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1 **C1q+ macrophages: passengers or drivers of cancer progression**

2 Margot Revel<sup>1</sup>, Catherine Sautès-Fridman<sup>1,2</sup>, Wolf Herman Fridman<sup>1,2</sup>, Lubka T. Roumenina<sup>1\*</sup>

3 <sup>1</sup> Centre de Recherche des Cordeliers, INSERM, Sorbonne Université, Université de Paris, F-  
4 75006 Paris, France.

5 <sup>2</sup> Equipe labellisée Ligue contre le Cancer, 2021-2

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7 \*Correspondence to: Lubka T. Roumenina, Ph.D.; Cordeliers Research Center, INSERM UMRS  
8 1138; 15 rue de l'Ecole de Medecine; 75006 Paris, France. Phone: 33-1-44-27-90-96/ Fax: 33-  
9 1-40-51-04-20, E-mail: [lubka.roumenina@sorbonne-universite.fr](mailto:lubka.roumenina@sorbonne-universite.fr), @roumenina

10

11 **Abstract**

12 The omics era made possible the quest for efficient markers for cancer progression and  
13 revealed that macrophage populations are much more complex than just the M1/M2  
14 dichotomy. Complement C1q pops up as a marker of a tolerogenic and immunosuppressive  
15 macrophage populations in both healthy and tumor tissues, but the specific role of C1q+  
16 tumor associated macrophages is poorly understood. C1q is co-expressed in healthy and  
17 tumor macrophages with HLA-DR, APOE and MRC1 (CD206), suggesting a resident origin of  
18 this population. Tumor associated macrophages expressing C1q correlate with T cell  
19 exhaustion and poor prognosis in numerous cancers. Herein, we discuss the plural roles of  
20 C1q in these macrophages and how it could drive cancer progression.

21 Keywords: C1q, tumor-associated macrophages, T cells exhaustion, complement system

22

## 23 **Highlights**

- 24 • Complement component C1q is a marker of a particular sub-population of tissue-  
25 resident macrophages and tumor associated macrophages (TAM), often expressing CD206,  
26 HLA-DR, SEPP1, FOLR2, APOE but not SPP1, as revealed by scRNAseq in different normal  
27 tissues and tumor types.
- 28 • In cancer presence of C1q+ TAM often correlates with poor prognosis.
- 29 • Presence of C1q+ TAM correlates with T cell exhaustion in cancer and immune  
30 tolerance induction in healthy tissue.
- 31 • C1q is the recognition molecule of the classical complement pathway, binding to  
32 immune complexes, pentraxins or other activators in the tumor microenvironment.
- 33 • C1q directly controls macrophage phenotype by interacting with surface receptors.
- 34 • C1q directly controls T cell phenotype by internalization, binding to mitochondria and  
35 regulation of mitochondrial metabolism.
- 36 • C1q is likely not only a biomarker of a TAM subpopulations but a driver of cancer  
37 progression.

38

## 39 **Outstanding Questions Box**

- 40 • Which mechanism does C1q regulate macrophage and T cell phenotypes in normal  
41 tissue and in tumor microenvironment? Does C1q act intracellularly in the producing  
42 macrophages or in an autocrine manner, at the cell surface?
- 43 • What is the exact origin of C1q+ macrophages?
- 44 • Could C1q, produced by other cells such as fibroblasts, regulate macrophages and T  
45 cells phenotype?
- 46 • What is the interplay between the complement cascade-mediated functions of C1q  
47 and its functions outside the cascade?
- 48 • What parallels can we draw between the break in immune tolerance and  
49 autoimmunity, driven by the congenital C1q deficiency and the immunosuppressive tumor  
50 microenvironment of the tumors that is rich in C1q+ macrophages?
- 51 • Can tumor-promoting C1q+ macrophages turn into an ally during immunotherapy  
52 with anti-checkpoint inhibitors?

53

## 54 **C1q+ TAM in the light of scRNAseq**

55 Searching for effective biomarkers to predict cancer progression is a holy grail in oncology. Big  
56 data is useful in this respect, but it is still challenging to find new biomarkers or fish out the  
57 most robust ones. Single cell RNA-seq can characterize the transcriptional state of individual  
58 cell types and allows one to define rare populations, otherwise lost in the bulk RNA-seq or  
59 undetectable by flow cytometry or CyTOF due to lack of prior knowledge of their existence.  
60 For a long time, macrophages had been divided into M1 (proinflammatory) or M2 (anti-  
61 inflammatory) populations. Such dichotomy seems too simplistic and outdated. Macrophages  
62 appear as a continuous spectrum of phenotypes between these two extreme populations.  
63 Thanks to single cell RNAseq analysis, scientists started to explore the macrophage universe,  
64 and two molecules have emerged: TREM2, a marker more often expressed by tumor  
65 infiltrating macrophages and associated with pro-tumorigenic actions<sup>1</sup>, and FOLR2, a marker  
66 that suggests tissue residency<sup>2</sup>. Numerous studies are now detailing another marker that is  
67 expressed on macrophages, Complement component 1q (C1q). C1Q+ macrophage  
68 populations in both healthy and tumor tissues (Figure 1) have been observed for many years,  
69 but a deeper understanding of its function is still lacking.

70 Recent studies suggest that C1q could be used as a marker of poor prognosis for various  
71 cancers. Transcriptomic data<sup>3</sup> and protein staining on tumor sections showed that a high  
72 presence of C1q+ macrophages<sup>3,4,5</sup> is associated with higher post-surgical recurrence in clear  
73 cell renal cell carcinoma (ccRCC)<sup>5</sup> as well as in hepatocellular carcinoma<sup>6</sup>, and breast cancer<sup>7</sup>.  
74 In osteosarcoma, the expression of C1Q, mostly by macrophages, negatively correlates with  
75 patient survival<sup>8</sup>. In pancreatic ductal adenocarcinoma (PDAC), C1q expression in primary  
76 tumors and hepatic metastasis is higher compared to normal tissue, and the presence of  
77 C1Q+M2-like macrophages is associated with worse prognosis<sup>9</sup>.

78

79 Despite its potential role as a biomarker, recent studies also suggest that C1q could drive  
80 tumor progression. Here we discuss recent evidence that suggests C1q+ macrophages play a  
81 major role in tumor immunity and cancer progression.

82

## 83 **C1q+ TAM – a bunch of gene correlations**

84 In healthy tissue, the C1Q+ macrophage population is characterized by the expression of  
85 C1QA, C1QB, C1QC, HLA-DRB1 and MRC1 (gene coding for the CD206 molecule)<sup>10</sup>. Different  
86 cancer types exhibit an increased C1Q+ Tumor Associated Macrophages (TAM) population  
87 expressing CD206, HLA-DR, SEPP1 and FOLR2<sup>11,12,4</sup>. C1Q+ macrophages also express APOE in  
88 renal cancer<sup>3</sup>, breast cancer<sup>7</sup>, and liver metastasis from colorectal cancer<sup>13</sup>. Nevertheless,  
89 there are exceptions. In renal cancer the C1Q+ TAM do not express FOLR2 but TREM2<sup>3</sup>, which  
90 is known to be associated with immunosuppression and poor prognosis in several cancers.  
91 Similarly, C1Q+ TAMs express TREM2 in liver metastases in patients and in mice with PDAC<sup>14</sup>.  
92 The C1Q+ TAM population has a strict mutually exclusive relationship with SPP1+ TAM<sup>15,16</sup>  
93 (Figure 2). This dichotomy is nearly perfect, although some studies have shown that  
94 C1Q+/SPP1+ associates with FOLR2+ and/or TREM2+ TAM<sup>17,7</sup>. In the colon, C1Q+  
95 macrophages can be found in both healthy and tumor tissue, whereas SPP1+ macrophages  
96 are only found in tumor tissue<sup>12</sup>. These macrophages can be deciphered by the transcriptional  
97 factors they express: C1Q+ TAM express mostly MAF/MAFB, while SPP1+ TAM express  
98 FOS/JUN for and CEBPB/ZEB2<sup>12</sup>. Although these cells likely originate from the same precursor,  
99 they have a very different evolutionary path (Box 1). C1q+ TAM may be linked to sex, as  
100 females with non-small cell lung cancer present with a higher number of C1Q+ TAMs, whereas  
101 male have higher number of SPP1+ macrpahages<sup>16</sup>.  
102 The ensemble of these gene expression correlations (positive correlation with APOE, HLA-DR,  
103 MRC1, FOLR2 or TREM2, and negative correlation with SPP1) draws a portrait of the C1Q+  
104 macrophages as a distinct immunosuppressive population. Indeed, in melanoma and basal  
105 cell skin carcinoma, C1Q+TAM are enriched in non-responders to immune checkpoint  
106 therapy<sup>17</sup>, suggesting that they play a role in regulating anti-tumor immunity. As such, single  
107 cell RNA sequencing (scRNA-seq) is unveiling the interplay between C1Q+ macrophages and  
108 other immune cells that might be responsible for the poor overall survival in cancer patients  
109 and lack of response to immunotherapies.

110

### 111 **C1q correlates with T cells exhaustion**

112

113 The presence of C1Q+ macrophages correlates with exhausted T cells, forming a dysfunctional  
114 immune circuit in ccRCC<sup>3-5</sup>. In colorectal cancer, C1Q+ TAM interact with T cell subsets.  
115 Analysis of ligand-receptors pairs revealed a significant enrichment of CXCL10-CXCR3 axis in

116 C1Q+ TAM, suggesting that production of CXCL10 by C1Q+ TAM binds to its receptor CXCR3,  
117 which is mostly present at the T cell surface. This finding highlights the potential role of C1Q+  
118 TAM in the recruitment and activation of the Th1 response<sup>12</sup> (Figure 3 point 1). In lung cancer  
119 a similar increase in CXCL-10 was described in C1Q+ TAM, in association with an enrichment  
120 of the transcription factors IRF1, IRF7 and STAT1<sup>16</sup>. IRF1 correlates with STAT1, HLA-DR, PD-1  
121 and LAG-3 in metastases of colorectal cancer<sup>18</sup>. Moreover, in ccRCC tumors, C1q+ cell density  
122 correlated with expression of inhibitory receptors PD-1 and LAG3 at the CD8+ T cells surface<sup>5</sup>,  
123 and these C1Q+ macrophages express additional immune checkpoint ligands, such as PD-L1  
124 and PDL-2<sup>4,5</sup>. In cervical cancer, patients with C1Q+ TAM also express high levels of immune  
125 checkpoint inhibitors including CD40L, CTLA4, LAG3, PD-1, and TIGIT<sup>19</sup> (Figure 3 point 2).  
126 Moreover, in mouse models of cancer, C1Q+ macrophages specifically express EB13, a subunit  
127 of the IL-35 cytokine, which allows their cross talk with intratumoral T cells and leads to their  
128 dysfunction when combined with the p35 subunit of IL-12<sup>20</sup>. This gene was already described  
129 as a promotor of CD8+ T cell exhaustion, when it is expressed by Tregs<sup>21</sup>. Interestingly, Tregs  
130 are also found at a higher proportion within tumors with C1Q+ TAM as compared with normal  
131 samples<sup>3</sup>.

132 The maturation of dendritic cells can impact on the expression levels of C1q, ultimately  
133 affecting T cell function. Indeed, immature dendritic cells express large amounts of C1q, but  
134 during dendritic cell maturation, which is driven by CXCL4, C1q gene is hyper-methylated and  
135 its expression decreases<sup>22</sup>. A murine model of subcutaneous injection of various murine tumor  
136 cell lines (lung cancer (LLC), colorectal cancer (MC38) or melanoma cancer (B16-F10)), showed  
137 that reduced C1q methylation (i.e. high C1q) promotes CD8+ T cell dysfunction and tumor  
138 progression<sup>20</sup>. In lung cancer and idiopathic pulmonary fibrosis, the methylation status of C1q  
139 decreases as compared to healthy tissue, leading to an increase in tumor associated C1Q  
140 expression, which is associated with poor prognosis<sup>23</sup>.

141 In cervical cancer, tumors with a gene signature of C1Q+ TAM are more infiltrated by immune  
142 cells and express more immune-checkpoint markers than tumors with a gene signature of  
143 SPP1+ TAM<sup>19</sup>. However, it is unclear which cells express these immune checkpoints. Compared  
144 to SPP1+ TAM, C1Q+ TAM also express higher levels of HLA-DR<sup>12</sup>, which could help C1Q+ TAM  
145 interact with immune cells.

146 Taken together, these lines of evidence show a clear correlation between C1q+ macrophages  
147 and the activation status of T cells, but is C1q a driver or a passenger in this process?

148

149

### 150 **C1q in TAM... So what?**

151 Are these C1q+ macrophages accompanying other factors responsible for poor prognosis or  
152 driving recurrence? Is C1q only a “marker” of these macrophages or does it play a direct role  
153 in their pro-tumoral effect? As C1q is a major factor involved in the complement system,  
154 recent studies are unveiling the role of the complement cascade in regulating tumor  
155 progression. However, C1q is a versatile molecule and has functions extending beyond the  
156 borders of the complement cascade. Below we describe emerging mechanisms by which C1q  
157 regulates tumor progression.

158

159

### 160 *C1q in the complement system*

161 C1q is the initiating protein of the complement cascade<sup>24</sup>. Complement is a part of the innate  
162 immune system and its best-known function is to defend the host against invading pathogens.  
163 C1q can trigger the classical pathway of the cascade when it binds to immune complexes,  
164 pentraxins or one of its other over 100 different ligands. The plasma source of the majority of  
165 complement proteins is the liver, but C1q is an exception, being secreted by tissue-resident  
166 macrophages<sup>24</sup>. To function within the cascade, C1q needs to associate with two serine  
167 proteases – C1r and C1s, which trigger the proteolytic cascade that results in the generation  
168 of proinflammatory anaphylatoxins C3a and C5a, the membrane attack complex C5b-9 and  
169 the opsonization of target cells. How could this cascade function in the context of cancer and  
170 what is the role of C1q in this scenario?

171 Transcriptomic analysis across different tumor types has revealed that cells of the tumor  
172 microenvironment or tumor cells themselves express components of the classical and  
173 alternative complement pathways, including C1q (Figure 1)<sup>24</sup>. It appears that their co-  
174 regulated overexpression is context-dependent, and their prognostic value is either favorable  
175 or poor or of undetermined significance in particular types of cancers. ccRCC falls in the  
176 “aggressive complement” group, where overexpression of these genes correlates with worse  
177 survival. *In situ* staining, scRNAseq analysis and *ex vivo* experiments revealed that the ccRCC  
178 tumors have a complement-rich environment, where some tumor cells produce C1r, C1s, C4



179 and C3 but need macrophage-derived C1q to activate the classical pathway on intratumoral  
180 IgG-containing immune complexes<sup>5,9</sup> (Figure 3 point 3). While complement is activated on  
181 most tumor cells and promotes cell death, it does not result in cell killing and even promotes  
182 tumor progression in some cancer types. This finding can be explained by the low intratumoral  
183 expression of the components of the terminal pathway<sup>3</sup> and the limited formation of a  
184 membrane attack complex due to expression of specific inhibitors at the tumor cell  
185 surface<sup>5,9,25</sup>. Thus, chronic inflammation mediated by C5a favors an immunosuppressive  
186 microenvironment and facilitates T cells exhaustion<sup>3,4</sup> (Figure 3 point 4). C1Q+ macrophages  
187 also expressed ApoE - a protein able to bind C1q and to activate the complement system<sup>26</sup>.  
188 Therefore, C1Q+ positive macrophages can induce tumor progression by triggering the  
189 complement cascade.

190

#### 191 *C1q and neoangiogenesis*

192 C1q can interact directly with endothelial cells (EC) to promote neoangiogenesis, via still  
193 unknown cell surface receptors or heparan sulfate<sup>27,5,28</sup>. C1q deposits can be found at the  
194 surface of EC in the absence of other complement factors such as C3 or C4. C1q can induce  
195 adhesion, spreading and expression of adhesion molecules by directly binding EC<sup>29</sup>. In  
196 addition, in a lesioned area, EC start to express C1q, which induces EC permeability,  
197 proliferation, migration and endothelial tube formation *in vitro*<sup>28</sup>. *In vivo*, C1q<sup>-/-</sup> mice show a  
198 disordered vascular network in subcutaneously implanted tumors. Other *in vitro* studies also  
199 indicate that the interaction of C1q with melanoma and PDAC cells promotes proliferation,  
200 migration and invasion of the tumor cells<sup>9,27</sup> (Figure 3 point 5).

201

#### 202 *C1q, immune tolerance, and T cell exhaustion*

203 In physiology C1q regulates human macrophage polarization via interactions with LAIR1, as a  
204 switch towards inflammation resolution to avoid autoimmunity<sup>30-32</sup> (Figure 3 point 6). Indeed,  
205 complete C1q deficiency, although very rare, is the strongest genetic predisposing factor to  
206 systemic autoimmunity<sup>33</sup>. C1q opsonizes apoptotic cells, enhances their uptake by  
207 macrophages and immature dendritic cells, modulates cytokine release, and promotes  
208 immune tolerance<sup>34-37</sup> (Figure 3 point 7). In this context macrophages and immature dendritic  
209 cells produce and secrete C1q, which will act in an autocrine manner<sup>38</sup>. In addition, C1Q+ cells

210 have been found to associate with tolerance in the fetal-maternal interface during pregnancy,  
211 which is reminiscent of that seen in cancer. An HLA-DR<sup>high</sup> group of cells, characterized by high  
212 expression of C1Q, APOE, various genes of lipid metabolism, EB13, IDO 1 and 2 (inducers of  
213 cell tolerance) and the immune-checkpoint PD-L1, limits T-cell expansion driven by fetal  
214 alloantigen and establishes an immune-tolerance to fetal allotransplant<sup>39</sup>. These processes  
215 are not well studied in cancer, but it is tempting to speculate that uptake of C1q-opsonized  
216 dying cancer cells may be perceived in a tolerogenic manner by the TAM, and that this C1q  
217 will re-orient their phenotype to hamper the immune response against tumor neo-antigens.

218

219

### 220 *C1q and cell metabolism*

221 Another function of C1q that is unrelated to the complement cascade is to play as a rheostat  
222 of the mitochondrial metabolism of CD8 T cells<sup>40</sup> (Figure 3 point 8). Extracellular C1q is  
223 internalized by CD8+ T cells and is found at the surface of mitochondria. Intracellularly C1q is  
224 involved in the upregulation of mitochondria biogenesis genes, leading to the differentiation  
225 of CD8+ T cells into memory T cells and not effector cells. By C1q its presence at the  
226 mitochondrial surface C1q dampens CD8+ T cell responses to self-antigens. Congenital  
227 deficiency of C1q is rare in humans, but it results in enhanced CD8+ T cell responses, becoming  
228 the strongest genetic predisposing factor for autoimmunity. C1q is not produced by T cells,  
229 but it is internalized from the extracellular milieu. Therefore, it is tempting to speculate that  
230 C1q, secreted by the C1Q+ macrophages in ccRCC downregulate the capacity of adjacent  
231 intratumoral CD8 T cells to respond to stimulation, thereby contributing to their exhausted  
232 phenotype. The intracellular role of C1q within macrophages has not been studied, but again,  
233 we speculate that it may affect their metabolism, and functional orientation towards an  
234 immunosuppressive phenotype, thereby inducing T cell exhaustion. In the context of  
235 atherosclerosis, C1q can modulate the cytokine expression of macrophages while they digest  
236 lipid proteins, leading to an M2-like polarization<sup>41</sup>. The MafB transcription factor, which is  
237 present in C1Q+TAM in colorectal cancer<sup>12</sup>, was also described to promote M2 polarization in  
238 atherosclerosis<sup>42</sup>.

239 These results raise the question whether pro-tumoral M2-like macrophages start to express  
240 C1q or does C1q allow the polarization of these cells into pro-tumoral macrophages?

241

## 242 **C1q in CAF, another actor of this story**

243 Like macrophages, recent studies have begun to distinguish different sub-types of fibroblasts  
244 and especially cancer-associated fibroblasts (CAF). They notably, showed that in breast and  
245 pancreatic cancer a subpopulation of CAF MHC-II+ led to an immunosuppressive tumor micro-  
246 environment<sup>43,44</sup>. In breast cancer, mesenchymal stem cells (MSC) can produce exosomes  
247 containing TGF- $\beta$  and C1q<sup>45</sup>. Fibroblasts are similar to MSC and can be considered as old  
248 MSC<sup>46</sup>. The exosomes from MSC but not from tumor cells drive the polarization of Monocytic  
249 myeloid derived suppressor cells (M-MDSC) into M2-macrophages overexpressing CD206, PD-  
250 L1 and MHC-II<sup>45</sup> (Figure 3 point 9). In contrast, in lung cancer, MHC-II+ CAF can activate the  
251 TCR of CD4+ T cells and rescue them from apoptosis via the C1q/C1qbp axis. Indeed, these  
252 CAFs have the ability to produce C1q that will be released and will interact with its receptor  
253 C1qbp present at the surface of CD4+ T cells<sup>47</sup> (Figure 3 point 10). These recent studies suggest  
254 that C1q excreted by CAF influence the tumor immune microenvironment, by both directly  
255 acting on T cells and on macrophage polarization. More studies are necessary to determine  
256 the role of CAF upstream of macrophage actions.

257

258

## 259 **Concluding remarks and future perspectives**

260 Analyses of scRNAseq data and the growing literature indicate that C1q+ TAM are key players  
261 in the tumor microenvironment, but many questions remain unanswered (see Outstanding  
262 Questions). Although formal proof is needed, it seems that C1q+ TAM are drivers of cancer  
263 progression with a direct pro-tumoral effect in the absence of immunotherapy. Interestingly,  
264 ccRCC tumors with mature tertiary lymphoid structures have IgG deposition on tumor cells  
265 and respond better to anti-checkpoint inhibitors<sup>48</sup>. The tertiary lymphoid structure signature  
266 contained C1q genes and APOE, and the macrophages likely contributed to the elimination of  
267 tumor cells and the mounting of an anti-tumoral immune response. Further studies are  
268 needed to determine how C1q and complement contribute to this process. It is still poorly  
269 understood if C1q acts in an autocrine manner, on the cell surface of the macrophages or  
270 intracellularly or as deposits on tumor cells. It is also unclear which cells C1q impacts on and  
271 how it controls immune activation vs tolerance and exhaustion. Understanding the  
272 mechanisms of action of C1q in the modulation of macrophages phenotype in health and

273 disease and whether C1q can be harnessed as a therapeutic target in combination with anti-  
274 cancer checkpoint inhibitors are perspectives for the future.

275 Beyond C1q, other complement proteins also impact on the TAM phenotype. Factor H  
276 differentiates CD14+ human monocytes into immunosuppressive macrophages in the context  
277 of breast but not renal cancer<sup>49,50</sup>. C5aR1 is also overexpressed on TAM, which exhibit an M2-  
278 like functional profile. C3aR deficiency is associated with reduced accumulation and functional  
279 skewing of TAM, increased T cell activation and response to anti-PD-1 therapy<sup>51</sup> in mouse  
280 models of sarcoma. In models of squamous carcinogenesis, C5aR1 inhibition improves  
281 chemotherapy efficacy by reprogramming macrophages to recruit cytotoxic CD8+ T cells<sup>52</sup>.  
282 Similar effects were observed in a mouse model of renal cancer<sup>53</sup>. Future studies should  
283 address how other complement proteins and activation fragments, such as FH or C3a and C5a  
284 impact on C1q+ TAM vs TAM lacking C1q expression.

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298

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300

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449 **Box 1. C1q TAM = tissue resident macrophages?**



450 The origins of C1Q+ TAM is not well understood, but a Single Cell RNAseq study across species  
451 (mouse, rat, pig and human) found a conserved gene signature of kidney resident  
452 macrophages composed of CD81, CD74 (coding for HLA-DR-gamma), APOE and C1QC<sup>54</sup>. A  
453 similar study in lung found conserved genes between human and mice macrophages,  
454 including APOE, MRC1, C1QA, C1QB and C1QC<sup>55</sup>. These data agree with the ontogenic study  
455 of multiple tumor types, describing a high similarity between C1Q+ and FOLR2+ macrophages.  
456 FOLR2+ is associated with an embryonic origin of macrophages<sup>9,10,15</sup>. In healthy lung tissue,  
457 two types of macrophages can be distinguished, alveolar or interstitial. A sub-type of alveolar  
458 macrophages expresses both C1Q and APOE<sup>56</sup>. C1Q+ macrophages are larger in size and are  
459 identified as resident in patients with colorectal cancer liver metastasis<sup>13</sup> and in the peritoneal  
460 cavity of mice<sup>57</sup>. These large C1Q+ TAM confer poor prognosis<sup>13</sup>, and they have features of  
461 foamy cells, overexpress genes of cholesterol metabolism, scavenger receptors, as well as  
462 C1QA and C1QB.

463 C1Q+ TAM express, in a conserved way in multiple cancer types, HLA-DR, APOE and MRC1,  
464 suggesting an embryonic, tissue resident origin. However, a recent study showed that C1Q  
465 and APOE are commonly expressed by TAM in breast cancer<sup>7</sup> and resident TAM express  
466 FOLR2+, while infiltrating TAM express TREM2+, opening new questions on the origin of the  
467 C1q+ TAM in particular contexts.

468 A transcriptional trajectory study using the example of colorectal cancer<sup>12</sup> shows that C1Q+  
469 TAM and SPP1+ TAM populations could arise from a common precursor: CD14-expressing  
470 monocytes, which differentiate towards FCN1+ monocyte-like cells and different macrophage  
471 populations. One of them are the SPP1+, while the other overexpresses IL-1 $\beta$ + and gives rise  
472 to the sub-population of C1q+ TAM. In healthy tissue, C1q regulates the macrophages  
473 polarization during the uptake of apoptotic cells by inhibiting NLRP3 gene expression, that  
474 suppresses the IL-1  $\beta$  cleavage<sup>36</sup>. It is interesting to note that the FCN1 is defense collagen  
475 and a close relative to C1q, called ficolin 1, activator of the lectin complement pathway<sup>24</sup>.  
476 Another complement protein, C5aR1, regulate the IL-1  $\beta$  production un macrophages<sup>58</sup>.  
477 Whether a cross-talk exist between these complement proteins/pathways in macrophages,  
478 being they TRM or TAM, is still unknown.

479 Finally, future work will show whether there is a difference in the C1q+ macrophages from  
480 normal tissue (the ones that generate the plasma C1q pool,  
481 <https://www.proteinatlas.org/ENSG00000173372-C1QA/celltype>) and the ones in tumors.

482

483

## 484 **Figure legends**

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486

487 **Figure 1:** Heatmap representing the expression of C1QB gene in different immune or non-  
488 immune cells depending on the cancer type. Data is retrieved from the Tumor Immune Single-  
489 cell Hub (TISCH) scRNA-seq database<sup>59</sup>. In light yellow are indicated positions for which data  
490 is not available, in blue low gene expression of C1QB and in red high gene expression of C1QB.  
491 Similar data were obtained for C1QA and C1QC.

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493

494 **Figure 2:** Single Cell Analysis from the TISCH database. Gene expression of SPP1, C1QA, C1QB  
495 and C1QC of Colorectal Carcinoma tumor (accession number GSE146771). On the left, the cell  
496 characterization of the Single Cell RNAseq analysis. The upper panel is a large view of the gene  
497 expression. The lower panel is a zoom in on the Monocytes/Macrophages population.

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500 **Figure 3:** Proposed mechanisms by which C1q+ TAM drive cancer progression. C1q is produced  
501 by sub-populations of TAM and TRM. 1) The C1Q+ TAM secrete the chemokine CXCL-10 that  
502 binds to its receptor CXCR3 at the surface of CD8+ and CD4+ T cells, especially Th1 T cells. This  
503 binding will activate and recruit T cells inside the tumor. 2) By its expression of immune  
504 checkpoint inhibitors, C1Q+ TAM interact with T cell immune checkpoints, favoring T cell  
505 exhaustion. 3) C1q impacts on tumor progression by activating the complement cascade in  
506 the extracellular space, which will generate the anaphylatoxins C3a and C5a, and 4) promote  
507 chronic inflammation. 5) The C1q molecule can interact directly with endothelial cells to  
508 promote neo-angiogenesis needed for the tumor growth. 6) In healthy tissue, C1q is produced  
509 by TRM and immature dendritic cells (imDC) to maintain homeostasis and opsonize apoptotic  
510 cells. 7) The C1q molecule can also act in an autocrine way on macrophages, by interacting  
511 with LAIR1 to regulate their polarization. 8) CD8+ T cells can internalize C1q, which will interact  
512 with mitochondria to control the CD8 metabolism. 9) The secretion of C1q-containing  
513 exosomes by fibroblasts leads to M2 polarization of macrophages. 10) MHC-II+ cancer

514 associated fibroblasts also produce C1q, which binds to its receptor C1qbp at the surface of  
515 CD4+ T cells to activate and to rescue them from apoptosis. Figure generated with  
516 biorender.com





