


Comparison of the sensitivity of patient-reported outcomes for detecting the benefit of biologics in severe asthma

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Abstract

Background: The sensitivity of patient-reported outcomes (PROs) to detect the effects of treatment change depends on the match between the change in items of the PRO and the change that takes place in a sample of people. The aim of this study is to compare the sensitivity of different PROs in detecting changes following the initiation of biologic treatment in asthma. **Methods:** Patients starting a biologic treatment as part of clinical care completed the Asthma Control Questionnaire (ACQ-6), the Severe Asthma Questionnaire (SAQ and SAQ-global scores) and the EQ5D (EQ-5D-5L and EQ5D-VAS) at baseline. They completed the ACQ-6, SAQ, SAQ-global and a retrospective global rating of change (GROc) scale at weeks 4, 8 and 16 and completed the EQ-5D-5L and EQ5D-VAS at week 16. The SAQ-global and EQ5D-VAS differ but both are single item 100-point questions. Sensitivity was measured by Cohen's D effect size at each of the three time points. **Results:** 110 patients were recruited. Depending on the time of assessment, effect size varied between 0.45 and 0.64 for the SAQ, between 0.50 and 0.77 for the SAQ-global; between 0.45 and 0.69 for ACQ-6; between 0.91 and 1.22 for GROc; 0.32 for EQ-5D-5L and 0.49 for EQ5D-VAS. **Conclusion:** The sensitivity to change of a questionnaire varies with the time of measurement. The three asthma-specific prospective measures (SAQ, SAQ-global and ACQ-6) have similar sensitivity to change. The single-item EQ5D-VAS was less sensitive than the asthma specific measures and less sensitive than the single-item SAQ-global. The EQ-5D-5L was least sensitive.

Keywords

Asthma, quality of life, measurement, Severe Asthma Questionnaire, Asthma Control Questionnaire-6, EQ5D

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Introduction

Patient-reported outcomes (PROs) form an increasingly important part of treatment evaluation because they measure the impact of treatment from the patient's perspective. In this article, we compare four different types of PRO in their ability to detect change over time in severe asthma following initiation of biologic therapy: a symptom questionnaire measuring asthma control, two quality of life questionnaires and a questionnaire of perceived change. A PRO is sensitive to change to the extent that the items of the questionnaire

capture the changes that occur in a specific population for a specific treatment.¹ Each of the four types of questionnaire captures different kinds of change and therefore may not be equally sensitive to treatment.

Asthma symptoms are, in part, associated with increased airflow obstruction leading to a lower forced expiratory volume in 1 s (FEV₁). The Asthma Control Questionnaire (ACQ-6) is a validated questionnaire that measures asthma control by the severity of asthma symptoms and shows a strong correlation with lung function.^{2,3} The ACQ-6 should



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therefore provide a sensitive measure of improvements in lung function produced by pharmacological treatment.

Quality of life (QoL) is affected by symptoms but also by other factors such as personality,⁴ economic status⁵ and social support.⁶ Small changes in symptoms could create large changes in QoL and the relevant questionnaires, as long as the QoL questionnaire measures aspects of life that change following improved asthma control.

There are two types of quality of life questionnaire, disease specific and generic. As their names suggest, disease specific questionnaires measure only deficits experienced in a particular disease whereas generic questionnaires measure deficits across a range of diseases. The severe asthma questionnaire (SAQ)⁷ is designed to measure the QoL deficits that occur in severe asthma and therefore may be more sensitive to change compared to a generic questionnaire such as the EQ5D,⁸ which is known to poorly represent the deficits of QoL in severe asthma⁹ and therefore should be less sensitive to change.

Measures of perceived change, such as global rating of change (GRoC), require people to evaluate how much they feel they have improved. These are seldom used as a primary outcome measures because of concerns about the accuracy of retrospective recall and other possible biases.¹⁰ Retrospective recall of health events is known to be poor, including an underestimation of asthma symptoms,¹¹ and people create implicit narratives of their treatment that can create an over-estimation of the benefit of treatment.¹⁰

The aim of this study is to compare the sensitivity to change as measured by effect size over several time points for four questionnaires: SAQ, ACQ-6, EQ5D and GRoC.

Methods

Design

An open label multi-centre sensitivity to change study, using the SAQ, other questionnaire measures and clinic data.

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Participants

Participants were recruited from six UK specialist severe asthma centres (Royal Devon & Exeter, University Hospitals Plymouth NHS Trust, North Bristol NHS Trust, University Hospitals Birmingham, Royal Brompton and Harefield Hospitals, Manchester University NHS Foundation Trust) as part of usual care. All patients had undertaken a full multidisciplinary assessment according to UK severe asthma guidelines.¹² Inclusion criteria: patients with severe asthma (GINA step 4 & 5) commencing a biologic treatment in normal clinical care following NICE guidelines.

Participants were asked to complete questionnaires at baseline and at routine biologic administration clinic visits (0, 4, 8, 12 and 16 weeks).

Questionnaires

Severe Asthma Questionnaire: The SAQ is a validated specific scale for severe asthma (10) comprising 16 items measuring difficulties caused by asthma in 16 domains over a 2 week period.⁷ Responses are given on a 7-point scale that are averaged to form the total SAQ score and three subscales, SAQ-MyLife, SAQ-MyMind, and SAQ-My Body.¹³ In addition, the SAQ provides a SAQ-global score based on a single 100-point Borg-type scale. Higher scores indicate better quality of life. The minimum clinically important difference (MCID) is 0.5 for the SAQ and 11 for the SAQ-global.

Asthma Control Questionnaire-6: The ACQ-6 has six items, of which four measure symptoms, one measures short acting bronchodilator use and one activity limitations over a 7 day period. Responses are given on a 7-point scale that averaged to form the ACQ-6 score. Lower scores indicate better asthma control, and a score of 0–0.75 is classified as well controlled asthma.² The MCID is 0.5.

The Euroqol (EQ-5D-5L): The Euroqol is a generic health status questionnaire comprising five dimensions measured on a single day.⁸ We used the index values of this scale rather than the average values, as index values are used in resourcing decisions. Index values were calculated using the 2012 value set for England.¹⁴

Global rating of change Questionnaire: Patients were asked to rate the GRoC in terms of improvement by circling a statement ‘which best describes how you feel since starting your new treatment for your asthma’. The GRoC used in this study comprises 11 response options, no change, a little better, somewhat better, moderately better, a good deal better and a great deal better, with deterioration represented by the same quantifiers but using the word ‘worse’.¹⁰

Procedure

After providing informed consent, patients completed baseline questionnaires (SAQ and ACQ-6) in the clinic

before their first treatment. Patients completed questionnaires (SAQ, ACQ-6, GRoC) during subsequent routine clinic visits as part of usual care at 4 weeks, 8 weeks, 12 weeks and 16 weeks after starting treatment. Patients receiving benralizumab attended at 0, 4, 8 and 16 weeks and so did not attend clinic at 12 weeks due to the transition to an eight weekly dosing schedule. The EQ-5D-5L and FEV₁ was assessed at baseline and at 16 weeks. Clinical records were used to provide demographic details. Exacerbations requiring OCS, healthcare utilisation and prednisolone dose (if on maintenance OCS) were documented at each clinic visit.

Ethical approval

This study received ethical approvals from the Research Ethics Committee/Health Research Authority (REC reference: 19/WA/0011, IRAS project ID: 250167) and was sponsored by University Hospitals Plymouth NHS Trust.

Statistics

We present the data for baseline, 4, 8 and 16 weeks only as the numbers attending week 12 were reduced as those receiving benralizumab did not attend. The COVID pandemic coincided with the start of data collection, so data at 16 weeks was reduced because many of the health staff who were collecting data were reassigned to COVID-related duties. For this reason, we present two analyses, one intention to treat and one per protocol, and we used t-tests to compare the per protocol patients with the intention to treat after removing the per protocol patients from the intention to treat sample. There is no imputation of missing data for the intention to treat sample: we report all valid data but only valid data. Although the sample size is reduced at later time points, the use of the two methods of analysis has the advantage of providing additional calculations of sensitivity to change.

Effect size was Cohen's D calculated from the mean difference in score between baseline and follow-up divided by the pooled standard deviation, except in the case of GRoC where Cohen's D was calculated from the mean score (i.e. the difference between the score and zero) divided by the standard deviation of the GRoC score. Cohen's D provides a standardised measure of change between questionnaires and therefore permits comparison of questionnaires that have different scaling properties. Conventional interpretation of Cohen's D is that an effect size of 0.2 is considered small, 0.5 is medium and 0.8 large.¹⁵ We used Spearman's ranked correlations.

Results

One hundred and 10 patients were enrolled in the study of whom 62 were female and 48 male. Any patient who

completed one or more questionnaires at any time point (0, 4, 8, 16 weeks) was included in the intention to treat sample. Twenty-two patients completed all questionnaires at weeks 0, 4, 8 and 16 of whom 12 were female and 10 male, and these patients form the per protocol sample. Of the 22 per protocol patients, 18 came from one severe asthma centre. Demographic and clinical characteristics of the intention to treat sample are shown in Table 1.

Table 2 shows the mean and standard deviations for the SAQ, SAQ-global and ACQ-6 at baseline, 4, 8 and 16 weeks, for the EQ-5D-5L at baseline and 16 weeks and for the GRoC at 4, 8, and 16 weeks. The data are shown separately for intention to treat and per protocol, and there was no significant difference between the intention to treat and per protocol group at any time point.

Table 3 shows the effect sizes at the three time points after baseline for all questionnaires as a function of the intention to treat and per protocol. Of the six comparisons made between the asthma specific questionnaires, the SAQ-global is slightly better than the other two for four comparisons and the SAQ slightly better than the other two for two comparisons. Overall, though variable the relative effect sizes are similar over time and between groups. However, the effect sizes of the EQ-5D-5L and EQ5D-VAS are lower than the disease specific scale for all comparisons.

Table 4 shows the baseline and follow-up scores for the subscales of the SAQ as a function of the intention to treat and per protocol and shows similar performance across all comparisons.

Table 5 shows the effect sizes at the three time points after baseline for the subscales of the SAQ as a function of the intention to treat and per protocol and shows similar performance across all comparisons.

Table 6 shows the correlations between the questionnaires and objective measures at baseline.

Discussion

The sensitivity of a questionnaire to change depends on the items of the questionnaire, on the treatment and on the population studied.¹ In this study, sensitivity was assessed at three time points. The dropout rate for this study was high due to the pandemic, and as dropout can be non-random, we provided two analysis, intention to treat and per protocol. Although there was no significant difference, there is a trend for those in the per protocol to have higher effect sizes compared to those in the intention to treat sample. For both the intention to treat and per protocol groups, the MCID for the SAQ, SAQ-global and ACQ-6 was achieved as early as week 4.

As we performed a per protocol analysis in addition to an intention to treat analysis, it is possible to compare the relative effect sizes of the questionnaires in different samples.

Table 1. Patient characteristics, means (95% confidence intervals).

	Intention to treat sample		Per protocol sample	
	n	Mean (CI)	n	Mean
Age, years	106	50 (47–53)	22	51 (44–59)
Female, (%)	59 (56)			
FEV1, %predicted	103	66.82 (62.69–70.96)	22	65.83 (57.32–74.34)
Caucasian, (%)	91 (86)		20 (91)	
BMI, kg/m ²	104	29.49 (28.06–30.92)	22	28.20 (25.85–30.54)
Maintenance prednisolone dose, mg/day	60	12.43 (9.64–15.21)	11	10.70 (5.90–5.43)
Exacerbations in the last 12 months requiring OCS	106	5.43 (4.54–6.31)	22	6.50 (4.23–8.77)
Emergency department visits	106	2.08 (1.09–3.06)	22	2.27 (0.58–5.13)
Hospital admissions	106	1.25 (0.48–2.01)	22	0.64 (0.16–1.26)
Cumulative prednisolone dose, mg/yr	105	4062 (3238–4885)	22	3766 (2472–5060)
Receiving biologic treatment, n (%)	106 (96)			
Omalizumab (%)	16 (15)		4 (18)	
Mepolizumab (%)	26 (25)		10 (45)	
Benralizumab (%)	62 (59)		8 (36)	
Reslizumab (%)	2 (2)		0	

Table 2. Means and standard deviations for questionnaires for the intention to treat group (n) and the per protocol group (n = 22).

	Baseline		4 weeks		8 weeks		16 weeks	
	ITT	PP	ITT	PP	ITT	PP	ITT	PP
SAQ	3.72 (106)	3.58	4.38 (102)	4.40	4.90 (73)	4.54	4.57 (48)	4.72
	1.46	1.21	1.54	1.45	1.51	1.52	1.43	1.35
SAQ-Global	46.13 (106)	47.72	57.19 (105)	59.31	60.40 (73)	67.45	63.15 (47)	63.64
	23.67	19.13	22.72	19.90	24.70	18.35	23.79	19.22
ACQ-6	3.13 (90)	3.07	2.53 (85)	2.30	2.35 (59)	2.02	2.23 (41)	2.17
	1.40	1.08	1.50	1.24	1.46	1.12	1.53	1.28
EQ5D index	0.69 (103)	0.71					0.76 (41)	0.80
	0.23	0.19					0.21	0.18
EQ5D-VAS	54.76 (101)	56.82					65.20 (36)	70.65*
	22.21	19.43					23.35	18.43
GRoC			1.83 (103)	2.09	2.25 (72)	3.00	2.84 (45)	3.18
			2.01	2.07	2.40	1.66	2.36	1.68

ACQ: Asthma Control Questionnaire; SAQ: Severe Asthma Questionnaire; GRoC: global rating of change; ITT: intention to treat; PP: per protocol.

Our results reveal that the relative effect size between questionnaires varies as a function of the population and as a function of time. Sensitivity to change is not an absolute property of a scale but something that varies over time and with the population. Previous comparisons of the sensitivity of quality of life questionnaires have provided comparisons at only one time point and with only one group.^{16,17} The variability of effect size as a function of group and time should be recognised when comparing between questionnaires.

Our results provide six comparisons of effect size. Some questionnaires are consistently better or worse than others. Some questionnaires are sometimes better or worse depending on which comparison is made.

The results show that that the relative sensitivity, as measured by effect size, of the different questionnaires changes to some extent between the six different comparisons, showing that comparisons of sensitivity drawn from only one group and at one time may not be generalisable. There is no evidence of important differences in sensitivity between the SAQ, SAQ-global and ACQ-6. The SAQ-global is slightly more sensitive on four of the six comparisons but, given the variability in relative effect sizes observed between these questionnaires, there is no clear evidence of difference between them.

The effect size of the EQ-5D-5L is consistently less than all the other questionnaires in all comparisons. The effect size of the EQ-5D-5L is about half that of the SAQ, SAQ-global

Table 3. Effect sizes (Cohen's D) for change in questionnaire scores at three follow-up time points for intention to treat and per protocol samples.

	4 weeks		8 weeks		16 weeks	
	ITT	PP	ITT	PP	ITT	PP
SAQ	0.44	0.61	0.38	0.70	0.60	0.89
SAQ-Global	0.48	0.59	0.60	1.06	0.72	0.83
ACQ-6	0.41	0.58	0.54	0.95	0.61	0.76
EQ5D					0.32	0.49
EQ5D-VAS					0.49	0.73
GROc	0.91	1.01	0.94	1.81	1.20	1.89

ACQ: Asthma Control Questionnaire; SAQ: Severe Asthma Questionnaire; GROc: global rating of change; ITT: intention to treat; PP: per protocol.

Table 4. Means and standard deviations for SAQ subscale scores for the intention to treat group (n) and the per protocol group (n = 22).

	Baseline		4 weeks		8 weeks		16 weeks	
	ITT	PP	ITT	PP	ITT	PP	ITT	PP
SAQ-My Life	3.80 (104)	3.75	4.45 (100)	4.48	4.37 (71)	4.70	4.68 (47)	4.82
	1.60	1.37	1.70	1.59	1.70	1.54	1.56	1.33
SAQ-My Mind	3.87 (104)	3.63	4.49 (100)	4.50	4.29 (71)	4.42	4.69 (47)	4.82
	1.74	1.45	1.72	1.62	1.73	1.78	1.58	1.65
SAQ-My Body	3.42 (104)	3.09	4.06 (100)	4.06	4.01 (71)	4.27	4.28 (47)	4.38
	1.48	1.27	1.58	1.53	1.43	1.52	1.50	1.50

SAQ: Severe Asthma Questionnaire; ITT: intention to treat; PP: per protocol.

Table 5. Effect sizes (Cohen's D) for change in SAQ subscale scores at three follow-up time points.

	4 weeks		8 weeks		16 weeks	
	ITT	PP	ITT	PP	ITT	PP
SAQ-My Life	0.39	0.49	0.35	0.65	0.56	0.79
SAQ-My Mind	0.36	0.57	0.24	0.49	0.49	0.77
SAQ-My Body	0.42	0.69	0.41	0.84	0.58	0.97

SAQ: Severe Asthma Questionnaire; ITT: intention to treat; PP: per protocol.

and ACQ-6, and it is therefore safe to conclude that the EQ-5D-5L is the least sensitive of the questionnaires studied. The comparative insensitivity of the EQ-5D-5L is to be expected from a scale that only partially captures the QoL deficits of severe asthma on a single day,⁸ and findings are also consistent with evidence that this scale is comparatively insensitive to rehabilitation.¹⁸

The sensitivity of the EQ5D-VAS was slightly better than the EQ-5D-5L, but not as sensitive as other scales and, in particular, less sensitive than the SAQ-global. Although the EQ5D-VAS and SAQ-global are single item scales varying between 0 and 100, they differ in two respects. The EQ5D-VAS asks patients to rate health; the SAQ-global asks

patients to rate quality of life specifically in relation to asthma. The EQ-5D-VAS is a scale with only the end points specified. The SAQ-global is a Borg scale with additional quantifiers placed at empirically derived points along the scale. Borg scales have been shown to be more reliable than visual analogue scales.¹⁹ Our data shows that they may also be more sensitive.

The SAQ has three subscales made up from groupings of the 16 items that make up the SAQ. There was no consistent difference in sensitivity for these three subscales. The My Body subscale was the most sensitive of the three at all three times points for the per protocol group but not for the intention to treat subscale, where it was most sensitive only at week 8. These findings illustrate how small differences in sensitivity can arise from differences in population and time point, again showing that sensitivity is not an absolute property of a questionnaire, but relative to the population and time of assessment.

Although all three subscales of the SAQ are sensitive to the effects of treatment, only the My Life subscale correlates with baseline FEV₁%. A larger data set has also shown that the correlation between FEV₁% and the My Life subscale is larger than My Life and the My Body subscales.¹³ As patients' judgements of quality of life are determined by numerous factors (including dispositional mood, lung

Table 6. Correlation coefficients (ρ) between questionnaire scores at baseline and objective measures.

	FEV ₁ percent predicted	Exacerbations in the last 12 months requiring OCS
SAQ	0.25*	−0.34**
SAQ-global	0.27**	−0.27**
SAQ-My Life	0.30**	−0.35**
SAQ-My Mind	0.14	−0.27**
SAQ-My Body	0.13	−0.29**
ACQ-6	−0.31**	0.32*
EQ-5D-5L	0.28**	−0.28**
EQ5D-VAS	0.34**	−0.30

ACQ: Asthma Control Questionnaire; SAQ: Severe Asthma Questionnaire.

function, effects of treatment), these data suggest that the My Life and My Body subscales are more affected by factors other than lung function. Examination of other correlations with other PROs demonstrated no relationship between sensitivity to change and FEV₁%. The conclusion from these data is that the use of QoL as an outcome variable provides different information from change in lung function and that both PROs and objective measures should be used in clinical trials as they provide different kinds of information about the effects of treatment.

Our study measured change in an open label study rather than comparative change between placebo and active treatment. We found that a GRoC had the highest sensitivity to change, but it does not follow that it is most sensitive in comparing placebo with active treatment. The reason is that retrospective measures are known to be affected by biases, including that of implicit theory.²⁰ When patients receive a new treatment, they form a narrative that the treatment is likely to be effective, and their response, for both placebo and treatment, is therefore affected by this narrative.^{21,22}

The high sensitivity of the SAQ-global but not the EQ5D-VAS shows that single item scales can sometimes be highly sensitive. The SAQ-global is derived from an earlier scale, the Global Quality of Life Scale.¹⁸ Multi-item scales require the patient to consider the components of quality of life. These components are then aggregated most commonly with no weighting, but sometimes as in the case of the EQ5D with weightings from people who are not ill. As a result, the components of the multi-item quality of life scale are not aggregated using the patient's own individual utilities or weights. By contrast, the SAQ-global requires patients do this aggregation themselves, something that will take place fast, automatically and with implicit rather than conscious consideration of events in a person's life.²³ Our findings indicate that this fast, implicit process can be no worse than that provided by longer questionnaires in detecting change.

Conclusion

The sensitivity of the ACQ-6, SAQ and SAQ-global to the effect of starting biologic treatment is similar. The EQ5D-VAS was less sensitive and the EQ-5D-5L was least sensitive to treatment. The comparative sensitivity of PROs varies slightly as a function of population and time of assessment, so comparisons with other data sets should be treated with caution.

Our study shows that in clinical practice, a rapid increase in quality of life can be detected by the SAQ and SAQ-global. The easy use of the SAQ-global makes it a suitable tool if time is limited. The EQ-5D should not be used in clinical practice to evaluate change in quality of life in severe asthma as it is comparatively insensitive to change, and the use of this questionnaire in economic decision making may lead to an underestimation of the benefit of biologic treatment.

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References

- Hyland ME. Selection of items and avoidance of bias in quality of life scales. *Pharmacoeconomics* 1992; 1(3): 182–190.
- Juniper EF, O'byrne PM, Guyatt GH, et al. Development and validation of a questionnaire to measure asthma control. *Eur Respir J* 1999; 14: 902–907.
- Thamrin C, Nydegger R, Stern G, et al. Associations between fluctuations in lung function and asthma control in two

- populations with differing asthma severity. *Thorax* 2011; 66: 1036–1042.
4. Huang IC, Lee JL, Ketheeswaran P, et al. Does personality affect health-related quality of life? A systematic review. *PLoS One* 2017; 12(3): e0173806.
 5. Niedzwiedz CL, Katikireddi SV, Pell JP, et al. Life course socio-economic position and quality of life in adulthood: a systematic review of life course models. *BMC Public Health* 2012; 12(1): 628.
 6. Helgeson VS. Social support and quality of life. *Qual Life Res* 2003; 12(Suppl 1): 25–31.
 7. Hyland ME, Jones RC, Lanario JW, et al. The construction and validation of the Severe Asthma Questionnaire. *Eur Respir J* 2018; 52: 1800618.
 8. Janssen MF, Pickard AS, Golicki D, et al. Measurement properties of the EQ-5D-5L compared to the EQ-5D-3L across eight patient groups: a multi-country study. *Qual Life Res* 2013; 22(7): 1717–1727.
 9. Whalley D, Globe G, Crawford R, et al. Is the EQ-5D fit for purpose in asthma? Acceptability and content validity from the patient perspective. *Health Qual Life Outcomes* 2018; 16(1): 160–164.
 10. Kamper SJ, Maher CG and Mackay G. Global rating of change scales: a review of strengths and weaknesses and considerations for design. *J Man Manipulative Ther* 2009; 17(3): 163–170.
 11. Hyland ME, Kenyon CA, Allen R, et al. Diary keeping in asthma: comparison of written and electronic methods. *BMJ* 1993; 306(6876): 487–489.
 12. NHS England. Specialised respiratory services (adult)–severe Asthma, 2018.
 13. Lanario JW, Hyland ME, Menzies-Gow A, et al. Validation of subscales of the severe asthma questionnaire (SAQ) using exploratory factor analysis (EFA). *Health Qual Life Outcomes* 2020; 18(1): 336–340.
 14. Devlin NJ, Shah KK, Feng Y, et al. Valuing health-related quality of life: an EQ-5D-5L value set for England. *Health Econ* 2018; 27(1): 7–22.
 15. Cohen J. A power primer. *Psychol Bull* 1992; 112(1): 155–159, DOI: [10.1037/0033-2909.112.1.155](https://doi.org/10.1037/0033-2909.112.1.155).
 16. Rutten-van Mölken MP, Custers F, Van Doorslaer EK, et al. Comparison of performance of four instruments in evaluating the effects of salmeterol on asthma quality of life. *Eur Respir J* 1995; 8(6): 888–898.
 17. Sanjuás C, Alonso J, Prieto L, et al. Health-related quality of life in asthma: a comparison between the St George’s respiratory Questionnaire and the Asthma Quality of Life Questionnaire. *Qual Life Res* 2002; 11(8): 729–738.
 18. Szentes BL, Schultz K, Nowak D, et al. How does the EQ-5D-5L perform in asthma patients compared with an asthma-specific quality of life questionnaire? *BMC Pulm Med* 2020; 20(1): 168–171.
 19. Hyland ME and Sodergren SC. Development of a new type of global quality of life scale, and comparison of performance and preference for 12 global scales. *Qual Life Res* 1996; 5(5): 469–480.
 20. Norman G. Hi! How are you? response shift, implicit theories and differing epistemologies. *Qual Life Res* 2003; 12: 239–249.
 21. Robinson J, Khan N, Fusco L, et al. Why are there discrepancies between depressed patients’ global Rating of change and scores on the Patient Health Questionnaire depression module? a qualitative study of primary care in England. *BMJ open* 2017; 7(4): e014519.
 22. Schmitt J and Di Fabio RP. The validity of prospective and retrospective global change criterion measures. *Arch Phys Med Rehabil* 2005; 86(12): 2270–2276.
 23. Kahneman D. *Thinking, fast and slow*. New York: Macmillan, 2011.