Case Report

Hypersensitivity Pneumonitis Associated with Red-Vented Bulbul: A New Encounter of Bird Related Hypersensitivity Pneumonitis

W. D. N. L. Amarasinghe,1 R. Jayasekara,1 B. D. W. Jayamanne,2 T. D. K. Nalaka,3,4 W. A. D. L. Amarasiri,1 R. Punchihewa,5 and A. Fernando1

1Central Chest Clinic, Colombo, Sri Lanka
2Department of Public Health, Faculty of Medicine, University of Kelaniya, Sri Lanka
3Department of Radiology, National Hospital of Sri Lanka, Sri Lanka
4Department of Pathology, National Hospital for Respiratory Diseases, Welisara, Sri Lanka

Correspondence should be addressed to W. D. N. L. Amarasinghe; amaranadee@gmail.com

Received 25 June 2019; Revised 7 August 2019; Accepted 17 September 2019; Published 9 December 2019

Academic Editor: Akif Turna

Copyright © 2019 W. D. N. L. Amarasinghe et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

1. Introduction

Hypersensitivity pneumonitis (HP) is an immunologically-mediated inflammatory lung disease caused by repetitive inhalation of antigens in a susceptible host [1–3]. Potential causative agents are grouped as microbes, animal proteins, and chemicals [1, 4, 5]. Bird related HP is becoming the commonest form caused by high- and low-molecular-weight proteins found in feathers, faeces, and other animal products commonly of pigeons, parrots, parakeets, love birds, cockatoos, budgerigars, and fowl [4, 6–8]. We report two cases that presented to the Central Chest Clinic, Sri Lanka with a multidisciplinary diagnosis of subacute HP associated with red-vented bulbuls. There have been no previously published reports of HP associated with red-vented bulbuls or any other visiting birds. Hence, this is the first paper to study this association.

2. Case Presentations

Both patients were retired professionals from semiurban areas in Colombo who denied exposure to any organic or inorganic particles, which can cause HP, within the last two years. However, repeated questioning revealed that there had been close contact with red-vented bulbuls which frequented their homes.
houses. The birds nested in chandeliers hung directly above the sofas in their living rooms where these patients spent most of their leisure times. Though there had been debris falling on to the sofa as the birds moved within the nests above, they had not attempted to remove the nests as they enjoyed their presence. The living rooms had been kept closed during the periods when the birds reproduced, to protect the eggs and nestlings. Additionally, both patients had closely handled the nestlings until they fledged. This confirmed that there had been continuous exposure to antigens of red vented bulbuls for around 8–10 hours per day for 10–18 months with a proximity of 2–3 meters in a closed or partially closed environment, in these patients.

2.1. Case 1. A 65-year female presented to the clinic with insidious-onset persistent and progressive dyspnoea, cough, and wheezing for six months. She was a diagnosed diabetic and had a history of allergic rhinitis. She revealed close contact with red-vented bulbul birds for around one-and-half years. Erythrocyte sedimentation rate (ESR) was 41 mm/h. Chest radiography showed patchy opacifications in right mid zone and reticular nodular shadows in mid zones of both lung fields (Figure 1). Saturation at rest was 95% and there was 7% desaturation during the 6-minute walk test where she walked a distance of 480 meters. Spirometry and body plethysmography showed severe restriction and air trapping (FVC 49.5%, total lung capacity-TLC-76%, residual volume-RV-110%, RV/TLC-158%), with reduction in diffusing capacity with patchy parenchymal involvement (carbon monoxide diffusing capacity-DLCO-67%, carbon monoxide transfer coefficient-KCO-99%). HRCT showed centrilobular nodules of ground glass density in upper zones, basal ground glass opacities, and significant lobular air trapping (Figure 2). Video-assisted thoracoscopic lung biopsy showed patchy and focal interstitial thickening with lymphocytic infiltrate, minimal fibrosis, and several noncaseating granulomata within the interstitium (Figure 3).

2.2. Case 2. A 61-year-old male presented with insidious-onset, persistent and progressive dyspnoea and dry cough for four months. At the onset, he had low grade fever and constitutional symptom which subsided over a few weeks. He was a known hypertensive. He revealed close contact with red-vented bulbuls for around ten months. ESR was 78 mm/h. Chest radiography showed reticular nodular shadows in mid and lower zones of both lung fields (Figure 4). Spirometry showed severe restriction (FVC 52.8%, TLC-56%, RV-65%) with a normal diffusing capacity (DLCO-86%, KCO-180%). HRCT showed centrilobular nodules of ground glass density in all three zones and significant lobular air trapping (Figure 5). Bronchial wash cytology revealed inflammatory cells (249/cumm) with predominant lymphocytes (80%). Transbronchial lung biopsy showed thickened alveolar septae with lympho-histiocytic infiltrate and occasional neutrophils and eosinophils. Some of the alveolar spaces contained foamy histiocytes. Fibrosis was not evident.

A diagnosis of subacute HP associated with exposure to red vented bulbuls was made for both cases at a multidisciplinary meeting for interstitial lung diseases held at the National hospital of Sri Lanka. The patient of case 1 achieved good clinical and lung function improvement (FVC 91.9%) by about two months and that of case 2 showed clinical and lung function improvement (FVC 72%) around the third month of treatment with oral prednisolone (0.5 mg/kg) tail down regimen and the avoidance of the offending exposure. Both patients did not have any recurrences following avoidance of...
exposure to red-vented bulbuls and achieved normal FVC values in spirometry assessment with treatment of oral prednisolone within one-year follow-up period.

3. Discussion

Bird related HP (Bird fancier’s lung) is increasingly becoming prevalent around the world and shows worse outcome than other forms [8, 9]. Bird related HP can be caused by high- and low-molecular-weight proteins (<5μm) found in feathers, faeces, and other animal products [10].

Suspicion of an association between symptoms and contact with a provoking antigen is the first step in the diagnostic process of HP, combined with the measurement of serological markers and specific IgG antibody levels, radiological findings, lung function assessment, bronchoalveolar lavage, and lung biopsy for the complete workup [4, 12]. The identification of causal antigen is impossible in about 30–60% of cases [1]. Species specific antibody are commercially available for pigeons, parrots, parakeets, cockatoos, and multiple domestic poultry species at present, and there are emerging avian associations as in this report, to be experimented [1, 11, 13]. However, laboratory measures of exposure, such as precipitin tests, specific inhalation challenge, and lymphocyte proliferation tests have failed to achieve consensus among international experts as important in diagnosing HP according to international modified Delphi survey, possibly due to the limited information on their test characteristics, lack of standardization, or limited availability [14].

Air space involvement of HP in the lung parenchyma presents as patchy ground glass opacities and/or centrilobular nodules in HRCT [15]. Shunting of blood away from poorly ventilated regions manifests as mosaic attenuation. Persistence of hypoattenuated areas in expiratory CT films indicates air-trapping [8]. Spirometry may demonstrate a restrictive lung disease pattern, with moderate to severe reduction of FVC and DLCO [10]. Lung biopsy may show cellular bronchiolitis, diffuse lymphocytic interstitial infiltration, and non-caseating granulomas [10, 16].
Proper history plays a key role in HP as early avoidance of causative associations invariably brings complete or partial recovery whereas delay would cause irreversible lung fibrosis. Multidisciplinary approach is highly recommended when the diagnosis is challenging [6].

The red-vented bulbul (Pycnonotus cafer) also known as Kondakurulla in Sinhala, is a resident breeder across the Indian subcontinent including Sri Lanka, and has established itself in several Pacific islands, parts of the United Arab Emirates, United States, and possibly New Zealand, not limiting to a part of the world [17–19]. They build nests in bushes at a height of around 2-3 meters and opportunistically invade houses specially using lamp shades and chandeliers for nesting, leaving an invariable risk exposure to the household [20, 21].

To our knowledge, all published cases of bird related HP are associated with birds raised as pets or for farming. An important fact to be highlighted is that when a clinician questions regarding risk exposure to birds, the patient would give a negative answer initially if they have not raised birds purposefully. The opportunity to remove the exposure would be missed in such instances. Hence, the present observation highlights the importance of specific questioning regarding this kind of casual exposures to visiting birds within living places.

Birds nesting and breeding inside houses is considered to bring prosperity to the household in some cultural beliefs. Hence updating public on harmful aspects of these associations would also be important.

Disclosure

Written informed consent was obtained for publication of this case report and accompanying images from relevant authority.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

References

[1] R. Nogueira, N. Melo, H. N. Bastos et al., "Hypersensitivity pneumonitis: antigen diversity and disease implications," *Palmonology*, vol. 15, no. 2, pp. 97–108, 2019.

[2] A. N. Musa, K. N. Hirmizi, and M. A. Ibrahim, "Hypersensitivity pneumonitis–a case series," *Respirology*, vol. 22, no. S3, pp. 256–256, 2017.

[3] A. Matsuo, T. Horiiuchi, K. Sonenara, and Y. Wada, "The first case report of hypersensitivity pneumonitis progressed to pulmonary fibrosis induced by Lyophyllum aggregatum," *Respirology*, vol. 22, no. S3, pp. 266–266, 2017.

[4] T. Ismail, C. Mcsharry, and G. Boyd, "Extrinsic allergic alveolitis," *Respirology*, vol. 11, no. 3, pp. 262–268, 2006.

[5] A. S. Lee, H. E. Jo, and T. J. Corte, "Hypersensitivity pneumonitis: a protein and challenging disease," *Respirology*, vol. 22, no. 8, pp. 1489–1490, 2017.

[6] M. Ueyama, H. Yutani, S. Terada et al., "Usefulness of lymphocyte stimulation test against pigeon serum in management of patients with chronic hypersensitivity pneumonitis," *Respirology*, vol. 22, no. S3, pp. 256–256, 2017.

[7] S. S. A. Aljah, Z. A. Bakar, H. Lockman, G. J. T. John, and A. R. A. Mutalif, "Hypersensitivity pneumonitis case series," *Respirology*, vol. 22, no. S3, pp. 256–257, 2017.

[8] L. Winkler, Z. Patel, and R. Gue, "Pigeon breeder’s lung: a case report of hypersensitivity pneumonitis," *West Virginia Medical Journal OA*, 2017.

[9] F. Morell, A. Roger, L. Reyes, M. J. Cruz, C. Murio, and X. Murto, "Bird Fancier's Lung," *Medicine*, vol. 87, no. 2, pp. 110–130, 2008.

[10] M. Selman, A. Pardo, and T. E. King, "Hypersensitivity Pneumonitis," *American Journal of Respiratory and Critical Care Medicine*, vol. 186, no. 4, pp. 314–324, 2012.

[11] M. Funke and J. M. Fellrath, "Hypersensitivity pneumonitis secondary to lovebirds: a new cause of bird fancier’s disease," *European Respiratory Journal*, vol. 32, no. 2, pp. 517–521, 2008.

[12] S. J. Bourke, J. C. Dalphin, G. Boyd, C. Mcsharry, C. L. Baldwin, and J. E. Calvert, "Hypersensitivity pneumonitis: current concepts," *European Respiratory Journal*, vol. 18, no. 32 suppl, pp. 81s–92s, 2001.

[13] C. Mcsharry, G. M. Dye, T. Ismail, K. Anderson, E. M. Spiers, and G. Boyd, "Quantifying serum antibody in bird fanciers’ hypersensitivity pneumonitis," *BMC Pulmonary Medicine*, vol. 6, no. 1, 2006.

[14] J. Morisset, K. A. Johannson, K. D. Jones et al., "Identification of diagnostic criteria for chronic hypersensitivity pneumonitis an international modified delphi survey," *American Journal of Respiratory and Critical Care Medicine*, vol. 197, no. 8, pp. 1036–1044, 2018.

[15] M. Remy-Jardin, J. Remy, B. Wallaert, and N. I. Müller, "Subacute and chronic bird breeder hypersensitivity pneumonitis: sequential evaluation with CT and correlation with lung function tests and bronchoalveolar lavage," *Radiology*, vol. 189, no. 1, pp. 111–118, 1993.

[16] Y. Ohtani, S. Saiki, M. Kitaichi et al., "Chronic bird fancier’s lung: histopathological and clinical correlation. an application of the 2002 ATS/ERS consensus classification of the idiopathic interstitial pneumonias," *Thorax*, vol. 60, no. 8, pp. 665–671, 2005.

[17] D. Bhatt and A. Kumar, "Foraging ecology of red-vented bulbul in Haridwar India," *Forktail*, vol. 2001, no. 17, pp. 109–110, 2001.

[18] Birdlife international 2018, "Pycnonotus cafer. The IUCN Red list of threatened species," 2018, e.T22712695A132102224. [Cited 13 June 2019.]

[19] Bushana, "Biodiversity of Sri Lanka: Red-vented Bulbul (Pycnonotus cafer) [Internet], Biodiversity of Sri Lanka," 2019, https://biodiversityofsrilanka.blogspot.com/2013/01/kondakurilled-vented-bulbul.html

[20] R. Somasiri, "The Island. The Island," 2012, [cited 2018 Jun 18] http://www.island.lk/index.php?page_cat=article-details&page=article-details&code_title=45008.

[21] C. M. Inglis, "Curious site for nest of the Bengal Redvented Bulbul (Molpastes haemarrhous bengalensis)," *Journal of the Bombay Natural History Society*, vol. 28, no. 3–4, pp. 1135–1136, 1922, 40.