The nature of multinucleated cells in the nasopharynx

M. Y. ALI

From the Department of Pathology, University of Singapore, General Hospital, Singapore 3, Malaysia

SYNOPSIS Exfoliated multinucleated epithelial cells were shown to be present in nasopharyngeal smears from patients with moniliasis, tuberculous lesions, and non-specific chronic inflammations of the nasopharynx. Multinucleated cells found in the nasopharynx were cytologically similar to the so-called measles giant cells. Because of the association of these cells with a number of chronic inflammatory lesions, their function is probably related to phagocytosis. In evidence thereof, a few multinucleated cells, presumably of epithelial origin, showed phagocytosed particles within their cytoplasm. This view is also based on corroborative evidence given by a number of workers that epithelial cells in the respiratory mucosa and in the endometrium were shown to assume phagocytic functions comparable to those of free histiocytes.

The epithelial lining of the human nasopharynx is composed of squamous, ciliated, and transitional epithelia. On average, approximately 40% of the nasopharyngeal mucosa is covered by pseudo-stratified columnar ciliated epithelium (Ali, 1965). The ciliated epithelium of the nasopharynx is, morphologically, similar to that lining the trachea and bronchi but has a tendency to be taller (Papanicolaou, 1954). The most characteristic cytological features of the ciliated columnar epithelium are the cilia, the heavy cuticular border, the round to oval granular nucleus, and the perinuclear cytoplasmic vacuolation. In addition, Papanicolaou (1954) also stated that binucleation and multinucleation are not infrequent; cells of this type which have lost their ciliated border may be easily misinterpreted as malignant. Furthermore, he explained that multinucleation may be observed in malignant as well as in benign cells but, in general, its specific diagnostic significance has not as yet been determined. Ide, Suntzeff, and Cowdry (1959), however, regarded the multinucleated cells in the trachea and bronchi as atypical cells.

During the course of cytological examination of nasopharyngeal smears for cytodagnosis of carcinoma, multinucleated cells (ciliated and non-ciliated) were frequently encountered. Numerous other structural changes in the ciliated epithelia, mainly degenerative, similar to those described by Bryan and Bryan (1953, 1959), were often found in the nasopharyngeal smears. Cytologically, the appearance of these degenerate cells was not confused with any of the features of malignant cells. However, at the beginning of this study, multinucleated cells posed some difficulty in interpretation. Although the presence of multinucleated epithelial cells in the bronchial mucosa has been recognized, there was no mention of their occurrence in the nasopharyngeal mucosa. Moreover, since the function of these cells is not well understood, a closer study of the conditions in which they prevail appears to be mandatory.

The purpose of this communication is to report the frequent occurrence of multinucleated cells amongst other cells exfoliated from the human nasopharynx and to present the conditions associated with their presence. The genesis and the possible function of these cells are briefly discussed.

MATERIALS AND METHODS

Nasopharyngeal swabs were taken from 138 patients suspected of having carcinoma of the nasopharynx and from 25 apparently healthy individuals. Two smears, on non-albuminised microscopic slides, were prepared from each swab and immediately fixed in equal parts of ether and 95% alcohol. Nasopharyngeal biopsies were then taken from the 138 patients and nasopharyngeal mucosa from 100 necropsied medico-legal cases was utilized for the study of the normal histology.

The cytological smears and paraffin sections of the biopsy and necropsy material were all stained with haematoxylin and eosin. The same stain was used for the histological and cytological preparation to facilitate comparison of various cell types in each.

The cytological preparations were examined for the presence of malignant cells and each specimen was com-

Received for publication 5 February 1965.
pared with the corresponding histological slides. The smears taken from apparently normal persons were also compared with the histological appearances in the necropsy material. Binucleated and multinucleated cells, when found in the smears, were recorded on index cards bearing the relevant clinical and histological data. Thus the conditions associated with the presence of multinucleated cells were distinguished. Acute inflammatory cells clear exudate. Associated with the presence of multinucleated conditions patients with inflammation. The remainder of cases were divided into three groups according to the corresponding histological diagnoses: those showing nasopharyngeal carcinoma (with or without mucosal ulceration), benign conditions, and normal controls (see the Table below). In the first group, there were 79 smears from histologically proven cases of nasopharyngeal carcinoma. Clinically and histologically 44 of these cases showed no ulceration of the nasopharynx. None of the smears from the 44 cases showed the presence of multinucleated cells. Each of the remaining 35 cases showed protruding and ulcerative nasopharyngeal tumours. Smears from all these were positive for malignant cells but only three showed the presence of multinucleated cells. The second group, 59 smears, was representative of various inflammatory lesions, including granulomas and vasomotor rhinitis. In 17 of the 59 smears, abundant binucleate and multinucleate cells were found. The conditions associated with the prevalence of these cells were tuberculous granulomas (Fig. 1a), moniliasis, and non-specific chronic inflammation. The remainder of the smears, from 42 patients with benign lesions, showed acute suppurative conditions. Multinucleated cells were not found in any of these smears. Finally, none of the 25 control smears showed the presence of binucleate or multinucleate cells.

RESULTS

The smears were divided into three groups according to the corresponding histological diagnosis: those showing nasopharyngeal carcinoma (with or without mucosal ulceration), benign conditions, and normal controls (see the Table below). In the first group, there were 79 smears from histologically proven cases of nasopharyngeal carcinoma. Clinically and histologically 44 of these cases showed no ulceration of the nasopharynx. None of the smears from the 44 cases showed the presence of multinucleated cells. Each of the remaining 35 cases showed protruding and ulcerative nasopharyngeal tumours. Smears from all these were positive for malignant cells but only three showed the presence of multinucleated cells. The second group, 59 smears, was representative of various inflammatory lesions, including granulomas and vasomotor rhinitis. In 17 of the 59 smears, abundant binucleate and multinucleate cells were found. The conditions associated with the prevalence of these cells were tuberculous granulomas (Fig. 1a), moniliasis, and non-specific chronic inflammation. The remainder of the smears, from 42 patients with benign lesions, showed acute suppurative conditions. Multinucleated cells were not found in any of these smears. Finally, none of the 25 control smears showed the presence of binucleate or multinucleate cells.

Multinucleated epithelial cells were difficult to identify in histological sections due to crowding of the nuclei in the pseudo-stratified ciliated epithelia. However, in cytological preparations two types of multinucleated cells were found. Binucleated and multinucleated ciliated cells with up to 12 nuclei,

![Figure 1a](http://jcp.bmj.com/)

**FIG. 1a.** Histological section of a tuberculous lesion in the nasopharynx showing chronic inflammatory cells at its ulcer crater. Haematoxylin and eosin × 150.

![Figure 1b](http://jcp.bmj.com/)

**FIG. 1b.** Lymphocytes and binucleate macrophage in the cytological smear from the ulcer illustrated in Fig. 1a. Haematoxylin and eosin × 500.

### TABLE

<table>
<thead>
<tr>
<th>Source of Material</th>
<th>No. of Smears Examined</th>
<th>Smears Showing Multinucleated Cells</th>
<th>Average Frequency (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Detailed Record</td>
<td>Total</td>
<td>No.</td>
</tr>
<tr>
<td>Normal (controls)</td>
<td>25</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Carcinoma</td>
<td>79</td>
<td>3</td>
<td>3.8</td>
</tr>
<tr>
<td>Ulcerating</td>
<td>35</td>
<td>3</td>
<td>8.5</td>
</tr>
<tr>
<td>Non-ulcerating</td>
<td>44</td>
<td>0</td>
<td>—</td>
</tr>
<tr>
<td>Inflammations</td>
<td>59</td>
<td>17</td>
<td>28.8</td>
</tr>
<tr>
<td>Chronic</td>
<td>25</td>
<td>17</td>
<td>68.0</td>
</tr>
<tr>
<td>Acute</td>
<td>34</td>
<td>0</td>
<td>—</td>
</tr>
</tbody>
</table>

**The nature of multinucleated cells in the nasopharynx** 425
the largest measuring approximately 20 microns in width, were the commonest types encountered (Figs. 2 and 3). The other type had no identifiable cilia, was elliptical in shape, and often showed phagocytosed particles in the cytoplasm (Fig. 4a). The shape and size of the nuclei of these cells and their abundant cytoplasm, but for the lack of cilia, showed a morphological resemblance to the multinucleated ciliated cells (Fig. 4b).

**DISCUSSION**

Ciliated epithelial cells are often shed as a result of infections of the respiratory mucosa. Degenerative changes in the ciliated epithelia and multinucleated cells diagnostic of viral infections have been shown by Bryan and Bryan (1959). The nasopharynx, being a part of the upper respiratory tract, would naturally show all the structural changes that might develop as a result of irritation and infection.

The occurrence of numerous multinucleated giant cells in the pharyngeal mucosa in the prodromal stage of measles was reported by Warthin (1931). However, Algana (1911) had earlier recognized these cells in the tonsils and the respiratory epithelium in necropsy specimens from cases of measles. The presence of multinucleated cells in the lymphoid tissue suggested that they were derived from lymphocytes or reticulum cells (Warthin, 1931). However, Semsroth (1939) reported that epithelial multinucleated giant cells with cytoplasmic inclusion bodies were found in the respiratory mucosa in a fatal case of prodromal measles. The formation of these cells, he suggested, was by fusion of adjoining cells which might or might not retain their cilia. Also, Masugi and Minami (1938) illustrated with clarity that the multinucleated cells in the respiratory mucosa have arisen from fusion of respiratory epithelial cells.

The measles giant cells, having acquired that name, were also demonstrated in cytological smears of nasal secretions from cases of measles by Bryan and Bryan (1959). In their description, they accepted the epithelial origin of these cells and mentioned that cilia were occasionally seen. Recently, Mottet and Szanton (1961) confirmed that the presence of the multinucleated cells in nasal secretions was diagnostic of measles. However, they realized that these cells might occur in some chronic nasal infections but there was little likelihood that such patients would be clinically confused with cases of measles. The finding of multinucleated epithelial cells in bronchial smears, earlier described by Papanicolaou (1954), has recently been shown to be associated with pulmonary tuberculosis and significantly frequent in cases of combined cancer and tuberculous disease of
The nature of multinucleated cells in the nasopharynx

The authors postulated that the function of the multinucleated cells, similar to that of macrophages, is related to phagocytosis. Morphological studies alone were not altogether conclusive in establishing the mode of formation of the multinucleated epithelial cells. The epithelial origin of the so-called virus-induced multinucleated giant cells was revealed by Enders and Peebles (1954). They showed that epithelial cells in tissue cultures fused to form multinucleated cells as a direct result of invasion by measles virus. Papanicolaou and Maddi (1958, 1959) also produced convincing evidence to show that epithelial phagocytes were formed in tissue cultures of endometrial epithelium, particularly in the presence of cellular detritus resulting from endometrial infection or, in some instances, from contamination with Monilia. The evidence presented in favour of the view that endometrial cells assumed phagocytic function comparable to that of the free histiocytes confirmed the earlier observations of Duthie (1930) in which he found evidence of phagocytosis by the ciliated bronchial epithelium. On the other hand, cells concerned with phagocytic activities in higher vertebrates were always considered to be of mesenchymal origin (Mudd, McCutcheon, and Lucke, 1934) and the majority of workers believed that epithelial cells were non-phagocytic (Robertson, 1941).

Based on the criteria in the literature reviewed above, multinucleated epithelial cells were found in chronic and viral infections of the respiratory tract. The findings in the present study showed that the presence of multinucleated cells in smears from the nasopharynx was definitely related to chronic and granulomatous lesions. These cells, resembling the measles giant cells, were undoubtedly of epithelial origin. But it was not always possible, on a morphological basis, to differentiate between the non-ciliated giant cells and other types of multinucleated macrophages. However, since epithelial cells have been shown to assume phagocytic function, there is at least presumptive evidence to show that their presence in chronic inflammatory conditions is related to phagocytosis. Giant cells of histiocytic and monocytic origins (Aronson and Elberg, 1962; Wassermann, 1963), also formed by fusion of cells, are not of the same size as the multinucleated epithelial cells. Multinucleated cells in which phagocytosed particles were identifiable had no cilia but were morphologically similar to the other ciliated giant cells in size, shape, and cytological details.

REFERENCES

——, ——— (1959). Ibid., 78, 156.
The nature of multinucleated cells in the nasopharynx

M. Y. Ali

doi: 10.1136/jcp.18.4.424

Updated information and services can be found at:
http://jcp.bmj.com/content/18/4/424

**Email alerting service**

*These include:*

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to: http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to: http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to: http://group.bmj.com/subscribe/