Neutrophils fan cancer’s flames

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A new study published in this issue of The EMBO Journal looks into the cross-talk between inflammation and cancer (Antonio et al., 2015). By using a zebrafish melanoma model, the authors reveal that neutrophils recruited at the wound site directly interact with cells undergoing oncogenic transformation and provide them with a paracrine proliferative support. Importantly, the authors demonstrate the clinical relevance of this association, showing that neutrophil infiltration has an independent prognostic value for human melanoma. This study reinforces the notion that inflammation flames carcinogenesis, which might have important implications for the improvement of antitumour therapies.

Genetic mutations are the initial spark of tumour initiation and the driving force in tumour progression; however, dramatic changes in the microenvironment accompany and fuel cancer cell growth. Indeed, a variety of infiltrating cell types impact the whole tumour activity and contribute to generate cancer cell heterogeneity which is required for tumour progression (Kreso & Dick, 2014). Inflammatory cells are one of the main components within the tumour microenvironment having a recognized pivotal role in tumour promotion, progression and metastatic spreading (Grivennikov & Karin, 2010). Hence, cancer-associated inflammation represents an important target for the development of novel anticancer therapies.

Neutrophils are innate immune cells and essential during tissue damage and wound healing processes. They are quickly mobilized to the damaged tissue and are required to protect against invading pathogens, build and modulate immune responses. In recent years, growing evidence shows neutrophil involvement in various physiologic processes and their functional relevance during tumourigenesis has become an active field of investigation. In cancer patients, high tumour-associated neutrophil density correlates with unfavourable outcome; therefore, mechanistic studies on the role of neutrophils in tumorigenesis are critical (Shen et al., 2014). Interestingly, neutrophils are reported to display both antitumour and pro-tumour responses according to different signals within the tumour microenvironment (Fridlender et al., 2009). Indeed, three phenotypically distinct sub-pools of neutrophils with conflicting functions have been identified in the circulation of tumour-bearing mice and cancer patients, indicating high neutrophil plasticity (Sagiv et al., 2015). The main antitumour activity of neutrophils is linked to their cytotoxicity, an example of which was recently reported in melanoma where a sub-pool of tumour-associated neutrophils promotes cancer cell killing via nitric oxide release (Finisguerra et al., 2015). Conversely, neutrophils have been extensively reported to promote tumour growth by influencing the tumour microenvironment using mechanism such as the promotion of angiogenesis (Nozawa et al., 2006; Shojaei et al., 2008) or the creation of a “safe” milieu where more immature neutrophils alongside with macrophages suppress antitumour immune responses (Katoh et al., 2013). However, evidence of neutrophil plasticity predicts that many novel neutrophil-mediated functions are yet to be discovered.

In this issue of The EMBO Journal, Antonio and colleagues made an important contribution to the debated role of neutrophils in tumorigenesis by unravelling their novel direct pro-tumorigenic activity during wound healing in a zebrafish model of melanoma (Antonio et al., 2015). Here, the expression of a RasG12V oncogene in skin leads to the development of invasive melanoma. Using this model, which enables in vivo imaging to study the dynamics of cell growth, the laboratory has previously shown that myeloid cells can provide direct trophic support to neoplastic cells during tumour initiation (Feng et al., 2012). Antonio and colleagues now report that, in the context of wound healing, neutrophils, but not macrophages, boost pre-neoplastic cell’s proliferative activity via a direct paracrine mechanism (Fig 1).

Post-surgery wound healing-associated inflammation might influence the tumorigenic ability of cancer cells remaining in the tissue; therefore, the investigation of tumour-promoting effects of wound healing is especially relevant in the context of tumour relapses after surgical tumour removal. By taking advantage of in vivo imaging of zebrafish larvae, Antonio et al. (2015) mimicked the consequences of cancer surgery through generation of a wound and directly investigated the consequences of the recruitment of inflammatory cells on the neoplastic cells in the wound proximity. Results showed that hydrogen peroxide released by the damaged tissue attracted a great number of neutrophils, but, in contrast to the physiologic response where neutrophils remained in the proximity of the wound, in the presence of the near pre-neoplastic cells, neutrophils were diverted from the wound site and engaged in direct interaction with growing cancer cells. Importantly, results indicated that this interaction provides pre-neoplastic cells with a proliferative advantage leading to an increase in clones size. Moreover, the authors provide evidence that the direct neutrophil-mediated pro-tumorigenic activity described in zebrafish might be relevant for human melanoma. They report that neutrophils are predominantly infiltrating human melanomas with ulcerative lesions and their
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presence correlates with increased proliferation. Significantly, Antonio et al (2015) show that neutrophil infiltration in melanoma is functionally important for disease progression since it refines the known prognostic value of ulceration and represents an independent prognostic indicator.

Collectively, the work by Antonio and colleagues corroborates the contribution of neutrophils to the tumorigenic process by defining a pro-tumorigenic role directly targeting cancer cells. The authors found immune-cell-derived PGE2 to be part of the paracrine mediators of wound-induced pre-neoplastic proliferation in zebrafish (Fig 1), but results show that other signals are also involved. What would be fascinating to define are the precise mechanisms of cancer cell–neutrophil interaction. On the ground of Antonio et al’s findings, future studies need to concentrate on the dissection of these neutrophil-mediated pro-tumour activities, which will provide novel targets to improve the efficacy of antitumour therapies.

References