacept and tolerogenic cell-based therapy). Treatment effectiveness was estimated with input from researchers leading the development and/or clinical trials of the relevant agents.

The survey included measures of sociodemographic variables, the Single-Item Health Literacy Screener [33] and the three-item version of the Subjective Numeracy

TABLE 1 Discrete choice task background scenario, attributes and levels

| Choice task background: When answering, we ask you to imagine that you are experiencing some joint symptoms (joint pain and stiffness) that are starting to impact on your daily activities and that the results of a blood test show you have a 60% chance of developing rheumatoid arthritis in the next 2 years. Your doctor suggests that you consider taking one of the following treatments for 1 year. In this case, would you prefer treatment A, treatment B or no treatment? | | |
|--|------------------------------------|-----------------------|
| Treatment attribute | Levels | |
| Chance of developing RA reduced from 60% to | 10% | 10 in 100 |
| | 20% | 20 in 100 |
| | 30% | 30 in 100 |
| | 40% | 40 in 100 |
| How the treatment is taken | A shallow injection under the skin | |
| | A drip into the vein | |
| | One or two tablets | |
| How often the medication has to be taken | Daily | |
| | Weekly | |
| | Monthly | |
| | Every 6 months | |
| Chance of reversible mild side effects (e.g. nausea, skin rashes or muscle pain) | 2% | 2 in 100 people |
| | 5% | 5 in 100 people |
| | 10% | 10 in 100 people |
| Chance of a serious infection (e.g. pneumonia) | 0% | 0 in 100 people |
| | 1% | 1 in 100 people |
| | 5% | 5 in 100 people |
| Chance of a serious, potentially irreversible side effect (e.g. brain inflammation, lymphoma or retinopathy) | 0.001% | 1 in 100 000 people |
| | 0.02% | 20 in 100 000 people |
| | 0.1% | 100 in 100 000 people |

Scale [34] and participants assessed their lifetime risk of developing RA using a five-item Likert-type scale ('very unlikely' to 'very likely'). Upon completion of the survey, all respondents were provided with information about RA for those who are at risk developed by an international team of patient partners, rheumatologists and researchers as part of a previous project [35].

Data analysis

The outcomes of interest were treatment preferences and preference heterogeneity, relative preference weights for levels of treatment attributes, estimated risk equivalents [maximum acceptable risk (MAR) for changes in treatment effectiveness] and choice frequency of fixed treatment profiles. Results were considered statistically significant if $P \leq 0.05$. Analyses were carried out using SPSS version 27.0 (IBM, Armonk, NY, USA) and NLogit (Econometric Software, Plainview, NY, USA). Participants who exhibited non-attendance to the survey were removed from the analysis, including those who completed the survey in 5 min and those who took 5–9 min to answer the survey, answered two or more of three background comprehension questions incorrectly and showed 'flat-lining' behaviour (always choosing 'treatment A' or always choosing 'treatment B') in the DCE choice tasks.

Panel latent class models (LCMs) were used on a pooled dataset combining data across countries. All attributes were considered non-linear and recoded using effects codes [36, 37]. Based on model fit tests [Akaike information criterion (AIC), log likelihood], the most suitable model was selected (models ranging from one to six classes were tested). See [Supplementary Data S1](#), available at *Rheumatology* online, for the final utility equation.

A class assignment model was fitted to assess the contribution of participant characteristics (i.e. country, perceived risk of developing RA, smoking, age, gender, health literacy, numeracy and educational level). A significant variable in the class assignment model indicates that this variable contributes to the class assignment (e.g. if the coefficient of the health literacy variable is positive and significant for class 1, participants with a higher health literacy score are more likely to belong to class 1).

Relative importance scores for the attributes relative to the most important attribute were calculated based on the results of the LCMs, separately for all classes. The class-adjusted relative importance was calculated by computing the relative importance score of all attributes in each class separately, after which they were weighted according to class assignment probability.

The MAR for a 40% reduction in the chance of getting RA (60% to 20%) was calculated for each of the three risk attributes according to the equation:

$$\text{MAR} = -\frac{(\beta_{\text{chance of developing RA}})}{(\beta_{\text{risk}})}.$$

For these calculations an LCA model was used, assuming linearity for all attributes included. The MAR was calculated separately for all classes after which they were

weighted according to class assignment probability. Predicted choice $[1/(1+\exp^{-y})]$ between the four hypothetical treatment profiles with fixed profiles and no treatment was tabulated using LCM estimates for each class.

Patient and public involvement

A panel of eight patient research partners [based in the UK ($n=4$), Germany ($n=2$), The Netherlands ($n=1$) and Sweden ($n=1$)] contributed to the development of research objectives, selection of attributes and levels, development of choice tasks and survey content and pretesting.

Results

Responses of 982 individuals (332 males, 650 females) in the UK, 984 (325 males, 656 females, 2 gender not reported) in Germany and 993 (340 males, 652 females, 1 other) in Romania were included. Participant characteristics are summarized in [Table 2](#). Health literacy scores were significantly lower in the sample in Romania compared with samples in Germany and the UK. Subjective numeracy scores were significantly higher in the sample in Romania compared with those in both Germany and the UK. Participants in Germany perceived their chances of developing RA to be significantly lower than those in either the UK or Romania.

Five classes were fitted in the LCM (see [Table 3](#) for model coefficients). The average probabilities of belonging to classes 1–5 were 51%, 10%, 26%, 4% and 10%, respectively. In all classes, participants preferred treatments that were more effective and had less risk of serious infection and serious side effects. For classes 1, 2 and 3, preventive treatment was preferred over no treatment, whereas in classes 4 and 5, no treatment was preferred. The relative importance of treatment attributes in each class is summarized in [Fig. 1](#). Treatment effectiveness was the most important attribute in all classes except for class 4, where the method of administration was more important. The least important attribute was mild side effects in all classes except for class 2, where the method of administration was least important. In classes 1, 3 and 4, oral tablets were preferred over injection or infusion, while in class 2, this attribute was not significant. In class 5, infusion was preferred over oral tablets and tablets over injections. In the same class, the frequency of administration did not significantly impact on preferences, while in the other classes, the least frequent attribute level of every 6 months was preferred. A preference for the lowest level of mild side effects was found in classes 1–3, with no significant impact on preferences in classes 4 and 5.

The model fit significantly improved when including country, health literacy, numeracy and perceived risk of developing RA in the class assignment model. Participants from Germany were less likely to belong to class 1, whereas participants from Romania were most likely to belong to class 5. Participants with low health literacy were more likely to belong to class 3, whereas

TABLE 3 Parameter estimates from the LCM of preferences for preventive treatment of RA

| Attribute | Level | Class 1 | | Class 2 | | Class 3 | | Class 4 | | Class 5 | |
|--------------------------------------|--------------------|-----------|------|-----------|------|----------|------|----------|------|----------|------|
| | | Estimate | S.E. | Estimate | S.E. | Estimate | S.E. | Estimate | S.E. | Estimate | S.E. |
| No treatment (opt out) | | −4.14*** | 0.08 | −3.81*** | 0.12 | −0.12*** | 0.02 | 1.08*** | 0.07 | 3.58*** | 0.06 |
| Chance of developing RA | 40% (ref) | −0.42*** | 0.04 | −3.31*** | 0.12 | −0.29*** | 0.02 | −0.47*** | 0.09 | −0.52*** | 0.15 |
| | 30% | −0.10*** | 0.02 | −0.79*** | 0.07 | −0.05** | 0.02 | 0.03 | 0.12 | −0.20 | 0.16 |
| | 20% | 0.13*** | 0.02 | 0.87*** | 0.07 | 0.11*** | 0.02 | 0.09 | 0.10 | 0.34** | 0.13 |
| | 10% | 0.40*** | 0.02 | 3.24*** | 0.13 | 0.23*** | 0.02 | 0.36*** | 0.09 | 0.39*** | 0.12 |
| Mode of administration | Tablet (ref) | 0.38*** | 0.04 | −0.11 | 0.10 | 0.20*** | 0.03 | 3.17*** | 0.18 | 0.04 | 0.18 |
| | Injection | −0.06*** | 0.01 | 0.00 | 0.04 | 0.02 | 0.02 | −1.30*** | 0.10 | −0.35*** | 0.11 |
| | Infusion | −0.32*** | 0.02 | 0.12 | 0.07 | −0.22*** | 0.02 | −1.87*** | 0.14 | 0.31** | 0.15 |
| Frequency of administration | Daily (ref) | −0.21*** | 0.06 | −0.32*** | 0.10 | −0.22*** | 0.04 | −0.37** | 0.18 | 0.20 | 0.22 |
| | Weekly | −0.17*** | 0.02 | −0.08 | 0.06 | −0.13*** | 0.02 | −0.61** | 0.15 | −0.11 | 0.15 |
| | Monthly | 0.09*** | 0.02 | −0.08 | 0.07 | 0.14*** | 0.03 | −0.19 | 0.19 | −0.17 | 0.18 |
| | Every 6 months | 0.29*** | 0.02 | 0.47*** | 0.06 | 0.21*** | 0.03 | 1.17*** | 0.15 | 0.07 | 0.16 |
| Chance of mild side effects | 2% (ref) | 0.14*** | 0.03 | 0.27*** | 0.04 | 0.05*** | 0.02 | −0.03 | 0.08 | 0.03 | 0.10 |
| | 5% | 0.03*** | 0.01 | 0.03 | 0.04 | 0.02 | 0.02 | 0.13 | 0.09 | −0.11 | 0.13 |
| | 10% | −0.17*** | 0.01 | −0.30*** | 0.04 | −0.07*** | 0.02 | −0.10 | 0.08 | 0.08 | 0.10 |
| Chance of serious infection | 0% (ref) | 0.22*** | 0.02 | 0.55*** | 0.05 | 0.21*** | 0.02 | 0.00 | 0.10 | 0.36*** | 0.09 |
| | 1% | 0.06*** | 0.02 | 0.07 | 0.05 | −0.02 | 0.02 | 0.02 | 0.09 | −0.11 | 0.12 |
| | 5% | −0.28*** | 0.01 | −0.62*** | 0.05 | −0.19*** | 0.02 | −0.02 | 0.08 | −0.25** | 0.11 |
| Chance of serious side effects | 1 in 100 000 (ref) | 0.32*** | 0.03 | 0.67*** | 0.05 | 0.20*** | 0.01 | −0.12 | 0.09 | 0.22** | 0.09 |
| | 20 in 100 000 | −0.04*** | 0.01 | 0.02 | 0.03 | −0.09*** | 0.02 | 0.36*** | 0.09 | −0.16 | 0.11 |
| | 100 in 100 000 | −0.28*** | 0.01 | −0.69*** | 0.05 | −0.11*** | 0.02 | −0.24*** | 0.09 | −0.06 | 0.11 |
| Average class probability | | 0.51 | | 0.10 | | 0.26 | | 0.04 | | 0.10 | |
| Class membership model ^a | | | | | | | | | | | |
| Constant | | 0.98*** | 0.34 | −11.06*** | 1.02 | 1.39*** | 0.35 | −2.72*** | 0.65 | | |
| Country | Germany | −0.59*** | 0.15 | 0.17 | 0.19 | −0.11 | 0.16 | −0.04 | 0.25 | – | – |
| | Romania | −0.69*** | 0.14 | −0.34* | 0.19 | −1.00*** | 0.16 | −0.82*** | 0.27 | – | – |
| High health literacy | | −0.15 | 0.15 | −0.11 | 0.23 | −0.52*** | 0.16 | 0.35 | 0.30 | – | – |
| High numeracy | | 0.73*** | 0.13 | 6.00*** | 0.56 | 0.47*** | 0.14 | 0.89*** | 0.22 | – | – |
| High perceived risk of developing RA | | 0.08 | 0.14 | −0.99*** | 0.20 | −0.16 | 0.15 | −0.16 | 0.24 | – | – |
| Model fit measures | Loglikelihood | −48050.65 | | | | | | | | | |
| | AIC | 96299.3 | | | | | | | | | |

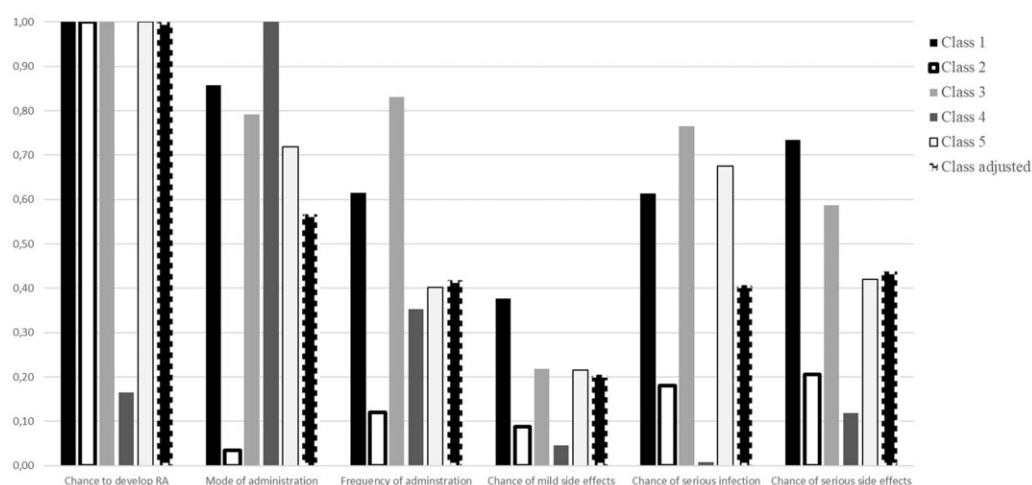
^aClass 5 is the reference category. * $P < 0.10$, ** $P < 0.05$, *** $P < 0.001$.

most important determinant of participants' choices overall. The importance of effectiveness aligns with two of three previous choice-based studies of preferences for preventive treatment [22, 24]. A third study found that serious side effects were more important than effectiveness, although most participants in that study were patients being treated for established RA [23]. Previous qualitative studies have suggested that positive viewpoints of preventive therapies are associated with high expectations of their effectiveness [38–41], highlighting the need to increase understanding of the true likely effectiveness of interventions to reduce RA risk.

A previous stated-choice study of preferences for preventive treatment of RA found that the probability of mild side effects and the method of treatment administration did not significantly determine participants' choice [22], which does not align with the current findings. This

discrepancy across studies could be due to differences in the definition of attributes and levels, preference method chosen and/or the study samples.

Preference heterogeneity was predicted by participant characteristics. Respondents with lower health literacy were more likely to prefer no treatment and more likely to focus on mode and frequency of administration. Others who preferred no treatment were also sensitive to the chance of serious side effects or infection and more likely to report low numeracy. Participants in Romania were more likely to belong to a class where no treatment was preferred than those in Germany and the UK. Another group whose choices were focused on treatment effectiveness to reduce RA risk were more likely to have high perceived risk of developing RA. These findings suggest that developing effective communication tools tailored to individuals' needs and illness beliefs is

Fig. 1 Relative importance of treatment attributes stratified by class in the LCM of preferences for preventive treatments for RA**TABLE 4** Choice frequencies and predicted choices per class (based on LCM output) for additional choice task with fixed attribute levels^a

| Characteristics | No treatment | Treatment A | Treatment B | Treatment C | Treatment D |
|--|--------------|----------------------------------|---------------------|-----------------------|----------------------------------|
| Chance of developing RA reduced from 60% to ... | 60 in 100 | 20 in 100 | 40 in 100 | 30 in 100 | 30 in 100 |
| How the treatment is taken | Not taken | Shallow injection under the skin | One or two tablets | One or two tablets | Shallow injection under the skin |
| How often the treatment has to be taken | Never | Weekly | Daily | Daily | Monthly |
| Chance of a mild side effect | 0 | 5 in 100 people | 10 in 100 people | 10 in 100 people | 10 in 100 people |
| Chance of a serious infection due to treatment | 0 | 1 in 100 people | 0 in 100 people | 0 in 100 people | 1 in 100 people |
| Chance of a serious side effect that is potentially irreversible | 0 | 20 in 100 000 people | 1 in 100 000 people | 100 in 100 000 people | 1 in 100 000 people |
| Choice frequency, <i>n</i> (%) | 425 (14) | 818 (28) | 606 (20) | 468 (16) | 642 (22) |
| Predicted uptake, % | | | | | |
| Class 1 | 0 | 23 | 27 | 21 | 28 |
| Class 2 | 1 | 73 | 2 | 6 | 19 |
| Class 3 | 18 | 18 | 20 | 19 | 25 |
| Class 4 | 12 | 1 | 35 | 51 | 1 |
| Class 5 | 90 | 2 | 4 | 4 | 1 |
| Class adjusted | 14 | 24 | 18 | 18 | 14 |

^aTreatment choices were unlabelled (i.e. participants chose between treatment A, B, C and D or no treatment). Attribute levels in each treatment profile reflected fixed profiles that reflected drug characteristics and known risks of agents under investigation as preventive treatments for RA (abatacept, atorvastatin, HCQ and tolerogenic cell-based therapy, respectively). Treatment effectiveness was estimated with input from researchers leading the development and/or clinical trials of the relevant agents.

essential to facilitate shared decision making by at-risk individuals about taking part in prevention studies or accepting future treatments in clinical settings. The finding that country explained the probability of class

membership suggests that treatment preferences are not generalizable across countries and further investigation of treatment preferences across a wide range of countries and the sociocultural and other (e.g. healthcare

system related) factors that drive preference variability across countries is needed.

Predicted uptake of treatment profiles intended to estimate treatments under consideration for RA prevention was similar across treatments but differed across sub-groups identified by the LCM. Participants with high perceived risk of RA were more likely to belong to the group predicted to prefer treatment profiles estimating biologic and cell-based therapy. Predicted uptake was higher than in two previous studies [23, 24] that included healthcare professional endorsement as a treatment attribute. In this study, this attribute was held constant in the choice task scenario, as it was felt that few people would accept a therapy their doctor did not endorse. Uptake estimates are strongly influenced by subjective attribute levels assigned to treatment profiles. In this study, these were chosen with input from investigators leading prevention trials. A treatment profile chosen to reflect abatacept was most often chosen, reflecting the assignment of the highest level of treatment effectiveness in the fixed choice task. Two previous studies concluded that biologic treatments are unlikely to be acceptable for RA prevention [23, 24]. However, the treatment profiles used in these studies to describe biologic therapies estimated them to be less effective at reducing the risk of RA than other treatments. They also included rituximab, which is administered via an infusion, i.e. the least preferred route for most participants. Rituximab, as a single infusion of 1000mg, has been shown to delay, but not prevent RA development [6] and thus was not included in the current study. Further investigation is warranted to establish preferences for preventive treatment benefits other than effectiveness to reduce risk, including delayed disease development and impact on early symptoms that may precede the diagnosis of RA.

The current study focused on preferences for preventive pharmacological interventions, as there are/have been several clinical trials of drugs to assess their ability to reduce the risk of RA development. Preventive lifestyle interventions were not addressed here. However, there is a rationale for certain preventive behavioural/lifestyle approaches, including smoking cessation [42], diet, exercise and weight management. Combined behavioural and pharmacological strategies may be effective in future clinical applications [43]. There is also converging evidence across qualitative studies that lifestyle interventions would be preferred over preventive drug therapies [20]. The efficacy of lifestyle approaches to prevent RA development [44] and preferences for such interventions are therefore important areas for future studies.

This study has several strengths, including large samples across three countries, robust preference elicitation methodology and input from clinical researchers, methodological experts and patient partners. The treatment attributes and levels included were based on extensive literature review, qualitative investigation and expert consultation, in line with best practice [28, 29, 45]. As with all preference elicitation studies, the results are generalizable to treatments well represented by these attributes

and levels and not to others. For example, alternative methods of treatment administration were not addressed in the present study.

A limitation is that participants were members of the general population asked to assume a hypothetical risk of RA. Previous research has highlighted common public misperceptions regarding RA that are likely to impact beliefs about the need for preventive treatment and associated decision making [46, 47]. We addressed this possibility by close collaboration with patient partners during the development of background information for survey participants and the inclusion of comprehension questions to check participant understanding of this material. Further research is needed to directly assess preferences of individuals with an elevated risk of RA, including those with RA-related autoantibodies and inflammatory-type joint symptoms. A further limitation is that participants were members of online panels who received a small payment to complete the survey and may not be fully representative of the general population. This was addressed to some extent by detailed analysis of data quality and exclusion of responses that did not meet our quality criteria. However, further investigation is needed to establish differences in the treatment preferences of survey panel members compared with samples recruited via alternative methods. This is being explored in a separate analysis of the current case study [25]. The use of an online survey further limits recruitment to those with access to web-based technology and a degree of literacy that may limit the ability to generalize our findings to other populations. Although this study benefits from input from patient partners across a range of European countries, no patient partner from Romania contributed to the survey design, although they were included in a previous related project that informed the clinical objectives of this study [48].

In conclusion, this study establishes that effective treatments to reduce the risk of RA are acceptable, informs the selection of acceptable candidate therapies and guides the design of prevention trials and the development of tailored risk communication tools for those considering preventive treatment.

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Data availability statement

Data are available upon reasonable request.

Supplementary data

Supplementary data are available at *Rheumatology* online.

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