

Expert Consensus on the Tapering of Oral Corticosteroids for the

Treatment of Asthma: a Delphi Study (100/100 characters)

Carey M. Suehs¹, Andrew Menzies-Gow², David Price^{3,4}, Eugene R. Bleecker⁵,
Giorgio Walter Canonica⁶, Mark Gurnell⁷, Arnaud Bourdin^{1,8} on behalf of the
Oral Corticosteroids Tapering Delphi Expert Panel*

¹Department of Respiratory Diseases, University of Montpellier, Centre Hospitalier Universitaire Montpellier, Montpellier, France; ²Royal Brompton and Harefield National Health Service Foundation Trust, London, United Kingdom; ³Observational and Pragmatic Research Institute, Singapore; ⁴Centre of Academic Primary Care, Division of Applied Health Sciences, University of Aberdeen, Aberdeen, United Kingdom; ⁵University of Arizona Department of Medicine, Tucson, Arizona; ⁶Personalised Medicine, Asthma and Allergy Center, Humanitas University and IRCCS Research Hospital, Milan, Italy; ⁷Wellcome Trust – Medical Research Council Institute of Metabolic Science and Cambridge National Institute for Health Research Biomedical Research Centre, University of Cambridge and Addenbrooke's Hospital, Cambridge, United Kingdom; ⁸PhyMedExp, University of Montpellier, Institut National de la Santé et de la Recherche Médicale, Centre National de la Recherche Scientifique, Centre Hospitalier Universitaire Montpellier, Montpellier, France

*A complete list of members of the Oral Corticosteroids Tapering Delphi Expert Panel may be found in the online supplement

Degrees and ORCID identifiers

Carey M. Suehs (PhD): 0000-0002-2175-3496
Andrew Menzies-Gow (MB BS, PhD): 0000-0001-9707-4986
David Price (FRCGP): 0000-0002-9728-9992
Eugene R. Bleecker (MD): 0000-0002-4767-3494
Giorgio Walter Canonica (MD): 0000-0001-8467-2557
Mark Gurnell (MD, PhD): 0000-0001-5745-6832
Arnaud Bourdin (MD, PhD): 0000-0002-4645-5209

Correspondence and requests for reprints should be addressed to: Carey M Suehs, Ph.D., Department of Respiratory Diseases, Hôpital Arnaud de Villeneuve, 371, av. du Doyen Gaston Giraud, F-34295 Montpellier Cedex 5, France.

E-mail: c-suehs@chu-montpellier.fr

Tel: +33 4.67.33.35.03

Fax: +33 4.67.33.58.27

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What is the current scientific knowledge on this subject?

Cumulative oral corticosteroid treatment for asthma is associated with costly and burdensome side effects and comorbidities. ‘OCS stewardship’ is advocated to protect patients from inappropriate OCS use and its consequences. The advent of effective OCS-sparing biological therapies also fosters new opportunities for tapering. Currently, evidence-based guidelines for OCS use, tapering, and associated comorbidity screening in asthma are lacking.

What does this study add to the field?

In the absence of clinical data to develop evidence-based guidelines, this modified Delphi consensus study brought together experts with relevant knowledge and clinical experience to generate a high-quality expert consensus statement on OCS use and tapering. The recommendations thus generated support minimizing OCS use in as much as possible. A cumulative yearly dose of 0.5 or 1g prednisolone equivalents would be indicative of poor asthma control. They also provide a first step towards development of an OCS tapering algorithm, as well as a minimum OCS adverse event screening list. Little consensus was achieved concerning the assessment and management of adrenal insufficiency, supporting a need for future related research in this specific domain. Finally, the experts strongly support shared decision making during OCS tapering.

Abstract

Rationale: There is a need to minimize oral corticosteroid use in patients with asthma to prevent their costly and burdensome adverse effects. Current guidelines do not provide recommendations for oral corticosteroid tapering in patients with asthma.

Objectives: To develop expert consensus on oral corticosteroid tapering among international experts.

Methods: A modified Delphi method was used to develop expert consensus statements relating to oral corticosteroid use, tapering, adverse effects, adrenal insufficiency, and patient-physician shared decision-making. Initial statements proposed by experts were categorized, filtered for repetition, and presented back to experts over three ranking rounds to obtain consensus ($\geq 70\%$ agreement).

Measurements and Main Results: 131 international experts participated in the study and 296 statements were ranked. Numerous recommendations and guidance regarding appropriate oral corticosteroid use were established. Experts agreed that oral corticosteroid tapering should be attempted in all patients with asthma receiving maintenance oral corticosteroid therapy, with personalization of tapering rhythm and speed. The importance of recognizing individual adverse effects was also established; however, a unified approach to the assessment of adrenal insufficiency was not reached. Shared decision-making was considered an important goal during the tapering process.

Conclusion: In this Delphi study expert consensus statements were generated on oral corticosteroid use, tapering, adverse effects screening, and shared decision-making, which may be used to inform clinical practice. Areas of non-consensus were identified, highlighting

uncertainty among the experts around some aspects of oral corticosteroid use in asthma, such as adrenal insufficiency, which underscores the need for further research in these domains.

Abstract word count: 247/250

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1 Introduction

2 Asthma is a chronic inflammatory disease, characterized by reversible airway obstruction and
3 airway hyperresponsiveness (1), which affects ~339 million individuals worldwide (2).
4 Approximately 5–10% of the overall asthma population have severe asthma (3), defined as
5 uncontrolled asthma despite adherence to maximal optimized therapy and treatment of
6 contributory factors (4). Severe asthma is associated with greater asthma-related morbidity,
7 increased healthcare costs, more frequent exacerbations, and greater oral corticosteroid
8 (OCS) use compared with mild/moderate asthma (5–8).

9 Early use of OCS in emergency department asthma treatment reduces hospital
10 admission rates (9), supporting its routine guideline-recommended use for asthma
11 exacerbations (4). Indeed, during acute exacerbations OCS have been observed to provide
12 rapid benefit (10). Nevertheless, such benefits may be dose- or duration-dependent, and the
13 current guidance remains somewhat empirical. Long-term, low-dose OCS add-on therapy is
14 restricted to Global Initiative for Asthma (GINA) Step 5 and positioned after trials of other
15 more preferential add-on treatments (e.g. tiotropium and biologicals), with consideration of
16 side effects (4). However, long-term OCS therapy continues to be widely used in severe
17 asthma, with global usage estimated at 20–60% (11).

18 Recent studies across multiple therapy areas demonstrate that cumulative OCS use
19 (including long-term and intermittent use) is associated with a dose- and duration-dependent
20 risk of potentially serious adverse effects including osteoporosis, hypertension, diabetes
21 mellitus, cataracts, fractures, obesity, and gastrointestinal disorders(6, 11–13). Risk of
22 adverse effects is evident at relatively low cumulative and mean daily OCS doses (12).
23 Furthermore, long-term OCS use is associated with increased risk of mortality, reduced
24 quality of life, and increased healthcare resource utilization and costs (5, 6, 14–16).

25 The costly and burdensome adverse effects associated with OCS use have prompted
26 international respiratory experts to call for a structured ‘OCS stewardship’ approach to
27 protect patients from inappropriate OCS use and its consequences (16, 17). Tapering has been
28 strengthened by the availability of effective OCS-sparing biological therapies; however, the
29 process should still be approached with caution to prevent symptom recurrence and to avoid
30 risking unrecognized adrenal insufficiency (12, 18). Reporting on successful OCS tapering
31 protocols is most often indirect (i.e. the tapering algorithm is not the subject of study per se)
32 and results in a diverse selection of study-specific algorithms (19–26), whose detail varies
33 significantly between published studies. Current recommendations (4, 27) do not provide
34 guidance on the choice of OCS tapering protocol or otherwise how to taper. From a clinical
35 perspective, the lack of asthma-specific guidelines on OCS tapering and the systematic
36 screening of adverse events represents a key barrier to reducing OCS use (16). In the absence
37 of clinical data to develop evidence-based guidelines, this modified Delphi consensus study
38 aimed to bring together experts with relevant knowledge and clinical experience to generate a
39 high-quality expert consensus statement on OCS use and tapering.

40

41 **Methods**

42 **Study Design**

43 An international panel of experts participated in a four-round Delphi study to develop a
44 systematic consensus on OCS tapering. The protocol was approved by the Institutional
45 Review Board, University Hospital of Montpellier (reference number: 2019 IRB-MTP 04-12)
46 and was registered on clinicaltrials.gov (NCT03934801). Surveys were administered
47 anonymously to the expert panel using SurveyMonkey online software

48 (www.surveymonkey.co.uk). Statistical analyses were performed using the R programming
49 environment version 3.6.1 (28).

50 **Participants and Expert Recruitment**

51 The study steering committee (ERB, AB, GWC, MG, AMG, DP) provided initial
52 recommendations of experts (based on their professional/association networks) to be invited
53 to enroll in the study and eligible/responding experts were asked to recommend additional
54 experts in the field. Pulmonologists/respiratory disease specialists, allergists,
55 endocrinologists, pediatricians, rheumatologists, and patient advocacy organization
56 representatives were eligible for study enrollment. Clinicians were required to manage
57 patients on a weekly basis and have clinical experience in managing disease following OCS
58 tapering/withdrawal to ensure a high-level knowledge in OCS management. Patient advocacy
59 organization representatives were required to represent an asthma patient advocacy group.
60 Experts were excluded if they were currently, or due to be (in the following 12 months)
61 employed by, or had ownership in a pharmaceutical company. Participants were encouraged
62 to provide complete responses to all survey rounds and reminders were delivered daily.

63 **Round 1: Expert Demographics and Brainstorming**

64 Participants completed an electronically administered questionnaire to provide demographic
65 information, including age, sex, qualifications, practice environment, specialty, years since
66 training completion, time spent caring for patients treated with OCS, and number of patients
67 seen per year. To initiate the brainstorming process, the questionnaire included open-ended
68 questions to generate an initial list of statements pertaining to six categories: appropriate OCS
69 use, OCS tapering, addressing adverse effects, adrenal insufficiency, patient-physician shared
70 decision-making, and other aspects they felt to be important. Experts were informed that all

71 OCS dosages should be expressed as prednisone-equivalent dosages, as reported in GINA
72 guidelines (4). Raw statements (which refer to adult patients unless otherwise indicated) were
73 categorized, filtered to avoid repetition, and amended for clarity (if necessary) to generate a
74 final list of statements for ranking. The demographics/brainstorming and ranking
75 questionnaires are available on the Open Science Framework platform (<https://osf.io/wrdbu/>).

76 **Rounds 2, 3, and 4: Ranking**

77 The final list of statements was presented to experts for ranking using a pre-defined Likert
78 scale ranging from ‘strongly disagree’ (−2 points) to ‘strongly agree’ (+2 points). Experts
79 were also asked to select specific responses for treatment duration, threshold values, and
80 assessment frequencies. A stopping rule was enforced for a given statement when $\geq 70\%$ of
81 experts indicated ‘agree’ or ‘strongly agree’ (positive consensus), or when $\geq 70\%$ indicated
82 ‘disagree’ or ‘strongly disagree’ (negative consensus) during any round. For statements
83 requiring a specific response consensus was defined as 70% of experts providing an identical
84 response. Items that achieved consensus were not re-presented in subsequent ranking rounds.

85

86 **Results**

87 **Participants**

88 Of the 363 experts invited to participate in this Delphi study, 169 were enrolled in the expert
89 panel and 131 completed at least one of the four survey rounds (Figure 1A). Participant
90 attrition rates during the ranking process were low; of the 108 experts who participated in the
91 first ranking round, 96 proceeded to complete all three rounds of ranking (Figure 1B). Most
92 experts were pulmonologists (73%) or allergists (18%); however, a wide range of specialties

93 were represented in the study. Demographics and professional characteristics of the expert
94 panel are provided in Table 1.

95 **Ranking Results**

96 The initial brainstorming survey was completed over a 2-month period (April–May 2019) and
97 three rounds of ranking surveys (rounds 2–4) were completed between 31 August and
98 26 September 2019. Ninety-one experts provided at least one brainstorming statement and
99 1447 statements were generated in total. Raw statements were categorized and filtered to
100 avoid repetition resulting in a final list of 296 statements. The following sections summarize
101 key points of consensus, but do not cover all 296 items presented to the experts. Full ranking
102 results for all 296 statements are available in the online supplement (pp 1–21).

103 **Section 1: Appropriate OCS Use for the Treatment of Asthma**

104 Over 95% of experts agreed or strongly agreed with the following statement: *‘In general, our*
105 *goal should be not to use OCS. When nevertheless required, dose and duration should be*
106 *minimized.’*

107 **Short-term OCS use:** Positive consensus was reached for five out of six statements
108 regarding appropriate short-term OCS use (online supplement p 1; 1.2.a–f). Short-term OCS
109 use (<15 days) was deemed appropriate in patients experiencing acute non-resolving or life-
110 threatening exacerbations and in patients experiencing eosinophilic or allergic exacerbations.
111 Experts also agreed that short-term OCS use was appropriate within an asthma management
112 plan or to avoid hospitalization. No consensus was reached on whether short-term OCS use
113 was appropriate to palliate unavailability of hospitalization services. Experts agreed that 5–7
114 days constitutes the usual maximal duration for a short course of OCS for treatment of an
115 exacerbation and that the optimal dosage of a short course of OCS should be 0.5 mg/kg/day.

116 Items that remained controversial included whether dosages for short courses of OCS for
117 treatment of an exacerbation should remain stable and whether the need for individual
118 tailoring of OCS dose would render systematic application of ‘ideal’ doses unlikely.

119 **Maintenance OCS use:** Nine statements were proposed regarding appropriate use of
120 OCS as a maintenance (long-term) treatment, with five statements reaching consensus (online
121 supplement pp 1–2; 1.6.a–i). Maintenance OCS use was considered appropriate in patients
122 with severe asthma experiencing inadequate control despite optimization of GINA Step 5
123 treatments, or when adverse effects that could not be managed by another treatment presented
124 during a tapering attempt. Consensus was also reached on eight of 13 statements
125 characterizing an adequate response to long-term OCS use (online supplement pp 2–3; 1.9.a–
126 m). In situations where OCS maintenance treatment is appropriate, experts considered ≤ 5
127 mg/day to be an acceptable dose (Figure 2).

128 Maintenance OCS use remained controversial in the context of adrenal insufficiency
129 and to reduce overall OCS exposure. There was no consensus on whether maintenance OCS
130 use is appropriate based on a patient’s T2 phenotype.

131 Over 90% of experts agreed or strongly agreed that the annual cumulative OCS dose
132 should be monitored as a marker of asthma control. Over 75% of experts selected a threshold
133 of 0.5 g or 1 g as the annual cumulative OCS dose indicative of poor control in ranking
134 round 3 (Figure 3).

135 It was agreed that biological therapies are useful OCS-sparing agents, and patients
136 should be systematically assessed for suitability for biological therapy. Daily OCS dose may
137 represent a reliable marker for the evaluation of biological treatment response (online
138 supplement p 5; 1.16.g; 1.17.a–c).

139 **Section 2: OCS Tapering**

140 Two general statements reached positive consensus in the first round of ranking: 1) *‘Tapering*
141 *(down to a minimal efficacious dose or complete weaning, if possible) should be attempted in*
142 *all asthma patients receiving maintenance OCS therapy, regardless of comorbidities’*; 2)
143 *‘The rhythm and speed of OCS tapering requires individualization for each patient.’*

144 Multiple statements reached positive consensus on when it may be appropriate to
145 attempt OCS tapering, and when cautious slow attempts of tapering and complete OCS
146 cessation are appropriate (Table 2). Tapering was deemed appropriate in multiple cases
147 (online supplement p 5; 2.2.a–f) including: if the intensity or duration of OCS use is a cause
148 for concern, if a patient exhibits OCS-related adverse effects or a lack of response to OCS,
149 holds a reasonable likelihood of hypothalamic-pituitary-adrenal-axis recovery, or experiences
150 improved asthma control following initiation of biological therapy. Tapering was also
151 deemed appropriate in patients experiencing asthma control with OCS maintenance therapy
152 for a minimum agreed-upon time; however, the duration of the minimum length of time
153 remained controversial.

154 Tapering attempts were deemed inappropriate in patients with eosinophilic
155 granulomatosis with polyangiitis or allergic bronchopulmonary aspergillosis that relapses
156 during tapering (online supplement p 6; 2.4.b–c). Further statements that remained
157 controversial included tapering in patients who demonstrated potentially harmful effects
158 during previous tapering attempts and whether tapering should be attempted in patients with
159 adrenal insufficiency (online supplement pp 5–6; 2.4.a,d).

160 Experts agreed that OCS tapering should incorporate some aspect of individualization
161 and multiple factors were considered that may influence the rhythm and speed of OCS

162 tapering (online supplement p 6; 2.5.a–g); such factors included: duration of previous
163 maintenance OCS treatment, history and future risk of adverse effects, and type of adverse
164 effect present. Three statements that remained controversial concerned the speed of OCS
165 tapering in patients with a fast/slow response to OCS, whether OCS tapering should be
166 guided by biomarkers at each weaning step, and whether the speed of tapering should be
167 dependent on the known rapidity of action of the OCS-sparing drug introduced.

168 Five statements concerning characteristics of an acceptable OCS tapering algorithm
169 reached positive consensus, and three statements remained controversial (Table 3). Experts
170 agreed that biologicals should play an important role in OCS tapering and that failure to
171 achieve a $\geq 50\%$ OCS dose reduction indicates failure of the biological and may warrant
172 switching strategies (online supplement p 9; 2.11.c,e); furthermore, when writing
173 prescriptions, the option to reduce dose should be considered (online supplement p 9; 2.12.c).

174 **Section 3: Addressing OCS-Related Adverse Effects**

175 All five general statements concerning adverse effects reached positive consensus in the first
176 round of ranking (online supplement p 9; 3.1.a–e). Experts agreed that patients receiving
177 OCS were at greater risk of adverse effects compared with patients receiving no OCS, and
178 adverse effects should always be addressed, but should not preclude attempting to taper OCS
179 to the lowest possible dose.

180 Experts reached positive consensus on two of three adverse effect subsets for whom
181 OCS tapering should be a priority (online supplement pp 10–11; 3.4.a–c). A positive
182 consensus was achieved in the first round of ranking for seven of ten elements that should be
183 included in a minimum checklist for adverse effect screening in patients receiving OCS
184 therapy, and three statements remained controversial (Table 4).

185 **Section 4: Managing Adrenal Insufficiency**

186 The majority of statements (44/55 [80%]) concerning adrenal insufficiency failed to reach
187 consensus following three ranking rounds. Many statements that remained controversial
188 concerned the sub-populations in which adrenal insufficiency should be assessed (online
189 supplement pp 13–14; 4.3.a–f, 4.4.a–d). Experts agreed that adrenal insufficiency should be
190 assessed in individuals on regular, long-term OCS therapy. Additionally, a positive consensus
191 was almost reached (69% agreement) on statements indicating that adrenal insufficiency
192 should be assessed in patients exceeding an OCS dose of >2 g per year or >four repeated
193 OCS short courses per year.

194 Experts agreed that adrenal insufficiency is inadequately assessed (online supplement
195 p 16; 4.11.a) and should be assessed regularly (online supplement p 121; 4.1.a) and when
196 OCS tapering has failed in OCS-treated patients (online supplement p 14; 4.5.b). Experts also
197 agreed that signs of adrenal insufficiency should be symptomatically treated as much as
198 possible during the tapering process and should not be viewed as a reason to give up on
199 tapering altogether (online supplement p 12; 4.1.b). Experts agreed that adrenal insufficiency
200 should be assessed using fasting morning cortisol and in case of intermediate results, follow
201 up with a (short) tetracosactide/cosyntropin (e.g. Synacthen®) test (online supplement p 15;
202 4.9.c). An additional general statement regarding whether hydrocortisone replacement is
203 preferred to continued prednisolone almost reached positive consensus, with 65% of experts
204 agreeing with the statement and 8% disagreeing; the remaining percentage remained neutral
205 on the subject (online supplement p 12; 4.1.c).

206 Consensus was reached on the need for the treating respiratory physician to assess for
207 adrenal insufficiency in patients with severe asthma, and that management of adrenal

208 insufficiency in patients with severe asthma should involve an endocrinologist or a
209 multidisciplinary approach (online supplement p 20; 6.1.c,d).

210 **Section 5: Patient-Physician Shared Decision-Making**

211 Experts agreed that shared decision-making should be a systematic practice and self-
212 management should be limited to individuals with good levels of comprehension (online
213 supplement p 17; 5.1.a,d). Eight statements achieved positive consensus on the importance of
214 shared decision-making (online supplement p 17; 5.2.a–h) and 14 statements reached positive
215 consensus concerning the content to be included in the shared decision-making process
216 (online supplement pp 17–18; 5.3.a–n).

217 **Section 6: Miscellaneous**

218 Experts agreed that primary care physicians prescribing at least three courses of OCS/year to
219 a patient should consider specialist referral (online supplement p 20; 6.2.a). Experts also
220 achieved positive consensus on 16/17 statements concerning future research of OCS tapering
221 (online supplement pp 20-21; 6.3.a–q). The only subject that remained controversial
222 concerning future work was the efficacy of internet-provided algorithms for delivering
223 symptom-driven OCS tapering guidance to asthma patients.

224

225 **Discussion**

226 This Delphi study generated expert consensus and recommendations on numerous statements
227 concerning appropriate OCS use, OCS tapering, adverse effects, patient-physician shared
228 decision-making, and future research domains. Consensus was reached on general statements
229 concerning adrenal insufficiency; however, beyond generalities, consensus was not reached.

230 Hence, improving the assessment of adrenal insufficiency was one of multiple domains
231 identified as requiring future research.

232 To our knowledge, no existing asthma-specific guidelines are currently available to
233 guide OCS tapering in clinical practice. Consensus stated that tapering should be attempted in
234 all asthma patients receiving maintenance OCS therapy, regardless of comorbidities;
235 however, speed and rhythm of tapering should be individualized. Furthermore, expert
236 consensus was reached on characteristics of an acceptable OCS tapering algorithm (Table 3),
237 which constitutes a first step towards the development of OCS tapering algorithms for use in
238 clinical practice. These consensus and related information are summarized in Figure 4.

239 Successful OCS tapering algorithms have been reported in the past (19–25, 29, 30), but
240 vary greatly in content and reporting quality. Currently, the most detailed and recent OCS
241 tapering algorithm is being tested in the eagerly awaited PONENTE study (26). Certain
242 previous studies also demonstrate that prescribing treatment guided by eosinophil levels can
243 improve control, whilst simultaneously resulting in some corticosteroid sparing (31–33).
244 Current GINA guidelines suggest OCS dose adjustment may be supported by internet-based
245 monitoring of symptom control and exhaled nitric oxide; however, the latter contributed little
246 to algorithm decisions, in favor of ACQ scores (34). In the current study, only asthma control
247 questionnaires reached positive consensus as a useful tool during OCS tapering. The need for
248 laboratory tests or at-home lung function measurements may render many biomarker
249 approaches impractical for patients and clinicians. In addition, GINA recommends gradually
250 decreasing or stopping OCS in patients with a good response to biological therapies.
251 Successful corticosteroid reduction following initiation of biological therapies, using pre-set
252 tapering protocols, has been demonstrated in multiple studies (18). However, the latter are
253 often short-term in nature with little focus on adrenal function assessments, and the full

254 potential of tapering was therefore not achieved/documentated. As the use of biological
255 therapies increases, studies evaluating OCS tapering regimens on a longer basis, which can
256 be personalized based on factors such as baseline OCS dosage and level of asthma control,
257 will become increasingly important (e.g. the PONENTE study) (26). The current consensus
258 statement provides broader guidance on when and how to taper OCS in patients with asthma
259 (Figure 4), regardless of whether a biological therapy has been initiated.

260 Regarding appropriate OCS use, experts felt that long-term use is not appropriate in
261 situations where other treatment options are available. However, if no alternative treatment
262 options are available, experts considered ≤ 5 mg/day to be an acceptable dose. This threshold
263 is considerably lower than the definition in current GINA guidelines, which defines low-dose
264 maintenance OCS as ≤ 7.5 mg/day (4) and may result from the way the question was designed
265 to span the range of thresholds mentioned during the brain storming phase of the study. The
266 reader should note that non-consensus fractions of experts are willing to use 10 mg/day doses
267 and higher, suggesting that there is considerable non-guideline-conforming OCS usage in the
268 field, even among experts. The low consensus threshold at 5mg may also reflect the
269 increasing importance of biologics in the domain and the resulting opportunities for tapering
270 down to the lowest efficacious dose possible or complete cessation. Regardless, the reader
271 should also keep in mind that a 5mg/day OCS dose amounts to a cumulative dose exceeding
272 1.8 g/year.

273 In this study, when experts were asked to consider cumulative OCS doses, they voted
274 that 0.5 or 1 g/year would be indicative of poor asthma control. This would correspond to
275 approximately 3.5-7 months of maintenance treatment at 5 mg/day. A previous study by Price
276 et al demonstrated that diabetes associated with OCS use emerged at lifetime cumulative
277 systemic corticosteroid exposures of 0.5–<1 g, with most other adverse events emerging at

278 1.0 to <2.5 g (12). Furthermore, a 2020 study stated that a yearly cumulative OCS dose above
279 1 g should be considered unacceptable in severe asthma and indicates the need for specialist
280 referral (35). Even a short term use, which amounts to a median of 20 mg per day for
281 approximately 6-days in a large database study, is associated with an increase in sepsis,
282 venous thromboembolism, and fracture in the next 30 days (36). These studies highlight the
283 need for earlier specialist referral and earlier consideration of OCS-sparing strategies in
284 patients receiving OCS.

285 Biological therapies were a common subject among the experts and the initiation of a
286 successful biological therapy was the highest-ranked situation appropriate for initiating OCS
287 tapering (Table 2). The reader should keep in mind that there are other important reasons for
288 initiating tapering, such as side effects or non-response (Table 2). Key criteria for success of
289 a biological therapy include maintenance of asthma control, reduction in exacerbations, and
290 decrease in dose of OCS (27, 37). However, there is no clear guidance on the magnitude of
291 OCS reduction that constitutes success or failure of a biological therapy. In this study,
292 consensus stated that failure to achieve $\geq 50\%$ reduction in OCS indicates failure of the
293 biological therapy and may warrant a switch in strategy. The guidance provided here will
294 support clinical decision-making.

295 Items included on the minimal checklist for adverse effect screening (Table 4) have
296 been well documented in the literature among individuals receiving OCS. Early detection of
297 adverse effects has been shown to be important in the treatment and management of OCS-
298 related complications; the items on the checklist provide a basis for adverse effect screening
299 in clinical practice (6, 11, 12, 38). This checklist further underlines the importance of adverse
300 effect prevention measures, including calcium and vitamin D supplementation and

301 appropriate prescribing of bisphosphonates for osteoporosis, optimizing ICS dose and
302 medication adherence. The latter may additionally allow further reduction in OCS dose.

303 Previous studies have shown that adrenal insufficiency is common among frequent
304 users of OCS following tapering (39); however, lack of clear guidance for clinicians on how
305 to manage adrenal insufficiency may hinder OCS reduction in patients with severe asthma
306 (16). Experts agreed on the need to regularly assess for adrenal insufficiency, and that fasting
307 morning cortisol tests may be used (followed up with a (short) tetracosactide/cosyntropin test
308 in case of intermediate results). Experts also highlighted the need for a process to be in place
309 for referral to an endocrinologist alongside further research and potential education in this
310 domain. The majority of experts agreed use of hydrocortisone replacement therapy is
311 preferential to continued prednisolone use to aid the tapering process in the case of adrenal
312 insufficiency; however, consensus was not reached. The lack of consensus on this point is not
313 surprising given that the optimal strategy for glucocorticoid replacement in patients with
314 adrenal insufficiency remains controversial in the literature. In the UK, hydrocortisone is the
315 first-line treatment in management of adrenal insufficiency, followed by prednisolone (40).
316 Prednisolone is less expensive and some experts contend it may mimic the circadian rhythm
317 more closely than the standard thrice-daily hydrocortisone therapy; however, prednisolone
318 may also be associated with increased relative risk of cardiovascular disease (40–42). Results
319 of ongoing head-to-head studies will improve understanding regarding this issue (43).

320 Shared decision-making in OCS tapering was viewed positively by the experts. The
321 consensus was that although the OCS-tapering process should be primarily driven by the
322 physician, patients should contribute to the decision-making process and be educated on OCS
323 use and tapering. Patient's perceptions are frequently ambivalent towards OCS and how they
324 navigated previous tapering attempts should be taken into account. This is in line with

325 emerging evidence showing that shared decision-making is becoming more common in
326 asthma management and has been shown to improve patient adherence, outcomes, and
327 satisfaction with care (44). Shared decision making tools/platforms to facilitate this process
328 (e.g. 43, 44) require further development and validation for general asthma populations.

329 The strengths of this study include participation of 131 experts across a range of
330 specialisms, ensuring that a wide breadth of knowledge and relevant expertise was
331 represented among the expert panel. Results from this study also benefit from the anonymity
332 of expert responses, alongside a clear, *a priori* definition of consensus criteria and controlled
333 feedback. Importantly, a lack of participant attrition was observed throughout all three
334 ranking rounds, increasing the validity of the consensus by avoiding suppression of minority
335 opinions and minimizing potential for bias (47). A limitation of the study was the large
336 number of raw statements that needed to be reduced and summarized; therefore, statements
337 presented to experts were not fully representative of all the raw statements.

338 This Delphi consensus study provides expert consensus statements around OCS use and
339 tapering, which may be used to inform clinical practice and optimize management of patients
340 with severe asthma. The recommendations also provide a first step towards development of
341 an OCS tapering algorithm and support the ongoing OCS stewardship effort by international
342 respiratory experts to reduce the harm from inappropriate OCS use and its consequences.
343 While consensus was generated on numerous statements, many remained controversial,
344 highlighting the existing uncertainty, even among international experts, around certain
345 aspects of OCS use in asthma, such as assessment and management of adrenal insufficiency.
346 These findings underscore the need for further research to inform clinical practice and drive
347 future evidence-based guideline development.

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References

1. Holgate ST, Wenzel S, Postma DS, Weiss ST, Renz H, Sly PD. Asthma. *Nat Rev Dis Primers* 2015;1:15025.
2. Global Asthma Network. *The Global Asthma Report 2018*. 2018. at <<http://www.globalasthmareport.org>>.
3. Chung KF, Wenzel SE, Brozek JL, Bush A, Castro M, Sterk PJ, Adcock IM, Bateman ED, Bel EH, Bleecker ER, Boulet L-P, Brightling C, Chanez P, Dahlen S-E, Djukanovic R, Frey U, Gaga M, Gibson P, Hamid Q, Jajour NN, Mauad T, Sorkness RL, Teague WG. International ERS/ATS guidelines on definition, evaluation and treatment of severe asthma. *Eur Respir J* 2014;43:343–373.
4. Global Initiative for Asthma. *Global Strategy for Asthma Management and Prevention*. 2020. at <<https://ginasthma.org>>.
5. Canonica GW, Colombo GL, Bruno GM, Di Matteo S, Martinotti C, Blasi F, Bucca C, Crimi N, Paggiaro P, Pelaia G, Passalacqua G, Senna G, Heffler E, SANI Network. Shadow cost of oral corticosteroids-related adverse events: A pharmaco-economic evaluation applied to real-life data from the Severe Asthma Network in Italy (SANI) registry. *World Allergy Organ J* 2019;12:100007.
6. Ekström M, Nwaru BI, Hasvold P, Wiklund F, Telg G, Janson C. Oral corticosteroid use, morbidity and mortality in asthma: A nationwide prospective cohort study in Sweden. *Allergy* 2019;74:2181–2190.
7. Suruki RY, Daugherty JB, Boudiaf N, Albers FC. The frequency of asthma exacerbations and healthcare utilization in patients with asthma from the UK and USA. *BMC Pulm Med* 2017;17:74.

8. Nagase H, Adachi M, Matsunaga K, Yoshida A, Okoba T, Hayashi N, Emoto K, Tohda Y. Prevalence, disease burden, and treatment reality of patients with severe, uncontrolled asthma in Japan. *Allergol Int* 2020;69:53–60.
9. Rowe BH, Spooner C, Ducharme FM, Bretzlaff JA, Bota GW. Early emergency department treatment of acute asthma with systemic corticosteroids. *Cochrane Database Syst Rev* 2001;CD002178.doi:10.1002/14651858.CD002178.
10. Tattersfield AE, Postma DS, Barnes PJ, Svensson K, Bauer CA, O’Byrne PM, Löfdahl CG, Pauwels RA, Ullman A. Exacerbations of asthma: a descriptive study of 425 severe exacerbations. The FACET International Study Group. *Am J Respir Crit Care Med* 1999;160:594–599.
11. Bleeker ER, Menzies-Gow AN, Price DB, Bourdin A, Sweet S, Martin AL, Alacqua M, Tran TN. Systematic Literature Review of Systemic Corticosteroid Use for Asthma Management. *Am J Respir Crit Care Med* 2020;201:276–293.
12. Price DB, Trudo F, Voorham J, Xu X, Kerkhof M, Ling Zhi Jie J, Tran TN. Adverse outcomes from initiation of systemic corticosteroids for asthma: long-term observational study. *J Asthma Allergy* 2018;11:193–204.
13. Liu D, Ahmet A, Ward L, Krishnamoorthy P, Mandelcorn ED, Leigh R, Brown JP, Cohen A, Kim H. A practical guide to the monitoring and management of the complications of systemic corticosteroid therapy. *Allergy Asthma Clin Immunol* 2013;9:30.
14. Lee H, Ryu J, Nam E, Chung SJ, Yeo Y, Park DW, Park TS, Moon J-Y, Kim T-H, Sohn JW, Yoon HJ, Kim S-H. Increased mortality in patients with corticosteroid-dependent asthma: a nationwide population-based study. *Eur Respir J* 2019;54:.

15. Hyland ME, Whalley B, Jones RC, Masoli M. A qualitative study of the impact of severe asthma and its treatment showing that treatment burden is neglected in existing asthma assessment scales. *Qual Life Res* 2015;24:631–639.
16. Chung LP, Upham JW, Bardin PG, Hew M. Rational oral corticosteroid use in adult severe asthma: A narrative review. *Respirology* 2020;25:161–172.
17. McBrien CN, Menzies-Gow A. Time to FOCUS on oral corticosteroid stewardship in asthma management. *Respirology* 2019;24:304–305.
18. Bourdin A, Husereau D, Molinari N, Golam S, Siddiqui MK, Lindner L, Xu X. Matching-adjusted comparison of oral corticosteroid reduction in asthma: Systematic review of biologics. *Clin Exp Allergy* 2020;50:442–452.
19. Cameron SJ, Cooper EJ, Crompton GK, Hoare MV, Grant IW. Substitution of beclomethasone aerosol for oral prednisolone in the treatment of chronic asthma. *Br Med J* 1973;4:205–207.
20. Bel EH, Wenzel SE, Thompson PJ, Prazma CM, Keene ON, Yancey SW, Ortega HG, Pavord ID, SIRIUS Investigators. Oral glucocorticoid-sparing effect of mepolizumab in eosinophilic asthma. *N Engl J Med* 2014;371:1189–1197.
21. Nair P, Wenzel S, Rabe KF, Bourdin A, Lugogo NL, Kuna P, Barker P, Sproule S, Ponnarambil S, Goldman M, ZONDA Trial Investigators. Oral Glucocorticoid-Sparing Effect of Benralizumab in Severe Asthma. *N Engl J Med* 2017;376:2448–2458.
22. Vogelmeier C, Kardos P, Hofmann T, Canisius S, Scheuch G, Muellinger B, Nocker K, Menz G, Rabe KF. Nebulised budesonide using a novel device in patients with oral steroid-dependent asthma. *Eur Respir J* 2015;45:1273–1282.
23. Braunstahl G-J, Chlumský J, Peachey G, Chen C-W. Reduction in oral corticosteroid use in patients receiving omalizumab for allergic asthma in the real-world setting. *Allergy Asthma Clin Immunol* 2013;9:47.

24. Lacronique J, Renon D, Georges D, Henry-Amar M, Marsac J. High-dose beclomethasone: oral steroid-sparing effect in severe asthmatic patients. *Eur Respir J* 1991;4:807–812.
25. Milgrom H, Fick RB, Su JQ, Reimann JD, Bush RK, Watrous ML, Metzger WJ. Treatment of allergic asthma with monoclonal anti-IgE antibody. rhuMAb-E25 Study Group. *N Engl J Med* 1999;341:1966–1973.
26. Menzies-Gow A, Corren J, Bel EH, Maspero J, Heaney LG, Gurnell M, Wessman P, Martin UJ, Siddiqui S, Garcia Gil E. Corticosteroid tapering with benralizumab treatment for eosinophilic asthma: PONENTE Trial. *ERJ Open Res* 2019;5:.
27. Chipps BE, Bacharier LB, Murphy KR, Lang D, Farrar JR, Rank M, Oppenheimer J, Zeiger RS. The Asthma Controller Step-down Yardstick. *Ann Allergy Asthma Immunol* 2019;122:241-262.e4.
28. R Core Team. *R: A Language and Environment for Statistical Computing*. Vienna, Austria: R Foundation for Statistical Computing; 2016. at <<https://www.R-project.org/>>.
29. Mullarkey MF, Lammert JK, Blumenstein BA. Long-term methotrexate treatment in corticosteroid-dependent asthma. *Ann Intern Med* 1990;112:577–581.
30. Nelson HS, Busse WW, deBoisblanc BP, Berger WE, Noonan MJ, Webb DR, Wolford JP, Mahajan PS, Hamedani AG, Shah T, Harding SM. Fluticasone propionate powder: oral corticosteroid-sparing effect and improved lung function and quality of life in patients with severe chronic asthma. *J Allergy Clin Immunol* 1999;103:267–275.
31. Chlumský J, Striz I, Terl M, Vondracek J. Strategy aimed at reduction of sputum eosinophils decreases exacerbation rate in patients with asthma. *J Int Med Res* 2006;34:129–139.

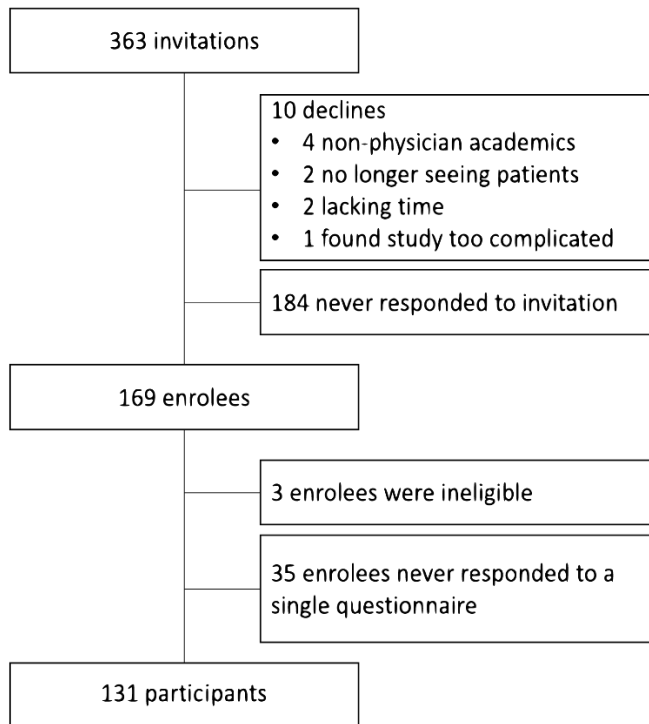
32. Green RH, Brightling CE, McKenna S, Hargadon B, Parker D, Bradding P, Wardlaw AJ, Pavord ID. Asthma exacerbations and sputum eosinophil counts: a randomised controlled trial. *Lancet* 2002;360:1715–1721.
33. Jayaram L, Pizzichini MM, Cook RJ, Boulet L-P, Lemièrè C, Pizzichini E, Cartier A, Hussack P, Goldsmith CH, Laviolette M, Parameswaran K, Hargreave FE. Determining asthma treatment by monitoring sputum cell counts: effect on exacerbations. *Eur Respir J* 2006;27:483–494.
34. Hashimoto S, Brinke AT, Roldaan AC, van Veen IH, Möller GM, Sont JK, Weersink EJM, van der Zee JS, Braunstahl G-J, Zwinderman AH, Sterk PJ, Bel EH. Internet-based tapering of oral corticosteroids in severe asthma: a pragmatic randomised controlled trial. *Thorax* 2011;66:514–520.
35. Bourdin A, Adcock I, Berger P, Bonniaud P, Chanson P, Chenivesse C, de Blic J, Deschildre A, Devillier P, Devouassoux G, Didier A, Garcia G, Magnan A, Martinat Y, Perez T, Roche N, Taillé C, Val P, Chanez P. How can we minimise the use of regular oral corticosteroids in asthma? *Eur Respir Rev* 2020;29:.
36. Waljee AK, Rogers MAM, Lin P, Singal AG, Stein JD, Marks RM, Ayanian JZ, Nallamothu BK. Short term use of oral corticosteroids and related harms among adults in the United States: population based cohort study. *BMJ* 2017;j1415.doi:10.1136/bmj.j1415.
37. Bousquet J, Brusselle G, Buhl R, Busse WW, Cruz AA, Djukanovic R, Domingo C, Hanania NA, Humbert M, Menzies Gow A, Phipatanakul W, Wahn U, Wechsler ME. Care pathways for the selection of a biologic in severe asthma. *Eur Respir J* 2017;50:.
38. Pisani P, Renna MD, Conversano F, Casciaro E, Muratore M, Quarta E, Paola MD, Casciaro S. Screening and early diagnosis of osteoporosis through X-ray and ultrasound based techniques. *World J Radiol* 2013;5:398–410.

39. Mortimer KJ, Tata LJ, Smith CJP, West J, Harrison TW, Tattersfield AE, Hubbard RB. Oral and inhaled corticosteroids and adrenal insufficiency: a case-control study. *Thorax* 2006;61:405–408.
40. Iqbal K, Halsby K, Murray RD, Carroll PV, Petermann R. Glucocorticoid management of adrenal insufficiency in the United Kingdom: assessment using real-world data. *Endocr Connect* 2019;8:20–31.
41. Williams EL, Choudhury S, Tan T, Meeran K. Prednisolone replacement therapy mimics the circadian rhythm more closely than other glucocorticoids. *The Journal of Applied Laboratory Medicine* 2016;1:152–161.
42. Quinkler M, Ekman B, Marelli C, Uddin S, Zelissen P, Murray RD, EU-AIR Investigators. Prednisolone is associated with a worse lipid profile than hydrocortisone in patients with adrenal insufficiency. *Endocr Connect* 2017;6:1–8.
43. ClinicalTrials.gov. Hydrocortisone vs prednisolone in AI (HYPER-AID). 2019;at <<https://clinicaltrials.gov/ct2/show/NCT03608943>>.
44. Blaiss MS, Steven GC, Bender B, Bukstein DA, Meltzer EO, Winders T. Shared decision making for the allergist. *Ann Allergy Asthma Immunol* 2019;122:463–470.
45. Tapp H, Shade L, Mahabaleshwarkar R, Taylor YJ, Ludden T, Dulin MF. Results from a pragmatic prospective cohort study: Shared decision making improves outcomes for children with asthma. *Journal of Asthma* 2017;54:392–402.
46. Fiks AG, Mayne SL, Karavite DJ, Suh A, O’Hara R, Localio AR, Ross M, Grundmeier RW. Parent-Reported Outcomes of a Shared Decision-Making Portal in Asthma: A Practice-Based RCT. *PEDIATRICS* 2015;135:e965–e973.
47. Gargon E, Crew R, Burnside G, Williamson PR. Higher number of items associated with significantly lower response rates in COS Delphi surveys. *Journal of clinical epidemiology* 2019;108:110–120.

Figure legends

Figure 1. (A) Study flow diagram. (B) Expert participation in three statement-ranking rounds.

A



B

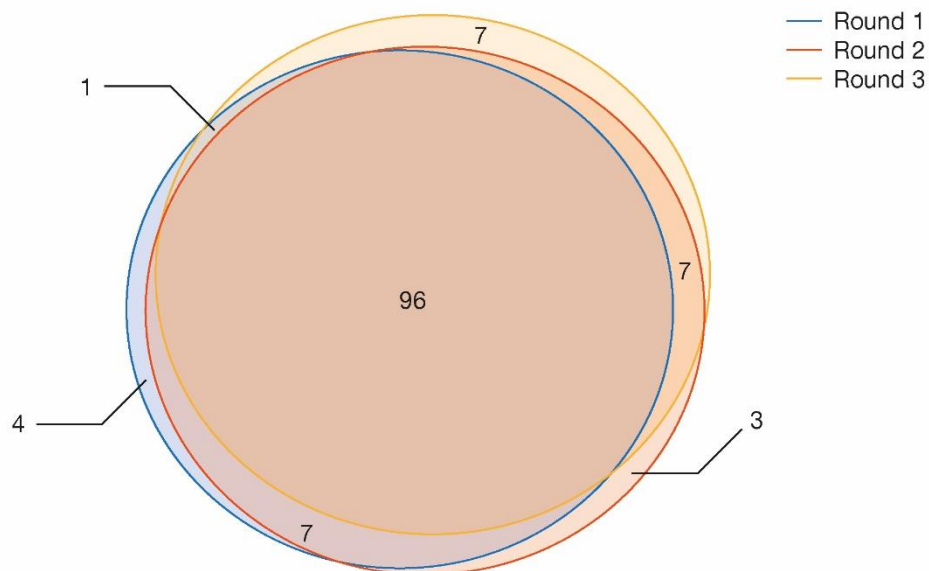


Figure 2. Percentage agreement among experts on acceptable doses for maintenance OCS treatment. OCS = oral corticosteroid.

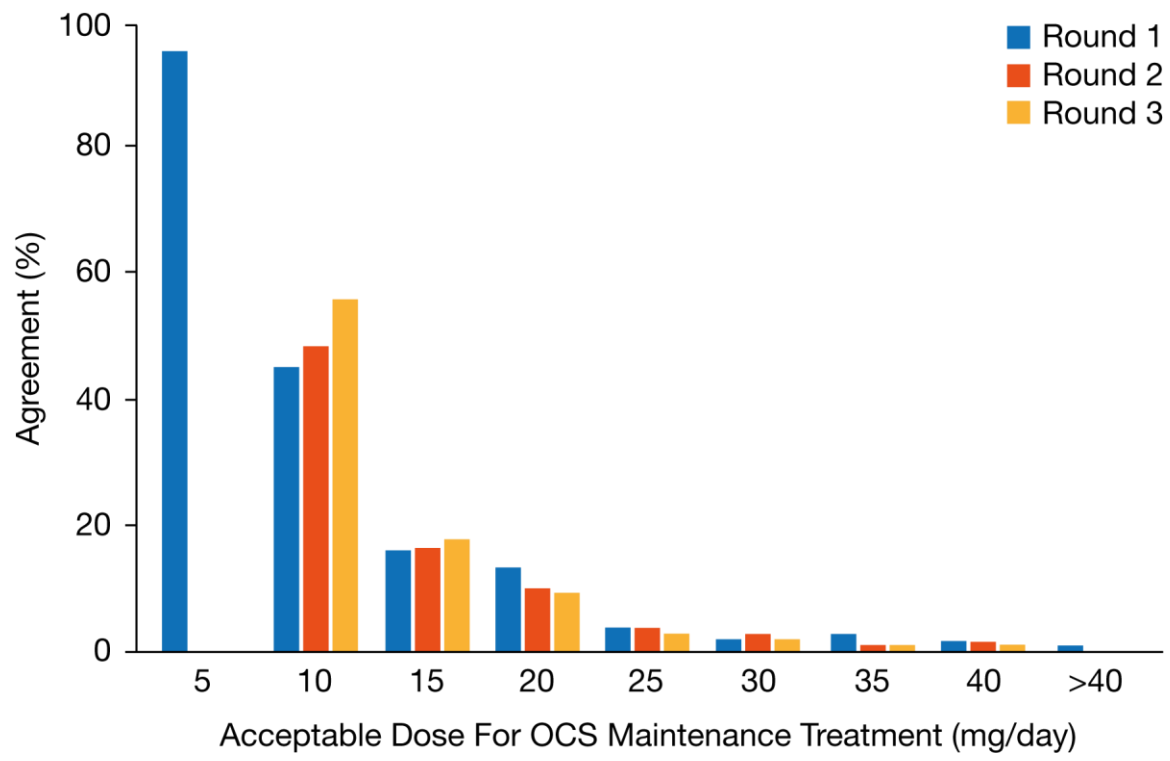


Figure 3. Percentage agreement among experts for threshold options indicating a yearly cumulative OCS dose that is suggestive of poor asthma control. NA = not applicable; OCS = oral corticosteroid.

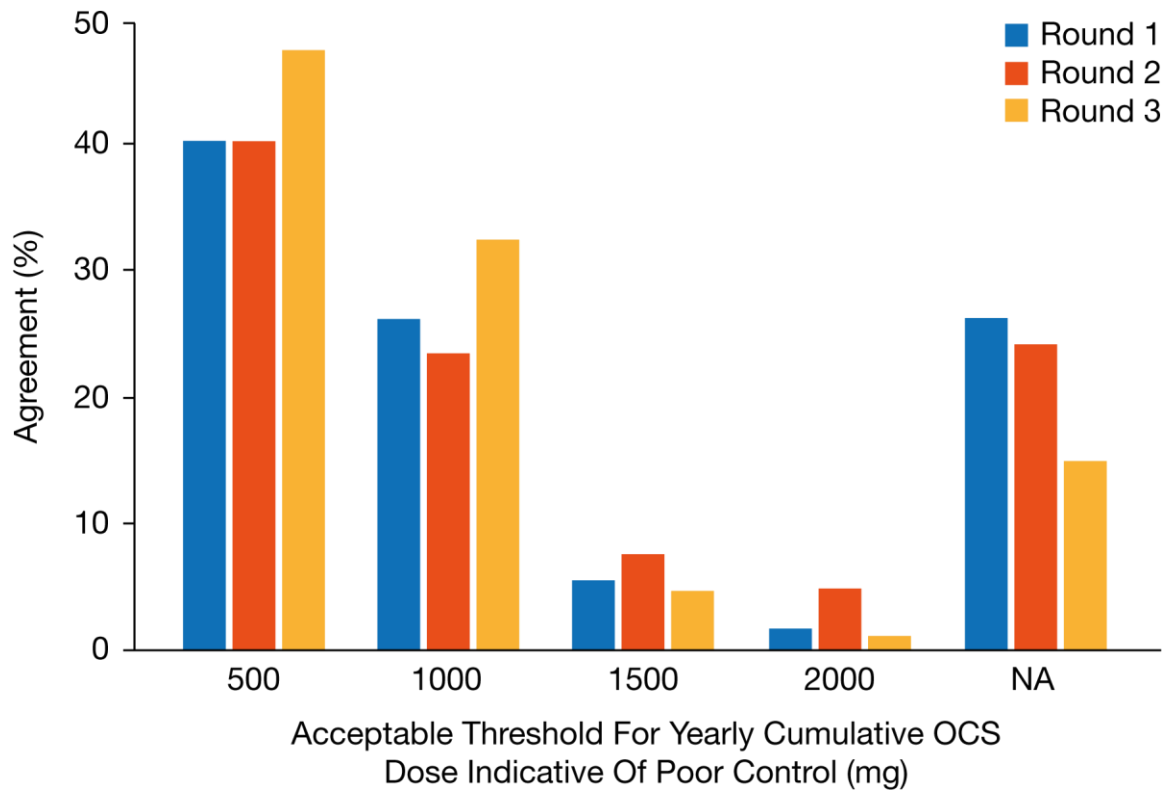
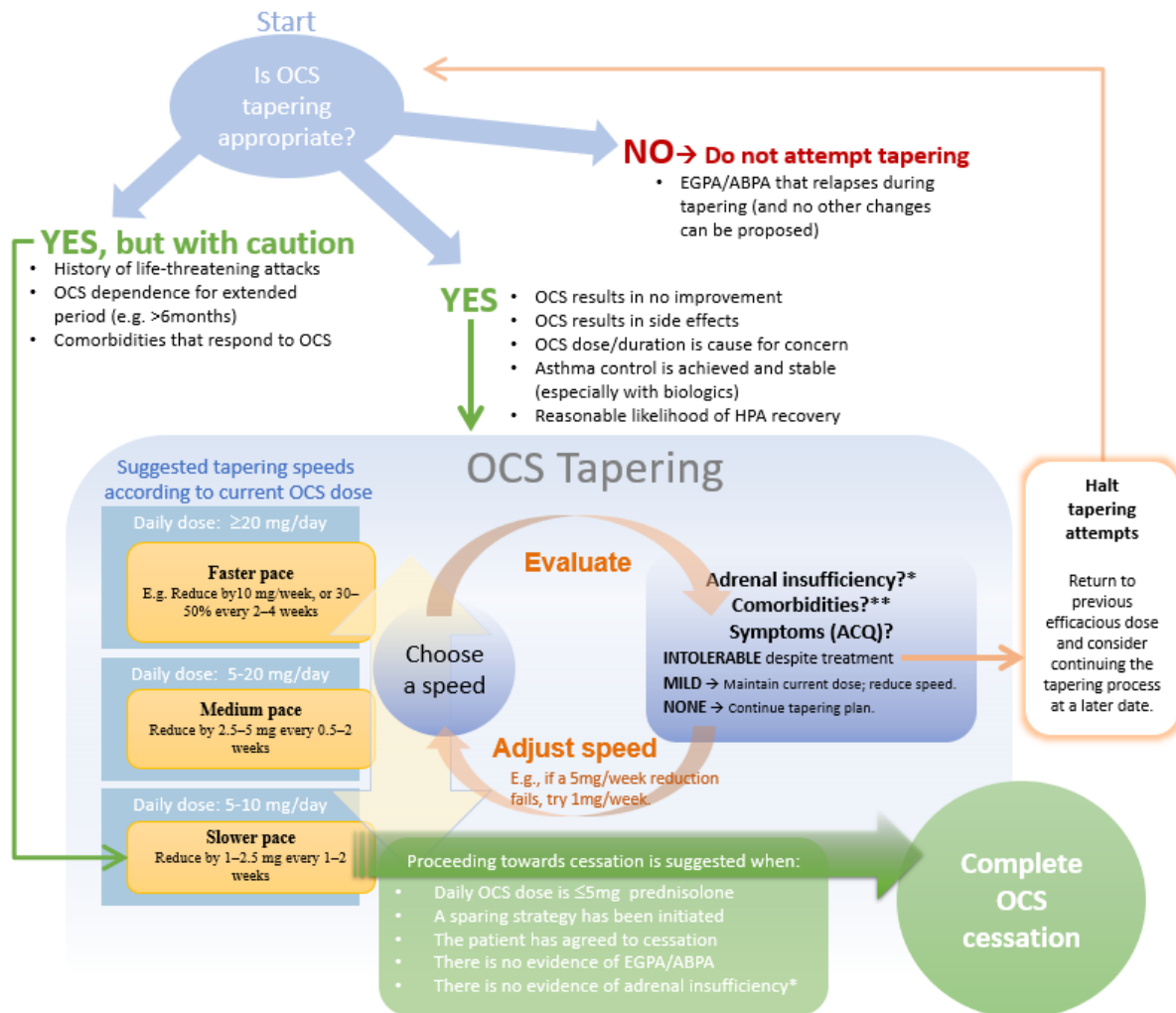


Figure 4. Graphic summary of consensus information on oral corticosteroid tapering.



*Adrenal insufficiency should be regularly assessed using fasting morning cortisol. In case of intermediate results, follow up with a (short) tetracosactide/cosyntropin test. Adrenal insufficiency management should be multidisciplinary, involving an endocrinologist.

**Comorbidity screening should include at least the following: glycemic control, bone density, blood pressure, cataracts and glaucoma, weight change, fracture risk score and

growth in pediatric populations. However, no consensus was achieved concerning the periodicity of comorbidity screening measures.

ABPA = allergic bronchopulmonary aspergillosis; ACQ = asthma control questionnaire; EGPA = eosinophilic granulomatosis with polyangiitis; HPA = hypothalamic–pituitary–adrenal axis; OCS = oral corticosteroids.

Table 1. Expert Panel Demographic Data

| Variable | Sample size (<i>n</i>) | Centrality |
|------------------------------|-------------------------------|-------------------|
| Age | 131 | 50.6 ± 9.64 |
| Sex (female) | 35/131 | 26.72% |
| Academic qualification | 131 | |
| MD (or equivalent) | 129 | 98.47% |
| PhD | 71 | 54.20% |
| Masters | 8 | 6.11% |
| Practice environment | 131 | |
| University hospital | 117 | 89.31% |
| Private practice | 11 | 8.40% |
| Academic environment | 37 | 28.24% |
| Patient care environment | 13 | 9.92% |
| Medical practice environment | 14 | 10.69% |
| Other | 7 | 5.34% |
| Specialties | 131 | |
| Allergist | 24 | 18.32% |
| Endocrinologist/Metabolic | 8 | 6.11% |
| Pediatrician | 1 | 0.76% |

| | | |
|---|-----|------------------|
| Patient advocacy organization representatives | 2 | 1.53% |
| Pulmonologist/Respiratory disease specialist | 95 | 72.52% |
| Rheumatologist | 1 | 0.76% |
| Years since completion of training | 131 | 19 (10 to 27) |
| Approximate % of work spent in caring for patients treated with OCS | 131 | 15 (5 to 30) |
| How often tapering is attempted in OCS patients | 131 | |
| NA (patient advocacy organization representative) | 2 | 1.53% |
| Occasionally | 4 | 3.05% |
| Frequently | 48 | 36.64% |
| Systematically | 77 | 58.78 |
| Participation in studies with aim of OCS tapering | 80 | 61.07% |
| Concerning OCS | | |
| Protocols, no. | 131 | 2 (1 to 4) |
| Scientific articles, no. | 131 | 2 (0 to 5) |
| Patients seen per year, no. | 131 | 50 (25 to 100) |
| Concerning asthma | | |
| Protocols, no. | 131 | 10 (4 to 20) |
| Scientific articles, no. | 131 | 30 (6 to 60) |
| Patients seen per year, no. | 131 | 300 (100 to 500) |

In all

| | | |
|-----------------------------|-----|-------------------|
| Protocols, no. | 131 | 20 (10 to 40) |
| Scientific articles, no. | 131 | 67 (25 to 132) |
| Patients seen per year, no. | 131 | 600 (400 to 1200) |

Definition of abbreviations: NA = not applicable; OCS = oral corticosteroid.

Table 2. Consensus Statements on OCS Tapering

| | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank* |
|--|-----------------------------|--------------------|-------------------|-----------------|--------------------------|----------------------------|
| Proceeding towards a tapering attempt is particularly appropriate when: | | | | | | |
| Biological treatment has been initiated and results in asthma control | 0.00 | 0.95 | 0.95 | 25.71 | 72.38 | 1.70 |
| The patient does not appear to respond to OCS treatment | 0.00 | 0.95 | 0.95 | 35.24 | 62.86 | 1.60 |
| A patient exhibits symptoms/comorbidities likely linked to OCS | 0.00 | 1.90 | 2.86 | 41.90 | 53.33 | 1.47 |
| Patients on maintenance OCS have gained control (for a minimum agreed-upon time) | 0.00 | 0.00 | 3.81 | 59.05 | 37.14 | 1.33 |
| The intensity or duration of OCS treatment gives reason for concern | 0.00 | 0.00 | 3.81 | 59.05 | 37.14 | 1.33 |
| There is reasonable likelihood of hypothalamic-pituitary-adrenal axis recovery | 0.00 | 1.90 | 11.43 | 54.29 | 32.38 | 1.17 |

Tapering should not be attempted in patients who:

| | | | | | | |
|--|------|------|-------|-------|-------|------|
| Have EGPA that relapses during tapering (and no other changes can be proposed) | 0.95 | 3.81 | 12.38 | 66.67 | 16.19 | 0.93 |
|--|------|------|-------|-------|-------|------|

| | | | | | | |
|--|------|------|-------|-------|------|------|
| Have ABPA that relapses during tapering (and no other changes can be proposed) | 0.00 | 9.52 | 19.05 | 61.90 | 9.52 | 0.71 |
|--|------|------|-------|-------|------|------|

Cautious slow tapering is particularly appropriate for patients who:

| | | | | | | |
|-----------------------------------|------|------|------|-------|-------|------|
| Have had life-threatening attacks | 0.95 | 3.81 | 3.81 | 60.00 | 31.43 | 1.17 |
|-----------------------------------|------|------|------|-------|-------|------|

| | | | | | | |
|---|------|------|------|-------|-------|------|
| Have been dependent on systemic steroids for an extended period (e.g. 6 months or more) | 0.00 | 2.86 | 6.67 | 63.81 | 26.67 | 1.14 |
|---|------|------|------|-------|-------|------|

| | | | | | | |
|--|------|------|------|-------|-------|------|
| Have comorbidities that respond to OCS | 0.00 | 3.81 | 9.52 | 70.48 | 16.19 | 0.99 |
|--|------|------|------|-------|-------|------|

Complete OCS cessation (weaning) can be implemented:

| | | | | | | |
|--|------|------|------|-------|-------|------|
| Following a short course of OCS treatment that lasted for 5–7 days | 0.95 | 1.90 | 1.90 | 44.76 | 50.48 | 1.42 |
|--|------|------|------|-------|-------|------|

| | | | | | | |
|--|------|------|------|-------|-------|------|
| Following a short course of OCS treatment if patients are on inhaled anti-inflammatory therapy | 1.90 | 1.90 | 2.86 | 48.57 | 44.76 | 1.32 |
|--|------|------|------|-------|-------|------|

| | | | | | | |
|--|------|-------|-------|-------|-------|------|
| When a sparing strategy has been initiated | 0.95 | 2.86 | 14.29 | 54.29 | 27.62 | 1.05 |
| When there is no evidence of adrenal insufficiency | 0.95 | 6.67 | 13.33 | 59.05 | 20.00 | 0.90 |
| When the patient has agreed to cessation | 1.90 | 4.76 | 20.00 | 50.48 | 22.86 | 0.88 |
| When there is no evidence of EGPA or ABPA | 0.00 | 7.62 | 19.05 | 56.19 | 17.14 | 0.83 |
| When the OCS dose is ≤ 5 mg prednisolone | 0.95 | 15.24 | 13.33 | 53.33 | 17.14 | 0.70 |

Definition of abbreviations: ABPA = allergic bronchopulmonary aspergillosis; EGPA, eosinophilic granulomatosis with polyangiitis; OCS = oral corticosteroid.

*Note that statements are ordered by mean rank score.

Table 3. Consensus Statements Concerning Development of an OCS Tapering Algorithm

| Positive consensus | Controversial |
|--|--|
| <ul style="list-style-type: none">• The initial tapering of high OCS doses (e.g. >20 mg/day) can proceed at a faster pace (e.g. –10 mg/week, or 30–50% reductions every 2–4 weeks)• OCS tapering should be gradual, with 2.5–5 mg steps every 0.5–2 weeks until an agreed-upon threshold is achieved (e.g. 5–10 mg/day), and then proceeds at a slower pace (1–2.5 mg every 1–2 weeks)• When a reduction in OCS by 5 mg weekly fails, a slower and lower dose reduction of 1 mg/week should be attempted• If mild symptoms occur, maintain the current dosage; they are likely to resolve as endogenous axis recovery occurs• If intolerable symptoms occur, return to the previous (efficacious) dose, and then later consider re-attempting tapering at a slower pace | <ul style="list-style-type: none">• In general, the speed of tapering should not exceed a reduction of 5 mg/week• OCS tapering should incorporate every-other-day OCS reductions (especially prior to discontinuation) to allow recovery of the endogenous axis• OCS tapering should be gradual by reducing the OCS dose by 30–50% every 2–4 weeks |

Definition of abbreviations: OCS = oral corticosteroid.

Table 4. Minimal Checklist for Adverse Effect Screening

| Positive consensus | Controversial |
|---|---|
| <ul style="list-style-type: none">• Growth (pediatric population)• Glycemic control• Bone density• Blood pressure• Cataracts and glaucoma• Weight change• Fracture risk score (e.g. FRAX) | <ul style="list-style-type: none">• Cardiovascular risk score• Lipid panel• Fluid retention |

Definition of abbreviations: FRAX = Fracture Risk Assessment Tool.

Adverse effects are not ordered/hierarchized, and should be given equal consideration.

Online Data Supplement:

Expert consensus on the tapering of oral corticosteroids for the treatment of asthma: a Delphi study

Carey M. Suehs, Andrew Menzies-Gow, David Price, Eugene R. Bleeker, Giorgio Walter Canonica, Mark Gurnell, Arnaud Bourdin on behalf of the Oral Corticosteroids Tapering Delphi Expert Panel*

Results from the three rounds of ranking in the OCS Tapering Delphi project (for items ranked with a five-point Likert scale). For each item, the round and sample size are given along with the percentage of experts that chose a given rank. Darker shades of green signify greater percentage consensus. The weighted mean rank and consensus category are given (positive = blue; negative = red; controversial = white).

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| In general, our goal should be to not use OCS. When nevertheless required, dose and duration should be minimized. | 1 . 1 . a | 1 | 108 | 2.78 | 0.93 | 0.93 | 19.44 | 75.93 | 1.65 | Positive |
| Short-term (e.g. <15 days) OCS use is appropriate in asthma patients during acute, non-resolutive exacerbation. | 1 . 2 . a | 1 | 108 | 0 | 0.93 | 5.56 | 54.63 | 38.89 | 1.31 | Positive |
| Short-term (e.g. <15 days) OCS use is appropriate in asthma patients during acute, life-threatening, exacerbation. | 1 . 2 . b | 1 | 108 | 0 | 0 | 0 | 22.22 | 77.78 | 1.78 | Positive |
| Short-term (e.g. <15 days) OCS use is appropriate in asthma patients during eosinophilic or allergic exacerbation. | 1 . 2 . c | 1 | 108 | 0 | 3.7 | 12.04 | 46.3 | 37.96 | 1.19 | Positive |
| Short-term (e.g. <15 days) OCS use is appropriate in asthma patients in the context of an asthma management plan. | 1 . 2 . d | 2 | 113 | 4.63 | 10.19 | 15.74 | 45.37 | 24.07 | 0.74 | Controversial |
| Short-term (e.g. <15 days) OCS use is appropriate in asthma patients to avoid hospitalization. | 1 . 2 . e | 1 | 108 | 2.78 | 11.11 | 10.19 | 48.15 | 27.78 | 0.87 | Positive |
| Short-term (e.g. <15 days) OCS use is appropriate in asthma patients to palliate the unavailability of hospitalization services. | 1 | 108 | 19.44 | 34.26 | 24.07 | 19.44 | 2.78 | -0.48 | Controversial | |
| | 2 | 113 | 16.81 | 32.74 | 18.58 | 27.43 | 4.42 | -0.3 | Controversial | |
| | 3 | 111 | 14.41 | 32.43 | 22.52 | 22.52 | 8.11 | -0.23 | Controversial | |
| Short-term (e.g. <15 days) OCS use is never appropriate in asthma patients. | 1 . 2 . g | 1 | 108 | 71.3 | 23.15 | 2.78 | 2.78 | 0 | -1.63 | Negative |
| As concerns dosages for short courses of OCS for the treatment of asthma exacerbations, individual tailoring is required to such an extent that the systematic application of "ideal" doses is unlikely. | 1 | 108 | 2.78 | 37.96 | 14.81 | 35.19 | 9.26 | 0.1 | Controversial | |
| | 2 | 112 | 5.36 | 31.25 | 19.64 | 38.39 | 5.36 | 0.07 | Controversial | |
| | 3 | 111 | 4.5 | 39.64 | 12.61 | 39.64 | 3.6 | -0.02 | Controversial | |
| Dosages for short courses of OCS for the treatment of asthma exacerbations should remain stable. | 1 | 108 | 1.85 | 21.3 | 20.37 | 45.37 | 11.11 | 0.43 | Controversial | |
| | 2 | 112 | 3.57 | 19.64 | 22.32 | 47.32 | 7.14 | 0.35 | Controversial | |
| | 3 | 111 | 0.9 | 16.22 | 19.82 | 52.25 | 10.81 | 0.56 | Controversial | |
| Dosages for short courses of OCS for the treatment of asthma exacerbations should be progressively escalated. | 1 . 5 . c | 1 | 108 | 26.85 | 53.7 | 8.33 | 7.41 | 3.7 | -0.93 | Negative |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics who are well controlled with a low dose of OCS (e.g. 5 mg/day or less of prednisone). | 1 . 6 . a | 1 | 108 | 15.74 | 54.63 | 12.96 | 13.89 | 2.78 | -0.67 | Negative |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics with inadequate control despite optimization of alternative (Step 5). | 1 | 108 | 5.56 | 14.81 | 13.89 | 49.26 | 6.48 | 0.46 | Controversial | |
| | 2 | 111 | 0.9 | 17.12 | 11.71 | 61.26 | 9.01 | 0.6 | Positive | |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics with poor inhaler compliance/technique. | 1 . 6 . c | 1 | 108 | 56.48 | 35.19 | 4.63 | 3.7 | 0 | -1.44 | Negative |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|---|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics with low-T2 phenotypes. | | 1 | 108 | 30.56 | 38.89 | 24.07 | 6.48 | 0 | -0.94 | Controversial |
| | 1 . 6 . d | 2 | 111 | 9.01 | 45.05 | 29.73 | 14.41 | 1.8 | -0.45 | Controversial |
| | | 3 | 111 | 13.51 | 52.25 | 24.32 | 9.01 | 0.9 | -0.68 | Controversial |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics with high T2 phenotypes/eosinophils. | | 1 | 108 | 18.52 | 29.63 | 19.44 | 31.48 | 0.93 | -0.33 | Controversial |
| | 1 . 6 . e | 2 | 111 | 6.31 | 36.94 | 24.32 | 30.63 | 1.8 | -0.15 | Controversial |
| | | 3 | 111 | 13.51 | 36.04 | 24.32 | 24.32 | 1.8 | -0.35 | Controversial |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics when it results in an overall reduction in OCS exposure (i.e. the total mg of OCS exposure per year; e.g. 5 mg/day is a 33% reduction when compared to 10) | | 1 | 108 | 3.7 | 22.22 | 30.56 | 36.11 | 7.41 | 0.21 | Controversial |
| | 1 . 6 . f | 2 | 111 | 0.9 | 14.41 | 22.52 | 38.56 | 3.6 | 0.5 | Controversial |
| | | 3 | 111 | 3.6 | 15.32 | 18.02 | 58.56 | 4.5 | 0.45 | Controversial |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics if when trying to taper OCS there is an adverse effect or comorbidity. | | 1 | 108 | 1.85 | 7.41 | 25 | 39.26 | 6.48 | 0.61 | Controversial |
| | 1 . 6 . g | 2 | 111 | 0 | 8.11 | 18.02 | 69.37 | 4.5 | 0.7 | Positive |
| Maintenance OCS therapy is appropriate (and does not require further improvement) in severe asthmatics with primary or secondary adrenal insufficiency. | | 1 | 108 | 6.48 | 11.11 | 12.96 | 36.48 | 12.96 | 0.58 | Controversial |
| | 1 . 6 . h | 2 | 111 | 3.6 | 12.61 | 18.02 | 52.25 | 13.51 | 0.59 | Controversial |
| | | 3 | 111 | 6.31 | 16.22 | 18.02 | 48.65 | 10.81 | 0.41 | Controversial |
| Maintenance OCS therapy is never appropriate in severe asthmatics. | 1 . 6 . i | 1 | 108 | 27.78 | 43.52 | 15.74 | 10.19 | 2.78 | -0.83 | Negative |
| As concerns maintenance OCS therapy, individual tailoring is required to such an extent that the systematic application of "ideal" doses is unlikely. | 1 . 8 . a | 1 | 108 | 1.85 | 16.67 | 13.89 | 42.59 | 25 | 0.72 | Controversial |
| As concerns maintenance OCS therapy, individual tailoring is required to such an extent that the systematic application of "ideal" doses is unlikely. | 1 . 8 . a | 2 | 111 | 0.9 | 17.12 | 9.91 | 56.76 | 15.32 | 0.68 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: normalization of lung function. | | 1 | 108 | 3.7 | 35.19 | 19.44 | 38.89 | 2.78 | 0.02 | Controversial |
| | 1 . 9 . a | 2 | 110 | 5.45 | 29.09 | 26.36 | 37.27 | 1.82 | 0.01 | Controversial |
| | | 3 | 109 | 1.83 | 30.28 | 20.18 | 47.71 | 0 | 0.14 | Controversial |
| An adequate response to long-term OCS in asthmatics can be characterized as: a stable peak flow during the last week of treatment. | | 1 | 108 | 4.63 | 24.07 | 19.44 | 49.07 | 2.78 | 0.21 | Controversial |
| | 1 . 9 . b | 2 | 110 | 0 | 30 | 30.91 | 38.18 | 0.91 | 0.1 | Controversial |
| | | 3 | 109 | 0.92 | 23.85 | 25.69 | 49.54 | 0 | 0.24 | Controversial |
| An adequate response to long-term OCS in asthmatics can be characterized as: suppression of blood eosinophils/other T2 biomarkers. | | 1 | 108 | 5.56 | 20.37 | 27.78 | 39.81 | 6.48 | 0.21 | Controversial |
| | 1 . 9 . c | 2 | 110 | 3.64 | 25.45 | 25.45 | 43.64 | 1.82 | 0.15 | Controversial |
| | | 3 | 109 | 1.83 | 26.61 | 20.18 | 50.46 | 0.92 | 0.22 | Controversial |
| An adequate response to long-term OCS in asthmatics can be characterized as: improvement in the Asthma Control Questionnaire score (MCID = 0.5) or the Asthma Control Test (ACT) (MCID = 5). | 1 . 9 . d | 1 | 108 | 0.93 | 6.48 | 7.41 | 72.22 | 12.96 | 0.9 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: decreasing the exacerbation rate to <2/year. | 1 . 9 . e | 1 | 108 | 0.93 | 4.63 | 12.96 | 63.89 | 17.59 | 0.93 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: decreasing the exacerbation rate by at least 30%. | | 1 | 108 | 0.93 | 24.07 | 32.41 | 37.04 | 5.56 | 0.22 | Controversial |
| | 1 . 9 . f | 2 | 110 | 0.91 | 26.36 | 16.36 | 49.09 | 7.27 | 0.35 | Controversial |
| | | 3 | 109 | 0 | 11.93 | 19.27 | 62.39 | 6.42 | 0.63 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|--------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| An adequate response to long-term OCS in asthmatics can be characterized as: decreasing the exacerbation rate by at least 50%. | 1.9.g | 1 | 108 | 0 | 7.41 | 14.81 | 57.41 | 20.37 | 0.91 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: decreasing hospitalizations for asthma to 0 per year. | 1.9.h | 1 | 108 | 0.93 | 5.56 | 11.11 | 56.48 | 25.93 | 1.01 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: a decreased need for rescue treatments. | 1.9.i | 1 | 108 | 0.93 | 12.96 | 19.44 | 56.48 | 10.19 | 0.62 | Controversial |
| | | 2 | 110 | 0.91 | 14.55 | 16.36 | 60.91 | 7.27 | 0.59 | Controversial |
| | | 3 | 109 | 0.92 | 9.17 | 12.84 | 72.48 | 4.59 | 0.71 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: when a clinical improvement is obtained that outweighs risks/harms. | 1.9.j | 1 | 108 | 0 | 3.7 | 12.96 | 60.19 | 23.15 | 1.03 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: improvement in asthma-related daily limitations/quality of life. | 1.9.k | 1 | 108 | 1.85 | 6.48 | 14.81 | 70.37 | 6.48 | 0.73 | Positive |
| An adequate response to long-term OCS in asthmatics can be characterized as: improvement in symptoms related to chronic sinusitis/nasal polyps. | 1.9.l | 1 | 108 | 2.78 | 19.44 | 33.33 | 41.67 | 2.78 | 0.22 | Controversial |
| | | 2 | 110 | 0.91 | 18.18 | 33.64 | 45.45 | 1.82 | 0.29 | Controversial |
| | | 3 | 109 | 0.92 | 16.51 | 29.36 | 52.29 | 0.92 | 0.36 | Controversial |
| An adequate response to long-term OCS in asthmatics can be characterized as: return to work (which would have been impossible without OCS). | 1.9.m | 1 | 108 | 0.93 | 2.78 | 16.67 | 68.52 | 11.11 | 0.86 | Positive |
| OCS may be used as a temporary measure in patients having recurrent eosinophilic asthma exacerbations whilst completing severe asthma assessments. | 1.10.a | 1 | 108 | 1.85 | 8.33 | 13.89 | 62.96 | 12.96 | 0.77 | Positive |
| The yearly cumulative dose of OCS should be monitored as a marker of asthma control. | 1.10.b | 1 | 108 | 0 | 2.78 | 6.48 | 63.89 | 26.85 | 1.15 | Positive |
| | | 2 | 108 | 0 | 17.59 | 25.93 | 48.15 | 8.33 | 0.47 | Controversial |
| | | 3 | 109 | 0.92 | 22.02 | 11.93 | 59.63 | 5.5 | 0.47 | Controversial |
| OCS therapy can be used to estimate the best obtainable improvement of asthma symptoms. | 1.10.c | 3 | 109 | 0.92 | 22.02 | 11.93 | 59.63 | 5.5 | 0.47 | Controversial |
| Short-term, prophylactic OCS use is appropriate in asthma patients when early signs/symptoms of significant exacerbation appear, if the patient is adherent with proper use of daily asthma therapy. | 1.10.d | 1 | 108 | 4.63 | 30.56 | 20.37 | 42.59 | 1.85 | 0.06 | Controversial |
| | | 2 | 108 | 4.63 | 27.78 | 22.22 | 42.59 | 2.78 | 0.11 | Controversial |
| | | 3 | 109 | 4.59 | 30.28 | 15.6 | 47.71 | 1.83 | 0.12 | Controversial |
| OCS can also be considered in patients with fixed airflow obstruction which becomes reversible on OCS (infrequent). | 1.10.e | 1 | 108 | 4.63 | 24.07 | 35.19 | 34.26 | 1.85 | 0.05 | Controversial |
| | | 2 | 108 | 1.85 | 26.85 | 29.63 | 40.74 | 0.93 | 0.12 | Controversial |
| | | 3 | 109 | 2.75 | 21.1 | 31.19 | 44.04 | 0.92 | 0.19 | Controversial |
| Asthma patients who have a second exacerbation within 6 weeks of a short "burst" prednisone-treated exacerbation should have a longer, tapering course of prednisone. | 1.11.a | 1 | 108 | 2.78 | 21.3 | 33.33 | 39.81 | 2.78 | 0.19 | Controversial |
| | | 2 | 108 | 2.78 | 28.7 | 18.52 | 46.3 | 3.7 | 0.19 | Controversial |
| | | 3 | 109 | 1.83 | 18.35 | 23.85 | 53.21 | 2.75 | 0.37 | Controversial |
| In adults and adolescents receiving maintenance OCS for asthma, the dose should be at least doubled to define an exacerbation. | 1.11.b | 1 | 108 | 2.78 | 20.37 | 18.52 | 52.78 | 5.56 | 0.38 | Controversial |
| | | 2 | 108 | 0.93 | 19.44 | 26.85 | 46.3 | 6.48 | 0.38 | Controversial |
| | | 3 | 109 | 2.75 | 11.93 | 16.51 | 62.39 | 6.42 | 0.58 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|------------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Patients hospitalized for asthma exacerbation and treated with systemic corticosteroids should be prescribed a short course (for example 5 days) of OCS upon discharge from the hospital. | 1 . 11 . c | 1 | 108 | 1.85 | 14.81 | 19.44 | 53.7 | 10.19 | 0.56 | Controversial |
| | | 2 | 108 | 0.93 | 22.22 | 18.52 | 51.85 | 6.48 | 0.41 | Controversial |
| | | 3 | 109 | 0 | 18.35 | 20.18 | 51.38 | 10.09 | 0.53 | Controversial |
| Prednisolone assays should be used in standard practice to verify OCS adherence. | 1 . 12 . a | 1 | 108 | 5.56 | 25 | 26.85 | 33.33 | 9.26 | 0.16 | Controversial |
| | | 2 | 108 | 6.48 | 34.26 | 29.63 | 24.07 | 5.56 | -0.12 | Controversial |
| | | 3 | 109 | 5.5 | 35.78 | 22.94 | 29.36 | 6.42 | -0.05 | Controversial |
| A 9AM cortisol test is sufficient for determining if a patient is OCS compliant. | 1 . 12 . b | 1 | 108 | 11.11 | 30.56 | 42.59 | 15.74 | 0 | -0.37 | Controversial |
| | | 2 | 108 | 13.89 | 32.41 | 36.11 | 16.67 | 0.93 | -0.42 | Controversial |
| | | 3 | 109 | 11.93 | 37.61 | 29.36 | 20.18 | 0.92 | -0.39 | Controversial |
| Long-acting or methylprednisolone injections are not necessary. | 1 . 13 . a | 1 | 108 | 0.93 | 13.89 | 17.59 | 36.11 | 31.48 | 0.83 | Controversial |
| | | 2 | 108 | 0 | 13.89 | 15.74 | 47.22 | 23.15 | 0.8 | Positive |
| Patients receiving frequent methylprednisolone injections for asthma treatment or exacerbations are at the same or similar risk of suffering side effects from steroids and developing adrenal insufficiency as those receiving OCS. | 1 . 13 . b | 1 | 108 | 2.78 | 14.81 | 7.41 | 35.19 | 39.81 | 0.94 | Positive |
| Long-acting or methylprednisolone injections are not superior to orally administered glucocorticoids. | 1 . 13 . c | 1 | 108 | 0 | 6.48 | 13.89 | 52.78 | 26.85 | 1 | Positive |
| Chronic OCS treatment of asthma in the pediatric age should be a rare exception. | 1 . 14 . a | 1 | 108 | 0 | 0 | 14.81 | 29.63 | 55.56 | 1.41 | Positive |
| OCS can lead to several systemic side-effects and growth deficits in pediatric patients. | 1 . 14 . b | 1 | 108 | 0 | 0 | 11.11 | 25 | 63.89 | 1.53 | Positive |
| Methotrexate is a useful steroid-sparing agent in asthma. | 1 . 16 . a | 1 | 108 | 25.93 | 35.19 | 29.63 | 9.26 | 0 | -0.78 | Controversial |
| | | 2 | 108 | 14.81 | 41.67 | 29.63 | 12.96 | 0.93 | -0.56 | Controversial |
| | | 3 | 109 | 14.68 | 48.62 | 26.61 | 8.26 | 1.83 | -0.66 | Controversial |
| Azathioprine is a useful steroid-sparing agent in asthma. | 1 . 16 . b | 1 | 108 | 25.93 | 37.04 | 31.48 | 5.56 | 0 | -0.83 | Controversial |
| | | 2 | 108 | 16.67 | 50.93 | 25.93 | 5.56 | 0.93 | -0.77 | Controversial |
| | | 3 | 109 | 15.6 | 56.88 | 21.1 | 6.42 | 0 | -0.82 | Negative |
| Mycophenolat mofetil is a useful steroid-sparing agent in asthma. | 1 . 16 . c | 1 | 108 | 25 | 35.19 | 33.33 | 6.48 | 0 | -0.79 | Controversial |
| | | 2 | 108 | 16.67 | 39.81 | 38.89 | 3.7 | 0.93 | -0.68 | Controversial |
| | | 3 | 109 | 14.68 | 47.71 | 33.94 | 3.67 | 0 | -0.73 | Controversial |
| Azithromycin is a useful steroid-sparing agent in asthma. | 1 . 16 . d | 1 | 108 | 5.56 | 25.93 | 31.48 | 36.11 | 0.93 | 0.01 | Controversial |
| | | 2 | 108 | 2.78 | 27.78 | 30.56 | 37.04 | 1.85 | 0.07 | Controversial |
| | | 3 | 109 | 2.75 | 26.61 | 32.11 | 35.78 | 2.75 | 0.09 | Controversial |
| The most useful OCS-sparing strategy is high-dose inhaled steroid in asthma. | 1 . 16 . e | 1 | 108 | 1.85 | 23.15 | 20.37 | 42.59 | 12.04 | 0.4 | Controversial |
| | | 2 | 108 | 3.7 | 23.15 | 24.07 | 39.81 | 9.26 | 0.28 | Controversial |
| | | 3 | 109 | 1.83 | 13.76 | 24.77 | 51.38 | 8.26 | 0.5 | Controversial |
| Bronchial thermoplasty is a useful steroid-sparing strategy in asthma. | 1 . 16 . f | 1 | 108 | 6.48 | 24.07 | 45.37 | 20.37 | 3.7 | -0.09 | Controversial |
| | | 2 | 108 | 3.7 | 21.3 | 41.67 | 31.48 | 1.85 | 0.06 | Controversial |
| | | 3 | 109 | 3.67 | 25.69 | 43.12 | 26.61 | 0.92 | -0.05 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|---|------------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Biologicals, such as IL5 and IL4Ra targeting drugs, are useful sparing agents in asthma. | 1 . 16 . g | 1 | 108 | 0.93 | 0 | 3.7 | 19.44 | 75.93 | 1.69 | Positive |
| There is a need for OCS-sparing agents. | 1 . 16 . h | 1 | 108 | 0 | 0 | 1.85 | 39.81 | 58.33 | 1.56 | Positive |
| Patients on maintenance OCS for severe asthma should be systematically assessed for suitability of biologicals. | 1 . 17 . a | 1 | 108 | 0 | 0 | 1.85 | 18.52 | 79.63 | 1.78 | Positive |
| OCS may be used as a provisional strategy for difficult to control, eosinophilic/T2 asthma until an effective biological treatment is available for the patient. | 1 . 17 . b | 1 | 108 | 1.85 | 2.78 | 9.26 | 67.59 | 18.52 | 0.98 | Positive |
| The daily dose of OCS treatment may represent a reliable marker for the evaluation of biological treatment response. | 1 . 17 . c | 1 | 108 | 0.93 | 7.41 | 12.04 | 49.07 | 30.56 | 1.01 | Positive |
| Patients should not have extra OCS at home because the risk of self treatment becoming a habit is too high. | | 1 | 108 | 8.33 | 50 | 13.89 | 19.44 | 8.33 | -0.31 | Controversial |
| | | 2 | 107 | 9.35 | 50.47 | 10.28 | 23.36 | 6.54 | -0.33 | Controversial |
| | 1 . 18 . a | 3 | 109 | 11.93 | 44.95 | 13.76 | 22.02 | 7.34 | -0.32 | Controversial |
| If OCS is to be used, preparations with lower adrenal suppression should be chosen at the lowest effective dose administered in the morning. | 1 . 18 . b | 1 | 108 | 0.93 | 0.93 | 14.81 | 58.33 | 25 | 1.06 | Positive |
| Establishing equivalence between ICS and OCS in children and in adults (systemic distribution of ICS) is of major importance. | 1 . 18 . c | 1 | 108 | 0 | 4.63 | 23.15 | 52.78 | 19.44 | 0.87 | Positive |
| Tapering (down to a minimal efficacious dose or complete weaning if possible) should be attempted in all asthma patients receiving maintenance OCS therapy, regardless of comorbidities. | 2 . 1 . a | 1 | 105 | 0 | 1.9 | 1.9 | 37.14 | 59.05 | 1.53 | Positive |
| The rhythm and speed of OCS tapering requires individualization for each patient. | 2 . 1 . b | 1 | 105 | 0 | 1.9 | 2.86 | 54.29 | 40.95 | 1.34 | Positive |
| Proceeding towards a tapering attempt is particularly appropriate when: patients on maintenance OCS have gained control (for a minimum, agreed-upon time). | 2 . 2 . a | 1 | 105 | 0 | 0 | 3.81 | 59.05 | 37.14 | 1.33 | Positive |
| Proceeding towards a tapering attempt is particularly appropriate when: biological treatment has been initiated and results in asthma control. | 2 . 2 . b | 1 | 105 | 0 | 0.95 | 0.95 | 25.71 | 72.38 | 1.7 | Positive |
| Proceeding towards a tapering attempt is particularly appropriate when: a patient exhibits symptoms/comorbidities likely linked to OCS. | 2 . 2 . c | 1 | 105 | 0 | 1.9 | 2.86 | 41.9 | 53.33 | 1.47 | Positive |
| Proceeding towards a tapering attempt is particularly appropriate when: there is a reasonable likelihood of hypothalamic-pituitary-adrenal axis recovery. | 2 . 2 . d | 1 | 105 | 0 | 1.9 | 11.43 | 54.29 | 32.38 | 1.17 | Positive |
| Proceeding towards a tapering attempt is particularly appropriate when: the intensity or duration of OCS treatment gives reason for concern. | 2 . 2 . e | 1 | 105 | 0 | 0 | 3.81 | 59.05 | 37.14 | 1.33 | Positive |
| Proceeding towards a tapering attempt is particularly appropriate when: the patient does not appear to respond to OCS treatment. | 2 . 2 . f | 1 | 105 | 0 | 0.95 | 0.95 | 35.24 | 62.86 | 1.6 | Positive |
| Tapering OCS should NOT be attempted in patients who: have demonstrated potentially harmful outcomes during previous weaning attempts (and all available medications have been appropriately initiated/tested). | | 1 | 105 | 1.9 | 18.1 | 18.1 | 56.19 | 5.71 | 0.46 | Controversial |
| | | 2 | 106 | 0.94 | 17.92 | 22.64 | 51.89 | 6.6 | 0.45 | Controversial |
| | 2 . 4 . a | 3 | 109 | 0.92 | 22.94 | 13.76 | 59.63 | 2.75 | 0.4 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|---|--------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Tapering OCS should NOT be attempted in patients who: have EGPA that relapses during tapering (and no other changes can be proposed). | 2.4.b | 1 | 105 | 0.95 | 3.81 | 12.38 | 66.67 | 16.19 | 0.93 | Positive |
| Tapering OCS should NOT be attempted in patients who: have ABPA that relapses during tapering (and no other changes can be proposed). | 2.4.c | 1 | 105 | 0 | 9.52 | 19.05 | 61.9 | 9.52 | 0.71 | Positive |
| Tapering OCS should NOT be attempted in patients who: have proven primary or secondary adrenal insufficiency. | | 1 | 105 | 4.76 | 11.43 | 22.86 | 52.38 | 8.57 | 0.49 | Controversial |
| | | 2 | 106 | 5.66 | 21.7 | 17.92 | 47.17 | 7.55 | 0.29 | Controversial |
| | 2.4.d | 3 | 109 | 4.59 | 29.36 | 18.35 | 42.2 | 5.5 | 0.15 | Controversial |
| Tapering OCS should NOT be attempted in patients who: have uncontrolled asthma. | | 1 | 105 | 3.81 | 17.14 | 20 | 47.62 | 11.43 | 0.46 | Controversial |
| | | 2 | 106 | 1.89 | 20.75 | 14.15 | 55.66 | 7.55 | 0.46 | Controversial |
| | 2.4.e | 3 | 109 | 0 | 21.1 | 13.76 | 55.96 | 9.17 | 0.53 | Controversial |
| Tapering OCS should NOT be attempted in patients who: have uncontrolled T2 high inflammation. | | 1 | 105 | 5.71 | 24.76 | 27.62 | 32.38 | 9.52 | 0.15 | Controversial |
| | | 2 | 106 | 4.72 | 27.36 | 29.25 | 30.19 | 8.49 | 0.1 | Controversial |
| | 2.4.f | 3 | 109 | 1.83 | 28.44 | 25.69 | 40.37 | 3.67 | 0.16 | Controversial |
| OCS tapering should be faster in patients who have been on maintenance OCS for shorter periods (less than 6 months for example). | 2.5.a | 1 | 105 | 0 | 17.14 | 12.38 | 56.19 | 14.29 | 0.68 | Positive |
| OCS tapering should be slower in patients who had a slow response to OCS (and vice-versa). | | 1 | 105 | 0.95 | 30.48 | 31.43 | 34.29 | 2.86 | 0.08 | Controversial |
| | | 2 | 106 | 2.83 | 34.91 | 29.25 | 31.13 | 1.89 | -0.06 | Controversial |
| | 2.5.b | 3 | 108 | 0.93 | 37.96 | 29.63 | 31.48 | 0 | -0.08 | Controversial |
| The speed of OCS tapering depends on the known rapidity of action of the sparing drug introduced. | | 1 | 105 | 0 | 18.1 | 23.81 | 44.76 | 13.33 | 0.53 | Controversial |
| | | 2 | 106 | 0.94 | 10.38 | 19.81 | 41.33 | 7.55 | 0.64 | Controversial |
| | 2.5.c | 3 | 108 | 0 | 17.59 | 20.37 | 54.63 | 7.41 | 0.52 | Controversial |
| The speed of OCS tapering depends on the history of and future risk for adverse events. | 2.5.d | 1 | 105 | 0 | 5.71 | 10.48 | 70.48 | 13.33 | 0.91 | Positive |
| The speed of OCS tapering depends on the type of comorbidity present (for EGPA, for example, tapering plans proposed in RCTs are used). | 2.5.e | 1 | 105 | 0 | 1.9 | 13.33 | 66.67 | 18.1 | 1.01 | Positive |
| OCS tapering should be based on patient collaboration and experience with side effects. | 2.5.f | 1 | 105 | 0 | 1.9 | 13.33 | 62.86 | 21.9 | 1.05 | Positive |
| OCS tapering should be guided by biomarkers at each weaning step. | | 1 | 105 | 0.95 | 28.57 | 36.19 | 32.38 | 1.9 | 0.06 | Controversial |
| | | 2 | 106 | 1.89 | 37.74 | 30.19 | 28.3 | 1.89 | -0.09 | Controversial |
| | 2.5.g | 3 | 108 | 0.93 | 50 | 21.3 | 25.93 | 1.85 | -0.22 | Controversial |
| OCS tapering should be gradual, by reducing the OCS dose by 30–50% every 24 weeks. | | 1 | 105 | 0.95 | 28.57 | 29.52 | 40 | 0.95 | 0.11 | Controversial |
| | | 2 | 106 | 1.89 | 27.36 | 20.75 | 48.11 | 1.89 | 0.21 | Controversial |
| | 2.6.a | 3 | 107 | 0 | 19.63 | 12.15 | 67.29 | 0.93 | 0.5 | Controversial |
| OCS tapering should be gradual, with 2.5–5 mg steps every 0.5–2 weeks until an agreed-upon threshold is achieved (e.g. 5–10 mg/day), and then proceeds at a slower pace (1–2.5 mg every 1–2 weeks). | 2.6.b | 1 | 105 | 0 | 3.81 | 13.33 | 72.38 | 10.48 | 0.9 | Positive |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|---|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| In general, the speed of tapering should not exceed a reduction of 5 mg per week. | 2 . 6 . c | 1 | 105 | 0 | 26.67 | 24.76 | 39.05 | 9.52 | 0.31 | Controversial |
| | | 2 | 106 | 1.89 | 23.58 | 17.92 | 50.94 | 5.66 | 0.35 | Controversial |
| | | 3 | 107 | 0 | 30.84 | 14.95 | 48.6 | 5.61 | 0.29 | Controversial |
| The initial tapering of high OCS doses (e.g. >20 mg per day) can proceed at a faster pace (e.g. -10 mg per week, or 30–50% reductions every 2–4 weeks). | 2 . 6 . d | 1 | 105 | 0.95 | 16.19 | 19.05 | 55.24 | 8.57 | 0.54 | Controversial |
| | | 2 | 106 | 0 | 13.21 | 16.98 | 55.98 | 2.83 | 0.59 | Controversial |
| | | 3 | 107 | 1.87 | 8.41 | 14.95 | 70.09 | 4.67 | 0.67 | Positive |
| When a reduction in OCS by 5 mg weekly fails, a slower and lower dose reduction of 1 mg per week should be attempted. | 2 . 6 . e | 1 | 105 | 0 | 5.71 | 12.38 | 72.38 | 9.52 | 0.86 | Positive |
| | | 3 | 107 | 0 | 5.71 | 12.38 | 72.38 | 9.52 | 0.86 | Positive |
| OCS tapering should incorporate every-other-day OCS reductions (especially prior to discontinuation) to allow recovery of the endogenous axis. | 2 . 6 . f | 1 | 105 | 1.9 | 16.19 | 27.62 | 44.76 | 9.52 | 0.44 | Controversial |
| | | 2 | 106 | 2.83 | 19.81 | 28.3 | 46.23 | 2.83 | 0.26 | Controversial |
| | | 3 | 107 | 2.8 | 15.89 | 26.17 | 50.47 | 4.67 | 0.38 | Controversial |
| If intolerable symptoms occur, return to the previous (efficacious) dose, and then later consider re-attempting tapering at a slower pace. | 2 . 6 . g | 1 | 105 | 0 | 0 | 3.81 | 75.24 | 20.95 | 1.17 | Positive |
| | | 3 | 107 | 0 | 0 | 3.81 | 75.24 | 20.95 | 1.17 | Positive |
| If mild symptoms occur, maintain the current dosage; they are likely to resolve as endogenous axis recovery occurs. | 2 . 6 . h | 1 | 105 | 0 | 6.67 | 23.81 | 69.52 | 3.81 | 0.67 | Controversial |
| | | 2 | 106 | 0.94 | 7.55 | 23.58 | 68.87 | 1.89 | 0.6 | Controversial |
| | | 3 | 107 | 0 | 7.48 | 19.63 | 69.16 | 3.74 | 0.69 | Positive |
| A tapering trial should end when: biomarkers trend toward abnormal. | 2 . 7 . a | 1 | 105 | 2.86 | 28.57 | 43.81 | 24.76 | 0 | -0.1 | Controversial |
| | | 2 | 106 | 1.89 | 44.34 | 28.3 | 23.58 | 1.89 | -0.21 | Controversial |
| | | 3 | 107 | 1.87 | 44.86 | 23.36 | 28.97 | 0.93 | -0.18 | Controversial |
| A tapering trial should end when: symptoms trend toward loss of control (retain lowest dose that maintains clinical benefit). | 2 . 7 . b | 1 | 105 | 0 | 5.71 | 2.86 | 81.9 | 9.52 | 0.95 | Positive |
| | | 3 | 107 | 0 | 5.71 | 2.86 | 81.9 | 9.52 | 0.95 | Positive |
| A tapering trial should end when: the patient is not motivated to continue. | 2 . 7 . c | 1 | 105 | 4.76 | 40.95 | 25.71 | 24.76 | 3.81 | -0.18 | Controversial |
| | | 2 | 106 | 5.66 | 43.4 | 29.25 | 18.87 | 2.83 | -0.3 | Controversial |
| | | 3 | 107 | 1.87 | 43.93 | 24.3 | 28.04 | 1.87 | -0.16 | Controversial |
| Peak expiratory flow is a useful biomarker during OCS tapering. | 2 . 8 . a | 1 | 105 | 3.81 | 30.48 | 35.24 | 25.71 | 4.76 | -0.03 | Controversial |
| | | 2 | 106 | 2.83 | 32.08 | 23.58 | 39.62 | 1.89 | 0.06 | Controversial |
| | | 3 | 106 | 0.94 | 33.02 | 16.04 | 49.06 | 0.94 | 0.16 | Controversial |
| Forced expiratory volume in 1 second (spirometry) is a useful biomarker during OCS tapering. | 2 . 8 . b | 1 | 105 | 1.9 | 23.81 | 22.86 | 44.76 | 6.67 | 0.3 | Controversial |
| | | 2 | 106 | 0 | 26.42 | 22.64 | 46.23 | 4.72 | 0.29 | Controversial |
| | | 3 | 106 | 0.94 | 26.42 | 16.98 | 50.94 | 4.72 | 0.32 | Controversial |
| Fraction exhaled nitric oxide is a useful biomarker during OCS tapering. | 2 . 8 . c | 1 | 105 | 0 | 11.43 | 29.52 | 53.33 | 5.71 | 0.53 | Controversial |
| | | 2 | 106 | 0 | 15.09 | 20.75 | 55.66 | 8.49 | 0.58 | Controversial |
| | | 3 | 106 | 0.94 | 14.15 | 24.53 | 54.72 | 5.66 | 0.5 | Controversial |
| Peripheral eosinophils are a useful biomarker during OCS tapering. | 2 . 8 . d | 1 | 105 | 0 | 13.33 | 27.62 | 50.48 | 8.57 | 0.54 | Controversial |
| | | 2 | 106 | 1.89 | 18.87 | 24.53 | 48.11 | 6.6 | 0.39 | Controversial |
| | | 3 | 106 | 0 | 15.09 | 23.58 | 55.66 | 5.66 | 0.52 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|------------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Sputum eosinophils are a useful biomarker during OCS tapering. | 2 . 8 . e | 1 | 105 | 2.86 | 20 | 32.38 | 35.24 | 9.52 | 0.29 | Controversial |
| | | 2 | 106 | 6.6 | 21.7 | 23.58 | 42.45 | 5.66 | 0.19 | Controversial |
| | | 3 | 106 | 5.66 | 19.81 | 32.08 | 38.68 | 3.77 | 0.15 | Controversial |
| Bronchoalveolar lavage fluid (BAL) eosinophils are a useful biomarker during OCS tapering. | 2 . 8 . f | 1 | 105 | 15.24 | 34.29 | 37.14 | 11.43 | 1.9 | -0.5 | Controversial |
| | | 2 | 106 | 14.15 | 44.34 | 28.3 | 12.26 | 0.94 | -0.58 | Controversial |
| | | 3 | 106 | 16.98 | 43.4 | 27.36 | 10.38 | 1.89 | -0.63 | Controversial |
| Asthma control questionnaires (ACT, ACQ) are a useful biomarker during OCS tapering. | 2 . 8 . g | 1 | 105 | 0.95 | 4.76 | 12.38 | 61.9 | 20 | 0.95 | Positive |
| Adrenal insufficiency assessments are a useful biomarker during OCS tapering. | 2 . 8 . h | 1 | 105 | 2.86 | 12.38 | 20.95 | 54.29 | 9.52 | 0.55 | Controversial |
| | | 2 | 106 | 4.72 | 19.81 | 25.47 | 44.34 | 5.66 | 0.26 | Controversial |
| | | 3 | 106 | 4.72 | 22.64 | 22.64 | 45.28 | 4.72 | 0.23 | Controversial |
| Biomarker guidance is useless or too troublesome during OCS tapering. | 2 . 8 . i | 1 | 105 | 24.76 | 31.43 | 24.76 | 15.24 | 3.81 | -0.58 | Controversial |
| | | 2 | 106 | 21.7 | 36.79 | 22.64 | 16.04 | 2.83 | -0.58 | Controversial |
| | | 3 | 106 | 11.32 | 51.89 | 21.7 | 13.21 | 1.89 | -0.58 | Controversial |
| Cautious, slow tapering is particularly appropriate for patients who: have comorbidities that respond to OCS. | 2 . 9 . a | 1 | 105 | 0 | 3.81 | 9.52 | 70.48 | 16.19 | 0.99 | Positive |
| Cautious, slow tapering is particularly appropriate for patients who: have had life-threatening attacks. | 2 . 9 . b | 1 | 105 | 0.95 | 3.81 | 3.81 | 60 | 31.43 | 1.17 | Positive |
| Cautious, slow tapering is particularly appropriate for patients who: have been dependent on systemic steroids for an extended period of time (e.g. 6 months or more). | 2 . 9 . c | 1 | 105 | 0 | 2.86 | 6.67 | 63.81 | 26.67 | 1.14 | Positive |
| Complete OCS cessation (weaning) can be implemented: when the OCS dose is less than or equal to 5 mg prednisolone. | 2 . 10 . a | 1 | 105 | 0.95 | 15.24 | 13.33 | 53.33 | 17.14 | 0.7 | Positive |
| Complete OCS cessation (weaning) can be implemented: following a short course of OCS treatment that lasted for 5–7 days. | 2 . 10 . b | 1 | 105 | 0.95 | 1.9 | 1.9 | 44.76 | 50.48 | 1.42 | Positive |
| Complete OCS cessation (weaning) can be implemented: following a short course of OCS treatment if patients are on inhaled anti inflammatory therapy. | 2 . 10 . c | 1 | 105 | 1.9 | 1.9 | 2.86 | 48.57 | 44.76 | 1.32 | Positive |
| Complete OCS cessation (weaning) can be implemented: when no severe exacerbations have occurred during the last 4 weeks. | 2 . 10 . d | 1 | 105 | 0 | 18.1 | 27.62 | 41.9 | 12.38 | 0.49 | Controversial |
| | | 2 | 106 | 1.89 | 22.64 | 16.98 | 51.89 | 6.6 | 0.39 | Controversial |
| | | 3 | 106 | 0.94 | 22.64 | 15.09 | 58.49 | 2.83 | 0.4 | Controversial |
| Complete OCS cessation (weaning) can be implemented: when there is no evidence of adrenal insufficiency. | 2 . 10 . e | 1 | 105 | 0.95 | 6.67 | 13.33 | 59.05 | 20 | 0.9 | Positive |
| Complete OCS cessation (weaning) can be implemented: when there is no evidence of EGPA or ABPA. | 2 . 10 . f | 1 | 105 | 0 | 7.62 | 19.05 | 56.19 | 17.14 | 0.83 | Positive |
| Complete OCS cessation (weaning) can be implemented: when a sparing strategy has been initiated. | 2 . 10 . g | 1 | 105 | 0.95 | 2.86 | 14.29 | 54.29 | 27.62 | 1.05 | Positive |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|------------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Complete OCS cessation (weaning) can be implemented: when the patient has agreed to cessation. | 2 . 10 . h | 1 | 105 | 1.9 | 4.76 | 20 | 50.48 | 22.86 | 0.88 | Positive |
| Pulmonary rehabilitation can be helpful before OCS tapering to improve physical activity and decrease dyspnea. It can facilitate OCS tapering. | | 1 | 105 | 0 | 14.29 | 34.29 | 44.76 | 6.67 | 0.44 | Controversial |
| | | 2 | 106 | 0 | 16.98 | 19.81 | 50.94 | 12.26 | 0.58 | Controversial |
| | 2 . 11 . a | 3 | 106 | 1.89 | 11.32 | 20.75 | 53.77 | 12.26 | 0.63 | Controversial |
| OCS tapering should be re-attempted every time a new biological treatment for eosinophilic asthma patients becomes available. | 2 . 11 . b | 1 | 105 | 0 | 2.86 | 7.62 | 57.14 | 32.38 | 1.19 | Positive |
| Biological therapies have become an essential support for OCS tapering. | 2 . 11 . c | 1 | 105 | 0.95 | 0.95 | 4.76 | 29.52 | 63.81 | 1.54 | Positive |
| Following the initiation of a biological therapy, if weaning is not achieved within 12 months, consider switching to a different biological. | 2 . 11 . d | 1 | 105 | 0 | 5.71 | 10.48 | 57.14 | 26.67 | 1.05 | Positive |
| Not achieving a >50% reduction in OCS dose (or a tolerable daily dose) is a failure for a given biological therapy that may mandate switching strategies. | 2 . 11 . e | 1 | 105 | 0 | 5.71 | 20 | 59.05 | 15.24 | 0.84 | Positive |
| Thermoplasty needs to be considered when OCS tapering fails and no other alternative is indicated (biologicals etc). | | 1 | 105 | 1.9 | 15.24 | 39.05 | 38.1 | 5.71 | 0.3 | Controversial |
| | | 2 | 106 | 0 | 13.21 | 29.25 | 50.94 | 6.6 | 0.51 | Controversial |
| | 2 . 11 . f | 3 | 106 | 0.94 | 10.38 | 34.91 | 44.34 | 9.43 | 0.51 | Controversial |
| Poor adherence and inhaler technique should be actively sought and managed to facilitate OCS tapering. | 2 . 12 . a | 1 | 105 | 0 | 0.95 | 0.95 | 32.38 | 65.71 | 1.63 | Positive |
| Monitoring during OCS tapering can be based on symptoms in almost all patients. | | 1 | 105 | 0.95 | 23.81 | 16.19 | 44.76 | 14.29 | 0.48 | Controversial |
| | | 2 | 106 | 4.72 | 27.36 | 7.55 | 51.89 | 8.49 | 0.32 | Controversial |
| | 2 . 12 . b | 3 | 106 | 0.94 | 23.58 | 14.15 | 55.66 | 5.66 | 0.42 | Controversial |
| OCS should be used at a minimum dose, so whenever writing a prescription for OCS, the option of reducing the dose should always be considered. | 2 . 12 . c | 1 | 105 | 0 | 2.86 | 7.62 | 58.1 | 31.43 | 1.18 | Positive |
| Comorbidities should be addressed at all times (not just during tapering). | 3 . 1 . a | 1 | 103 | 0 | 0 | 1.94 | 42.72 | 55.34 | 1.53 | Positive |
| Asthma patients receiving OCS therapy are at a higher risk of complications compared to those without OCS exposure. | 3 . 1 . b | 1 | 103 | 0 | 0 | 2.91 | 29.13 | 67.96 | 1.65 | Positive |
| OCS tapering becomes a primary outcome/goal of asthma management when a patient is affected by OCS-related comorbidities. | 3 . 1 . c | 1 | 103 | 0 | 2.91 | 4.85 | 28.16 | 64.08 | 1.53 | Positive |
| The evaluation of comorbidities is mandatory prior to tapering OCS. | 3 . 1 . d | 1 | 103 | 0 | 3.88 | 7.77 | 53.4 | 34.95 | 1.19 | Positive |
| In general, the presence of comorbidities should not preclude attempting to taper down to the lowest efficacious dose or complete withdrawal (if possible). | 3 . 1 . e | 1 | 103 | 0.97 | 1.94 | 0.97 | 60.19 | 35.92 | 1.28 | Positive |
| Comorbidities to address prior to or when initiating tapering: those that require or respond well to OCS treatment (immune diseases, vasculitis, adrenal suppression, etc) | 3 . 2 . a | 1 | 103 | 0 | 0.97 | 2.91 | 55.34 | 40.78 | 1.36 | Positive |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|---|--------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Comorbidities to address prior to or when initiating tapering: respiratory comorbidities or those that may cause (or mimic) asthma (rhinosinusitis, nasal polyposis, GERD, bronchiectasis, vocal cord dysfunction, inducible laryngeal obstruction, dysfunctional breathing, etc). | 3.2.b | 1 | 103 | 0 | 3.88 | 4.85 | 60.19 | 31.07 | 1.18 | Positive |
| Comorbidities to address prior to or when initiating tapering: chronic non-communicable diseases often exacerbated by (or even caused by) OCS use (hyperglycemia/diabetes, metabolic disease, cardiovascular diseases, high blood pressure, glaucoma, cataract, osteoporosis, etc). | 3.2.c | 1 | 103 | 0 | 1.94 | 6.8 | 54.37 | 36.89 | 1.26 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: glycemic control/HbA1c. | 3.3.a | 1 | 103 | 0 | 0.97 | 0.97 | 54.37 | 43.69 | 1.41 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: blood pressure. | 3.3.b | 1 | 103 | 0 | 0.97 | 6.8 | 58.25 | 33.98 | 1.25 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: fluid retention. | 3.3.c | 1 | 103 | 0 | 7.77 | 27.18 | 54.37 | 10.68 | 0.68 | Controversial |
| | | 2 | 106 | 1.89 | 24.53 | 21.7 | 51.89 | 0 | 0.24 | Controversial |
| | | 3 | 106 | 1.89 | 18.87 | 26.42 | 50.94 | 1.89 | 0.32 | Controversial |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: cardiovascular risk score (e.g. CHADS2). | 3.3.d | 1 | 103 | 0 | 6.8 | 25.24 | 47.57 | 20.39 | 0.82 | Controversial |
| | | 2 | 106 | 1.89 | 24.53 | 27.36 | 44.34 | 1.89 | 0.2 | Controversial |
| | | 3 | 106 | 2.83 | 21.7 | 25.47 | 45.28 | 4.72 | 0.27 | Controversial |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: lipid panel. | 3.3.e | 1 | 103 | 0 | 7.77 | 29.13 | 48.54 | 14.56 | 0.7 | Controversial |
| | | 2 | 106 | 0.94 | 23.58 | 24.53 | 50 | 0.94 | 0.26 | Controversial |
| | | 3 | 106 | 2.83 | 25.47 | 22.64 | 47.17 | 1.89 | 0.2 | Controversial |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: fracture risk score (e.g. FRAX). | 3.3.f | 1 | 103 | 0 | 0 | 17.48 | 50.49 | 32.04 | 1.15 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: bone density. | 3.3.g | 1 | 103 | 0 | 0 | 4.85 | 49.51 | 45.63 | 1.41 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: cataracts and glaucoma. | 3.3.h | 1 | 103 | 0 | 2.91 | 10.68 | 50.49 | 35.92 | 1.19 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: growth (pediatric population). | 3.3.i | 1 | 103 | 0 | 0 | 3.88 | 42.72 | 53.4 | 1.5 | Positive |
| The minimum checklist for comorbidity screening in the OCS-treated population should include: weight change. | 3.3.j | 1 | 103 | 0 | 0.97 | 6.8 | 65.05 | 27.18 | 1.18 | Positive |
| Comorbidity subsets for whom OCS tapering is a priority: those with evidence of a clinically significant OCS adverse effect. | 3.4.a | 1 | 103 | 0 | 0 | 0.97 | 38.83 | 60.19 | 1.59 | Positive |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Comorbidity subsets for whom OCS tapering is a priority: those with chronic non-communicable diseases often exacerbated by (or even caused by) OCS use (glucose metabolism, metabolic disease, cardiovascular diseases, high blood pressure, glaucoma, cataract, osteoporosis, etc). | 3 . 4 . b | 1 | 103 | 0 | 0 | 0.97 | 45.63 | 53.4 | 1.52 | Positive |
| | | 1 | 103 | 0 | 5.83 | 25.24 | 48.54 | 20.39 | 0.83 | Controversial |
| | | 2 | 106 | 0.94 | 17.92 | 26.42 | 48.11 | 6.6 | 0.42 | Controversial |
| Comorbidity subsets for whom OCS tapering is a priority: those with a non-T2 phenotype. | 3 . 4 . c | 3 | 105 | 0.95 | 15.24 | 21.9 | 54.29 | 7.62 | 0.52 | Controversial |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... | 3 . 4 . d | 1 | 103 | 0 | 0 | 10.68 | 63.11 | 26.21 | 1.16 | Positive |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... such as age (youth). | 3 . 4 . e | 1 | 103 | 0 | 0 | 10.68 | 50.49 | 38.83 | 1.28 | Positive |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... such as age (elderly). | 3 . 4 . f | 1 | 103 | 0 | 3.88 | 16.5 | 54.37 | 25.24 | 1.01 | Positive |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... such as post-menopausal women. | 3 . 4 . g | 1 | 103 | 0 | 8.74 | 19.42 | 51.46 | 20.39 | 0.83 | Positive |
| | | 1 | 103 | 0 | 8.74 | 32.04 | 45.63 | 13.59 | 0.64 | Controversial |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... such as gender (female). | 3 . 4 . h | 2 | 106 | 0.94 | 14.15 | 16.98 | 41.37 | 6.6 | 0.58 | Controversial |
| | | 3 | 105 | 0 | 8.57 | 25.71 | 60 | 5.71 | 0.63 | Controversial |
| | | 1 | 103 | 0 | 11.65 | 30.1 | 47.57 | 10.68 | 0.57 | Controversial |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... such as vitamin D deficiency. | 3 . 4 . i | 2 | 106 | 0.94 | 15.09 | 27.36 | 52.83 | 3.77 | 0.43 | Controversial |
| | | 3 | 105 | 0.95 | 8.57 | 34.29 | 53.33 | 2.86 | 0.49 | Controversial |
| Comorbidity subsets for whom OCS tapering is a priority: those with important risk factors associated with increased OCS-susceptibility... such as known PDGF-D gene polymorphism. | 3 . 4 . j | 1 | 103 | 0 | 7.77 | 31.17 | 23.3 | 7.77 | 0.31 | Controversial |
| | | 2 | 106 | 1.89 | 13.21 | 54.72 | 27.36 | 2.83 | 0.16 | Controversial |
| | | 3 | 105 | 0.95 | 8.57 | 62.86 | 23.81 | 3.81 | 0.21 | Controversial |
| | | 1 | 103 | 0.97 | 35.92 | 32.04 | 25.24 | 5.83 | -0.01 | Controversial |
| Obese patients should have a polysomnography test prior to tapering. | 3 . 6 . a | 2 | 106 | 7.55 | 51.89 | 15.09 | 18.87 | 6.6 | -0.35 | Controversial |
| | | 3 | 105 | 1.9 | 60 | 22.86 | 11.43 | 3.81 | -0.45 | Controversial |
| Obesity should be aggressively managed with dietary advice and, where suitable and safe, consideration of bariatric surgery. | 3 . 6 . b | 1 | 103 | 0 | 0.97 | 6.8 | 63.11 | 29.13 | 1.2 | Positive |
| | | 1 | 103 | 0 | 15.53 | 29.13 | 50.49 | 4.85 | 0.45 | Controversial |
| The risk of triggering a bipolar disorder in predisposed patients on continuous OCS treatment should be discussed with a psychiatrist. | 3 . 7 . a | 2 | 106 | 2.83 | 25.47 | 24.53 | 42.45 | 4.72 | 0.21 | Controversial |
| | | 3 | 105 | 1.9 | 24.76 | 23.81 | 45.71 | 3.81 | 0.25 | Controversial |
| OCS addiction requires assessment of patient psychological profiles. | 3 . 7 . b | 1 | 103 | 0 | 7.77 | 21.36 | 63.11 | 7.77 | 0.71 | Positive |
| | | 1 | 103 | 5.83 | 36.89 | 31.07 | 25.24 | 0.97 | -0.21 | Controversial |
| All patients over 65 years with severe asthma Step 5 and cardiac failure, should begin tapering only in case of stable cardiac disease. | 3 . 8 . a | 2 | 106 | 4.72 | 45.28 | 16.98 | 31.13 | 1.89 | -0.2 | Controversial |
| | | 3 | 105 | 4.76 | 38.1 | 20.95 | 33.33 | 2.86 | -0.09 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| In OCS patients with cardiovascular diseases, a coronarography should be performed even if the patient has no symptoms. | 3 . 8 . b | 1 | 103 | 11.65 | 49.51 | 28.16 | 9.71 | 0.97 | -0.61 | Controversial |
| In OCS patients with cardiovascular diseases, a coronarography should be performed even if the patient has no symptoms. | 3 . 8 . b | 2 | 106 | 17.92 | 56.6 | 16.98 | 8.49 | 0 | -0.84 | Negative |
| Patients >75 years of age with uncontrolled, Step 4–5 asthma and cardiac disease should have a cardiology evaluation prior to tapering. | 3 . 8 . c | 1 | 103 | 4.85 | 38.83 | 26.21 | 26.21 | 3.88 | -0.15 | Controversial |
| | | 2 | 106 | 4.72 | 33.02 | 23.58 | 34.91 | 3.77 | 0 | Controversial |
| | | 3 | 105 | 5.71 | 45.71 | 17.14 | 27.62 | 3.81 | -0.22 | Controversial |
| For GINA Step 5 patients, fungal disease must be ruled out in the first weeks of OCS treatment. | 3 . 9 . a | 1 | 103 | 0 | 17.48 | 33.01 | 45.63 | 3.88 | 0.36 | Controversial |
| | | 2 | 106 | 0 | 23.58 | 25.47 | 41.51 | 9.43 | 0.37 | Controversial |
| | | 3 | 105 | 0.95 | 20 | 21.9 | 51.43 | 5.71 | 0.41 | Controversial |
| OCS tapering should occur prior to cataract surgery. | 3 . 9 . b | 1 | 103 | 0.97 | 25.24 | 29.13 | 42.72 | 1.94 | 0.19 | Controversial |
| | | 2 | 106 | 0.94 | 26.42 | 36.79 | 33.02 | 2.83 | 0.1 | Controversial |
| | | 3 | 105 | 0 | 20 | 41.9 | 34.29 | 3.81 | 0.22 | Controversial |
| In patients with EGPA, tapering must be performed in collaboration with a rheumatologist. | 3 . 9 . c | 1 | 103 | 3.88 | 21.36 | 21.36 | 48.54 | 4.85 | 0.29 | Controversial |
| | | 2 | 106 | 4.72 | 22.64 | 23.58 | 41.51 | 7.55 | 0.25 | Controversial |
| | | 3 | 105 | 2.86 | 25.71 | 20.95 | 44.76 | 5.71 | 0.25 | Controversial |
| For patients treated with DDAVP (desmopressin), sodium levels should be monitored during tapering to avoid significant hyponatremia. | 3 . 9 . d | 1 | 103 | 0 | 2.91 | 46.6 | 44.66 | 5.83 | 0.53 | Controversial |
| | | 2 | 106 | 0 | 4.72 | 46.23 | 43.4 | 5.66 | 0.5 | Controversial |
| | | 3 | 105 | 0 | 4.76 | 50.48 | 41.9 | 2.86 | 0.43 | Controversial |
| ACOS/COPD rule-out should be performed for patients with a history of tobacco use or biomass exposure. | 3 . 9 . e | 1 | 103 | 0.97 | 15.53 | 20.39 | 57.38 | 5.83 | 0.51 | Controversial |
| | | 2 | 106 | 3.77 | 19.81 | 19.81 | 51.89 | 4.72 | 0.34 | Controversial |
| | | 3 | 105 | 1.9 | 21.9 | 8.57 | 60 | 7.62 | 0.5 | Controversial |
| The cost of OCS side effects should be more properly invested in more effective treatments such as biologicals. | 3 . 9 . f | 1 | 103 | 0 | 0.97 | 10.68 | 50.49 | 37.86 | 1.25 | Positive |
| OCS tapering may be necessary for assessing the possibility of EGPA or other systemic vasculitis. | 3 . 9 . g | 1 | 103 | 1.94 | 3.88 | 22.33 | 65.05 | 6.8 | 0.71 | Positive |
| Adrenal insufficiency among OCS-treated asthma patients should be regularly assessed. | 4 . 1 . a | 1 | 101 | 1.98 | 11.88 | 12.87 | 57.43 | 15.84 | 0.73 | Positive |
| In as much as possible during the tapering process, troublesome signs (such as aches and pains) of adrenal insufficiency should be symptomatically treated and not viewed as a reason to give up on tapering altogether. | 4 . 1 . b | 1 | 101 | 0.99 | 8.91 | 14.85 | 62.38 | 12.87 | 0.77 | Positive |
| In case of adrenal insufficiency during tapering, hydrocortisone replacement is preferred to continued prednisolone, and may ease the tapering process. | 4 . 1 . c | 1 | 101 | 1.98 | 3.96 | 28.71 | 46.53 | 18.81 | 0.76 | Controversial |
| | | 2 | 106 | 1.89 | 7.55 | 27.36 | 50 | 13.21 | 0.65 | Controversial |
| | | 3 | 105 | 1.9 | 5.71 | 27.62 | 55.24 | 9.52 | 0.65 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Adrenal insufficiency should be assessed: systematically when the daily dose of OCS is tapered down to an agreed-upon threshold... | 4 . 2 . a | 1 | 101 | 1.98 | 9.9 | 12.87 | 55.45 | 19.8 | 0.81 | Positive |
| Adrenal insufficiency should be assessed: systematically when the daily dose of OCS is tapered down to an agreed-upon threshold... such as 3 mg/day. | 4 . 2 . b | 1 | 101 | 4.95 | 31.68 | 35.64 | 18.81 | 8.91 | -0.05 | Controversial |
| | | 2 | 105 | 2.86 | 34.29 | 27.62 | 28.57 | 6.67 | 0.02 | Controversial |
| | | 3 | 105 | 4.76 | 33.33 | 24.76 | 33.33 | 3.81 | -0.02 | Controversial |
| Adrenal insufficiency should be assessed: systematically when the daily dose of OCS is tapered down to an agreed-upon threshold... such as 5 mg/day. | 4 . 2 . c | 1 | 101 | 3.96 | 12.87 | 24.75 | 46.53 | 11.88 | 0.5 | Controversial |
| | | 2 | 105 | 1.9 | 18.1 | 15.24 | 52.38 | 12.38 | 0.55 | Controversial |
| | | 3 | 105 | 3.81 | 13.33 | 19.05 | 54.29 | 9.52 | 0.52 | Controversial |
| Adrenal insufficiency should be assessed: systematically when the daily dose of OCS is tapered down to an agreed-upon threshold... such as 7.5 mg/day. | 4 . 2 . d | 1 | 101 | 3.96 | 31.68 | 33.66 | 22.77 | 7.92 | -0.01 | Controversial |
| | | 2 | 105 | 4.76 | 34.29 | 36.19 | 18.1 | 6.67 | -0.12 | Controversial |
| | | 3 | 105 | 5.71 | 36.19 | 24.76 | 27.62 | 5.71 | -0.09 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... | 4 . 3 . a | 1 | 101 | 3.96 | 17.82 | 19.8 | 55.45 | 2.97 | 0.36 | Controversial |
| | | 2 | 105 | 3.81 | 22.86 | 11.43 | 51.43 | 10.48 | 0.42 | Controversial |
| | | 3 | 105 | 3.81 | 21.9 | 10.48 | 58.1 | 5.71 | 0.4 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those on regular, long-term OCS therapy. | 4 . 3 . b | 1 | 101 | 0.99 | 4.95 | 9.9 | 62.38 | 21.78 | 0.99 | Positive |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those exceeding a cumulative yearly dose of 500 mg OCS. | 4 . 3 . c | 1 | 101 | 1.98 | 23.76 | 37.62 | 32.67 | 3.96 | 0.13 | Controversial |
| | | 2 | 105 | 2.86 | 36.19 | 25.71 | 30.48 | 4.76 | -0.02 | Controversial |
| | | 3 | 105 | 0 | 39.05 | 32.38 | 26.67 | 1.9 | -0.09 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those exceeding a cumulative yearly dose of 1 g OCS. | 4 . 3 . d | 1 | 101 | 1.98 | 19.8 | 26.73 | 38.61 | 12.87 | 0.41 | Controversial |
| | | 2 | 105 | 1.9 | 25.71 | 24.76 | 39.05 | 8.57 | 0.27 | Controversial |
| | | 3 | 105 | 0 | 20 | 30.48 | 43.81 | 5.71 | 0.35 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those exceeding a cumulative yearly dose of 2 g OCS. | 4 . 3 . e | 1 | 101 | 1.98 | 13.86 | 20.79 | 41.58 | 21.78 | 0.67 | Controversial |
| | | 2 | 105 | 0.95 | 17.14 | 19.05 | 44.76 | 18.1 | 0.62 | Controversial |
| | | 3 | 105 | 0 | 16.19 | 19.05 | 48.57 | 16.19 | 0.65 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those exceeding a cumulative yearly dose of >2 g OCS. | 4 . 3 . f | 1 | 101 | 1.98 | 8.91 | 21.78 | 39.6 | 27.72 | 0.82 | Controversial |
| | | 2 | 105 | 0.95 | 13.33 | 17.14 | 45.71 | 22.86 | 0.76 | Controversial |
| | | 3 | 105 | 0.95 | 14.29 | 16.19 | 47.62 | 20.95 | 0.73 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those who have had two repeated short courses of OCS in a given year. | 4 . 4 . a | 1 | 101 | 8.91 | 33.47 | 28.71 | 8.91 | 0 | -0.62 | Controversial |
| | | 2 | 105 | 11.43 | 55.34 | 26.67 | 5.71 | 0.95 | -0.7 | Controversial |
| | | 3 | 105 | 7.62 | 60.95 | 19.05 | 12.38 | 0 | -0.64 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those who have had three repeated short courses of OCS in a given year. | 4 . 4 . b | 1 | 101 | 6.93 | 46.53 | 28.71 | 16.83 | 0.99 | -0.42 | Controversial |
| | | 2 | 105 | 10.48 | 40 | 28.57 | 18.1 | 2.86 | -0.37 | Controversial |
| | | 3 | 105 | 5.71 | 48.57 | 21.9 | 22.86 | 0.95 | -0.35 | Controversial |
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those who have had four repeated short courses of OCS in a given year. | 4 . 4 . c | 1 | 101 | 5.94 | 28.71 | 26.73 | 29.7 | 8.91 | 0.07 | Controversial |
| | | 2 | 105 | 3.81 | 28.57 | 18.1 | 41.9 | 7.62 | 0.21 | Controversial |
| | | 3 | 105 | 3.81 | 28.57 | 20.95 | 40.95 | 5.71 | 0.16 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Adrenal insufficiency should be assessed only in selected sub-populations... such as those who have had >4 repeated short courses of OCS in a given year. | 4 . 4 . d | 1 | 101 | 2.97 | 12.87 | 20.79 | 43.56 | 19.8 | 0.64 | Controversial |
| | | 2 | 105 | 1.9 | 12.38 | 16.19 | 48.57 | 20.95 | 0.74 | Controversial |
| | | 3 | 105 | 1.9 | 13.33 | 15.24 | 54.29 | 15.24 | 0.68 | Controversial |
| Adrenal insufficiency should be assessed when signs/symptoms of adrenal insufficiency appear. | 4 . 5 . a | 1 | 101 | 0.99 | 6.93 | 5.94 | 43.56 | 42.57 | 1.2 | Positive |
| Adrenal insufficiency should be assessed when OCS tapering trials are unsuccessful. | 4 . 5 . b | 1 | 101 | 0 | 14.85 | 10.89 | 52.48 | 21.78 | 0.81 | Positive |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: 0.25 mg/kg/d. | 4 . 6 . a | 1 | 101 | 1.98 | 21.78 | 33.45 | 18.81 | 1.98 | -0.03 | Controversial |
| | | 2 | 105 | 0.95 | 28.57 | 50.48 | 16.19 | 3.81 | -0.07 | Controversial |
| | | 3 | 105 | 1.9 | 32.38 | 42.86 | 20.95 | 1.9 | -0.11 | Controversial |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: 0.50 mg/kg/d. | 4 . 6 . b | 1 | 101 | 3.96 | 21.78 | 43.56 | 29.7 | 0.99 | 0.02 | Controversial |
| | | 2 | 105 | 3.81 | 29.52 | 48.57 | 16.19 | 1.9 | -0.17 | Controversial |
| | | 3 | 105 | 1.9 | 26.67 | 43.81 | 26.67 | 0.95 | -0.02 | Controversial |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: 15–20 mg/day | 4 . 6 . c | 1 | 101 | 0.99 | 21.78 | 32.67 | 38.61 | 5.94 | 0.27 | Controversial |
| | | 2 | 105 | 2.86 | 16.19 | 41.9 | 31.43 | 7.62 | 0.25 | Controversial |
| | | 3 | 105 | 1.9 | 17.14 | 43.81 | 31.43 | 5.71 | 0.22 | Controversial |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: 30 mg/day in men and 20 mg/day in women. | 4 . 6 . d | 1 | 101 | 4.95 | 18.81 | 46.53 | 28.71 | 0.99 | 0.02 | Controversial |
| | | 2 | 105 | 4.76 | 24.76 | 44.76 | 23.81 | 1.9 | -0.07 | Controversial |
| | | 3 | 105 | 2.86 | 25.71 | 41.9 | 26.67 | 2.86 | 0.01 | Controversial |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: doubling in cases of stress/sick days. | 4 . 6 . e | 1 | 101 | 0.99 | 5.94 | 16.83 | 56.44 | 19.8 | 0.88 | Positive |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: one intake per day. | 4 . 6 . f | 1 | 101 | 4.95 | 32.67 | 33.66 | 24.75 | 3.96 | -0.1 | Controversial |
| | | 2 | 105 | 6.67 | 36.19 | 27.62 | 23.81 | 5.71 | -0.14 | Controversial |
| | | 3 | 105 | 9.52 | 39.05 | 26.67 | 21.9 | 2.86 | -0.3 | Controversial |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: two intakes per day. | 4 . 6 . g | 1 | 101 | 2.97 | 23.76 | 31.68 | 36.63 | 4.95 | 0.17 | Controversial |
| | | 2 | 105 | 1.9 | 23.81 | 36.19 | 31.43 | 6.67 | 0.17 | Controversial |
| | | 3 | 105 | 2.86 | 14.29 | 32.38 | 45.71 | 4.76 | 0.35 | Controversial |
| In case of adrenal insufficiency during tapering, OCS should be switched to physiological doses of hydrocortisone with the following characteristics: three intakes per day. | 4 . 6 . h | 1 | 101 | 4.95 | 40.59 | 31.68 | 17.82 | 4.95 | -0.23 | Controversial |
| | | 2 | 105 | 2.86 | 41.9 | 32.38 | 19.05 | 3.81 | -0.21 | Controversial |
| | | 3 | 105 | 6.67 | 34.29 | 33.33 | 22.86 | 2.86 | -0.19 | Controversial |
| Hydrocortisone is not obligatory; OCS can be maintained at 2–4 mg once daily (starting at 4 mg) | 4 . 7 . a | 1 | 101 | 5.94 | 19.8 | 29.7 | 41.58 | 2.97 | 0.16 | Controversial |
| | | 2 | 105 | 6.67 | 21.9 | 25.71 | 41.9 | 3.81 | 0.14 | Controversial |
| | | 3 | 105 | 3.81 | 22.86 | 29.52 | 40 | 3.81 | 0.17 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|---|-----------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Hydrocortisone is not obligatory; OCS can be maintained at 5 mg once daily. | 4 . 7 . b | 1 | 101 | 7.92 | 19.8 | 21.78 | 48.51 | 1.98 | 0.17 | Controversial |
| | | 2 | 105 | 7.62 | 18.1 | 23.81 | 46.67 | 3.81 | 0.21 | Controversial |
| | | 3 | 105 | 4.76 | 17.14 | 25.71 | 49.52 | 2.86 | 0.29 | Controversial |
| Hydrocortisone is not obligatory; OCS can be maintained at 7.5 mg once daily. | 4 . 7 . c | 1 | 101 | 13.86 | 29.7 | 29.7 | 25.74 | 0.99 | -0.3 | Controversial |
| | | 2 | 105 | 11.43 | 42.86 | 26.67 | 17.14 | 1.9 | -0.45 | Controversial |
| | | 3 | 105 | 10.48 | 37.14 | 26.67 | 23.81 | 1.9 | -0.3 | Controversial |
| Switching to hydrocortisone should be performed: as soon as adrenal insufficiency is diagnosed. | 4 . 8 . a | 1 | 101 | 2.97 | 12.87 | 38.61 | 37.62 | 7.92 | 0.35 | Controversial |
| | | 2 | 105 | 1.9 | 28.57 | 30.48 | 34.29 | 4.76 | 0.11 | Controversial |
| | | 3 | 105 | 3.81 | 23.81 | 25.71 | 42.86 | 3.81 | 0.19 | Controversial |
| Switching to hydrocortisone should be performed: when the patient has been weaned down to 5 mg OCS (and signs of adrenal insufficiency are present). | 4 . 8 . b | 1 | 101 | 3.96 | 7.92 | 26.73 | 55.45 | 5.94 | 0.51 | Controversial |
| | | 2 | 105 | 2.86 | 10.48 | 29.52 | 53.33 | 3.81 | 0.45 | Controversial |
| | | 3 | 105 | 2.86 | 10.48 | 23.81 | 61.9 | 0.95 | 0.48 | Controversial |
| Switching to hydrocortisone should be performed: when the patient has been weaned down to 5 mg OCS (regardless of adrenal insufficiency assessments). | 4 . 8 . c | 1 | 101 | 5.94 | 50.5 | 32.67 | 10.89 | 0 | -0.51 | Controversial |
| | | 2 | 105 | 5.71 | 47.62 | 33.33 | 8.57 | 4.76 | -0.41 | Controversial |
| | | 3 | 105 | 7.62 | 46.67 | 24.76 | 20 | 0.95 | -0.4 | Controversial |
| Switching to hydrocortisone should be performed: when the patient has been weaned down to 7 mg OCS (and signs of adrenal insufficiency are present). | 4 . 8 . d | 1 | 101 | 3.96 | 24.75 | 35.64 | 27.72 | 7.92 | 0.11 | Controversial |
| | | 2 | 105 | 3.81 | 35.24 | 32.38 | 23.81 | 4.76 | -0.1 | Controversial |
| | | 3 | 105 | 5.71 | 35.24 | 23.81 | 33.33 | 1.9 | -0.1 | Controversial |
| Switching to hydrocortisone should be performed: when the patient has been weaned down to 7 mg OCS (regardless of adrenal insufficiency assessments). | 4 . 8 . e | 1 | 101 | 6.93 | 55.45 | 31.68 | 2.97 | 2.97 | -0.6 | Controversial |
| | | 2 | 105 | 6.67 | 50.48 | 34.29 | 4.76 | 3.81 | -0.51 | Controversial |
| | | 3 | 105 | 9.52 | 54.29 | 24.76 | 9.52 | 1.9 | -0.6 | Controversial |
| Switching to hydrocortisone is not obligatory/important when managing adrenal insufficiency. | 4 . 8 . f | 1 | 101 | 8.91 | 36.63 | 31.68 | 18.81 | 3.96 | -0.28 | Controversial |
| | | 2 | 105 | 10.48 | 29.52 | 31.43 | 20.95 | 7.62 | -0.14 | Controversial |
| | | 3 | 105 | 15.24 | 30.48 | 33.33 | 16.19 | 4.76 | -0.35 | Controversial |
| Adrenal insufficiency should be assessed: using only a fasting morning cortisol. | 4 . 9 . a | 1 | 101 | 3.96 | 44.55 | 27.72 | 21.78 | 1.98 | -0.27 | Controversial |
| | | 2 | 105 | 7.62 | 42.86 | 22.86 | 23.81 | 2.86 | -0.29 | Controversial |
| | | 3 | 105 | 4.76 | 46.67 | 21.9 | 23.81 | 2.86 | -0.27 | Controversial |
| Adrenal insufficiency should be assessed: using only a (short) Synacthen test. | 4 . 9 . b | 1 | 101 | 0.99 | 32.67 | 35.64 | 24.75 | 5.94 | 0.02 | Controversial |
| | | 2 | 105 | 4.76 | 34.29 | 28.57 | 25.71 | 6.67 | -0.05 | Controversial |
| | | 3 | 105 | 1.9 | 40.95 | 24.76 | 29.52 | 2.86 | -0.1 | Controversial |
| Adrenal insufficiency should be assessed: using fasting morning cortisol, and in case of intermediate results, follow up with a (short) Synacthen test. | 4 . 9 . c | 1 | 101 | 0.99 | 6.93 | 19.8 | 55.45 | 16.83 | 0.8 | Positive |
| Adrenal insufficiency assessments should be interpreted with caution; current laboratory tests require improvement in terms of sensitivity and specificity. | 4 . 9 . d | 1 | 101 | 0 | 6.93 | 31.68 | 54.46 | 6.93 | 0.61 | Controversial |
| | | 2 | 105 | 2.86 | 13.33 | 29.52 | 47.62 | 6.67 | 0.42 | Controversial |
| | | 3 | 105 | 0 | 13.33 | 25.71 | 54.29 | 6.67 | 0.54 | Controversial |
| Adrenal insufficiency should be assessed: never; patients should be systematically substituted during tapering irrespective of any test. | 4 . 9 . e | 1 | 101 | 17.82 | 56.44 | 22.77 | 2.97 | 0 | -0.89 | Negative |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|------------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Adrenal insufficiency should be assessed: never; patients should be substituted during tapering only according to signs/symptoms. | 4 . 9 . f | 1 | 101 | 20.79 | 50.5 | 19.8 | 7.92 | 0.99 | -0.82 | Negative |
| Adrenal insufficiency is insufficiently assessed or under-recognized. | 4 . 11 . a | 1 | 101 | 0.99 | 1.98 | 15.84 | 54.46 | 26.73 | 1.04 | Positive |
| Steroid withdrawal syndrome (symptoms of glucocorticoid deficiency in the setting of a proven normal hypothalamic-pituitary-adrenal axis) occurs more often than adrenal insufficiency. | 4 . 11 . b | 1 | 101 | 0 | 2.97 | 38.61 | 51.49 | 6.93 | 0.62 | Controversial |
| | | 2 | 105 | 0 | 10.48 | 31.43 | 50.48 | 7.62 | 0.55 | Controversial |
| | | 3 | 105 | 0 | 7.62 | 29.52 | 57.14 | 5.71 | 0.61 | Controversial |
| Administration of exogenous glucocorticoids even in small doses for only a few days leads to a measurable suppression of the hypothalamic-pituitary-adrenal axis. | 4 . 11 . c | 1 | 101 | 0.99 | 15.84 | 24.75 | 53.47 | 4.95 | 0.46 | Controversial |
| | | 2 | 105 | 1.9 | 24.76 | 22.86 | 44.76 | 5.71 | 0.28 | Controversial |
| | | 3 | 105 | 0.95 | 17.14 | 28.57 | 49.52 | 3.81 | 0.38 | Controversial |
| OCS treatment may not suppress the hypothalamic-pituitary-adrenal axis at all, or it may cause central suppression and adrenal gland atrophy of varying degrees. | 4 . 11 . d | 1 | 101 | 0.99 | 13.86 | 29.7 | 49.5 | 5.94 | 0.46 | Controversial |
| | | 2 | 105 | 1.9 | 10.48 | 33.33 | 45.71 | 8.57 | 0.49 | Controversial |
| | | 3 | 105 | 0.95 | 6.67 | 34.29 | 52.38 | 5.71 | 0.55 | Controversial |
| A correct OCS tapering regime does not require frequent assessments of adrenal insufficiency. | 4 . 12 . a | 1 | 101 | 2.97 | 31.68 | 18.81 | 43.56 | 2.97 | 0.12 | Controversial |
| | | 2 | 105 | 9.52 | 32.38 | 19.05 | 35.24 | 3.81 | -0.09 | Controversial |
| | | 3 | 105 | 4.76 | 27.62 | 20 | 43.81 | 3.81 | 0.14 | Controversial |
| Reduce the dose of glucocorticoid replacement to the minimum dose possible. This should be judged on hydrocortisone day curves (if on hydrocortisone), or prednisolone day curves/8- hour prednisolone levels. | 4 . 12 . b | 1 | 101 | 2.97 | 19.8 | 49.5 | 26.73 | 0.99 | 0.03 | Controversial |
| | | 2 | 105 | 3.81 | 22.86 | 51.43 | 20.95 | 0.95 | -0.08 | Controversial |
| | | 3 | 105 | 1.9 | 25.71 | 48.57 | 22.86 | 0.95 | -0.05 | Controversial |
| If systemic effects (e.g. arthritis pain) occur during OCS tapering, patients are advised to slow down the tapering pace because the complaints will disappear after some time. | 4 . 12 . c | 1 | 101 | 0 | 11.88 | 24.75 | 59.37 | 3.96 | 0.55 | Controversial |
| | | 2 | 105 | 0.95 | 10.48 | 22.86 | 59.05 | 6.67 | 0.6 | Controversial |
| | | 3 | 105 | 0 | 10.48 | 24.76 | 61.9 | 2.86 | 0.57 | Controversial |
| If adrenal insufficiency occurs during tapering, first increase OCS, and then later re-attempt tapering at a slower pace. | 4 . 12 . d | 1 | 101 | 2.97 | 13.86 | 19.8 | 57.43 | 5.94 | 0.5 | Controversial |
| If adrenal insufficiency occurs during tapering, first increase OCS, and then later re-attempt tapering at a slower pace. | 4 . 12 . d | 2 | 105 | 2.86 | 9.52 | 15.24 | 65.71 | 6.67 | 0.64 | Positive |
| When symptoms occur, stop further tapering until they resolve (this can take weeks/months), and then continue. | 4 . 12 . e | 1 | 101 | 0 | 19.8 | 23.76 | 53.45 | 0.99 | 0.38 | Controversial |
| | | 2 | 105 | 1.9 | 23.81 | 16.19 | 56.19 | 1.9 | 0.32 | Controversial |
| | | 3 | 105 | 0.95 | 15.24 | 24.76 | 55.24 | 3.81 | 0.46 | Controversial |
| An undetectable eosinophil count may be a sign of glucocorticoid excess. | 4 . 13 . a | 1 | 101 | 4.95 | 41.58 | 22.77 | 28.71 | 1.98 | -0.19 | Controversial |
| | | 2 | 105 | 8.57 | 29.52 | 26.67 | 33.33 | 1.9 | -0.1 | Controversial |
| | | 3 | 105 | 6.67 | 31.43 | 22.86 | 38.1 | 0.95 | -0.05 | Controversial |
| The interpretation of short Synacthen test results should take into account the effect of inhaled glucocorticoids. | 4 . 13 . b | 1 | 101 | 0.99 | 21.78 | 32.67 | 37.62 | 6.93 | 0.28 | Controversial |
| | | 2 | 105 | 0 | 24.76 | 29.52 | 37.14 | 8.57 | 0.3 | Controversial |
| | | 3 | 105 | 0 | 19.05 | 35.24 | 40.95 | 4.76 | 0.31 | Controversial |
| Patients who fail their first short Synacthen test with a 30-min cortisol of <350 nmol/L or 12 g/dL, should be counselled that there is a 50% chance of lifelong replacement therapy. | 4 . 13 . c | 1 | 101 | 2.97 | 14.85 | 30.69 | 18.81 | 0 | -0.02 | Controversial |
| | | 2 | 105 | 0.95 | 12.38 | 40.95 | 22.86 | 2.86 | 0.14 | Controversial |
| | | 3 | 105 | 0.95 | 14.29 | 60 | 23.81 | 0.95 | 0.1 | Controversial |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|------------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Patients with a subsequent morning cortisol of <200 nmol/L should be informed that there is a >90% chance that they will need lifelong steroids. | | 1 | 101 | 2.97 | 15.84 | 60.95 | 17.82 | 0.99 | -0.02 | Controversial |
| | | 2 | 105 | 0.95 | 17.14 | 60.95 | 15.24 | 1.9 | 0 | Controversial |
| | 4 . 13 . d | 3 | 105 | 1.9 | 11.43 | 60.95 | 24.76 | 0.95 | 0.11 | Controversial |
| Patient-physician shared decision-making for OCS tapering should be a systematic practice. | 5 . 1 . a | 1 | 101 | 0 | 1.98 | 4.95 | 52.48 | 40.59 | 1.32 | Positive |
| In most cases, the decision to taper OCS treatment is not shared, but taken alone by the clinician. | | 1 | 101 | 8.91 | 39.6 | 8.91 | 38.61 | 3.96 | -0.11 | Controversial |
| | | 2 | 105 | 13.33 | 39.05 | 9.52 | 32.38 | 5.71 | -0.22 | Controversial |
| | 5 . 1 . b | 3 | 105 | 6.67 | 50.48 | 11.43 | 26.67 | 4.76 | -0.28 | Controversial |
| The self-management of OCS treatments should be discouraged. | | 1 | 101 | 1.98 | 25.74 | 10.89 | 38.61 | 22.77 | 0.54 | Controversial |
| | | 2 | 105 | 2.86 | 31.43 | 13.33 | 35.24 | 17.14 | 0.32 | Controversial |
| | 5 . 1 . c | 3 | 105 | 1.9 | 33.33 | 14.29 | 40.95 | 9.52 | 0.23 | Controversial |
| The self-management of OCS tapering should be limited to patients with a good level of comprehension. | 5 . 1 . d | 1 | 101 | 1.98 | 14.85 | 11.88 | 57.43 | 13.86 | 0.66 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: it educates the patient on the benefits/risks associated with OCS use. | 5 . 2 . a | 1 | 101 | 0 | 0 | 0.99 | 58.42 | 40.59 | 1.4 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: it allows the patients to understand the purpose of OCS tapering. | 5 . 2 . b | 1 | 101 | 0 | 0 | 0 | 66.34 | 33.66 | 1.34 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: it provides necessary support and guidance to the patient. | 5 . 2 . c | 1 | 101 | 0 | 0 | 3.96 | 65.35 | 30.69 | 1.27 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: it can increase the chances of success; improve outcomes. | 5 . 2 . d | 1 | 101 | 0 | 1.98 | 1.98 | 61.39 | 34.65 | 1.29 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: ambivalent attitudes towards tapering are frequent. | 5 . 2 . e | 1 | 101 | 0 | 6.93 | 13.86 | 59.41 | 19.8 | 0.92 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: "aches and pains" during OCS withdrawal can occur, and planning how to manage them is likely to improve withdrawal progress. | 5 . 2 . f | 1 | 101 | 0.99 | 0.99 | 3.96 | 62.38 | 31.68 | 1.23 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: patient engagement/empowerment in the process can optimize the outcome. | 5 . 2 . g | 1 | 101 | 0 | 0 | 1.98 | 62.38 | 35.64 | 1.34 | Positive |
| Patient-physician shared decision-making for OCS tapering is important because: patients are often expected to self-medicate at home. | 5 . 2 . h | 1 | 101 | 1.98 | 4.95 | 15.84 | 63.37 | 13.86 | 0.82 | Positive |
| Patient-physician shared decision-making should include: a decision aid including full disclosure of short- and long-term exacerbation/adverse events profile. | 5 . 3 . a | 1 | 101 | 0 | 1.98 | 11.88 | 66.34 | 19.8 | 1.04 | Positive |
| Patient-physician shared decision-making should include: patient education on the benefits/risks associated with OCS use. | 5 . 3 . b | 1 | 101 | 0 | 0 | 0 | 65.35 | 34.65 | 1.35 | Positive |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|--------|-------|-------------|----------------------|----------------|----------------|----------------|-------------------|--------------------|---------------------------|
| Patient-physician shared decision-making should include: the benefits/risks associated with OCS tapering and why it is important. | 5.3.c | 1 | 101 | 0 | 0 | 0 | 56.44 | 43.56 | 1.44 | Positive |
| Patient-physician shared decision-making should include: the dangers of abrupt tapering /OCS discontinuation. | 5.3.d | 1 | 101 | 0 | 0 | 0 | 56.44 | 43.56 | 1.44 | Positive |
| Patient-physician shared decision-making should include: the patient's thoughts (concerns, fears, hopes, expectations) and preferences. | 5.3.e | 1 | 101 | 0 | 0 | 3.96 | 60.4 | 35.64 | 1.32 | Positive |
| Patient-physician shared decision-making should include: symptoms that may occur due to weaning, how to recognize and manage them (including adrenal insufficiency). | 5.3.f | 1 | 101 | 0 | 0.99 | 0.99 | 60.4 | 37.62 | 1.35 | Positive |
| Patient-physician shared decision-making should include: multidisciplinary work (for example, collaboration between respiratory, endocrinology, and rheumatology experts). | 5.3.g | 1 | 101 | 0.99 | 3.96 | 12.87 | 49.5 | 32.67 | 1.09 | Positive |
| Patient-physician shared decision-making should include: a joint evaluation of the patient's global health status and/or quality of life. | 5.3.h | 1 | 101 | 0 | 3.96 | 11.88 | 66.34 | 17.82 | 0.98 | Positive |
| Patient-physician shared decision-making should include: using biomarkers for monitoring and individualization of the action plan. | 5.3.i | 2 | 105 | 0.99 0.95 | 10.89 12.38 | 26.73 16.19 | 53.47 60.95 | 7.92 9.52 | 0.56 0.66 | Controversial Positive |
| Patient-physician shared decision-making should include: steroid-sparing strategies and their benefits/risks. | 5.3.j | 1 | 101 | 0 | 0.99 | 5.94 | 61.39 | 31.68 | 1.24 | Positive |
| Patient-physician shared decision-making should include: clear, agreed-upon protocols/action plan on how tapering will be carried out and what to expect. | 5.3.k | 1 | 101 | 0 | 0.99 | 4.95 | 62.38 | 31.68 | 1.25 | Positive |
| Patient-physician shared decision-making should include: a warning regarding the consequences of not following the action plan. | 5.3.l | 1 | 101 | 0.99 | 2.97 | 7.92 | 70.3 | 17.82 | 1.01 | Positive |
| Patient-physician shared decision-making should include: a means of contacting the doctor/team so the patient can reach out and get support. | 5.3.m | 1 | 101 | 0 | 0.99 | 3.96 | 59.41 | 35.64 | 1.3 | Positive |
| Patient-physician shared decision-making should include: discussion with both patients and their families/caregivers. | 5.3.n | 1 | 101 | 0 | 1.98 | 11.88 | 59.41 | 26.73 | 1.11 | Positive |
| Advice for OCS self-managers: if possible, do not opt for regular OCS use. | 5.4.a | 1 | 101 | 0 | 1.98 | 7.92 | 51.49 | 38.61 | 1.27 | Positive |
| Advice for OCS self-managers: the lowest active dose of OCS for the shortest duration is preferable. | 5.4.b | 1 | 101 | 0 | 0 | 1.98 | 53.47 | 44.55 | 1.43 | Positive |
| Advice for OCS self-managers: closely monitor symptoms while tapering, including those of adrenal insufficiency. | 5.4.c | 1 | 101 | 0 | 0 | 5.94 | 59.41 | 34.65 | 1.29 | Positive |
| Advice for OCS self-managers: help the process of OCS tapering by overcoming minor discomfort related to it. | 5.4.d | 1 | 101 | 0.99 | 0 | 3.96 | 67.33 | 27.72 | 1.21 | Positive |
| Advice for OCS self-managers: respect your doctor's recommendations in as much as possible, and contact him/her (or team) when there is a problem. | 5.4.e | 1 | 101 | 0 | 0.99 | 3.96 | 63.37 | 31.68 | 1.26 | Positive |

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|---|--------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Advice for OCS self-managers: increase the OCS dose to the previous dose if a weaning step causes (intolerable) symptoms. | 5.4.f | 1 | 101 | 0 | 3.96 | 10.89 | 54.46 | 30.69 | 1.12 | Positive |
| | | 1 | 101 | 0.99 | 10.89 | 22.77 | 44.55 | 20.79 | 0.73 | Controversial |
| Advice for OCS self-managers: never use a dose lower than the agreed-up threshold (e.g. 7.5 mg) without substitution. | 5.4.g | 2 | 105 | 4.76 | 23.81 | 29.52 | 35.24 | 6.67 | 0.15 | Controversial |
| | | 3 | 105 | 0.95 | 33.33 | 23.81 | 37.14 | 4.76 | 0.11 | Controversial |
| Advice for OCS self-managers: always make dosage changes under medical supervision. | 5.4.h | 1 | 101 | 0 | 10.89 | 15.84 | 52.48 | 20.79 | 0.83 | Positive |
| Physicians should drive the decision-making when it comes to OCS tapering. | 5.5.a | 1 | 101 | 0 | 8.91 | 12.87 | 59.41 | 18.81 | 0.88 | Positive |
| | | 1 | 101 | 1.98 | 10.89 | 21.78 | 56.44 | 8.91 | 0.59 | Controversial |
| | | 2 | 105 | 0.95 | 27.62 | 17.14 | 48.57 | 5.71 | 0.3 | Controversial |
| Physicians should limit prescriptions to ensure that tapering is occurring. | 5.5.b | 3 | 105 | 1.9 | 19.05 | 21.9 | 56.19 | 0.95 | 0.35 | Controversial |
| | | 1 | 101 | 1.98 | 23.76 | 26.73 | 29.7 | 17.82 | 0.38 | Controversial |
| | | 2 | 105 | 2.86 | 30.48 | 14.29 | 41.9 | 10.48 | 0.27 | Controversial |
| The self-management of OCS treatments should be discouraged. | 5.5.c | 3 | 105 | 1.9 | 39.05 | 12.38 | 39.05 | 7.62 | 0.11 | Controversial |
| | | 1 | 101 | 4.95 | 41.58 | 10.89 | 35.64 | 6.93 | -0.02 | Controversial |
| | | 2 | 105 | 3.81 | 47.62 | 15.24 | 30.48 | 2.86 | -0.19 | Controversial |
| Forewarning patients of "aches and pains" during OCS withdrawal is likely to impede withdrawal progress. | 5.5.d | 3 | 105 | 2.86 | 49.52 | 11.43 | 33.33 | 2.86 | -0.16 | Controversial |
| | | 1 | 101 | 0 | 9.9 | 24.75 | 58.45 | 9.9 | 0.65 | Controversial |
| | | 2 | 105 | 0.95 | 7.62 | 25.71 | 59.05 | 6.67 | 0.63 | Controversial |
| When OCS tapering decisions are not taken mutually, this can lead to medical malpractice and litigation. | 5.5.e | 3 | 105 | 0 | 8.57 | 27.62 | 58.1 | 5.71 | 0.61 | Controversial |
| | | 1 | 101 | 3.96 | 24.75 | 35.64 | 31.68 | 3.96 | 0.07 | Controversial |
| | | 2 | 105 | 9.52 | 41.9 | 15.24 | 32.38 | 0.95 | -0.27 | Controversial |
| In some cases, you might need to have a consent form signed before patients start OCS treatment. | 5.5.f | 3 | 105 | 6.67 | 36.19 | 23.81 | 30.48 | 2.86 | -0.13 | Controversial |
| Many times, patients feel their safety depends on OCS and it takes a lot of effort to convince them to taper. | 5.5.g | 1 | 101 | 0 | 9.9 | 19.8 | 55.45 | 14.85 | 0.75 | Positive |
| The majority of patients want to reduce their OCS use and will actively participate in doing so. | 5.5.h | 1 | 101 | 0 | 2.97 | 10.89 | 62.38 | 23.76 | 1.07 | Positive |
| OCS tapering can be successful even if the patient doesn't think it will work. | 5.5.i | 1 | 101 | 0.99 | 4.95 | 16.83 | 62.38 | 14.85 | 0.85 | Positive |
| | | 1 | 101 | 4.95 | 36.63 | 26.73 | 25.74 | 5.94 | -0.09 | Controversial |
| | | 2 | 105 | 4.76 | 40.95 | 29.52 | 23.81 | 0.95 | -0.25 | Controversial |
| It is better to allow patients to control their own prednisolone doses to control symptoms than to give high dose bursts for exacerbations. | 5.5.j | 3 | 105 | 2.86 | 45.71 | 26.67 | 22.86 | 1.9 | -0.25 | Controversial |
| | | 1 | 101 | 0 | 13.86 | 35.64 | 43.56 | 6.93 | 0.44 | Controversial |
| | | 2 | 105 | 0 | 10.48 | 35.24 | 52.38 | 1.9 | 0.46 | Controversial |
| The patient generally has full confidence in his/her doctor and experiences tapering as a success on his/her illness. | 5.5.k | 3 | 105 | 0 | 14.29 | 30.48 | 48.57 | 6.67 | 0.48 | Controversial |
| | | 1 | 101 | 0.99 | 12.87 | 30.69 | 46.53 | 8.91 | 0.5 | Controversial |
| | | 2 | 105 | 0.95 | 10.48 | 26.67 | 56.19 | 5.71 | 0.55 | Controversial |
| The patient is usually the major player and follows an action plan with an easy contact with the multidisciplinary team. | 5.5.l | 3 | 105 | 1.9 | 15.24 | 20.95 | 57.14 | 4.76 | 0.48 | Controversial |

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|--|--------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| Physicians should be trained on how to coach patients during the tapering process. | 5.5.m | 1 | 101 | 0 | 0 | 8.91 | 72.28 | 18.81 | 1.1 | Positive |
| Patients should be educated with standard material (generated and endorsed e.g. by ERS) about the OCS therapy. | 5.5.n | 1 | 101 | 0 | 0.99 | 6.93 | 75.25 | 16.83 | 1.08 | Positive |
| Shared decision-making is made difficult by the level of individualization and adaptation required during OCS tapering. | | 1 | 101 | 1.98 | 20.79 | 15.84 | 51.49 | 9.9 | 0.47 | Controversial |
| | | 2 | 105 | 1.9 | 32.38 | 22.86 | 40 | 2.86 | 0.1 | Controversial |
| | 5.5.o | 3 | 105 | 0.95 | 38.1 | 20.95 | 38.1 | 1.9 | 0.02 | Controversial |
| Shared decision-making is dependent on the willingness and ability of both sides to interact. | 5.5.p | 1 | 101 | 0 | 0 | 3.96 | 69.31 | 26.73 | 1.23 | Positive |
| Patients are suffering a lot and a strong patient-doctor relationship is required to achieve a safe, optimum outcome from OCS tapering. | 5.5.q | 1 | 101 | 0.99 | 0 | 13.86 | 53.47 | 31.68 | 1.15 | Positive |
| All OCS-treated asthma patients should be referred to an expert center able to propose multidisciplinary assessment and access to innovations. | 6.1.a | 1 | 101 | 0 | 1.98 | 5.94 | 38.61 | 53.47 | 1.44 | Positive |
| Maintenance OCS for severe asthma should only be considered after evaluation by a severe asthma specialist (the definition of this specialist may vary from region to region). | 6.1.b | 1 | 101 | 0 | 2.97 | 4.95 | 32.67 | 59.41 | 1.49 | Positive |
| The respiratory physician treating severe asthma patients must assess for adrenal insufficiency. | 6.1.c | 1 | 101 | 0.99 | 6.93 | 13.86 | 49.5 | 28.71 | 0.98 | Positive |
| Adrenal insufficiency management in patients with severe asthma should involve an endocrinologist/multidisciplinary approach. | 6.1.d | 1 | 101 | 0 | 3.96 | 23.76 | 38.61 | 33.66 | 1.02 | Positive |
| Primary care physicians prescribing more than three courses of OCS to a patient with asthma in 1 year should consider a referral to a specialist. | 6.2.a | 1 | 101 | 0 | 0.99 | 0 | 27.72 | 71.29 | 1.69 | Positive |
| The primary care physician should be part of the multidisciplinary team. | 6.2.b | 1 | 101 | 0 | 3.96 | 16.83 | 55.45 | 23.76 | 0.99 | Positive |
| OCS use in asthma should also be discouraged at the primary care level. | | 1 | 101 | 6.93 | 28.71 | 6.93 | 36.63 | 20.79 | 0.36 | Controversial |
| | | 2 | 105 | 10.48 | 28.57 | 8.57 | 33.33 | 19.05 | 0.22 | Controversial |
| | 6.2.c | 3 | 105 | 7.62 | 29.52 | 9.52 | 34.29 | 19.05 | 0.28 | Controversial |
| The following is an important subject of future research: improving the delivery of asthma care. | 6.3.a | 1 | 101 | 0 | 0.99 | 5.94 | 54.46 | 38.61 | 1.31 | Positive |
| The following is an important subject of future research: integration and dissemination of how to use predictive biomarkers in clinical practice. | 6.3.b | 1 | 101 | 0 | 2.97 | 8.91 | 57.43 | 30.69 | 1.16 | Positive |
| The following is an important subject of future research: improving the use of biological treatments in asthma. | 6.3.c | 1 | 101 | 0 | 0 | 4.95 | 38.61 | 56.44 | 1.51 | Positive |
| The following is an important subject of future research: while striving to obtain a balance between over and under-treatment with OCS, patients often experience adverse quality of life. How best to manage this requires future research. | 6.3.d | 1 | 101 | 0 | 0 | 19.8 | 55.45 | 24.75 | 1.05 | Positive |

| Statement | Number | Round | Sample size | Strongly disagree, % | Disagree, % | Neutral, % | Agree, % | Strongly agree, % | Weighted mean rank | Consensus |
|--|--------|-------|-------------|----------------------|-------------|------------|----------|-------------------|--------------------|---------------|
| The following is an important subject of future research: whether hydrocortisone supplementation is less harmful than prednisone should be established. | 6.3.e | 1 | 101 | 0.99 | 2.97 | 15.84 | 50.5 | 29.7 | 1.05 | Positive |
| The following is an important subject of future research: The impact of shared decision-making on important outcomes. | 6.3.f | 1 | 101 | 0 | 0.99 | 20.79 | 52.48 | 25.74 | 1.03 | Positive |
| The following is an important subject of future research: OCS tapering regime algorithms and optimization. | 6.3.g | 1 | 101 | 0.99 | 0.99 | 5.94 | 50.5 | 41.58 | 1.31 | Positive |
| The following is an important subject of future research: real-life, cost-benefit/effectiveness evaluations for steroid-sparing strategies taking into account side-effects and comorbidities, quality of life, and the societal costs of maintenance OCS. | 6.3.h | 1 | 101 | 0 | 0 | 6.93 | 40.59 | 52.48 | 1.46 | Positive |
| The following is an important subject of future research: direct comparisons between biologicals, especially anti-IL-5. | 6.3.i | 1 | 101 | 0.99 | 0.99 | 10.89 | 38.61 | 48.51 | 1.33 | Positive |
| The following is an important subject of future research: strategic ways to reduce OCS use for the overall at-risk populations. | 6.3.j | 1 | 101 | 0 | 0.99 | 2.97 | 58.42 | 37.62 | 1.33 | Positive |
| The following is an important subject of future research: methods for determining OCS starting doses. | 6.3.k | 1 | 101 | 0 | 7.92 | 17.82 | 55.45 | 18.81 | 0.85 | Positive |
| The following is an important subject of future research: the role of the endocrinologist and when referral should occur. | 6.3.l | 1 | 101 | 0 | 4.95 | 11.88 | 64.36 | 18.81 | 0.97 | Positive |
| The following is an important subject of future research: improving the assessment of adrenal insufficiency. | 6.3.m | 1 | 101 | 0.99 | 0 | 7.92 | 54.46 | 36.63 | 1.26 | Positive |
| The following is an important subject of future research: the efficacy of internet-provided algorithms for delivering symptom-driven OCS tapering guidance to asthma patients. | 6.3.n | 1 | 101 | 0.99 | 9.9 | 22.77 | 47.52 | 18.81 | 0.73 | Controversial |
| | | 2 | 105 | 0 | 13.33 | 22.86 | 50.48 | 13.33 | 0.64 | Controversial |
| | | 3 | 105 | 0 | 11.43 | 20.95 | 59.05 | 8.57 | 0.65 | Controversial |
| The following is an important subject of future research: how should OCS tapering be addressed in countries where there is limited access to biological treatments? | 6.3.o | 1 | 101 | 0 | 2.97 | 8.91 | 58.42 | 29.7 | 1.15 | Positive |
| The following is an important subject of future research: what aspect/phenotype of asthma is being treated by OCS that the currently available biological therapies are not treating? | 6.3.p | 1 | 101 | 0 | 0.99 | 7.92 | 42.57 | 48.51 | 1.39 | Positive |
| The following is an important subject of future research: in the context of successful OCS weaning subsequent to the initiation of a biological, what kind of follow-up should be proposed? | 6.3.q | 1 | 101 | 0 | 1.98 | 15.84 | 56.44 | 25.74 | 1.06 | Positive |

ABPA = allergic bronchopulmonary aspergillosis; ACOS = Asthma-COPD overlap syndrome; ACQ = Asthma Control Questionnaire; ACT = Asthma Control Test; BAL = bronchoalveolar lavage fluid; COPD = chronic obstructive pulmonary disease; DDAVP = desmopressin; EGPA = eosinophilic granulomatosis with polyangiitis; ERS = European Respiratory Society; FRAX = Fracture Risk Assessment Tool; GERD = gastroesophageal reflux disease; GINA = Global Initiative for Asthma; HbA1c = hemoglobin A1c; ICS = inhaled corticosteroid; IL = interleukin; MCID = minimal clinically important difference; OCS = oral corticosteroid; PDGF-D = platelet-derived growth factor D

The OCS Tapering Delphi Expert Panel
(Alphabetical order)

- **Al-Ahmad, Mona**
 - Microbiology Department, Faculty of Medicine, Kuwait University, Kuwait
- **Babu, K Suresh**
 - Queen Alexandra Hospital, Portsmouth, Hampshire, PO6 3LY, United Kingdom
- **Bakakos, Petros**
 - 11 Kononos St, Athens, Attiki, 11634, Greece
- **Ball, Stephen**
 - Manchester University Foundation Trust, Endocrinology Department, Oxford Road, Manchester, M13 9WL, United Kingdom
- **Bel, Elisabeth**
 - Department of Pulmonology, F5-168, Amsterdam UMC, Univ. of Amsterdam, Meibergdreef 9, Amsterdam, 1105AZ, Netherlands
- **Bjermer, Leif**
 - Department of Lung and Allergology, Skane University Hospital, Lund, 22185, Sweden
- **Blanc, François-Xavier**
 - Nantes University, Nantes, 44093, France
- **Blasi, Francesco**
 - Internal Medicine Department, Respiratory Unit and Adult Cystic Fibrosis Center, Fondazione IRCCS Cà Granda Ospedale Maggiore Policlinico, Milano, and Department of Pathophysiology and Transplantation, Università degli Studi di Milano, Italy
- **Bourdin, Arnaud**
 - Department of Respiratory Diseases, University of Montpellier, CHU Montpellier, Montpellier, France
- **Brown, Thomas**
 - Portsmouth Hospitals NHS Trust, Queen Alexandra Hospital, Southwick Hill Road, Cosham, Portsmouth, Hampshire, PO63LY, United Kingdom
- **Brussino, Luisa**
 - Department of Medical Science, University of Torino-Allergy and Immunology Unit, Mauriziano Hospital, C.so Re Umberto 109, Torino, 10100, Italy
- **Burhan, Hassan**
 - Royal Liverpool University Hospital, Prescot Street, Link 6Z, Royal Liverpool University Hospital, Prescot Street, Liverpool, L7 8XP, United Kingdom
- **Calvert, James**
 - North Bristol NHS Trust, Westbury on Trym, Bristol, Avon, BS10 5NB, United Kingdom
- **Caminati, Marco**
 - Department of Medicine, University of Verona & Asthma Center and Allergy Unit, Verona University Hospital, Verona, Italy
- **Campos Cerda, Ricardo**
 - Centro de Atención de Enfermedades Cardiopulmonares, 880 Garibaldi St, Office 9, 1st Floor, Guadalajara, Jalisco, 44200, Mexico
- **Canonica, Giorgio Walter**

- Personalised Medicine, Asthma and Allergy Center, Humanitas Research Hospital, Milan, Italy
- **Caruso, Cristiano**
 - Fondazione Policlinico Universitario A. Gemelli IRCCS, Rome, Italy
- **Cataldo, Didier**
 - University of Liege (GIGA-research center) and CHU of Liege, Tower of Pathology (B23), Hippocrates Avenue 13, Liege, 4000, Belgium
- **Chanez, Pascal**
 - Department of Respiratory Diseases, AP-HM, INSERM, INRA, C2VN Aix Marseille Université, Marseille, France
- **Chanson, Philippe**
 - Assistance Publique-Hôpitaux de Paris, Hôpital de Bicêtre, Service d'Endocrinologie et des Maladies de la Reproduction, Centre de Référence des Maladies Rares de l'Hypophyse; Université Paris-Saclay, Univ. Paris-Sud, Inserm, Signalisation Hormonale, Physiopathologie Endocrinienne et Métabolique, 94276, Le Kremlin-Bicêtre, France
- **Chapman, Ken**
 - Asthma & Airway Centre, University Health Network, University of Toronto, Room 7-451 East Wing, 399 Bathurst Street, Toronto, Ontario M5T 2S8, Canada
- **Chaudhuri, Rekha**
 - NHS Greater Glasgow and Clyde, Asthma/COPD Clinical Research Centre, Gartnavel General Hospital, Glasgow, G62 6QL, United Kingdom
- **Chenivresse, Cécile**
 - CHU Lille, Service de Pneumologie et Immuno-allergologie, Centre de référence constitutif pour les maladies pulmonaires rares, Univ. Lille, Inserm 1019, Centre Infection et Immunité de Lille, Institut Pasteur de Lille, Lille, France
- **Choudhury, Sirazum**
 - Imperial College London, Section of Investigative Medicine, 6th Floor Commonwealth Building, Du Cane Road, London, W12 0NN, United Kingdom
- **Christoff, George**
 - Medical University - Sofia, Faculty of Public Health, 8 "Bialo more" str, Sofia, 1527, Bulgaria
- **Chung, Li Ping**
 - Department of Respiratory Medicine, Fiona Stanley Hospital, Perth, Western Australia 6150, Australia
- **Clairelyne Dupin**
 - Assistance Publique-Hôpitaux de Paris, Hôpital Saint-Louis, 1 avenue Claude Vellefaux, 75475 Paris Cedex 10, France
- **Clifton, Ian**
 - St James's University Hospital, Beckett Street, Leeds, West Yorkshire, LS9 7TF, United Kingdom
- **Cochrane, Belinda**
 - Campbelltown Hospital, Therry Rd, Campbelltown, NSW, 2560, Australia
- **Colantuono, Stefania**
 - Allergy Unit, Fondazione Policlinico Gemelli, IRCCS; Department of Translational and Precision medicine, Sapienza University of Rome, Rome, Italy

- **Cosmi, Lorenzo**
 - University of Firenze, AOU Careggi, Largo Brambilla, Firenze 50100, Italy
- **Costello, Richard**
 - RCSI, Beaumont Hospital, Dublin 9, Ireland
- **Côté, Andréanne**
 - Institut Universitaire Cardiologie et Pneumologie de Québec, Laval University, 2725 ch Ste- Foy, Quebec, G1V 4V5, Canada
- **Crimi, Nunzio**
 - Via Etnea 676, Catania, 95125, Italy
- **Crooks, Michael G.**
 - Hull York Medical School, Academic Respiratory Medicine, Castle Hill Hospital, Cottingham, HU16 5JQ, United Kingdom
- **D'Amato, Maria**
 - Respiratory Department, Monaldi Hospital, Via D. Fontana, 134, Naples, Campania, 80128, Italy
- **De Gennaro, Mónica S.**
 - Fundacion CIDEA, Paraguay 2035 3Cuerpo 2SS, CABA, Buenos Aires, C1121ABE, Argentina
- **Debono, Miguel**
 - Sheffield Teaching Hospitals, Royal Hallamshire Hospital, Glossop Road, Sheffield, S10 2JF, United Kingdom
- **Del Giacco, Stefano**
 - Department of Medical Sciences and Public Health, University of Cagliari, Asse Didattico E1, Cittadella Universitaria, 09042 Monserrato (Cagliari), Italy
- **Dennison, Patrick**
 - University Hospitals Southampton Foundation Trust, Tremona road, Southampton, Hampshire, SO16 6YD, United Kingdom
- **Deschildre, Antoine**
 - CHU Lille, Pneumologie et Allergologie Pédiatriques, Hôpital Jeanne de Flandre, Avenue Avinée, F-59000 Lille, France
- **Detoraki, Aikaterini**
 - Azienda Ospedaliera Universitaria Federico II, Via Pansini 5, Naples, 80131, Italy
- **Devouassoux, Gilles**
 - University Claude Bernard Lyon 1; HCL, Service de Pneumologie, Bâtiment I 103 Grande Rue de la Croix Rousse, F-69004 Lyon, France
- **Didier, Alain**
 - Center for Pathophysiology Toulouse Purpan, INSERM U1043, CNRS UMR 5282, Toulouse III University and CHU Toulouse, France
- **Dorscheid, Del**
 - University of British Columbia, 166 - 1081 Burrard Street, Vancouver, BC, V6Z 1Y6, Canada
- **Fardon, Tom**
 - NHS Tayside, East Block, Ninewells Hospital; Department of Respiratory Research, University of Dundee, Dundee, DD1 9SY, Scotland, United Kingdom
- **Faruqi, Shoaib**

- The Hull University Teaching Hospital NHS Trust, Department of Respiratory Medicine, Castle Hill Hospital, Castle Road, Cottingham, East Yorkshire, HU16 5JQ, United Kingdom
- **FitzGerald, JM Mark**
 - The Lung Centre, Vancouver General Hospital, Institute for Heart and Lung Health, Gordon and Leslie Diamond Health Care, Vancouver, BC, Canada
- **Gaga, Mina**
 - Athens Chest Hospital, 152 Mesogion Ave, Athens 11527, Greece
- **Genova, Sonya**
 - UHATEM N.I.Pirogov, 21 Totleben blvd, Sofia, 1606, Bulgaria
- **Gibson, Peter**
 - University of Newcastle, Lookout RD, New Lambton Hts, Newcastle, NSW 2305, Australia
- **Gore, Robin**
 - Cambridge University NHS Foundation Trust, Box 40, Addenbrooke's Hospital, Cambridge, CB2 0QQ, United Kingdom
- **Guilleminault, Laurent**
 - Toulouse University Hospital Centre, Larrey Hospital, F-31059 Toulouse, France
- **Gurnell, Mark**
 - University of Cambridge & Addenbrooke's Hospital, Cambridge, United Kingdom
- **Hamerlijnck, Dominique**
 - Atini, Zeeburgerkade 540, Amsterdam, Noord-Holland, 1019HR, Netherlands
- **Hanania, Nicola**
 - Baylor College of Medicine, 7200 Cambridge, Suite 8A. 269, Houston, 77030 Texas, United States of America
- **Heaney, Liam**
 - Centre for Experimental Medicine, School of Medicine, Dentistry and Biomedical Sciences, Queen's University Belfast, Wellcome-Wolfson Institute for Experimental Medicine, 97 Lisburn Road, Belfast BT9 7BL, Northern Ireland, United Kingdom
- **Heffler, Enrico**
 - Humanitas University, Via Rita Levi Montalcini 4, Pieve Emanuele, MI, 20090, Italy
- **Hernandez Colin, Dante Daniel**
 - Instituto Jalisciense de Investigacion Clinica Penitenciaria, Centro 20, Guadalajara, Jalisco, 44100, México
- **Hew, Mark**
 - Alfred Health, 55 Commercial Road, Prahran, Melbourne, Victoria 3004, Australia
- **Hoyte, Flavia**
 - National Jewish Health, Division of Allergy and Immunology, Denver, CO, 80206, United States of America
- **Humbert, Marc**
 - Université Paris-Saclay, Inserm, Hôpital Bicêtre (Assistance Publique Hôpitaux de Paris), 78 rue du Général Leclerc, F-94270 Le Kremlin-Bicêtre, France
- **Idzko, Marco**
 - Department of Pneumology, Medical Clinic II, Medical University Vienna, 6L, Währingerstraße 18-20, Vienna, 1090, Austria
- **Jenkins, Christine**

- The George Institute for Global Health, PO Box M201, Missenden Rd, NSW, 2050, Australia
- **Kauppi, Paula**
 - HUH, Inflammation Center, Department of Allergy (Adult Unit), P.O. Box 160, 00029 HUH; Skin and Allergy Hospital, Meilahdentie 2, Helsinki, Finland
- **Kostikas, Konstantinos**
 - Respiratory Medicine Department, University of Ioannina, Ioannina, Greece
- **Kuna, Piotr**
 - Department of Internal Medicine, Asthma and Allergy, Medical University of Lodz, Poland
- **Kupczyk, Maciej**
 - Medical University of Lodz, Kopcinskiego 22, Lodz, 90-350, Poland
- **Kupryś-Lipińska, Izabela**
 - Medical University of Lodz, Department of Internal Medicine, Asthma and Allergy, 22 Kopcinskiego Str, Lodz 90-153, Poland
- **Labor, Marina**
 - University Hospital Centre Osijek, Huttlerova 4, Osijek 31000, Croatia
- **Langton, David**
 - Peninsula Health, 2 Hastings St, Frankston, Victoria, 3199, Australia
- **Latorre, Manuela**
 - Pulmonary Unit, NOA Hospital (Nuovo Ospedale Apuano), Massa, Italy
- **Lehtimäki, Lauri**
 - Faculty of Medicine and Health Technology, Tampere University, Tampere, Finland
- **Louis, Renaud**
 - CHU of Liege, University of Liege, GIGA I3 Research Group, Liege, 4000, Belgium
- **Loukides, Stylianos**
 - National and Kapodistrian University of Athens Medical School, 2nd Respiratory Department, Rimini 1 Xaidari 12462, Greece
- **Lugogo, Njira Lucia**
 - Department of Medicine, Division of Pulmonary and Critical Care Medicine, University of Michigan, 300 North Ingalls St Suite 2C40, Ann Arbor, Michigan, 48109, United States of America
- **Mahay, Guillaume**
 - Centre Hospitalier Universitaire de Rouen, Service de Pneumologie, Oncologie Thoracique, Soins Intensifs Respiratoires, 1 rue de Germont, F-760031 Rouen, France
- **Mahboub, Bassam**
 - DHA & University of Sharjah, PO box 4545, Dubai 4546, United Arab Emirates
- **Masoli, Matthew**
 - Royal Devon & Exeter Hospital, Respiratory Department, Barrack Road, Exeter, Devon, EX2 5DW, United Kingdom
- **Maspero, Jorge**
 - Fundacion Cidea, Paraguay 2035, Segundo Subsuelo, Buenos Aires, Caba C1121ABE, Argentina
- **Meeran, Karim**
 - Imperial College London, Department of Endocrinology, 9th floor, East Wing, Charing Cross campus, Fulham Palace Road, London W6 8RF, United Kingdom
- **Menzella, Francesco**

- Azienda USL di Reggio Emilia-IRCCS, Pneumology Unit, Santa Maria Nuova Hospital, Via Amendola 2, Reggio Emilia 42122, Italy
- **Menzies-Gow, Andrew**
 - Royal Brompton and Harefield NHS Foundation Trust, London, United Kingdom
- **Middleton, Peter**
 - Westmead Hospital, Hawkesbury Rd, Westmead, Sydney, NSW 2145, Australia
- **Milanese, Manlio**
 - Pulmonology Unit S. Corona Hospital, ASL2 Savonese, Via XXV Aprile 38, Pietra ligure, Savona 17027, Italy
- **Mitchell, Patrick D.**
 - Rockyview General Hospital, 7007 14 ST SW, Calgary, Alberta, T2V 1P9, Canada
- **Mohan, Arjun**
 - ECU Pulmonary Clinic & Severe Asthma Program, Division of Pulmonary and Critical Care Medicine, Brody School of Medicine-East Carolina University, 3E-111A, BSOM, Greenville, NC, 27834, United States of America
- **Paggiaro, Pierluigi**
 - University of Pisa, via Paradisa 2, Pisa, 56124, Italy
- **Papadopoulos, Nikolaos G.**
 - Division of Infection, Inflammation and Respiratory Medicine, University of Manchester, Manchester, United Kingdom; Allergy Dpt, 2nd Pediatric Clinic, National Kapodistrian University of Athens, Athens, Greece
- **Papaioannou, Andriana I.**
 - Attikon Hospital, Rimini 1, Chaidari, Athens, 12462 Greece
- **Pavord, Ian**
 - University of Oxford, Respiratory Medicine Unit, Nuffield Department of Medicine, Oxford OX3 7FZ, United Kingdom
- **Peché, Rudi**
 - CHU Charleroi site Vesale, Rue de Gozee 706, Montingny-l-Tilleul, Hainaut, 6110 Belgium
- **Pelaia, Corrado**
 - University Magna Graecia of Catanzaro, Viale Europa, Catanzaro, 88100, Italy
- **Pelaia, Girolamo**
 - University Magna Graecia of Catanzaro, Viale Europa, Catanzaro, 88100, Italy
- **Perng, Diahn-Warng**
 - Taipei Veterans General Hospital, 201 Shi-Pai Rd, Sec 2, Taipei, 11217 Taiwan
- **Pfeffer, Paul**
 - Barts Health NHS Trust, St Bartholomew's Hospital, West Smithfield, London EC1A 7BE, United Kingdom
- **Pilette, Charles**
 - Université catholique de Louvain, Cliniques Universitaires St-Luc and Institute of Experimental and Clinical Research, Department of Pulmonology, 10 Avenue Hippocrate, Brussels, B-1200, Belgium
- **Pison, Christophe**
 - UGA, Service Hospitalier Universitaire Pneumologie Physiologie, Pôle Thorax et Vaisseaux, Centre Hospitalier et Universitaire de Grenoble, Grenoble, CS10217, 38043 Grenoble Cedex 9, France
- **Plavec, Davor**
 - Srebrnjak Children's Hospital, Srebrnjak 100, Zagreb, 10000, Croatia; Medical Faculty, University JJ Strossmayer, Osijek, 31000, Croatia

- **Popov, Todor A**
 - University Hospital Sv. Ivan Rilski, 13, Urvich St., Sofia, 1612, Bulgaria
- **Popović-Grle, Sanja**
 - University Hospital Centre Zagreb, Clinic for Lung Diseases Jordanovac, Jordanovac 104, Zagreb, 10000; School of Medicine, University of Zagreb, Croatia
- **Powlson, Andrew S**
 - University of Cambridge, Institute of Metabolic Science, Addenbrooke's Hospital, Cambridge, CB2 0QQ, United Kingdom
- **Puggioni, Francesca**
 - Personalized Medicine, Asthma, and Allergy, Humanitas Clinical and Research Center, IRCCS, Milan, Italy
- **Reddel, Helen**
 - Woolcock Institute of Medical Research, University of Sydney, 431 Glebe Point Rd, Glebe, NSW 2037, Australia
- **Rhee, Chin Kook**
 - Seoul St. Mary's Hospital, The Catholic University of Korea, 222 Banpodaero, Seochogu, Seoul, South Korea
- **Roche, Nicolas**
 - Respiratory Medicine, Cochin Hospital (APHP - Centre), University Paris Descartes (UMR 1016 Institut Cochin), Université de Paris, Paris, France
- **Rupani, Hitasha**
 - Portsmouth Severe Asthma Service, Queen Alexandra Hospital, Southwick Hill Road, Portsmouth, Hampshire, PO6 5LY, United Kingdom
- **Sabroe, Ian**
 - Sheffield Teaching Hospitals NHS Foundation Trust, Royal Hallamshire Hospital, Glossop Road, Sheffield, S10 2JF, United Kingdom
- **Samitas, Konstantinos**
 - Athens Chest Hospital "SOTIRIA", Mesogion 152 Ave, Athens, Attiki 11527, Greece
- **Santus, Pierachille**
 - Department of Biomedical and Clinical Sciences, University of Milan, Via GB Grassi, 74, Milano, 20157, Italy
- **Schleich, Florence**
 - Rue Hubert Delvenne 1, Hody, 4172, Belgium
- **Selmi, Carlo**
 - Division of Rheumatology and Clinical Immunology, Humanitas Clinical and Research Center – IRCCS, Humanitas University, Via A. Manzoni 113, Rozzano, Milano, 20089, Italy
- **Senna, Gianenrico**
 - Allergy Unit – Asthma Center, University of Verona, Italy
- **Sherlock, Mark**
 - Department of Endocrinology, Beaumont Hospital, Beaumont, Dublin D9 and Royal College of Surgeons, Ireland
- **Siddiqui, Salman**
 - University of Leicester and Leicester NIHR Biomedical Research Centre (Respiratory Theme), Glenfield Hospital, Leicester, LE3 9QP, England, United Kingdom
- **Smith, Andrew**

- NHS Lanarkshire, University Hospital Wishaw, 50 Netherton Street, Wishaw, Lanarkshire, ML2 0DP, Scotland, United Kingdom
- **Spanevello, Antonio**
 - Dipartimento di Medicina e Chirurgia, Malattie dell'Apparato Respiratorio, Università degli Studi dell'Insubria, Varese – Como; Dipartimento di Medicina e Riabilitazione Cardio Respiratoria, U.O. di Pneumologia Riabilitativa, Istituti Clinici Scientifici Maugeri, IRCCS Tradate, Italia
- **Taillé, Camille**
 - Assistance Publique-Hôpitaux de Paris and Université de Paris, 46 rue Henri Huchard, F-75018 Paris Cedex, France
- **Taube, Christian**
 - Universitätsmedizin Essen- Ruhrlnadklinik, Tüschener Weg 40, Essen, 45239, Germany
- **ten Brinke, Anneke**
 - Medical Centre Leeuwarden, H Dunantweg 2, Leeuwarden, 8934AD, Netherlands
- **Tudoric, Neven**
 - Clinical Hospital Dubrava, Avenija Gojka Suska 6, Zagreb, 10000, Croatia
- **Tunsäter, Alf**
 - Department of Respiratory Medicine & Allergology, Skane University Hospital, Lund, 221 85, Sweden
- **Ulrik, Charlotte**
 - Head of Respiratory Research Unit & Severe Asthma Clinic Hvidovre, Department Of Respiratory Medicine, Hvidovre University Hospital, DK-2650 Hvidovre, Denmark
- **Upham, John**
 - Princess Alexandra Hospital, 199 Ipswich Road, Woolloongabba, Brisbane, Qld 4102, Australia
- **Van Ganse, Eric**
 - PharmacoEpidemiology Lyon (PELyon), Faculté d'Odontologie; HESPER, Claude Bernard Lyon1 University, Respiratory Medicine; Croix Rousse University Hospital, 69008 Lyon, France
- **Walker, Brandie**
 - Canada Division of Respirology, Department of Medicine, University of Calgary; Calgary COPD and Asthma Program, Alberta Health Services; Airways Working Group, Respiratory Health Strategic Clinical Network; 1007 Health Sciences Centre, 3330 Hospital Drive NW, Calgary, AB, T2N 4N1, Canada
- **Wang, Eileen**
 - National Jewish Health, 1400 Jackson Street, K624a, Denver, CO, 80238; University of Colorado, 13001 East 17th Place, Aurora, CO, 80045, United States of America
- **Wark, Peter**
 - Priority Research Centre for Healthy Lungs, Respiratory Medicine HMRI, Lookout Road, New Lambton, New South Wales 2305, Australia
- **Wechsler, Michael**
 - NJH Cohen Family Asthma Institute, Department of Medicine, National Jewish Health, 1400 Jackson St, Denver, CO 80206, United States of America
- **Winders, Tonya**
 - Allergy & Asthma Network, 8229 Boone Blvd, Ste 260, Vienna, VA, 22182, United States of America
- **Zervas, Eleftherios**

- 7th Resp. Department and Asthma Centre, Athens Chest Hospital, Mesogion Ave. 152, Athens, Attica 11527, Greece