



HAL
open science

Uveitis associated with cancer immunotherapy: long-term outcomes

Christine Fardeau, Mehdi Bencheqroun, Arielle Levy, Sophie Bonnin,
Marie-Adélaïde Ferchaud, Leïla Fardeau, Florence Coscas, Bahram Bodaghi,
Bénédicte Lebrun-Vignes

► **To cite this version:**

Christine Fardeau, Mehdi Bencheqroun, Arielle Levy, Sophie Bonnin, Marie-Adélaïde Ferchaud, et al..
Uveitis associated with cancer immunotherapy: long-term outcomes. *Immunotherapy*, 2021, 13 (18),
pp.1465-1481. 10.2217/imt-2021-0032 . hal-03543514

HAL Id: hal-03543514

<https://hal.sorbonne-universite.fr/hal-03543514>

Submitted on 26 Jan 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

1 Title page

2

3 Uveitis associated with cancer immunotherapy: long-term outcomes

4

5 Authors: Christine Fardeau¹, Mehdi Bencheqroun¹, Arielle Levy¹, Sophie Bonnin^{1,2}, Marie-
6 Adélaïde Ferchaud¹, Leila Fardeau³, Florence Coscas⁴, Bahram Bodaghi¹, Bénédicte Lebrun-
7 Vignes⁵.

8

9 Corresponding author: Dr. Christine Fardeau

10

11

12 ¹ Department of ophthalmology, Reference center for rare diseases, La Pitié-Salpêtrière
13 Hospital, Paris-Sorbonne University, 47-83 Boulevard de l'Hôpital, 75013 Paris, France.

14 ² Department of ophthalmology, Fondation Rothschild, Université de Paris, 25, rue Manin,
15 75019 Paris

16 ³ Institut de Démographie, Université Paris 1 - Panthéon Sorbonne.

17 ⁴ Centre Ophtalmologique de l'Odéon, 113 bd Saint Germain, Paris, Agora Académie.

18 ⁵ Department of Pharmacovigilance, La Pitié-Salpêtrière Hospital, Paris-Sorbonne University,
19 47-83 Boulevard de l'Hôpital, 75013 Paris, France.

20

21

22 Abbreviated title: Cancer immunotherapy-associated uveitis

23

24 **Key words**

25 Uveitis, macular edema, hypopion, cancer immunotherapy, rituximab, nivolumab,
26 ipilimumab, anti-BRAF, anti-MEK, ibritunib

27

28 **Abstract**

29 **Background :** to report the long-term outcome of uveitis associated with cancer
30 immunotherapy (CIT)

31 *Methods:* retrospective review of serial patients with CIT-associated uveitis treated using
32 various regimen.

33 *Results:* Eight included patients treated with rituximab (anti-CD20), nivolumab (anti-PD-1),
34 ipilimumab (anti-CTLA-4), vemurafenib and dabrafenib (anti-BRAF), trametinib (anti-MEK),
35 and ibritunib, showed uveitis with hypopion (1 patient), macular edema (5 patients) and
36 choroiditis (2 patients). Various regimen of corticosteroid therapy showed a favorable
37 ophthalmological outcome, whether the CIT was continuing or suspended.

38 **Discussion-Conclusion:** Local corticosteroids injections in combination with CIT could be
39 suggested as a first-line treatment. This could help to preserve the quality of life without
40 threatening the vital prognosis.

41

42 **LayAbstract:**

43

44 *Purpose:*To report the long-term outcome of intra-ocular inflammation (uveitis) associated
45 with cancer immunotherapy (CIT)

46 *Methods:* Serial patients complaining of blurred vision and painful eyes, showed intra-ocular
47 inflammation that was related to CIT, after infectious, inflammatory, and tumoral causes of
48 uveitis have been ruled out. The length of follow-up was more than 12 months for most
49 patients.

50 *Results:* Eight serial patients treated with rituximab (anti-CD20), nivolumab (anti-PD-1),
51 ipilimumab (anti-CTLA-4), vemurafenib and dabrafenib (anti-BRAF), trametinib (anti-MEK),
52 and ibritunib, showed intra-ocular inflammation with hypopion (1 patient), macular edema (5
53 patients) and choroiditis (2 patients). Various regimen of corticosteroid therapy showed a
54 favorable ophthalmological outcome, whether the CIT was continuing or suspended.

55 *Conclusions:* Local corticosteroids injections in combination with CIT could be suggested as
56 a first-line treatment. This could help to preserve the quality of life without threatening the
57 vital prognosis.

58 **Key words**

59 Uveitis, macular edema, hypopion, cancer immunotherapy, rituximab, nivolumab,
60 ipilimumab, anti-BRAF, anti-MEK, ibritunib

61 **Summary points :**

- 62 1. This study reported 8 cases of CIT-associated uveitis, among which 7 experienced a visual
63 loss and 5 had a macular edema, 2 had nodular choroiditis, 1 had hypopion.
- 64 2. Changes in macular edema, choroiditis, and hypopion were analysed for a long-term
65 follow-up superior to 12 months.
- 66 3. The underlying metastatic cancer were heterogenous, and immunotherapy drugs were
67 numerous. However the clinical ophthalmological uveitis related to CIT showed a similar
68 good response to the corticosteroid treatment, regardless of further CIT.
- 69 4. At 3 months, the short-term visual outcomes were favorable after treatment with systemic
70 or local injections of cortosteroids, whether the CIT was continuing or suspended.
- 71 5. At 12 months, CIT continuation was associated with uveitis activity, that showed sustained
72 good response to sub-Tenon's triamcinolone injections.
- 73 6. So the role of the ophthalmologist may be useful in (1) making quickly the diagnosis
74 relating uveitis to CIT, ruling out infectious, auto-immune and tumoral causes (2) explaining
75 the often favorable uveitis outcome using corticosteroid local injections (3) reassuring
76 patients and oncologists, worried about visual fonction (4) performing quickly
77 ophthalmological treatment to keep a good quality of life (5) following the patients to assess
78 positive and negative treatment effects (6) repeating the corticosteroid local injections if
79 needed (7) rediscussing the treatment in case of ophthalmological failure.
- 80

81

82

83

84

85

86

87 Immunotherapy
88• 2021 Oct 28.
89 doi: 10.2217/imt-2021-0032. Online ahead of print.

90 Uveitis associated with cancer immunotherapy: long-term outcomes

91 [Christine Fardeau](#)¹, [Mehdi Bencheqroun](#)¹, [Arielle Levy](#)¹, [Sophie Bonnin](#)^{1,2}, [Marie-Adélaïde](#)
92 [Ferchaud](#)¹, [Leila Fardeau](#)³, [Florence Coscas](#)⁴, [Bahram Bodaghi](#)¹, [Bénédicte Lebrun-Vignes](#)⁵

93 Affiliations collapse

94 Affiliations

- 95• ¹Department of Ophthalmology, Reference Center for Rare Diseases, La Pitié-Salpêtrière
96 Hospital, Paris-Sorbonne University, Paris, 75013, France.
97• ²Department of Ophthalmology, Fondation Rothschild, Université de Paris, Paris, 75019,
98 France.
99• ³Institut de Démographie, Université Paris 1 - Panthéon Sorbonne, Paris, France.
100• ⁴Centre Ophtalmologique de l'Odéon, Agora Académie, Paris, France.
101• ⁵Department of Pharmacovigilance, La Pitié-Salpêtrière Hospital, Paris-Sorbonne University,
102 Paris, 75013, France.
103 PMID: 34709074 DOI: [10.2217/imt-2021-0032](https://doi.org/10.2217/imt-2021-0032)

104 Abstract

105 **Background:** We report the long-term outcome of uveitis associated with cancer
106 immunotherapy (CIT). **Methods:** This retrospective review included serial patients with CIT-
107 associated uveitis treated using various regimen. **Results:** Eight patients treated with
108 rituximab (anti-CD20), nivolumab (anti-PD-1), ipilimumab (anti-CTLA-4), vemurafenib and
109 dabrafenib (anti-BRAF), trametinib (anti-MEK) and ibritunib showed uveitis with hypopion
110 (one patient), macular edema (five patients) and choroiditis (two patients). Various regimens
111 of corticosteroid therapy showed a favorable ophthalmological outcome, whether the CIT was
112 continuing or suspended. **Conclusion:** Local corticosteroid injections in combination with
113 CIT could be suggested as a first-line treatment. This could help to preserve the quality of life
114 without threatening the vital prognosis.

115 **Keywords:** anti-BRAF; anti-MEK; cancer immunotherapy; hypopion; ibritunib; ipilimumab;
116 macular edema; nivolumab; rituximab; uveitis.

117 Plain Language Summary

118 Lay abstract This study aims to report the long-term outcome of intra-ocular inflammation (uveitis)
119 associated with cancer immunotherapy (CIT). Serial patients complaining of blurred vision and painful
120 eyes showed intra-ocular inflammation that was related to CIT, after infectious, inflammatory and
121 tumoral causes of uveitis have been ruled out. The length of follow-up was more than 12 months for
122 most patients. Eight serial patients treated with rituximab (anti-CD20), nivolumab (anti-PD-1),
123 ipilimumab (anti-CTLA-4), vemurafenib and dabrafenib (anti-BRAF), trametinib (anti-MEK) and ibritunib
124 showed intra-ocular inflammation with hypopion (one patient), macular edema (five patients) and
125 choroiditis (two patients). Various regimens of corticosteroid therapy showed a favorable
126 ophthalmological outcome, whether the CIT was continuing or suspended. Local corticosteroid
127 injections in combination with CIT could be suggested as a first-line treatment. This could help to
128 preserve the quality of life without threatening the vital prognosis

129 **Introduction**

130 Uveitis can affect all layers of the eye and may be due to various etiologies: autoimmune,
131 infectious, iatrogenic or tumor such as masquerade syndrome. Macular edema is the leading
132 cause of visual loss in uveitis [1,2]. Recently, a new iatrogenic uveitis entity has been
133 associated with cancer immunotherapy (CIT). CIT inhibits T-cell inactivation driven by tumor
134 cells [3,4]. The promising results of CIT have led to an increasing number of indications
135 involving larger cohorts of patients [5]. For 10 years, about 50 novel agents have been
136 approved for the treatment of metastatic cancers, including (1) monoclonal antibodies directed
137 against tumor cell surface receptors (such as rituximab, an anti-CD20); (2) immune
138 checkpoint inhibitors (ICIs) such as anti-PD-1 (pembrolizumab, nivolumab), anti-PD-L1
139 (atezolimab, avelumimab) and anti-CTLA-4 (ipilimumab) monoclonal antibodies that block
140 the interaction between inhibitory T-cell receptors and their ligands; and (3) small-molecule
141 kinase inhibitors, in particular BRAF inhibitors (vemurafenib, dabrafenib) and MEK
142 inhibitors (trametinib) that inhibit the mitogen-activated protein kinase signaling pathway
143 involved in cell regulation.

144 Immune-related adverse events (irAEs) have been associated with ICIs, BRAF inhibitors and
145 MEK inhibitors in about 25% of patients, including skin (vitiligo, rash, pruritis) and liver
146 (auto-immune hepatitis) AEs as well as colitis and thyroiditis [6,7]. However, the severity of
147 the reported ocular AEs varies from mild (dry eyes and conjunctivitis) to serious (intraocular
148 and orbital inflammation).

149 Intraocular inflammation was mainly reported in the anterior chamber while a posterior
150 segment involvement may lead to Vogt-Koyanagi-Harada (VKH) disease, retinal vascular
151 occlusion, macular edema and papillitis [8-19].

152 The aim of this study was to report the long-term outcomes of patients with CIT-associated
153 uveitis, and discuss the various therapeutic options based on a literature review.

154
155 **Patients and methods**

156 Serial patients with CIT-associated intraocular inflammation seen between January 1, 2018
157 and December 31, 2019 were retrospectively included. Patients treated with CIT and reporting
158 eye problems were promptly referred to an ophthalmologist and then to a reference tertiary
159 uveitis center. A collegial therapeutic discussion about the therapeutic interventions involving
160 oncologists, and ophthalmologists was held in a multidisciplinary meeting.

161 This was a retrospective interventional case series on the approved indications of CIT in
162 France. Informed consent was obtained for all patients before performing retinal angiography,
163 aqueous humor sampling, and before treatment change. Ethic institutional committee approval
164 was obtained.

165 For each patient, the following data were collected: cancer history and disease progression,
166 previous and current treatments, and detailed ophthalmologic examinations performed at
167 baseline and during the follow-up.

168 All subjects underwent a comprehensive eye examination including the best-corrected visual
169 acuity (BCVA) on a decimal scale converted into LogMAR, a slit-lamp examination, and a
170 dilated fundus examination. Intraocular inflammation in the aqueous humor and vitreous was
171 scored using the SUN grading system[20]. Color fundus photographs, autofluorescence
172 photographs, fluorescein (FA) and indocyanine green (ICGA) angiography were performed
173 using CR2 plus (Canon, Tokyo, Japan) devices, the HRA2 (Heidelberg Engineering,
174 Heidelberg, Germany), and Optos (Optos Inc. Marlborough, MA 01752 USA) respectively.
175 Spectral-Domain Optical Coherence Tomography (SD-OCT) was performed using the
176 Spectralis (Heidelberg Engineering, Heidelberg, Germany). Flare measurements were
177 performed using the FC-2000® flare meter (KOWA, Tokyo, Japan), to objectively quantify
178 the inflammatory reaction in the aqueous humor [21]. Goldmann visual field examinations

179 were performed by orthoptists. A thorough work-up was performed to exclude infectious and
180 common ocular and systemic auto-immune causes of uveitis. Particular attention was paid to
181 the systemic causes such as sarcoidosis that can be induced by CIT [22]. So the work-up
182 included for all patients serogy for Human Immunodeficiency virus, syphilis, Lyme disease, B
183 and C hepatitis, and Quantiferon® test, along with blood count, ionogram, protein
184 electrophoresis, sedimentation rate, angiotensine convertase enzyme, lysozyme, antinuclear
185 and anti DNA. In case of clinical doubt aqueous humor sampling was performed for
186 IL10/IL6 rate to rule out intraocular lymphoma diagnosis. HLA A29 was tested to rule out
187 Birdshot retinochoroiditis which is an auto-immune posterior uveitis characterized by fundus
188 showing deep cream-coloured spots predominantly located in nasal inferior to the optic nerve
189 head. Chest scan and brain magnetic resonance imaging results were known thanks to the
190 patient's oncologic department. If induced-sarcoidosis was suspected, pulmonary function
191 tests and salivary glandular biopsy were performed. Alternative causes of autoimmune-
192 associated or infectious uveitis were excluded from the analysis.

193
194 AE severity was graded according to the Common Terminology Criteria for Adverse Events
195 (CTCAE) version 5.0 [23].

196 The tumor response to CIT was classified according to the revised RECIST guideline [24].

197 AEs were reported to the National and world Global pharmacovigilance after checking
198 accurately each case.

199 Clinical uveitis features along with OCT, FA, ICGA, and flare meter results were described at
200 baseline and during the follow-up.

201 Therapeutic interventions were collegially discussed and decided by consensus with the
202 oncologists. They included ICI discontinuation or continuation, ICI resumption or not if ICI
203 had been previously discontinued and/or the use of systemic or local corticosteroids. Local
204 corticosteroids were given as drops or subconjunctival dexamethasone injections. Delayed
205 local corticosteroids were given as sub-Tenon's triamcilonone injections. Systemic
206 corticosteroids were given as methylprednisone pulses for 3 days followed by oral prednisone
207 at 1mg/kg for 10 days and then tapered until a daily dose of 10 mg/kg.

208 Patients were followed one month after the first examination and at least quarterly thereafter
209 to assess the efficacy of the therapeutic intervention and the occurrence of AEs. The study
210 was conducted in compliance with the Good Clinical Practice and the tenets of the
211 Declaration of Helsinki. We focussed on chronic intraocular inflammation that is a rare irAEs.
212 However, this study gave us the opportunity to discuss the risk/benefit ratio in a context of
213 life-threatening disease.

214 **Statistical analysis.**

215 Descriptive statistical analysis was performed using STATA/SE 11.0 software (College
216 Station, Texas, US). Variables were compared using a U Mann-Whitney non-parametric test
217 for continuous variables and a p-value <0.05 was considered significant.

218

219 **Results**

220 Eight patients (4 women and 4 men) with intraocular inflammation associated with CIT given
221 for metastatic cancer were included (table 1). Patients median age was 68.5 years (IQ1 60.5,
222 IQ3 75.7; range: 58-83). The presence of neoplasia and its grading are described in table 1.

223 Treatments included rituximab (case 1), ibrutinib (case 2), nivolumab (cases 4, 5, 6, 8),
224 ipilimumab (case 8), dabrafenib + trametinib (case 3), and verumafenib (case 7). One patient
225 (case 3) received a combination of nivolumab, dabrafenib and trametinib for 6 months and
226 only dabrafenib and trametinib thereafter (table 1).

227

228 Three patients experienced additional extra-ocular irAEs, including interstitial pneumonitis
229 (cases 3, 6), cryoglobulinemia (case 3), and peripheral neuropathy (case 7) leading to CIT
230 discontinuation (table 1).

231 Uveitis occurred after a median treatment duration of 4.5 months (IQ1 3.7, IQ4 14.2; range: 1-
232 24 months). Upon admission, uveitis was predominant in the anterior segment in 3 patients
233 (cases 5, 7, 8), including in association with hypopyon in 1 case (figure 5), in the intermediate
234 segment in 2 patients (cases 1, 3) and 3 patients (cases 2, 4, 6) had panuveitis. The baseline
235 features and grading of uveitis are shown in table 2. None of the patients had a previous
236 history of uveitis.

237 The median baseline VA was 0.2 LogMAR (range: 0-2) in both eyes. One patient (case 1)
238 with a history of corneal scars was monophthalmic. The median flare was 26.2 photons/ms
239 (range: 7.9-315.3) in both eyes. The median intraocular pressure was 15 mmHg (range: 9-18).
240 Posterior synechias were present in 6 eyes of 4 patients (table 2).

241 Macular edema was present in 9 eyes of 5 patients, and 3 of these patients showed macular
242 cysts that were mainly located in the inner nuclear layer (cases 2, 3, 4, figure 3). The median
243 central foveal thickness was 260 microns (range: 248-500) in both eyes. Choroidal nodular
244 lesions were seen in 4 eyes of 2 patients (cases 2, 4); on ICGA, they appeared hypofluorescent
245 10 minutes after dye injection and disappeared after 30 minutes, suggesting a stronger
246 inflammatory choroidal involvement than suggested on the fundus examination (Figure 4).

247 No case of diplopia and ocular paralysis was found. No systemic auto-immune disease
248 induced by CIT was found; in particular the sarcoidosis work-up was negative.

249 The first-line therapeutic intervention was based on corticosteroids. Systemic corticosteroids
250 were given in 2 patients with a good ophthalmological response at 3 months in both cases
251 (case 2 and case 6, respectively). Corticosteroids were injected locally in 4 patients with a
252 good ophthalmological response at 3 months. CIT was discontinued in 4 patients: 2 due to
253 ocular AEs (cases 1, 5) and 2 due to extra-ocular AEs (cases 6, 7) (Table 1).

254 At 3 months, the median BCVA was 0.05 LogMAR (range: 0-2) in both eyes and was
255 significantly different compared to the mean baseline VA ($p = 0.02$). The median flare ($n=5$)
256 was 8.0 photons/ms (IQ1 3.1, IQ4 19.6; range: 2.1-54.3) in both eyes and was significantly
257 different compared to the baseline value ($p = 0.001$). The median central foveal thickness was
258 273 microns (IQ1 264, IQ4 366; range: 231-376) in both eyes and was significantly different
259 compared to the baseline value ($p < 0.001$).

260
261 The second-line therapeutic intervention consisted in CIT resumption at various timepoints:
262 (1) at 1 month, with a switch from vemurafenib to nivolumab in 1 patient (case 7), and from
263 ipilimumab to nivolumab in 1 patient (case 8); (2) at 6 months in 1 patient (case 5), associated
264 with an increase in tumor mass leading to a rapid death; and (3) at 3 years in 1 patient (case 1)
265 without uveitis recurrence in the short term.

266 Meanwhile, CIT was continued in 3 patients (cases 2, 3, 4) and did not seem to prevent a good
267 ophthalmological response at 3 months in these three cases. However, bilateral uveitis
268 remained active in 2 out of these 3 patients (cases 3, 4). Case 3 presented a good response to
269 sub-Tenon's triamcilonone injections with a sustained efficiency on the macular edema. Case
270 4 experienced a recurrence of bilateral uveitic macular edema, but unfortunately the patient
271 died before receiving a new ophthalmological intervention.

272

273 **Discussion**

274 This study reported 8 cases of CIT-associated uveitis, among which 7 experienced a visual
275 loss. Five had a macular edema, 2 had nodular choroiditis, and 1 had hypopyon. The follow-up
276 was at least 12 months, except for the last included patient case 8. So changes in ME,
277 choroiditis, and hypopyon had been analysed for a long-term period of time. First, at 3

278 months, the short-term visual outcomes were favorable after corticosteroids treatment using
279 either systemic or local injections way. CIT was continued in 3 patients, with no negative
280 effects on the ophthalmological response. At 12 months, CIT continuation was associated
281 with sustained uveitis activity in 2 patients: one died a few weeks later and the other one
282 received sub-Tenon's triamcinolone injections every 10 months.

283 The occurrence of CIT uveitis has been currently described in metastatic melanomas (4/8
284 patients) as in other advanced cancers, suggesting that the uveitis is associated with the CIT
285 rather than related to the immune response to the original cancer. Moreover the visual fields
286 of the current patients were preserved contrary to what is usually found in cancer associated
287 retinopathy and melanoma associated retinopathy which are subtypes of paraneoplastic
288 syndrome. Two current patients were investigated for antigenic retinal proteins known to be
289 remated to cancer-associated retinopathy, in particular arrestin and recoverin search was
290 negative.

291
292 The reported incidence of severe ocular irAEs is less than 1% [16,25]. A prevalence and an
293 incidence of irAEs of respectively 0.4% and 0.7 per 1,000 patients-months have been reported
294 in patients treated with anti-PD-L1 [10]. Another recent systematic review on the ocular
295 manifestations associated with ICIs has reported an estimated prevalence ranging between
296 0.3% and 0.6% [13].

297 Combining nivolumab and ipilimumab has been associated with a stronger antitumor effect,
298 but also with more irAEs, such as uveitis, with a frequency of 6% [26-28]. In a systematic
299 review of 234 patients, ipilimumab has also been strongly associated with irAEs, including
300 autoimmune complications in 10.3% of cases, 4.3% being classified as uveitis [29].

301 In a review on ocular toxicities induced by targeted anticancer agents, Fu et al. have found
302 that small-molecule drugs are associated with a higher incidence of ocular toxicities than
303 monoclonal antibodies (37.5% *versus* 28.6%) [30].

304
305 A mean age of 54 ± 10 years has been reported at the time of the diagnosis of uveitis in
306 patients treated with anti-CTLA-4 and anti-PD-1 antibodies (ipilimumab/pembrolizumab,
307 nivolumab/acetoluzumab, avelumab, durvalumab) [31]. In the current study the patients were
308 younger and this could be due to the various types of neoplasia included.

309
310 Uveitis occurred after a median treatment duration of 4.5 months, close to previously described
311 treatment durations. Indeed, in a review of 15 cases of uveitis associated with anti-CTLA-4
312 and anti-PD-1 antibodies, most cases occurred in the first 6 months following ICI initiation
313 [31]. However, shorter treatment durations have also been associated with the occurrence of
314 uveitis. In patients treated with nivolumab and pembrolizumab, 15 cases of uveitis have been
315 reported by Wei Wang et al. after a median treatment duration of 9 weeks [32]. After an
316 infusion of anti-PD-1 and anti-PD-L1 antibodies, the median period of time to ocular irAE
317 occurrence was 29 ± 41 days [10]. In our study, uveitis also occurred after treatment
318 discontinuation in one patient as previously described [10].

319
320 The CIT-associated uveitis is usually bilateral; no strictly unilateral cases have been reported.
321 They corresponded either to enucleated patients or corneal scars preventing the intraocular
322 evaluation, as for one patient of the current study [8,33].

323
324 The spectrum of CIT-associated uveitis ranged from anterior uveitis to panuveitis. Most of the
325 reported cases had anterior uveitis, as in the case series of 22 CIT-associated uveitis reported
326 by Whilst et al. in which half of the patients had anterior uveitis [8,16,34-40]. Posterior
327 segment inflammation showed various aspects, and most of them are summarized in table 3

328 [41-59]. Serous retinal detachment suggesting VKH-like disease has been reported with the
329 use of anti-CTLA-4 treatment [41,42] and nivolumab combined with antiBRAF [11,46].
330 Moreover, bilateral cystoid macular edema shown in 4 of the current cases has been
331 previously reported to be associated with nivolumab [14,15,45], BRAF [12,52] and MEK
332 inhibitors [8,53]. The cysts were mainly located in the inner nuclear layer in 3 out of the 4
333 current cases experiencing cystoid macular edema (cases 1, 3, 4), as previously shown in a
334 large case series [8]. The inner and outer layers were more rarely involved (case 1) [45]. ME
335 has been previously reported in patients treated with rituximab [57-59].

336 An association between ibrutinib, Bruton tyrosine kinase inhibitor, and macular edema has
337 been previously reported [56]. In the global pharmacovigilance database (Vigilyze), 46 cases
338 of uveitis associated with ibrutinib (3 in France) have been reported, and ibrutinib was the
339 only suspected drug in 44 cases. The disproportionality analysis (based on the IC025) was
340 significant, suggesting an over-representation of AEs compared to other drugs.

341
342 Optic disc edema, papillitis, and leakage of the optic nerve head have been reported, in
343 particular in patients treated with pembrolizumab [34], in patients with severe hypotony
344 treated with pembrolizumab [50], and in patients treated with nivolumab [15]. Choroidal
345 involvements seen as hypofluorescent spots on ICGA were found in 2 current patients and
346 previously reported in patients treated with ipilimumab, nivolumab and pembrolizumab
347 [16,43,44,47].

348
349 In a recent study of 22 cases of CIT-associated uveitis, the combination of trametinib and
350 dabrafenib has been most often associated with the occurrence of inflammatory macular
351 edema [8]. Among the ICIs, nivolumab has been associated with posterior pole inflammation
352 leading to macular edema or disc leakage, and peripheral leakage on FA [15,31,43,45].
353 Pembrolizumab has also been associated with macular edema and disc and peripheral leakage
354 [33,48].

355 In patients treated with anti-CTLA-4 antibodies, including ipilimumab and tremelimumab,
356 secondary uveitis has been frequently reported. Anti-CTLA-4 antibodies act on the lymph
357 nodes while anti-PD-1 antibodies act in the peripheral tissues [60]. This could explain why
358 CTLA-4-associated irAEs are more common than anti-PD-1/anti-PD-L1-associated irAEs
359 [61].

360
361 Uveitis was scored using the CTCAE version 5.0 that defines successive grades [23]. Grade 1
362 corresponds to anterior uveitis with trace cells, grade 2 to anterior uveitis with 1 to 2+ cells,
363 grade 3 to anterior uveitis with 3+ or more cells or intermediate/posterior/panuveitis and
364 grade 4 to a vision at 20/200 or worse in the affected eye. Thresholds at which treatment
365 should be discontinued have been defined based on irAE severity. In particular, ICI
366 discontinuation in the presence of any grade 3 or 4 irAE has been suggested, associated or not
367 with the use of systemic corticosteroids [61]. However, the application of the CTCAE grading
368 scale may be limited to ocular toxicity management. In our series, even grade 3 and 4 irAEs
369 responded well to dexamethasone subconjunctival injections, long-term dexamethasone
370 drops, and sub-Tenon's triamcinolone injections, avoiding the initiation of systemic
371 corticosteroids. Previous studies have described a good response to local corticosteroid
372 injections. For example, grade 4 panuveitis with bilateral serous retinal detachment in patients
373 treated with nivolumab has been successfully treated with late dexamethasone implant
374 injections (Ozurdex®) [32]. Recent studies have shown a similar successful control of
375 inflammation using local corticosteroid injections and drops in patients treated with CICs,
376 associated with targeted therapy [10,19].

377 Overall, a high sensitivity to corticosteroids has been observed in cases of anterior uveitis and
378 most cases of posterior uveitis and panuveitis reported associated with CIT [8,10,19].

379
380 The AEs of local corticosteroid injections are well known and include induced cataract and
381 glaucoma. For example, cataract occurrence has been reported at 6 months in 20% of uveitis
382 eyes treated by corticosteroid periocular injections [62]. However, cataract surgery is a well
383 codified procedure, especially in elderly patients. Moreover, the endophthalmitis rate has been
384 reduced using systematic intracameral cefuroxime injections at the end of the surgical
385 procedure [63]. That is why cataract may appear as an acceptable AE in elderly patients
386 undergoing surgery with a rapid visual improvement.

387 Regarding raised intraocular pressure, it reached ≥ 24 mmHg in one third of eyes 6 months
388 after periocular corticosteroid injections in patients treated for uveitis [62]. Thus, the
389 intraocular pressure should be monitored in the second month following local corticosteroid
390 injections.

391
392 **Systemic corticosteroids:** The efficacy of systemic corticosteroids has been reported in
393 uveitis secondary to CIT in various case reports, as in 2 of the current patients [11,41,
394 44,46,48,49]. Indeed, the classical management of macular edema associated with bilateral
395 uveitis may include systemic corticosteroids as a first-line therapy, associated with
396 immunosuppressive and immunomodulatory agents thereafter to spare the daily corticosteroid
397 dose needed to control uveitis, and a treatment algorithm has been previously suggested
398 [64,65]. However, it has been suggested that systemic corticosteroids could decrease the
399 overall response in patients treated with anti-PD-(L)1 antibodies for lung cancer and taking
400 more than 10 mg/kg of prednisone daily [66]. On the other hand, systemic corticosteroids
401 have been shown not to influence the antitumor response to anti-CTL-4 [67].

402
403 **CIT discontinuation:** In a recent study, ICIs were continued in 2 patients despite the
404 occurrence of ocular complications and their ocular irAEs were resolved at 1 and 6 months,
405 respectively [10]. In our series, CIT discontinuation for 6 months was associated with tumor
406 progression and a rapid death despite a late resumption in one patient.

407 Thus, CIT continuation while promptly managing uveitis with corticosteroids injections could
408 be discussed as a first-line therapy.

409
410 **Recurrence of uveitis:** A tendency to relapse was observed in 2 of our patients after CIT
411 restart, as previously reported in studies assessing various ICIs, in particular nivolumab,
412 pembrolizumab, and ipilimumab [8,19,32]. In case of posterior pole involvement,
413 corticosteroid injections could be repeated if well tolerated. Moreover, the long-term use of
414 dexamethasone drops has been suggested to effectively maintain a low level of inflammation
415 in the anterior chamber [19].

416
417 All previous recommendations suggest to have an immediate assessment performed by an
418 ophthalmologist in case of ophthalmological problem [61,68].

419 In the context of a first-line management, it could be justified to use first topical and peri-
420 ocular injections of corticosteroids while continuing CIT. Most of the current cases were good
421 responders to local corticosteroids, even patients with a severe vision loss, and patients with
422 inflammation of the macular area.

423 The use of corticosteroids as a first line is supported by: (1) the good response to
424 corticosteroids of the inflammation in the aqueous humor and the posterior pole, (2) the long
425 period of time to recurrence, (3) the advanced age of patients making acceptable the
426 occurrence of cataract as an AE, and (4) the possibility to avoid the use of systemic

427 corticosteroids that could negatively affect the expected response. Moreover, CIT
428 continuation did not seem to prevent a good visual prognosis. CIT discontinuation should be
429 discussed on a case-by-case basis, in case of poor response to the ophthalmological treatment.
430 Although the underlying metastatic cancer are heterogenous, and immunotherapy drugs are
431 numerous, the clinical ophthalmological uveitis related to CIT had shown a similar response
432 to the corticosteroid treatment, regardless of further CIT. So the role of the ophthalmologist
433 may be useful in (1) making quickly the diagnosis relating uveitis to CIT, ruling out
434 infectious, auto-immune and tumoral causes (2) explaining the often favorable uveitis
435 outcome using corticosteroid local injections (3) reassuring patients and oncologists, worried
436 about visual fonction (4) performing quickly ophthalmological treatment to keep a good
437 quality of life (5) following the patients to assess positive and negative treatment effects (6)
438 repeating the corticosteroid local injections if needed (7) rediscussing the treatment in case of
439 ophthalmological failure.

440
441 **This study has some limitations**, including the lack of power due to a small sample size.
442 Moreover, this was a real-life study in which various types of first-line ophthalmological
443 treatments were assessed.

444 The study strenght could be the ophthalmological follow-up that allowed assessing the long-
445 term results of the therapeutic options.

446
447 **In summary**, local corticosteriod injections for uveitis seem to be associated with favorable
448 visual outcomes while continuing CIT. They could be suggested as a first-line therapy for the
449 management of CIT-associated uveitis.

450 Ophthalmologists should explain the often favorable visual outcome of CIT-associated uveitis
451 when treated early with local corticosteroids, reassure patients and the care team, explain the
452 expected rapid visual improvement and the possible occurrence of AEs, including cataract and
453 glaucoma, and regularly check corticosteroid efficacy and AEs. In case of failure, they should
454 early adapt treatment after collegial discussion on a case-by-case basis.

455 456 **Conclusion**

457 The long-term ophthalmological and systemic assesements of patients with CIT-associated
458 uveitis described in this series together with literature data suggest to use local corticosteroid
459 injections as a first line while CIT is continued. The ophthalmological interventions should be
460 given early not to alter patients' quality of life. Indeed, preserving the quality of life could be
461 a new important challenge for the numerous patients treated with CIT, and represent a huge
462 progress in terms of survival outcomes. Uveitis treatment could maintain the quality of life
463 while not impairing the expected outcome of CIT on tumor mass increase. An adapted uveitis
464 management could improve the visual prognosis, without impairing the vital prognosis.

465
466
467 Disclosure: None of the authors has any financial/conflicting interests to disclose.

468 469 **References**

- 470 1. Accorinti M, Okada AA, Smith JR, Gilardi M. Epidemiology of macular edema in uveitis.
471 *Ocul. Immunol. Inflamm.* 27(2):169-180 (2019).
- 472 2. Lardenoye CW, van Kooij B, Rothova A. Impact of macular edema on visual acuity in
473 uveitis. *Ophthalmology* 113(8):1446–1449 (2006).
- 474 3. Velcheti V, Schalper K. Basic Overview of Current Immunotherapy Approaches in
475 Cancer. *Am. Soc. Clin. Oncol. Educ. Book* 35:298-308 (2016).

- 476 4. Pardoll DM. The blockade of immune checkpoints in cancer immunotherapy. *Nat. Rev.*
477 *Cancer* 12(4):252-264 (2012).
- 478 5. Motzer RJ, Penkov K, Haanen J, *et al.* Avelumab plus Axitinib versus Sunitinib for
479 Advanced Renal-Cell Carcinoma. *N. Engl. J. Med.* 380(12):1103-1115 (2019).
- 480 6. Boutros C, Tarhini A, Routier E, *et al.* Safety Profiles of anti-CTLA-4 and anti-PD-1
481 Antibodies Alone and in Combination. *Nat. Rev. Clin. Oncol.* 13(8), 473-486 (2016).
- 482 7. Michot JM, Bigenwald C, Champiat S, *et al.* Immune-related adverse events with immune
483 checkpoint blockade: a comprehensive review. *Eur. J. Cancer* 54:139–148 (2016). ** important
484 topic for treatments more and more used
- 485 8. Whist E, Symes RJ, Chang JH, *et al.* Uveitis caused by treatment for malignant melanoma:
486 a case series. *Retin. Cases Brief Rep.* 00:1-6 (2019). ** a case series well described
- 487 9. Guedj M, Quéant A, Funck-Brentano E, *et al.* Uveitis in patients with latestage cutaneous
488 melanoma treated with vemurafenib. *JAMA Ophthalmol.* 132(12):1421-1425 (2014).** good
489 ophthalmological description
490
- 491 10. Bitton K, Michot JM, Barreau E, *et al.* Prevalence and Clinical Patterns of Ocular
492 Complications Associated With Anti-PD-1/PD-L1 Anticancer Immunotherapy. *Am. J.*
493 *Ophthalmol.* 202:109-117 (2019).
- 494 11. Fujimura T, Kambayashi Y, Tanita K, *et al.* HLA-DRB1*04:05 in two cases of Vogt-
495 Koyanagi-Harada disease-like uveitis developing from an advanced melanoma patient treated
496 by sequential administration of nivolumab and dabrafenib/trametinib therapy. *J. Dermatol.*
497 45(6):735-737 (2018).
- 498 12. Gavric AU, Ocvirk J, Mekjavic PJ. Ocular changes in metastatic melanoma patients
499 treated with MEK inhibitor cobimetinib and BRAF inhibitor vemurafenib. *Radiol. Oncol.*
500 52(2):213–219 (2018).
501
- 502 13. Abdel-Rahman O, Oweira H, Petrausch U, *et al.* Immunerelated ocular toxicities in solid
503 tumor patients treated with immune checkpoint inhibitors: a systematic review. *Expert Rev.*
504 *Anticancer Ther.* 17(4):387–394 (2017).
- 505 14. Theillac C, Straub M, Breton AL, Thomas L, Dalle S. Bilateral uveitis and macular edema
506 induced by nivolumab: a case report. *BMC Ophthalmol.* 17(1):227 (2017).
- 507 15. Richardson DR, Ellis B, Mehmi I, Leys M. Bilateral uveitis associated with nivolumab
508 therapy for metastatic melanoma: a case report. *Int. J. Ophthalmol.* 10(7):1183-1186 (2017).
- 509 16. Conrady CD, Laroche M., Pecen P, Palestine A, Shakoor A, Singh A. Checkpoint
510 inhibitor-induced uveitis: a case series. *Graefes Arch. Clin. Exp. Ophthalmol.* 256(1):187-191
511 (2018). **good ophthalmological description
- 512 17. Robinson MR, Reed G, Csaky KG, Polis MA, Whitcup SM. Immune-recovery uveitis in
513 patients with cytomegalovirus retinitis taking highly active antiretroviral therapy. *Am. J.*
514 *Ophthalmol.* 130(1):49-56 (2000).
- 515 18. Tsui E, Gonzales JA. Retinal Vasculitis Associated with Ipilimumab. *Ocul. Immunol.*
516 *Inflamm.* 28(6):868-870-3 (2020).
- 517 19. Venkat AG, Arepalli S, Sharma S, *et al.* Local therapy for cancer therapy-associated
518 uveitis : a case series and review of the literature. *Br. J. Ophthalmol.* 104(5):703-711(2020). **
519 interesting discussion
- 520 20. Jabs DA, Nussenblatt RB, Rosenbaum JT. Standardization of Uveitis Nomenclature
521 (SUN) Working Group. Standardization of uveitis nomenclature for reporting clinical data.
522 Results of the First International Workshop. *Am. J. Ophthalmol.* 140(3):509-516 (2005).

- 523 21. Herborg CP Jr, Tugal-Tutkun I, Neri P, Pavésio C, Onal S, LeHoang P. Failure to integrate
524 quantitative measurement methods of ocular inflammation Hampers Clinical Practice and
525 Trials on New Therapies for Posterior Uveitis. *J. Ocul. Pharmacol. Ther.* 33(4):263-277
526 (2017).
- 527 22. Rodriguez EF, Lipson E, Suresh K, Cappelli LC, Monaco SE, Maleki Z. Immune
528 checkpoint blocker-related sarcoid-like granulomatous inflammation: a rare adverse event
529 detected in lymph node aspiration cytology of patients treated for advanced malignant
530 melanoma. *Hum. Pathol.* 91:69-76 (2019).
- 531 23. U.S. Department of Health and Human Services. National Cancer Institute. Common
532 terminology criteria for adverse events (CTCAE V. 5.0) (2019).
- 533 24. Eisenhauer EA, Therasse P, Bogaerts J, *et al.* New response evaluation criteria in solid
534 tumours: revised RECIST guideline (version 1.1). *Eur. J. Cancer* 45(2):228-47 (2009).
- 535 25. Dalvin L, Shields CL, Orloff M, Sato T, Shields JA. Checkpoint inhibitor immune
536 therapy: systemic indications and ophthalmic side effects. *Retina* 38:1063–1078 (2018).
- 537 26. Hammers HJ, Plimack ER, Infante JR, *et al.* Safety and Efficacy of Nivolumab in
538 Combination With Ipilimumab in Metastatic Renal Cell Carcinoma: The CheckMate 016
539 Study. *J. Clin. Oncol.* 35(34):3851-3858 (2017).** [prospective study](#)
- 540 27. Wolchok JD Kluger H, Callahan MK, *et al.* Nivolumab plus ipilimumab in advanced
541 melanoma. *N. Engl. J. Med.* 369(2):122–133 (2013).
- 542 28. Kwon ED, Drake CG, Scher HI, Fizazi K, *et al.* Ipilimumab versus placebo after
543 radiotherapy in patients with metastatic castration-resistant prostate cancer that had
544 progressed after docetaxel chemotherapy (CA184-043): a multicentre, randomised, double-
545 blind, phase 3 trial. *Lancet Oncol.* 15(7):700–712 (2014).
- 546 29. Abdel-Wahab N, Shah M, Suarez-Almazor ME, Nishikawa H. Adverse events associated
547 with immune checkpoint blockade in patients with cancer: a systematic review of case
548 reports. Nishikawa H, ed. *PLoS One.* 11 (2016).** [Review](#)
- 549 30. Fu C, Gombos DS, Lee J, *et al.* Ocular toxicities associated with targeted anticancer
550 agents: an analysis of clinical data with management suggestions. *Oncotarget* 8(35),58709-27
551 (2017). * [Review](#)
- 552 3. Sun MM, Levinson RD, Filipowicz A, *et al.* Uveitis in Patients Treated with CTLA-4 and
553 PD-1 Checkpoint Blockade Inhibition. *Ocul. Immunol.Inflamm.* 28(2):217-227 (2020).
- 554 32. Wei W, Wai-Ching L, Ling C. Cancer Immunology, Immunotherapy Recurrent grade 4
555 panuveitis with serous retinal detachment related to nivolumab treatment in a patient with
556 metastatic renal cell carcinoma. *Ophthalmology* 126(2):283-295 (2019).
- 557 33. Aaberg MT, Aaberg TMJ. Pembrolizumab administration associated with posterior
558 uveitis. *Retin. Cases Brief Rep.* 11(4):348–351 (2017).
- 559 34. Abu Samra K, Valdes-Navarro M, Lee S, Swan R, Foster CS, Anesi SD. A case of
560 bilateral uveitis and papillitis in a patient treated with pembrolizumab. *Eur. J.*
561 *Ophthalmol.*26(3):e46-8 (2016).
- 562 35. Arai T, Harada K, Usui Y, Irisawa R, Tsuboi R. Case of acute anterior uveitis and Vogt-
563 Koyanagi-Harada syndrome-like eruptions induced by nivolumab in a melanoma patient. *J.*
564 *Dermatol.* 44(8):975-976 (2017).
- 565 36. Basiliou A, Lloyd JC. Posterior subcapsular cataracts and hypotony secondary to severe
566 pembrolizumab induced uveitis: Case report. *Can. J. Ophthalmol.* 51(1):e4-6 (2016).
- 567 37. Kanno H, Ishida K, Yamada W, *et al.* Uveitis induced by programmed cell death protein 1
568 inhibitor therapy with nivolumab in metastatic melanoma patient. *J. Infect. Chemother.*
569 23(11):774-777 (2017).

- 570 38. Manusow JS, Khoja L, Pesin N, Joshua AM, Mandelcorn ED. Retinal vasculitis and
571 ocular vitreous metastasis following complete response to PD-1 inhibition in a patient with
572 metastatic cutaneous melanoma. *J. Immunother. Cancer* 2(1):41 (2014).
573
- 574 39. Roberts P, Fishman GA, Joshi K, Jampol LM. Chorioretinal Lesions in a Case of
575 Melanoma-Associated Retinopathy Treated With Pembrolizumab. *JAMA Ophthalmol.*
576 134(10):1184-8 (2016).
- 577 40. Taylor SC, Hrisomalos F, Linette GP, Rao PK. A case of recurrent bilateral uveitis
578 independently associated with dabrafenib and pembrolizumab therapy. *Am. J.*
579 *Ophthalmol. Case Rep.* 2:23-25 (2016).
- 580 41. Wong RK, Lee JK, Huang JJ. Bilateral drug (ipilimumab)-induced vitritis, choroiditis, and
581 serous retinal detachments suggestive of vogt-koyanagi-harada syndrome. *Retin. Cases Brief*
582 *Rep.* 6(4):423-426 (2012).
- 583 42. Crosson JN, Laird PW, Debiec M, Bergstrom CS, Lawson DH, Yeh S. Vogt-Koyanagi-
584 Harada-like syndrome after CTLA-4 inhibition with ipilimumab for metastatic melanoma. *J.*
585 *Immunother.* 38(2):80-84 (2015).
- 586 43. Fierz F, Meier F, Chaloupka K. Intraocular inflammation associated with new therapies
587 for cutaneous melanoma-case series and review. *Klin. Monatsbl. Augenheilkd* 233, 540-544
588 (2016).
- 589 44. Obata S, Saishin Y, Teramura K, Ohji M. Vogt-Koyanagi-Harada Disease-Like Uveitis
590 during Nivolumab (Anti-PD-1 Antibody) Treatment for Metastatic Cutaneous Malignant
591 Melanoma. *Case Rep. Ophthalmol.* 10(1):67-74 (2019).
- 592 45. De Velasco G, Bermas B, Choueiri TK. Autoimmune arthropathy and uveitis as
593 complications of programmed death 1 inhibitor treatment. *Arthritis Rheumatol.* 68(2), 556-
594 557 (2016).
- 595 46. Matsuo T, Yamasaki O. Vogt-Koyanagi-Harada disease-like posterior uveitis in the
596 course of nivolumab (anti-PD-1 antibody), interposed by vemurafenib (BRAF inhibitor), for
597 metastatic cutaneous malignant melanoma. *Clin. Case Rep.* 31(5):694-700 (2017).
- 598 47. Hanna KS. A Rare Case of Pembrolizumab-Induced Uveitis in a Patient with Metastatic
599 Melanoma. *Pharmacotherapy* 36(11) (2016).
- 600 48. Diem S, Keller F, Ruesch R, *et al.* Pembrolizumab-triggered Uveitis: An Additional
601 Surrogate Marker for Responders in Melanoma Immunotherapy? *J. Immunother.* 39(9):379-
602 382 (2016).
- 603 49. Bricout M, Petre A, Amini-Adle M, *et al.* Vogt-Koyanagi-Harada-like Syndrome
604 Complicating Pembrolizumab Treatment for Metastatic Melanoma. *J. Immunother.*
605 40(2):77-82 (2017).
- 606 50. Reid G, Lorigan P, Heimann H, Hovan M. Management of chronic hypotony following
607 bilateral uveitis in a patient treated with pembrolizumab for cutaneous metastatic melanoma.
608 *Ocul. Immunol. Inflamm.* 27(6):1012-1015 (2019).
- 609 51. Fonollosa A, Mesquida M, Adan A. Uveitic macular oedema after treatment with
610 vemurafenib. *Acta Ophthalmol.* 93(8):686-687 (2015).
- 611 52. Draganova D, Kerger J, Caspers L, Willermain F. Severe bilateral panuveitis during
612 melanoma treatment by Dabrafenib and Trametinib. *J. Ophthalmic. Inflamm. Infect.* 5:17
613 (2015).
- 614 53. Sarny S, Neumayer M, Kofler J, El-Shabrawi Y. Ocular toxicity due to Trametinib and
615 Dabrafenib. *BMC Ophthalmol.* 17(1):146 (2017).
- 616 54. Joshi L, Karydis A, Gemenetzi M, Shao EH, Taylor SR. Uveitis as a Result of MAP
617 Kinase Pathway Inhibition. *Case Rep. Ophthalmol.* 4(3):279-282 (2013).
- 618 55. Flaherty KT, Robert C, Hersey P, *et al.* Improved survival with MEK inhibition in BRAF-
619 mutated melanoma. *N. Engl. J. Med.* 367(2):107-114 (2012).

- 620 56. Mirgh SP, Ahmed R, Agrawal N, Bothra S, Mohan B, Garg A, Thekkudan SF, Khushoo
621 V, Bhurani D. Knowing the flip side of the coin: ibrutinib associated cystoid macular edema.
622 *Indian J. Hematol. Blood Transfus.* 36(1):208-210 (2020).
- 623 57. Bussone G, Kaswin G, de Menthon M, Delair E, Brézin AP, Guillevin L. Macular oedema
624 following rituximab infusion in two patients with Wegener's granulomatosis. *Clin. Exp.*
625 *Rheumatol.* 28(1 Suppl 57):90-2 (2010).
- 626 58. Gilca M, Merrill PT. Cystoid macular edema secondary to rituximab. *Retin. Cases Brief*
627 *Rep.* 13(2):111–114 (2019).
- 628 59. Juruc I, Juric Tomasevic B, Basic-Jukic N. Unique case of macular edema following
629 rituximab treatment for antibody-mediated rejection in kidney transplant recipient. *Ther.*
630 *Apher. Dial.* 25(3):361-362 (2021).
- 631 60. Fife BT, Bluestone JA. Control of peripheral T-cell tolerance and autoimmunity via the
632 CTLA-4 and PD-1 pathways. *Immunol. Rev.* 224:166–182 (2008).
- 633 61. Brahmer JR, Lacchetti C, Schneider BJ, *et al.* Management of immune-related adverse
634 events in patients treated with immune checkpoint inhibitor therapy: American Society of
635 clinical oncology clinical practice guideline. *J. Clin. Oncol.* 36(17):1714–1768 (2018).
- 636 62. Sen HN, Vitale S, Gangaputra SS *et al.* Periocular corticosteroid injections in uveitis
637 :effects and complications. *Ophthalmology* 121(11): 2275–2286 (2014).
- 638 63. Jabbarvand M, Hashemian H, Khodaparast M, Jouhari M, Tabatabaei A, Rezaei S:
639 Endophthalmitis Occurring after Cataract Surgery: Outcomes of More Than 480 000 Cataract
640 Surgeries, Epidemiologic Features, and Risk Factors. *Ophthalmology* 123(2):295-301 (2016).
- 641 64. Fardeau C, Champion E, Massamba N, LeHoang P. Uveitic macular edema. *Eye*
642 30(10):1277-1292 (2016).
- 643 65. Massa H, Papis SY, Adewoyin T, Vergados A, Patra S, Panos GD. Macular edema
644 associated with non-infectious uveitis: pathophysiology, etiology, prevalence, impact and
645 management challenges. *Clin. Ophthalmol.* 13:1761-1777 (2019).
- 646 66. Arbour KC, Mezquita L, Long N, *et al.* Impact of Baseline Steroids on Efficacy of
647 Programmed Cell Death-1 and Programmed Death-Ligand 1 Blockade in Patients With Non-
648 Small-Cell Lung Cancer. *J. Clin. Oncol.* 36(28):2872-2878 (2018).
- 649 67. Horvat TZ, Adel NG, Dang TO, *et al.* Immune-Related Adverse Events, Need for
650 Systemic Immunosuppression, and Effects on Survival and Time to Treatment Failure in
651 Patients With Melanoma Treated With Ipilimumab at Memorial Sloan Kettering Cancer
652 Center. *J. Clin. Oncol.* 33(28):3193-3198 (2015).
- 653 68. Puzanov I, Diab A, Abdallah K, *et al.* Managing toxicities associated with immune
654 checkpoint inhibitors: consensus recommendations from the Society for immunotherapy of
655 cancer (SITC) toxicity management Working group. *J. Immunother. Cancer* 5(1):95

656

657 **Figure captions**

658 Figure 1: Patient 1. 81-year old woman treated with rituximab for 4 months for chronic
659 lymphoid leucemia and referred for visual loss of her only functional eye (RE) due to macular
660 edema complicating intermediate uveitis. The left eye showed former corneal scars. 1A
661 OCT of the RE: cysts mainly located in the inner nuclear layer. 1B OCT of the RE: 1 month
662 after sub-Tenon's triamcinolone injection, the macular edema disappeared, the visual acuity
663 improved from 0.2 to 0 LogMAR.

664

665 Figure 2: Patient 2. An 88-year old man previously treated for Waldstrom disease,
666 experienced 3 years later a metastatic dermic-hypodermic sarcoma treated with ibrutinib and

667 cyclophosphamide. Eighteen months later, a bilateral visual loss was reported due to
668 panuveitis associated with bilateral macular edema and multifocal peripheral nodular
669 choroidal lesions. Both eyes showed similar lesions. RE : 2A wide field retinography showing
670 peripheral nodular lesions 2B FA 10 min after dye injection, showed a leakage in the central
671 macular area and papillitis 2C: ICGA 4 min after dye injection, showed numerous
672 hypofluorescent nodular lesions 2D they disappeared at 30 min, suggesting an inflammatory
673 activity although an effect of masking by the normal staining of the retinal pigmentary
674 epithelium at 10 minutes may interfere. 2E OCT showing macular cysts mainly located in the
675 inner nuclear layer. 2F the macular edema resolved after treatment with systemic
676 corticosteroids for 6 months in both eyes.

677
678

679 Figure 3: Patient 3. 61-year old man treated with nivolumab, dabrafenib, and trametinib for a
680 BRAF V600E mutation-positive melanoma. Intermediate uveitis complicated by macular
681 edema occurred 18 months after treatment discontinuation due to cryoglobulinemia and
682 pneumopathy. The peripheral isopters of the visual field remained normal and the etiological
683 work-up was negative. In particular, the aqueous humor testing showed a cytokine profile
684 consistent with an inflammatory or infectious process (IL-10 at 3 pg/mL and IL-6 at 105
685 pg/mL). The Herpes virus PCR was negative. Diagnostic vitrectomy showed the presence of
686 pigmented histiocytes, a few lymphocytes and the absence of malignant cells. The anti-
687 recoverin antibody assay was negative. Both eyes showed similar lesions. RE : 3A: wide-field
688 retinography. 3B: autofluorescence showing a shadowing of the inflammatory vitreous. 3C:
689 OCT showed macular edema cysts located mainly in the inner nuclear layer. 3D: A sub-
690 Tenon's triamcinolone injection was effective, the macular cysts disappeared and the visual
691 acuity improved from 0.2 to 0.1 LogMAR.

692

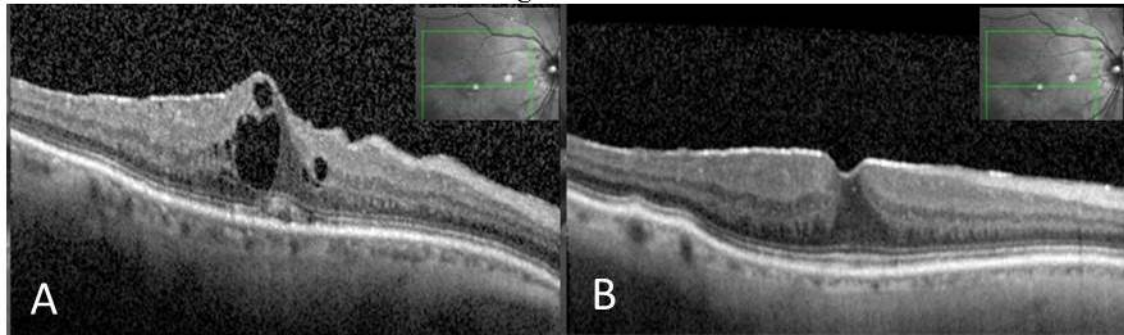
693 Figure 4: Patient 4. A 63-year old woman with KRAS-mutant bronchial adenocarcinoma (T3,
694 N3, M1b) was treated with nivolumab and bevacizumab for more than one year. She
695 experienced a bilateral visual loss due to bilateral panuveitis associated with macular edema
696 and focal choroidal nodular lesions. Both eyes showed similar lesions. RE 4A retinography
697 showing papillitis and peripheral nodular lesions. 4B: OCT horizontal scan showing cysts in
698 the inner nuclear layer. 4C: autofluorescence showing nodular hyperautofluorescent
699 lesions. 4D: FA showing a papillary and macular leakage and cyst filling. 4E: ICGA showing
700 numerous focal hypofluorescent nodular lesions, well visible at 10 minutes. 4F: they
701 disappeared during the late phase (30 min).

702

703 Figure 5: Patient 5. A 75-year old woman was treated with nivolumab for metastatic pleural
704 mesothelioma for 4 months when a bilateral visual loss occurred, due to acute anterior uveitis
705 with hypopyon, successfully treated with dexamethasone subconjunctival injections. In the
706 RE, hypopyon complicating acute anterior uveitis.

707

Figure 1

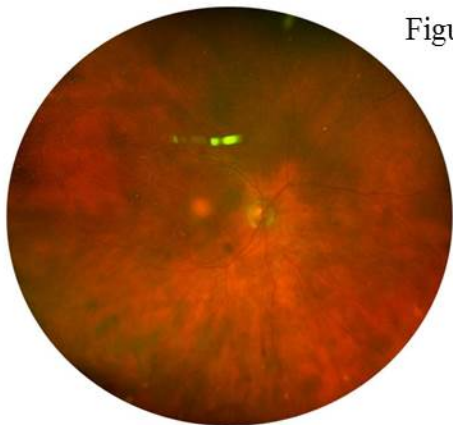


A

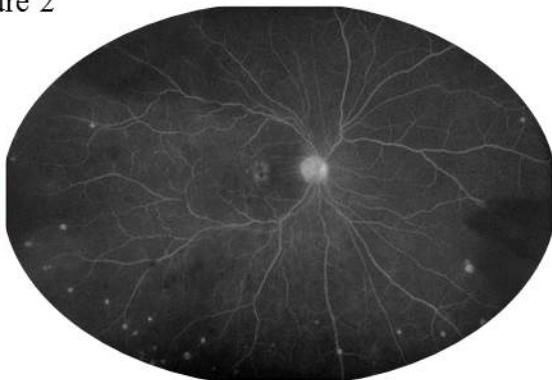
B

Figure 2

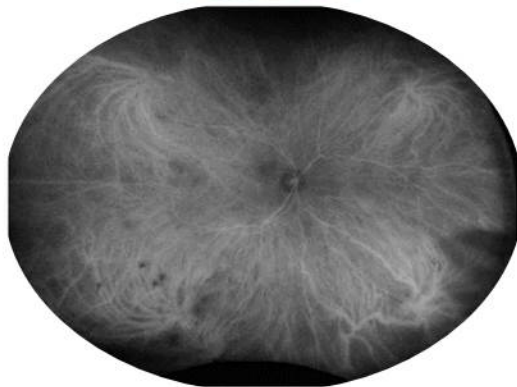
A



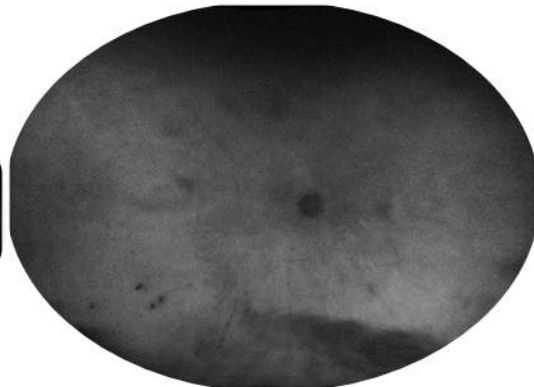
B



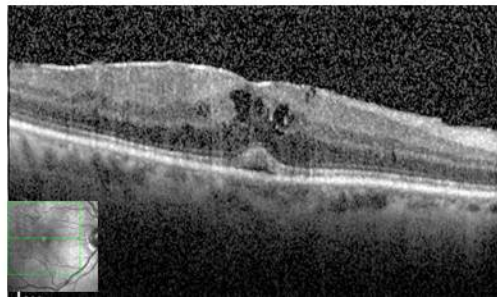
C



D



E



F

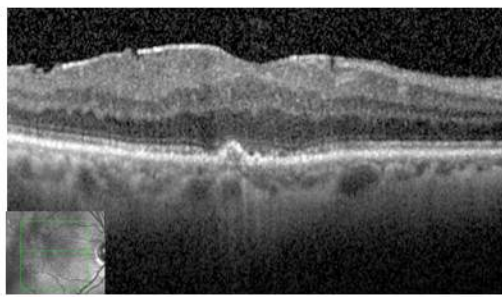


Figure 3

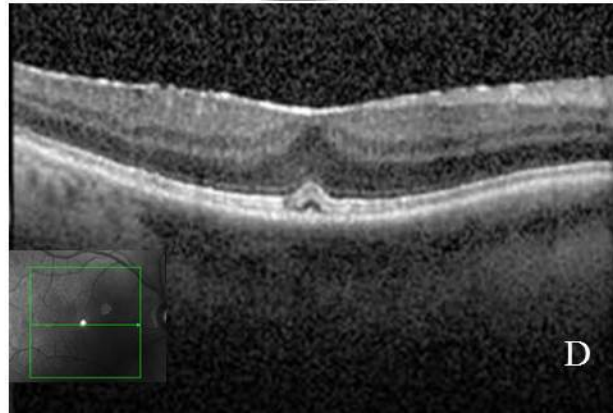
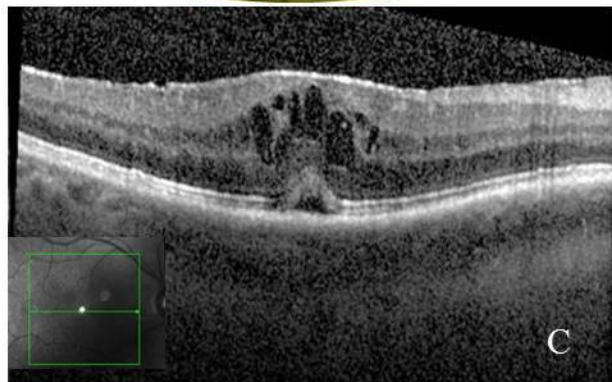
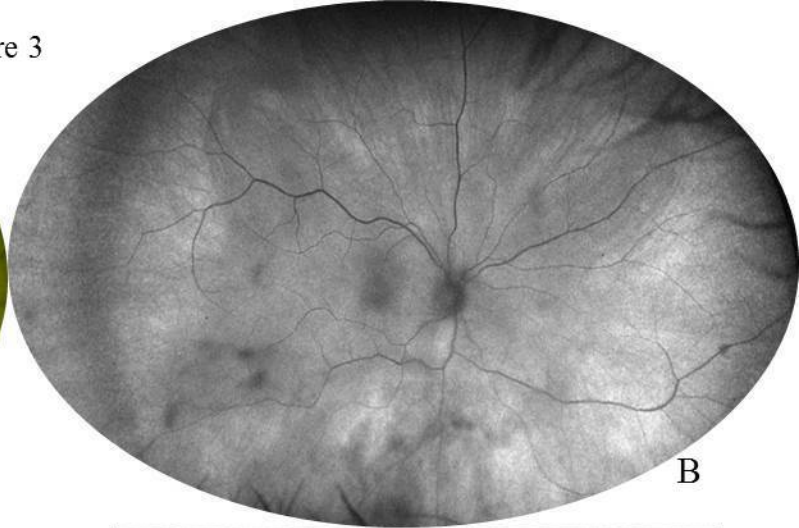
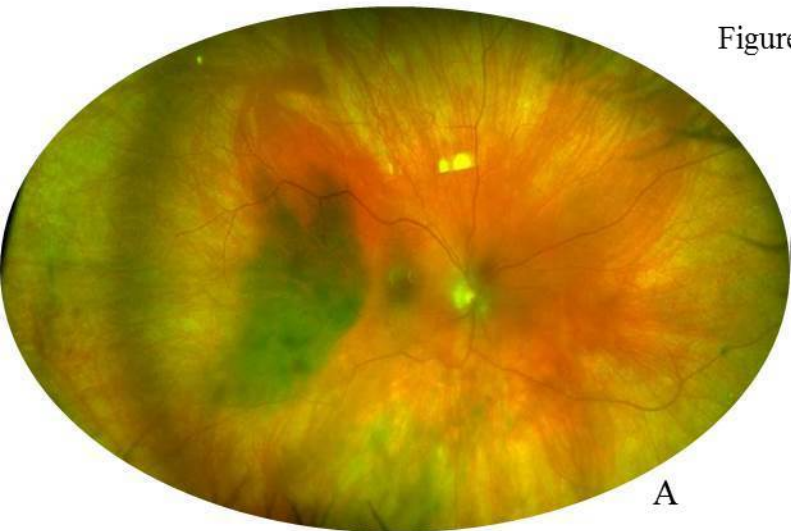


Figure 4

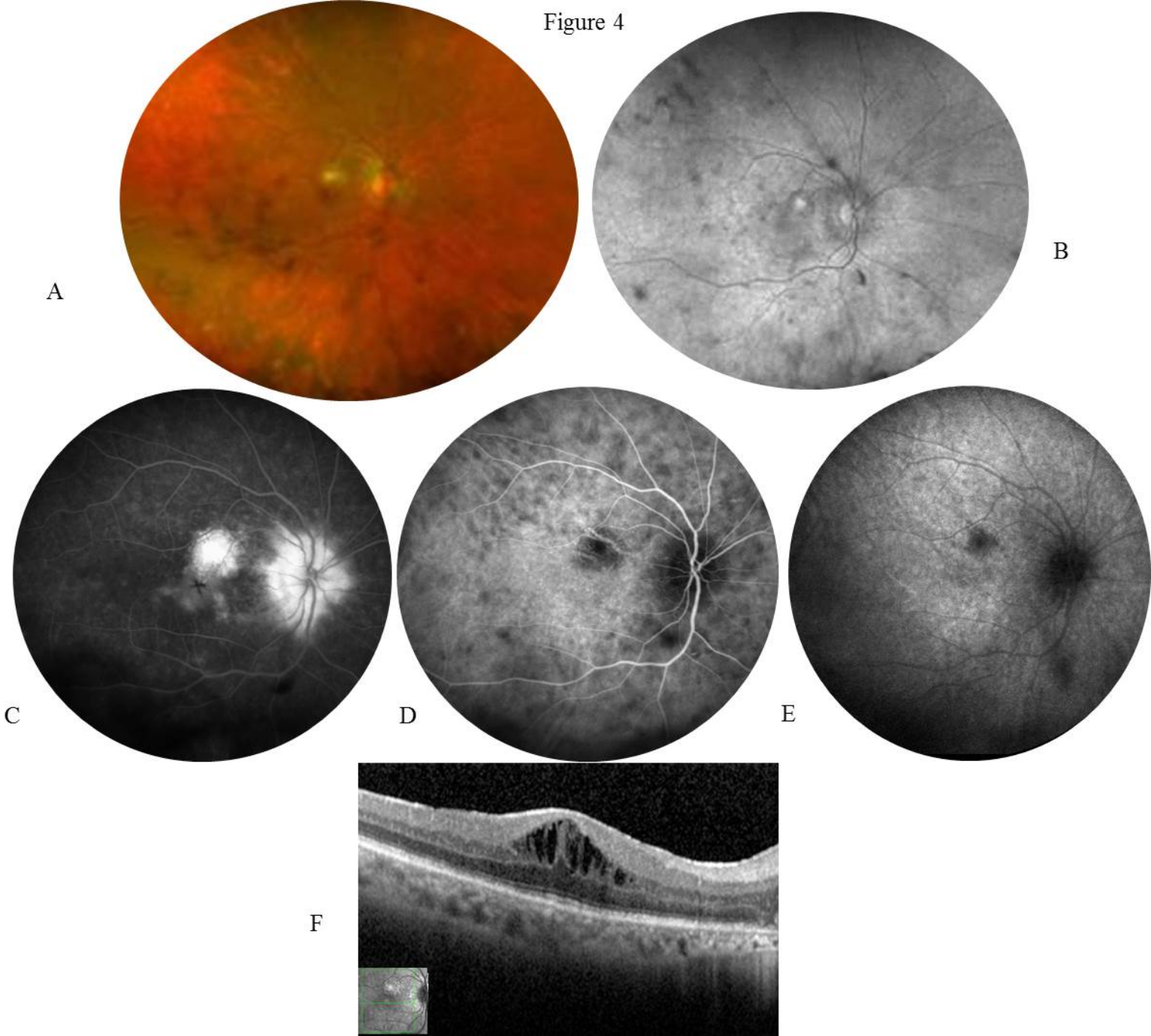
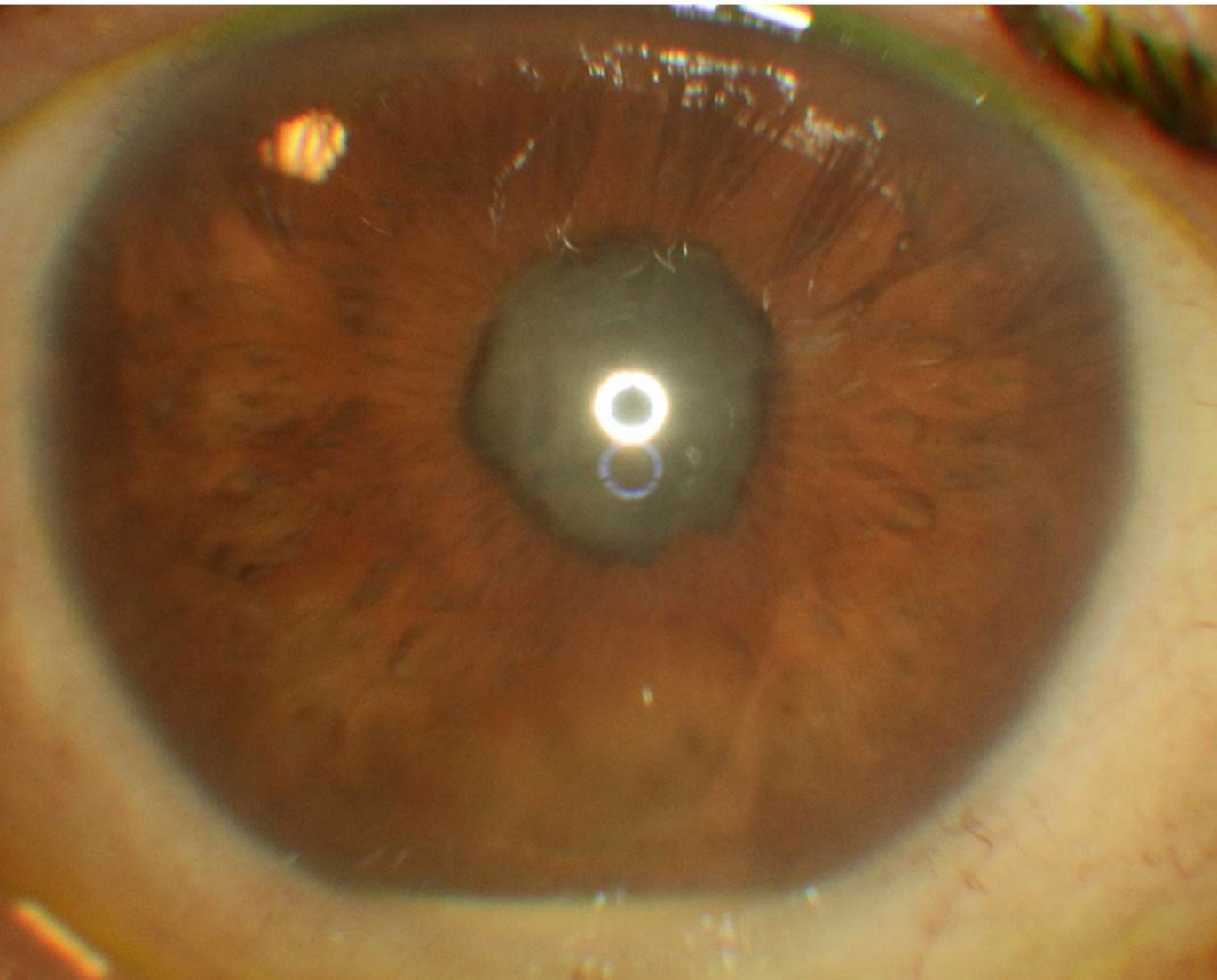


Figure 5



1st therapeutic intervention2nd intervention

Patient number, gender and age at uveitis onset	Neoplasia grading	CIT type	Time between ICI initiation and uveitis onset (months)	Oncological response	Baseline visual acuity in the RE/LE in decimals and LogMAR	Active uveitis features	CIT discontinuation	Ophthalmological treatments	Visual acuity at 3 months RE/LE LogMAR	Favorable ophthalmological outcome at 3 months	CIT resumption	Uveitis activity at 12 months	Final outcome at 12 months
1 F. 81	Chronic lymphoid leukemia	Rituximab (Anti-CD20)	4	CR	0.6/0.01 0.2/2	Bilateral intermediate Uveitis and ME	yes	Local corticosteroids Sub-Tenon's triamcinolone injection	0/2	Yes	Resumption after 3 years for leukemia recurrence	No	Alive CR
2 M. 72	Less differentiated sarcoma in Waldenström disease	Ibrutinib (Bruton tyrosine Kinase inhibitor)	18	PR	0.5/0.5 0.3 /0.3	Bilateral panuveitis choroiditis and ME	no	Systemic corticosteroids	0.2/0.1	Yes	ongoing	No	Alive PR
3 M. 61	Stage 4 BRAF V600E mutation-positive melanoma	Nivolumab (Anti-PD-1) + Dabrafenib (Anti-BRAF) + Trametinib (Anti-MEK)	24	PR	0.8/0.8 0.1/0.1	Bilateral intermediate uveitis and ME	no	Sub-Tenon's triamcinolone injection	0.2/0.2	yes	ongoing	Yes Good VA with triamcinolone injections	Alive CR
4 F. 65	T3 lung adenocarcinoma	Nivolumab (Anti-PD-1)	13	PR	0.5 /0.7 0.3/0.2	Bilateral panuveitis choroiditis and ME	no	Dexamethasone subconjunctival injections	0.1/0.1	yes	ongoing	Yes Bilateral ME	Death
5 F. 74	Pleural mesothelioma	Nivolumab (Anti-PD-1)	4	CR	0.01/0.01 2/2	Bilateral anterior uveitis Bilateral hypopion	yes	Local antibiotics and dexamethasone drops Dexamethasone subconjunctival injections	0/0	Yes	Resumption after 6 months for tumor mass increase	No	Death
6 F. 83	Melanoma	Nivolumab (Anti-PD-1)	5	CR	0.8/0.1 0.1/1	Bilateral posterior uveitis Papillary edema Contiguous ME	Yes for interstitial pneumonitis	Systemic corticosteroids	0/0.3	no	no	no	Alive PR
7 M. 58	BRAF V600E mutation-positive superficial spreading melanoma	Vemurafenib (Anti-BRAF)	5	CR	0.6/0.8 0.2/0.1	Bilateral anterior uveitis	Yes for peripheral neuropathy	Local corticosteroids Dexamethasone drops	0/0	yes	Yes with nivolumab	no	Alive CR
8 M.59	Stage 4 BRAF V600E mutation-positive melanoma	Nivolumab + Ipilimumab (Anti-CTLA-4)	1	PR	1.0/1.0 0/0	Bilateral anterior uveitis	Yes ipilimumab discontinuation	Local corticosteroids Dexamethasone drops	0/0	yes	Yes with nivolumab	No at 3 months	Alive PR at 3 months

CIT: Cancer ImmunoTherapy; PR: Partial response; CR: complete response; ME: macular edema; OPH:

Patient	Baseline uveitis features	Corneal precipitates	Post synechiae	Hypopyon	IOP >21 mmHg	Vitreous cells grading RE/LE	Flare Photon/ms RE/LE	Macular edema	Central foveal thickness Microns RE/LE	FA and ICGA findings	Uveitis grading CTCAE
1	Bilateral intermediate uveitis	fine	NO	NO	NO	+++	8.4/ND	YES	462/ND	Retinal peripheral venous leakage Papillitis	3
2	Bilateral panuveitis	Small granulomatous	NO	NO	NO	++	26.2/16.6	YES	441/483	Retinal edematous capillaritis Choroidal nodular lesions	3
3	Bilateral intermediate uveitis	fine	YES	NO	NO	++/+++	108.1/315.3	YES	500/399	Macular leakage	3
4	Bilateral panuveitis	Small granulomatous	YES	NO	NO	++/+++	44.9/50.4	YES	455/410	posterior pole leakage Choroidal nodular lesions	3
5	Bilateral anterior uveitis	fine	YES	YES	NO	++	152.3/64.7	NO	282/259	ND	4
6	Bilateral posterior uveitis	fine	NO	NO	NO	++	3.1/3.9	YES	295/360	Papillitis	4
7	Bilateral anterior uveitis	fine	YES	NO	NO	½+/½+	23.5/25.4	NO	275/269	ND	2
8	Bilateral anterior uveitis	fine	NO	NO	NO	½+/½+	nd	NO	248/250	Papillary edema	1

CTCAE: Common terminology criteria for adverse events; ND: not determined; IOP: intraocular pressure

Posterior segment involvement	Posterior segment involvement features	CIT drug(s)	Number of cases	Corticosteroid type (local/systemic/topical) (local=IVT , subTenonien injections)	CIT drug stop	Uveitis resolution after therapeutic intervention
Wong et al. 2012	VKH-like	ipilimumab	1	Systemic	Yes	Yes
Crosson et al. 2015	VKH-like	ipilimumab	1	None	No	ND
Fierz et al. 2016	Papillary edema Choroidal nodular lesions	ipilimumab	1	Systemic	Yes	No
Robinson et al. 2004	ME and papillitis	ipilimumab	1	Local	Yes	Yes
Tsui et al. 2019	ME and retinal vasculitis	ipilimumab	1	Local and systemic	Yes	Yes
Sun et al. 2019	CME Panuveitis	ipilimumab ipilimumab + nivolumab	2 2	Periocular triam/topical Local	ND ND	Yes / no Yes / no
Conrady et al. 2018	Subretinal fluid Multifocal choroiditis	ipilimumab + nivolumab	2	Systemic/local	Yes	Yes
Conrady et al. 2018	Central retinal artery occlusion Multifocal choroiditis	ipilimumab + nivolumab	1	Systemic	ND	No
Venkat et al. 2019	Papillitis and peripheral leakage on FA	ipilimumab + nivolumab	1	Local	Yes Ipilimumab discontinued	Yes
Obata et al. 2019	VKH-like Granulomatous corneal precipitates Choroidal granulomas	nivolumab	1	Local IVT	Yes	Yes
Arai et al. 2017	VKH-like	nivolumab	1	Local	Yes	
Theillac et al. 2017	ME, papillary edema Granulomatous corneal precipitates	nivolumab	1	Systemic	Yes	Yes
Wei et al. 2019	Serous retinal detachment	nivolumab	1	Local and systemic IVT DXM implant	Yes	Yes
De Velasco et al. 2016	CME	nivolumab	1	Intraocular steroids	No	Yes
Richardson et al. 2017	ME and papillitis	nivolumab	1	Local IVT	Yes	Yes
Conrady et al. 2018	Subretinal fluid Multifocal choroiditis	nivolumab	1	Local	ND	Yes
Sun et al. 2019	Panuveitis	nivolumab	1	Local	ND	Yes
Fujimura et al. 2018	VKH-like	Nivolumab + dabrafenib/trametinib	2	Systemic	Yes	Yes
Matsuo et al. 2017	VKH-like	Nivolumab + vemurafenib	1	Systemic	Yes	Yes
Bitton et al. 2019	Subretinal fluid	pembrolizumab	1	Systemic	Yes	Initial improvement and then relapse
Sun et al. 2019	Panuveitis	pembrolizumab	2	Local	ND	Yes / no
Conrady et al. 2018	ME	pembrolizumab	1	Local IVT	ND	Yes
Aaberg et al. 2017	Posterior uveitis, retinal vasculitis	pembrolizumab	1	Local IVT	No	Yes
Hanna et al. 2016	Panuveitis Multifocal choroiditis	pembrolizumab	1	Local and systemic	Yes	Yes
Diem et al. 2016	panuveitis	pembrolizumab	1	Systemic	Yes	Yes
Manusow et al. 2014	Retinal vasculitis	pembrolizumab	1	Local and systemic	No	Yes
Abu et al. 2016	Papillitis	pembrolizumab	1	Local	ND	Yes
Bricout et al. 2017	VKH-like	pembrolizumab	1	Systemic	ND	Yes
Reid et al. 2019	Optic disc edema hypotony	pembrolizumab	1	Systemic	Yes	No
Whist et al. 2019	hypotony	pembrolizumab	1	Systemic	Yes	No
Venkat et al. 2019	ME and peripheral leakage on FA	atezolizumab	1	Local IVT	Yes	Yes
Conrady et al. 2018	Retinal vasculitis	atezolizumab		Systemic		Yes
Guedj et al. 2014	ME	vemurafenib	1	Systemic	Yes	No
Fierz et al. 2016	Retinal vasculitis	vemurafenib	1	Local	Yes	Yes
Whist et al. 2019	ME	vemurafenib	1	Local	No	Yes
Fonollosa et al. 2015	ME	vemurafenib	1	ND	ND	Yes
Gavric et al. 2018	ME Subretinal fluid	Vemurafenib + cobimetinib	1 5	Topical	ND	ND
Whist et al. 2019	ME	Dabrafenib + trametinib	4	Systemic	ND	Yes
Draganova et al. 2015	Multiple subretinal detachment	Dabrafenib + trametinib	1		Yes	Yes
Sarny et al. 2017	ME and focal lesions	Dabrafenib + trametinib	1	ND	Yes	Yes
Joshi et al. 2013	Panuveitis	Dabrafenib + trametinib	1	ND	Yes	Yes
Flaherty et al. 2012	RPE detachment	trametinib	1		Yes	Yes
Mirgh et al. 2020	ME	ibrutinib	1	Local	ND	Yes
Bussone et al. 2010	ME	rituximab	2	Systemic	ND	ND
Gilca et al. 2019	ME	rituximab	1	Systemic followed by local	Yes	Yes
Juric et al. 2021	ME	Rituximab	1	Local	ND	yes

IVT: intravitreal therapy; Triam: triamcinolone; CME: cystoid macular edema; ME: macular edema; ND: not determined; RPE: retinal pigmentary epithelium; FA: fluorescein angiography; VKH: Vogt-Koyanagi-Harada disease; DXM dexamethasone