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Diethylcarbamazine and Non-Diethylcarbamazine Related Bancroftian Granuloma: An Immunohistochemical Study of Eosinophil Toxic Proteins

Jose Figueredo-Silva¹, Carmelita Cavalcanti², Luciano Tavares Montenegro³, Joaquim Norões⁴, ⁵, Gerusa Dreyer⁵, ⁶, ⁷

¹Núcleo de Ensino e Pesquisa em Patologia, Faculdade de Ciências Médicas, Universidade Estadual do Piauí, Piauí, Brazil; ²Laboratório de Imunopatologia Keizo Asami (LIKA), Universidade Federal de Pernambuco, Pernambuco, Brazil; ³Departamento de Patologia, Centro de Ciências da Saúde, Universidade Federal de Pernambuco, Pernambuco, Brazil; ⁴Serviço de Urologia, Departamento de Cirurgia, Centro de Ciências da Saúde, Universidade Federal de Pernambuco, Pernambuco, Brazil; ⁵Núcleo de Ensino Pesquisa e Assistência em Filariose (NEPAF), Hospital das Clínicas, Universidade Federal de Pernambuco, Pernambuco, Brazil; ⁶Centro de Pesquisas Aggeu Magalhães (CPqAM), Fundação Oswaldo Cruz (FIOCRUZ), Pernambuco, Brazil; ⁷NGO Amaury Coutinho Rua Conselheiro Portela, 665, Sala 120, Espinheiro, Recife, 50670-900, Pernambuco, Brazil

ABSTRACT

It has been suggested, mostly using in vitro experiments, that defenses against parasites involve mainly activated eosinophils and their toxic proteins, such as major basic protein (MBP), eosinophil cationic protein (ECP) and eosinophil peroxidase (EPO). Eosinophil degranulation has been described around degenerating onchocercal microfilariae in patients treated with diethylcarbamazine (DEC). In bancroftian filariasis, traditional histopathologic studies have shown remarkable numbers of eosinophils in granulomatous lesions associated with both DEC-induced and spontaneous death of adult Wuchereria bancrofti parasites. No immunohistochemical study targeting eosinophil degranulation has been previously performed in these granulomas, which are found mainly within intrascrotal lymphatic vessels. This investigation was undertaken in 22 (12 DEC-treated and 10 untreated) male patients in order to determine the immunohistochemical expressions of MBP, EPO and ECP in bancroftian granulomas, using the indirect method. Stained intact eosinophils, as well as granular, extra-cellular material positive for all three proteins, were found in all granulomas. The immunohistochemical patterns were similar in both DEC-treated and untreated cases, irrespective of microfilaremia, blood eosinophilia, and granuloma age. Positive intact cells were observed mostly at the periphery of the granulomas, whereas granular material predominated in central areas around dead or degenerating parasites. These results indicate that eosinophils accumulate in the granulomas and degranulate preferentially in close proximity to degenerating or dead adult parasites. In bancroftian granulomas, influx and degranulation of eosinophils are considered a consequence of parasite death, rather than its cause. (Int J Biomed Sci 2010; 6(2):111-119)

Keywords: bancroftian filariasis; eosinophils; eosinophil cationic protein; eosinophil peroxidase; filarial granuloma; immunohistochemistry; lymphatic filariasis; major basic protein; Wuchereria bancrofti

INTRODUCTION

The association between parasitic infections and eosinophilia in the blood and tissues of the host has been known for more than a century and is well-documented, being
Eosinophils in Bancroftian Granuloma

helminths the most common cause of blood eosinophilia in global terms (1). The exact role of eosinophils in host protection is still debated (2), but they seem primarily to be potent effector cells involved in the defense against infective larval stages of parasitic helminths. In contrast, living adult worms appear to be resistant to eosinophil attack (3). Since the in vivo mechanisms of action of human eosinophils have been difficult to establish, information has being gathered mostly from unnatural hosts as experimental models or in vitro studies. For instance, in vitro investigations have shown that eosinophils operate via ADCC (antibody-dependent cytotoxicity) directed against schistosomula (4) and microfilariae of Onchocerca volvulus (5).

Although ADCC, appears well suited for killing parasites too large to be phagocytized, it has been difficult to prove that it actually occurs in vivo (6). In fact, data supporting eosinophil function in vivo are mainly indirect since the evidence refers to interferences in interleukin-dependent responses (7). Several studies have also shown that eosinophil-derived cationic proteins, such as the major basic protein (MBP), the eosinophil cationic protein (ECP), and the eosinophil peroxidase (EPO) are particularly involved in cytotoxic activity (8-11). Nevertheless, these highly toxic eosinophil proteins are responsible for a considerable amount of the host inflammatory pathology that accompanies helminth infections (12).

Previous histological studies have shown close associations between tissue eosinophils and adult Wuchereria bancrofti parasites, damaged or dead, both during the course of diethylcarbamazine (DEC) treatment and naturally (13-19). Segments of coiled parasites in different stages of disruption are usually encircled by a granulomatous inflammatory reaction with remarkable numbers of eosinophils, as described in detail elsewhere (20). In such scenarios, eosinophil degranulation is expected to occur, but no studies have specifically investigated this. In men, spontaneous or drug-induced adult W. bancrofti parasite death is followed by the development of a palpable nodule, most commonly localized in intrascrotal lymphatic vessels (21, 22). The present study was undertaken to determine the immunohistochemical expression of MBP, EPO and ECP in granulomatous lesions in scrotal nodules removed from DEC-treated and non-treated patients.

MATERIAL AND METHODS

Tissue specimens
This retrospective study was conducted at the Center for Teaching, Research and Tertiary Referral for Bancroftian Filariasis (NEPAF) and the Laboratory of Immunopathology Keizo Asami-LIKA, both at Federal University of Pernambuco in Recife, Brazil. Tissue specimens for this study were randomly selected from the filariasis histopathology tissue bank. The material consisted of excised nodules that had been detected by physical examination of the intrascrotal contents; they were obtained during previous trials described elsewhere (17, 18, 23-26) and from patients participating in a diagnostic protocol used at NEPAF (J. Norões, unpublished). Before nodule excision, all patients signed an informed consent form. The study was conducted in accordance with the principles of the Declaration of Helsinki and the guidelines on Good Clinical Practice. Granuloma age, which could be determined only for patients treated with DEC, was defined as the interval between DEC intake and the day of excision biopsy. The DEC oral dose was 6mg/kg/day, given as a single dose or as a 12-day course.

Microfilaria counts
The microfilaria (Mf) density was obtained from venous blood collected between 23:00 and 01:00 o’clock and filtered through a 3μm polycarbonate membrane, stained, and examined by microscopy. Microfilaremia was defined as having no Mf detected in 11 mL of blood (if no Mf was detected in one mL, an additional 10 mL was obtained and examined within 7-15 days) (24). In treated patients the microfilaria density was investigated before DEC intake. In the untreated patients the first blood sample was collected one to three days after palpation of the nodule by the examining physician.

Blood absolute eosinophil counts
The peripheral eosinophil concentration was measured by Neubauer counting chamber. All patients receiving DEC blood eosinophilia was evaluated before and between 7 and 14 days after the first dose of DEC intake as described elsewhere (27). In patients with spontaneous nodules (i.e., those who had not received DEC), the eosinophil counts were measured when the nodule was first palpated by the examining physician and 7 to 14 days later.

Histopathology
The surgical specimens, fixed in buffered 10% formalin, were elastic, white-grayish, and either chord-like (varying from 2.5 × 0.5 cm to 4.5 × 0.4 cm) or ovoid (0.5 to 2.5 cm in its largest diameter). Several transverse slices from each specimen were routinely processed, and 5 μm-thick sections were stained with hematoxylin-eosin and
Masson’s trichrome, as previously described (19). Images were obtained with an Olympus CX31 microscope and an Olympus digital camera C-7070 (Tokyo, Japan).

**Immunohistochemistry for major basic protein (MBP), eosinophilic peroxidase (EPO) and eosinophilic cationic protein (ECP)**

Immunohistochemical staining was conducted using the standard indirect method. A set of six de-paraffinized 5 μm-thick sections of the same material used for the histopathological study was subjected to microwave for 20 minutes at maximum power as an antigen retrieval method. The endogenous peroxidases were blocked with a methanol solution containing 0.03% hydrogen peroxide and 1% sodium azide for 20 minutes. Tissue sections were then incubated overnight at 4°C with either normal swine serum (Dako Corporation, Carpinteria, CA) as a negative control, or with primary high-affinity chromatography-purified rabbit IgG antibodies to MBP, EPO, and ECP. Goat high-affinity anti-Fc-rabbit IgG conjugated to peroxidase (Jackson Immunoresearch Laboratories, Inc., West Grove, PA) was applied to the tissues for 60 minutes at 37°C. The slides were developed using diaminobenzidine and counterstained with 10% Harris hematoxylin. After each treatment stage and incubation, the slides were washed in phosphate buffered saline. Brown staining was considered an expression of immunoreactivity to MBP, EPO, and ECP. Interpretation of immunohistochemical results included both the location of the positive reaction and the staining pattern: intact, well-defined cells versus extra-cellular granular staining. The presence of extra-cellular positive granular material was interpreted as evidence of eosinophil degranulation. It was beyond to the scope of the study to assess quantitatively the immunohistochemical reaction for each protein studied.

**Statistics**

Descriptive statistics were used in the analysis of the data (Excel 2002). Pre and post eosinophil counts were log-transformed. Means of log-transformed counts were compared using the Student t test for paired samples.

**RESULTS**

**Patients**

The clinical, parasitological and antifilarial treatment schedules of the patients are shown in Table 1. Peripheral blood eosinophilia in relation to Mf counts and antifilarial treatment are shown in Table 2.

**Histopathology**

The tissue responses to the adult worm have been described in detail elsewhere (20). Briefly, the specimens consisted of lymphatic vessel segments with variable degrees of dilation. The vessels contained adult *W. bancrofti* in different stages of degeneration; progressing and healing lesions coexisted in the same tissue section. In all cases but one, a granulomatous response was seen around the damaged worms. The non-granulomatous pattern was observed in a patient seven days after beginning a 12-day course of DEC. In this case, a considerably dilated segment of the lymphatic showed cross-sections of adult worms with early signs of degeneration such as deformity and collapse of internal structures. Eosinophils, macrophages and lymphocytes infiltrated the vessel wall diffusely and were seen also entrapped in a fibrin-like material in the lumen, together with numerous free Mf (Fig. 1A). In all remaining treated and untreated cases, the lymphatic lumens were partially occluded by granulomas surrounding cross sections of dead adult worms in distinct stages of disintegration; the granulomas were composed of macrophages, lymphocytes, plasma cells and large numbers of eosinophils. In some granulomas, the damaged worms were circumscribed by a necrotic, acidophilic, granular material (Fig. 2A). The number of multinucleated giant cells and peripheral concentric fibrosis were accentuated in the more advanced cases, irrespective of antifilarial treatment and microfilariaemia status. Eosinophils were present in all cases, not only within the granuloma and interspersed with collagen fibers, but also spreading into the outlying areas, even in one case 90 days after DEC treatment; in this case, a segment of damage cuticle was surrounded by giant cells (Fig. 3A). In one case, 19 days after a single DEC dose, calcified and non-calcified segments of dead parasites coexisted in the same section. Neutrophils and microabscesses formation were not observed. In 18 cases, the adult worm gender could be recognized. In 17 cases only females were found, and in one case both male and female worms were present. All females were pregnant. In the remaining four cases, worm gender could not be established due to the advanced stage of disintegration. Eleven patients were Mf-negative in 11 mL of blood (Table 1) and the tissue specimens in seven of these showed free and/or intrauterine microfilariae.

**Immunohistochemistry**

In all tissue sections, intact eosinophils and granular, extra-cellular material were stained for the three proteins studied. In both DEC-treated and untreated cases, the
topographic distribution of labeled cells and extra-cellular granular material was similar. In the patient who received seven days of DEC treatment and did not have a well-established granulomatous reaction, stained intact cells and extra cellular granular material were abundant in the fibrin-like material around apparently degenerated extra-uterine Mf, localized within the lymphatic lumen (Fig. 1B, C, D). In all remaining cases, numerous labeled intact cells were present in the granulomas, running in concentric layers between other unmarked cellular components of the inflammatory response, and also extending to the peripheral tissues.Granular stained material was detected mainly around damaged adult worms (Fig. 2B, C, D), corresponding to necrotic area observed in con-

Table 1. Clinical and parasitological features and antifilarial treatment doses of the patients

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Total</th>
<th>DEC related</th>
<th>Not-DEC related</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number of patients (%)</td>
<td>22</td>
<td>12</td>
<td>10</td>
</tr>
<tr>
<td>Mean age (range), years</td>
<td>22.5 (15-34)</td>
<td>22.6 (19-34)</td>
<td>21.4 (15-32)</td>
</tr>
<tr>
<td>DEC dose</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Single dose</td>
<td></td>
<td>5</td>
<td>NA</td>
</tr>
<tr>
<td>12-day course</td>
<td></td>
<td>7</td>
<td>NA</td>
</tr>
<tr>
<td>Granuloma age (range), days</td>
<td>30.8 (7-90)</td>
<td>30.8 (7-90)</td>
<td>NA*</td>
</tr>
<tr>
<td>Site of the nodule</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Right</td>
<td>7</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>5</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Spermatic cord</td>
<td>11</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Infratesticular</td>
<td>1</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Paratesticular</td>
<td>0</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Geometric mean Mf density (range) per 1mL of blood</td>
<td>34.3 (0-4128)</td>
<td>90.7 (0-4128)</td>
<td>9.4 (0-1765)</td>
</tr>
<tr>
<td>Number of patients</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mf+/1mL</td>
<td>10</td>
<td>6</td>
<td>4</td>
</tr>
<tr>
<td>Mf+/1mL</td>
<td>1b</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Mf-/1mL</td>
<td>11</td>
<td>6</td>
<td>5</td>
</tr>
</tbody>
</table>

*It was not possible to provide the granuloma age, since the patients did not know when the nodule appeared (by self physical examination or by discomfort); *0.2 Mf/mL.

Table 2. Microfilarial density and eosinophil counts in peripheral blood of treated and untreated patients

<table>
<thead>
<tr>
<th>Relation to DEC treatment</th>
<th>Mf density</th>
<th>No. of patients</th>
<th>Mf density/mL</th>
<th>Eosinophils/mm³</th>
<th>p value*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Pre</td>
<td>Post</td>
</tr>
<tr>
<td>DEC-related</td>
<td>&lt; 100/mL</td>
<td>5a</td>
<td>6.5 (0-65)</td>
<td>360.3 (150-800)</td>
<td>352.4 (175-850)</td>
</tr>
<tr>
<td></td>
<td>&gt; 100/mL</td>
<td>7</td>
<td>597.4 (123-4128)</td>
<td>303.6 (50-600)</td>
<td>1205.4 (625-3800)</td>
</tr>
<tr>
<td>Not DEC-related ±</td>
<td>&lt; 100/mL</td>
<td>7b</td>
<td>0.8 (0-0.2)</td>
<td>280.2 (75-1650)</td>
<td>267.0 (50-1700)</td>
</tr>
<tr>
<td></td>
<td>&gt; 100/mL</td>
<td>3</td>
<td>833.5 (243-1765)</td>
<td>362.9 (300-425)</td>
<td>689.0 (475-950)</td>
</tr>
<tr>
<td>All patients</td>
<td>22</td>
<td>27.2 (0-4128)</td>
<td>315.3 (50-1650)</td>
<td>522.8 (50-3800)</td>
<td>0.015</td>
</tr>
</tbody>
</table>

*aTwo were Mf negative; *six were Mf negative; *Student t test (comparing log geometric means: post versus pre observations points); ± Pre and post indicate eosinophil counts measured when the nodule was first palpated by the examining physician and 7-14 days later, respectively.
Conventional histopathology (Fig. 2A). In several granulomas, worms in different stages of degradation coexisted. This was particularly exemplified in the nodule shown in figure 3; granular stained material was detected on the surface of intrauterine Mf. Even in the specimen collected 90 days after treatment (Fig. 3B), adult worms in advanced stage of disintegration were also found (Fig. 3C, D). In two cases (treated and non-treated), intact stained cells were found in the very act of attachment to the surface of dead parasites (Fig. 4A, B). Cellular and extra-cellular positive immunohistochemical staining for the three eosinophil proteins was scarce or even absent in relation to calcified worms segments, although intact cells were still present at the periphery of the granulomas (Fig. 4C). None of the control slides revealed any immunohistochemical staining for MBP (Fig. 4D), EPO, or ECP.

**DISCUSSION**

In our study, all three toxic proteins were localized in the same areas of the granuloma in both DEC-treated and
untreated cases, irrespective of microfilaremia and blood eosinophilia, and granuloma age. These findings seem to indicate that these parameters do not influence the role of eosinophils in the localized inflammatory process associated with the death of adult worms. The presence of labeled cells and granular, extra-cellular material suggests that eosinophils have accumulated and degranulated preferentially in close proximity to the adult parasites which are centrally located in the granulomas. Since eosinophils represent a hallmark of the inflammatory granulomatous reaction to degenerating adult *W. bancrofti*, it would be surprising if they did not degranulate. Indeed, it is believed that the acidophilic necrotic material observed by conventional histology (13-19) around degenerating parasites is composed, at least in part, of eosinophil-derived toxic granular material.

Positive immunohistochemical staining was seen in intact eosinophils and in extracellular granular material...
around intrauterine degenerating Mf. It is likely that eosinophils reached the Mf through a disrupted wall of the adult female, a finding suggested by the concomitant presence of Mf outside the parasite. One could speculate that the eosinophil influx and degranulation might be enhanced by DEC-damaged Mf. In fact, in lymph node (28) and skin (29, 30) material from patients with onchocerciasis, it has been demonstrated that eosinophils are found degranulating at the surface of Mf soon after DEC treatment. Of interest, we recently observed a female:male prevalence ratio of 4.5:1 in granulomatous tissues material (25); all females in the present series were pregnant.

Bancroftian adult filarial worms probably die from a variety of causes, involving both parasite- and host-related factors and drugs such as DEC; however, the exact pathways leading to their death are largely unexplained (31). Whatever the cause, there are several lines of evidence indicating that the presence of eosinophils and their toxic products in tissues harboring degenerating and dead adult worms is secondary to the parasite death rather than its cause. First, although our cases were selected randomly, the study included nodules obtained from patients with and without antifilarial treatment, and with a range of microfilaraemia and pre-treatment blood eosinophilia levels. Our study design allowed at least two different mechanisms of adult parasite death: DEC-related and natural conditions. The non-DEC related nodules in essence controlled for any interference of DEC on eosinophil homeostasis in microfilaremic patients such as recruitment, activation and degranulation (32). Irrespective of the clinical, parasitological and chemotherapeutic category of the patients, all the granulomas showed a similar immunohistochemical pattern for all three eosinophil proteins, making it unlikely that the death of adult *W. bancrofti* had been initiated by eosinophil-dependent or related mechanisms.

Second, absence of eosinophils in tissues containing living *W. bancrofti* adult worms, despite peripheral blood hyper eosinophilia due to concomitant intestinal helminthiases, has been reported (33). Third, post-treatment blood eosinophilia is a regular feature in response to ivermectin – a potent microfilaricidal drug without macrofilaricidal effects – in essentially all microfilaricemic patients regardless of the doses. The magnitude and kinetics of blood eosinophilia are proportional to pre-treatment Mf levels; blood eosinophilia peaks between 7 and 30 days after treatment, followed by a progressive return to normal levels thereafter (27). Despite this “acute” microfilaria-related blood eosinophilia, adult *W. bancrofti* worms remained alive, as evaluated by ultrasonography until several months after treatment (34).

Fourth, the complex and intriguing phenomenon by which the adult worms escape eosinophil attack is illustrated by some well-defined clinical conditions, such as tropical pulmonary eosinophilia. This uncommon extralymphatic manifestation of lymphatic filariasis is characterized by peripheral eosinophilia, and by high titers of IgE and antifilarial IgG4 antibodies (35). However, in spite of pre-treatment blood eosinophilia of more than 20,000/mm³, living adult worms can be seen by ultrasound in intra-scrotal lymphatics, and these can remain unaffected by aggressive DEC treatment (12mg/kg/day/30 days) (36). Thus, DEC is not 100% effective against adult worms and many of them are unaffected, even with increasing doses in both asymptomatic Mf-positive and Mf-negative individuals (26). Fifth, the so-called mixed reaction, in which alive and dead parasite share the same nest (19-21, 23, 26), demonstrates that some adult parasites are able to remain alive in spite of the strong inflammatory reaction directed against nearby damaged/dead worms. Of note, these mixed reaction have been observed in parasite nests from both DEC-treated and untreated patients. In our view, clarification of this remarkable phenomenon would help to unveil the mechanisms governing parasite resistance or susceptibility to the toxic products of eosinophils, antifilarial drugs, and natural conditions threatening the survival of adult worms. Sixth, in other filarial nematodes such as *O. volvulus*, there is ample evidence that DEC kills microfilaria and spares adult parasites, especially during the Mazzotti’s reaction when massive eosinophil recruitment and degranulation occur (30), which can threaten the patient’s life in cases of high parasite load. Localized lymphatic vessel inflammation with accumulation of eosinophils and subsequent degranulation was noted in a nodule formed as early as 7 days after beginning DEC treatment, when a clear granulomatous inflammatory reaction was not yet established. This finding explains why nodules can be detected by physical examination within seven day after the first dose of DEC intake (21, 26). Immunostained extra-cellular material and intact eosinophils were absent or scarce around remnants of *W. bancrofti* adult worms with calcification, a condition that supposedly “sterilizes” the parasite antigenic sources (25). Positive immunostaining was also observed in lesions 90 days after treatment, when non-calcified remnants of adult worms were barely discernable. The lifespan and the turnover of tissue-resident eosinophils are unknown. They usually undergo apoptosis after degranu-
loration and are replaced by newly arrived cells; they arrive at their tissue destination prepared for immediate action or prolonged tissue survival (37). In such situation, they might have some effector activity in maintaining control of tissue repair and remodeling, angiogenesis, fibroblast activation and collagen production (12).

A remarkable finding in our material is the virtual absence of neutrophils in the tissue sections, even in earlier cases. This finding is relevant in view of the attention that has been paid to the Wolbachia endosymbiont bacterium present in most filarial nematodes and its link with neutrophilic infiltrates after parasite death (38). On the other hand, despite the absence of Wolbachia (39), dying Loa loa adult worms cause significant inflammation with initial neutrophil accumulation and suppuration, followed by granulomatous reaction (40). In addition, the number, size and cellular composition of granulomas in lymphatic vessels or renal lymph nodes did not differ after removal of Wolbachia in gerbils chronically infected with B. pahangi, indicating that Wolbachia may not play a pivotal role in the development of the lymphatic lesions (41). It is clear that additional studies are needed on Wolbachia-host interactions in the development of W. bancrofti granulomas and their impact on pathogenesis of adult worm-related chronic conditions.

Our current and previous studies in bancroftian filariasis give additional support to the concept that, as far as adult helminths are concerned, tissue eosinophilia appears to be triggered after the sudden release of antigens from parasites dying either spontaneously or following chemotherapy (42). The present study has limitations: (a) a relatively small number of cases; (b) the lack of a quantitative approach for histochemistry patterns; (c) as a retrospective investigation, it was not possible to perform plasma cytokine and eosinophil-derived proteins profiles. Thus similar investigations using more refined techniques would be welcome; (d) due to the rarity of palpable filarial nodule formation in females and children (24), specimens from these population were not included and it is not possible to extrapolate the results to these groups. In summary, based on a pioneering, integrated clinical, laboratorial and immunohistochemical investigation, our findings indicate that eosinophils per se are not responsible for the initiation of lethal damage to adult worms, as much as intrascrotal filarial granulomas are concerned. Rather, eosinophil influx and degranulation in DEC-related and non DEC-related bancroftian granulomas likely represent a consequence of parasite death rather than its cause. However, eosinophils may play an important role in postmortem worm degradation as well as in granuloma evolution and modulation (12).

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CONFLICT OF INTEREST

I (Gerusa Dreyer) declare, on behalf of all authors, that we have no personal or professional interests that could be construed to have influenced the paper.

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