Original Article

Hansen's Neuritis Revisited – A Clinicopathological Study

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ABSTRACT

Introduction: Leprosy affecting the nerve solely or with concomitant skin lesions is not an uncommon condition in clinical practice. It is responsible for extensive morbidity and often poses a diagnostic challenge. This study aims to highlight the clinicopathological features of Hansen's neuritis (HN). **Materials and Methods:** In this retrospective study, cases of histologically diagnosed HN, from January 2010 to July 2017, were reviewed in the light of clinical features, treatment history, and outcome. **Results:** There were 18 cases of HN which accounted for 3.97% of total nerve biopsy samples ($n = 453$) and 0.02% of total histopathology samples ($n = 81,013$). The male: female ratio was 5:1 in the cases of HN. Age range was 20–79 years with a mean age of 42.4 years (standard deviation: ± 14.03). Among the HN cases, there were 13 cases of pure neuritic leprosy (61.1%). Mononeuritis multiplex was the most common finding in the nerve conduction study. Six (33.3%) cases exhibited histological features of borderline tuberculoid leprosy, followed by five (27.8%) cases of mid-borderline features, three (16.7%) cases each of borderline lepromatous and burnt-out HN, and one (5.6%) case of polar tuberculoid leprosy. Lepra bacilli were detected on Fite-Faraco stain in 44.4% cases. **Conclusion:** Diagnosis of HN depends on astute search for skin lesions, nerve thickening or tenderness, sensory or motor symptoms, histopathological examination, and demonstration of lepra bacilli.

KEYWORDS: Hansen's neuritis, lepra bacilli, mononeuritis multiplex, pure neuritic leprosy

INTRODUCTION

Hansen's disease is a chronic granulomatous disorder affecting the skin and nerves and is caused by *Mycobacterium leprae*. The earliest classification of this disease based on immunity status and response was given by Ridley and Jopling in 1966.^[1] Wade in 1952 first proposed the disease "neural leprosy," in which only peripheral nerves were involved. The technical committee of International Leprosy Congress in 1953, held in Madrid, accepted "neuritic leprosy" as one subtype among the major groups of leprosy.^[2] Neuritic leprosy is one of the important causes of inflammatory peripheral neuropathy worldwide, especially in the tropical countries. In the absence of skin involvement, pure neuritic leprosy (PNL) may clinically mimic vasculitic neuropathy. In this study, we describe the

clinicopathological profile of Hansen's neuritis (HN) diagnosed in our institute.

MATERIALS AND METHODS

The study was conducted in the Department of Pathology, Kasturba Medical College, Mangalore, Karnataka, India. Institutional Ethical Committee's permission was obtained before commencing this study. We retrieved clinicopathological data and slides of all the histologically diagnosed cases of HN from our archives, during the period from January 2010 to July 2017. The samples were received in the department from the parent institute, other medical colleges, and hospitals in and

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around Mangalore. All the cases were microscopically reviewed taking note of clinical details, including findings of nerve conduction study (NCS), which was collected from the medical records department and biopsy requisition forms. The corresponding skin biopsies were also reviewed.

Formalin-fixed, paraffin-embedded transverse and longitudinal sections of nerve segments were stained with hematoxylin and eosin, Masson trichrome, and Fite-Faraco (Fite) stains. Simultaneously, the transverse sections were also stained with Kulchitsky Pal (KPal) stain (for myelin), after secondarily fixing of the nerve biopsies in Fleming's solution. In the nerve and skin biopsies, the bacillary index (BI) was computed based on Ridley's scoring scheme.^[3] Microscopic pathology in the nerve was categorized into indeterminate^[4] and burnt-out/healed Hansen's categories as well as in addition to the five classes of Ridley and Jopling scheme.

All the demographic data, clinical findings, and results of microscopic examination were tabulated and analyzed by mean, standard deviation (SD), and percentages. The results were laid out in the form of graph and tables. Statistical methods could not apply in this study because of small sample size. Sincere attempts were made to collect the follow-up data in all the cases.

RESULTS

In the above-mentioned period, a total number of 81,013 samples were received in the pathology department for histopathological examination, of which there were 1982 skin biopsies (2.45% of total histopathology samples). Hansen's disease of the skin was diagnosed in 131 (6.6% of total skin biopsies and 0.16% of total histopathology cases). During the same time span, 453 nerve biopsies (0.56% of total histopathology samples) were received for etiological diagnosis of peripheral neuropathy. HN was diagnosed in 18 (3.97% of total nerve biopsy samples and 0.02% of total histopathology samples).

Sural nerve was biopsied in 16 cases, followed by right greater auricular nerve and cutaneous branch of left radial nerve in one case each. The male: female ratio was 5:1 in the cases of HN with 15 (83.3%) cases diagnosed in men. Age range was 20–79 years with a mean age of 42.4 years (SD $\pm$ 14.03). Maximum number of male patients presented in the age group of 41–50 years (5 cases, 27.8%), followed by three cases (16.7%) each in the age groups of 21–30 years and 31–40 years, two cases (11.1%) in 51–60 years, and one case (5.6%) each in the age groups of 11–20 years and 71–80 years. The three female cases belonged

to the age groups of 31–40 years, 41–50 years, and 51–60 years (5.6% each) [Table 1].

Skin lesions consistent with leprosy were present in 5 out of 18 cases (27.78%) of HN, and in other 2 cases, there was no information available on skin affection by lepra bacilli. Hence, there were 13 cases of PNL (61.1%) among the HN cases, and these accounted for 2.8% cases of all nerve biopsy ($n = 453$) samples. Case 1 had a single hypopigmented and hypoesthetic skin patch in left buttock, and Case 4 had multiple hypopigmented macules in the trunk and erythematous patches in both the legs. Histopathological examination of the skin biopsies in Cases 1 and 4 revealed borderline tuberculoid (BT) (BI: zero) and borderline lepromatous (BL) (BI: 5) morphology, respectively. The corresponding nerve biopsies in both these cases showed BB morphology. Cases 11, 16, and 18 also had skin lesions which were not biopsied. Age range in PNL was 20–56 years with a mean age of 40.6 years (SD $\pm$ 10.78). Maximum number of male patients was in the age group of 41–50 years (4 cases; 30.7%), followed by two cases (15.4%) each in the age groups of 21–30 years, 31–40 years, and 51–60 years, and one case (7.7%) in the age group of 11–20 years. One female case (7.7%) each presented in the age groups of 31–40 years and 41–50 years.

The patients had myriad signs and symptoms. Most of the cases (44.4%; 8 out of 18 cases) presented with features of sensory loss and motor weakness with variation in spatial and temporal distribution. In 4 cases (22.2%), one or multiple peripheral nerves were thickened. Three cases (16.7%) had nonhealing trophic ulcers in the foot. Two (11.1%) patients were alcoholics and one case (5.6%) had retroviral disease. Two (11.1%) cases each had clinical history of peripheral neuropathy and mononeuritis multiplex. Case 1 presented with lower motor neuron palsy of bilateral facial nerves of 5 days duration [Figure 1], involving only the upper fibers, and Case 18 had claw hand for 2 years. Rheumatoid factor positivity was noted in Case 6, and there was family history of leprosy in skin in Case 14. There were no clinical details available in Case 9 [Table 1].

The findings of NCS available in 13 (72.2%) cases revealed a common pattern of mononeuritis multiplex with severe degree of sensory more than motor axonopathy. In Case 14, NCS recorded axonal neuropathy involving the bilateral median, right superficial peroneal, saphenous, sural, and posterior tibial nerves. Seven cases had a clinical diagnosis of HN, of which three cases had differential diagnoses of Charcot-Marie-Tooth disease, vasculitic neuropathy and diabetic neuropathy. In one (5.6%) case each, the provisional diagnosis

Table 1: Clinicopathological summary of Hansen's neuritis cases

Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
1	25/ male	Inability to completely close eyes since 5 days. Examination revealed lower motor neuron paresis of bilateral facial nerves involving only the upper fibres. Bilateral greater auricular nerve, supraorbital nerve, ulnar and posterior tibial nerves were thickened	Single hypopigmented and hypoaesthetic skin patch in left buttock	Mononeuritis multiplex, sensory more than motor axonopathy	Hansen's neuritis	Right greater auricular nerve	BB; BI: 2 Complete loss of all myelinated fibres. Foamy histiocytes, interspersed with lymphocytes, plasma cells in endoneurium along with few poorly formed epithelioid granulomas. Perineurium and epineurium show dense lymphohistiocytic infiltrates and around blood vessels	BT; BI: 0 Pandermal epithelioid granulomas without Langhans giant cells	MDT for MB leprosy	Complete recovery
2	20/ male	Slowly progressive numbness over left medial malleolus, lateral aspect of left foot and antero-medial aspect of left leg of 2 years duration with slipping of footwear. Examination revealed diminished sensation of pain and temperature, absent left ankle reflex. No nerve thickening	Absent	Mononeuritis multiplex, sensory more than motor axonopathy	N/A	Left sural nerve	BB; BI: 0 Severe degree loss of all myelinated fibres. Endoneurium effaced by poorly formed epithelioid granulomas, few foamy macrophages and interspersed plasma cells. The perineurium show lymphocytic infiltrate. The epineurium show perivascular lymphocytic infiltrate		MDT for MB leprosy	Complete recovery
3	49/ female	Consistent with mononeuropathy multiplex. No nerve thickening	Absent	Mononeuritis multiplex, sensory more than motor axonopathy	Diabetic neuropathy or Hansen's neuritis	Right sural nerve	BT; BI: 0 Severe degree loss of all myelinated fibres. Epithelioid granulomas		MDT for MB leprosy	Complete recovery

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Table 1: Contd...

Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
4	44/ male	Reduced sensation left little finger of 5 years duration. Examination revealed bilateral asymmetric loss of pain and temperature sensation in legs. Bilateral ulnar and sural nerves were thickened	Multiple hypopigmented macules in the trunk and erythematous patches in both the legs	Mononeuritis multiplex, sensory more than motor axonopathy	Hansen's neuritis	Right sural nerve	involving the endo-, peri-, and epineurium. Dense perivascular lymphoplasmacytic infiltrate in the epineurium BB; BI: 5 Complete loss of all myelinated fibres. Endo-, peri- and epineurial ill-defined aggregates composed of histiocytes and lymphocytes. Moderate perivascular lymphoplasmacytic infiltrate in endo- and epineurium	BL; BI: 5 Lympho- and foamy histiocytic infiltrate around the upper dermal blood vessels and dermal nerves	MDT for MB leprosy	Complete recovery
5	30/ male	Trophic ulcer in left foot. Decreased sensation in upper limbs. No nerve thickening	Absent	Axonal neuropathy	N/A	Left sural nerve	BL; BI: 5 (globi present) Complete loss of all myelinated fibres. Moderate lymphohistiocytic infiltrate involving the endoneurium and perineurium. Perineurium is thickened. Dense perivascular lymphocytic infiltrate in the epineurium		MDT for MB leprosy	Complete recovery
6	45/ male	Peripheral neuropathy. Rheumatoid factor positive	Absent	N/A	Rheumatoid vasculitis	Right sural nerve	BL; BI: 4 Complete loss of all myelinated fibers. Endoneurium is infiltrated by numerous lymphocytes, foamy histiocytes, and few epithelioid histiocytes.		MDT for MB leprosy	N/A

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Table 1: Contd...

Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
7	50/ male	Right common peroneal nerve and left ulnar nerve palsy with thickening	Absent	Mononeuritis multiplex	N/A	Left sural nerve	<p>Perineurium is focally thickened. Peri- and epineurium has dense lymphocytic infiltrate and scattered histiocytes</p> <p>BB; BI: 2</p> <p>Complete loss of all myelinated fibers. Endoneurium showed perivascular, moderate -to-dense lymphohistiocytic infiltrate, and sparse neutrophilic infiltrate along with epithelioid granulomas and foamy histiocytes. Endoneurial capillaries are hyalinized with prominent endothelial cells. Perineurium is thickened with moderate lymphohistiocytic infiltrate. Epineurium had moderate perivascular lymphohistiocytic infiltrate and neovascularization</p>		MDT for MB leprosy	Complete recovery
8	38/ female	Peripheral neuropathy, nonhealing ulcer in right foot	Absent	N/A	N/A	Left sural nerve	<p>Burnt-out Hansen's neuritis; BI: 0</p> <p>Complete loss of all myelinated fibres. Dense collagenization in the endoneurium. Endo-, peri-, and epineurium had moderate perivascular</p>		N/A	N/A

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Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
9	42/ male	N/A	N/A	N/A	N/A	Right sural nerve	lymphoplasmacytic infiltrate. Focal epithelioid histiocytes aggregates were seen. Nutrient blood vessel showed subendothelial fibrosis and medial hypertrophy BT; BI: 0 Complete loss of all myelinated fibres. Moderate endo-, peri-, and epineurial lymphohistiocytic inflammation. Dense perivascular lymphocytic infiltrate with focal neovascularization in the epineurium. Nutrient blood vessel showed subendothelial fibrosis, medial hypertrophy, and calcification		N/A	N/A
10	51/ male	Right ulnar nerve thickening. Distal muscle weakness	Bilateral lower limb trophic ulcer	Mononeuritis multiplex	NA	Left sural nerve	BT; BI: 0 Complete loss of all myelinated fibres. Endoneurial epithelioid granulomas rimmed by lymphocytic and interspersed plasma cells. Prominent endoneurial capillaries Peri- and epineurium showed moderate to dense lymphoplasmacytic infiltrate and neovascularization		MDT for MB leprosy	Complete recovery

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Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
11	32/ male	Tingling, numbness and decreased sensation in both lower limbs and hands since 1 year	Reddish raised lesion over toes, fingers, and extremities since 2 months	Mononeuritis multiplex	Hansen's neuritis	Left radial cutaneous nerve	BL; BI: 6 (globi present) Asymmetric, moderate degree loss of myelinated nerve fibres. Partial effacement of fascicular architecture by moderate lymphohistiocytic infiltrate and foam cells in the endoneurium. Peri- and epineurium showed moderate to dense lymphohistiocytic and ill-defined epithelioid cell aggregates	Not done	MDT for MB leprosy	Complete recovery
12	38/ male	Alcoholic. Since 3 years progressive asymmetrical to symmetrical upper and lower limb weakness, and complete claw hand	Absent	N/A	Hansen's neuritis or Charcot Marie Tooth disease	Left sural nerve	BT; BI: 0 Severe degree loss of small and large myelinated nerve fibres, endo-, and perineurial, well-defined, epithelioid granulomas with lymphoplasmacytic infiltrate. Epineurium shows perivascular lymphoplasmacytic infiltrate		MDT for MB leprosy	Mild motor weakness persisting. Claw hand persisting
13	56/ male	Mononeuritis multiplex	NA	NA	Hansen's neuritis	Right sural nerve	BT; BI: 1 Complete loss of myelinated nerve fibres. Endoneurial epithelioid granulomas with lymphoplasmacytic rimming, and scattered foamy histiocytes.		MDT for MB leprosy	N/A

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Table 1: Contd...

Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
14	25/ male	2.5 months history of asymmetrical multiple sensory painful small fiber mononeuropathies. Right arm wasting History of skin Hansen's in the brother (details NA)	Absent	Bilateral median, right superficial peroneal, saphenous, sural and posterior tibial nerves: Axonal neuropathy	Hansen's neuritis or Vasculitic neuropathy	Right sural nerve	Peri- and epineurium exhibits dense shows lymphocytic infiltrate admixed with plasma cells and few histiocytes BT; BI: 0 Complete loss of myelinated nerve fibres and endoneurial fibrosis. Epithelioid granulomas admixed with few foamy histiocytes and rimmed by lymphocytes are noted in endo-, peri- and epineurial compartment. Epineurial compartment also shows moderate perivascular lymphocytic infiltrate		MDT for MB leprosy	Resolution of sensory symptoms
15	36/ male	Alcoholic. Right foot drop since 4 months. History of trauma in right leg 8 months back. Right dorsiflexion at ankle weak. External hallucis longus muscle weak on right side. Sensory loss in dorsum of right foot	Absent	Axonal neuropathy in right peroneal nerve	Vasculitic neuropathy	Right sural nerve	TT; BI: 0 Severe degree loss of myelinated fiber. Necrotizing epithelioid granulomas with Langhans giant cells involving the endo-, peri-, and epineurium. Granulomas are rimmed by dense lymphocytic infiltrate and plasma cells. Epineurial perivascular, dense lymphohistiocytic infiltrate		MDT for MB leprosy	Resolution of sensory symptoms. Significant improvement in foot drop

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Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
16	55/ female	Nonhealing ulcer in left foot since 10 years, and had received treatment for leprosy. Known case of retroviral disease	Previous skin lesion (details N/A)	Mononeuritis multiplex	N/A	Right sural nerve	Burnt-out Hansen's neuritis; BI: 0 Complete loss of small and large myelinated nerve fibres, marked endoneurial sclerosis, and mild lymphocytic infiltrate. Peri- and epineurial mild-to-moderate lymphocytic infiltrate. Epineurial neovas- cularization, subendothelial fibrosis and mild to moderate perivascular lymphocytic infiltrate	Not done	No treatment	N/A
17	48/ male	Paraesthesia of bilateral foot and left hand for 2-3 years, loss of sensation in right foot for 2-3 months, bilaterally weak ankle dorsiflexion, graded sensory loss bilaterally in both upper and lower limbs	Absent	Sensory more than motor axonopathy	Nutritional neuropathy	Left sural nerve	BB; BI: 6 (globi present) Asymmetric, moderate degree loss of myelinated nerve fibres. Endo- and perineurial poorly formed epithelioid granulomas, with lymphocytes and foam cells. Epineurial moderate perivascular lymphocytic infiltrate		MDT for MB leprosy	On-going treatment
18	79/ male	Known case of motor neuron disease. History of bilateral claw hand of 2 years duration	Hypopigmented skin lesion in left lower limb. Slit skin smear: Negative for lepra bacilli	Severe motor and sensory axonal polyn europathy	Healed Hansen's neuritis	Left sural nerve	Burnt-out Hansen's neuritis; BI: 0 Complete loss of small and large myelinated nerve fibres, marked endoneurial sclerosis and mild to moderate lymphocytic infiltrate. Perineurium is thickened and	Not done	No treatment	

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Table 1: Contd...

Case number	Age (years)/ sex	Clinical features	Skin involvement	NCS	Provisional clinical diagnosis	Site of biopsy	Histological findings in nerve biopsy	Skin biopsy findings	Treatment	Follow-up
							exhibits moderate lymphocytic infiltrate. Medial hypertrophy in nutrient vessel with moderate, perivascular lymphohistiocytic infiltrate			

BB: Mid-borderline, BI: Bacillary index, BL: Borderline lepromatous, BT: Borderline tuberculoid, N/A: Not available, MB: Multibacillary, MDT: Multidrug therapy, NCS: Nerve conduction study, TT: Tuberculoid

was burnt-out HN, vasculitic neuropathy, rheumatoid vasculitis, and nutritional neuropathy. Provisional clinical diagnosis was not available in seven (38.9%) cases [Table 1].

In the nerve biopsies examined, six (33.3%) cases exhibited histological features of BT leprosy [Figure 2], followed by five (27.8%) cases of mid-borderline (BB) features [Figure 1], three (16.7%) cases each of BL [Figure 3] and burnt-out HN [Figure 4], and one (5.6%) case of polar tuberculoid (TT) leprosy [Figure 5]. There was no case of polar lepromatous (LL) or indeterminate morphology among the cases studied. Among the HN cases which had concomitant skin lesions, there were two cases of BB (Case 1 and 4) and healed HN each (Case 16 and 18), and one case (Case 11) BL. There were six cases having BT histology, followed by three cases of BB, two cases each of BL, and one case each of burnt-out HN and TT morphology among the 13 PNL cases.

Complete loss of myelinated fiber loss was seen in 12 (66.7%) cases and among them four cases showed BT morphology, followed by three cases each of BB and burnt-out morphology, and two cases of BL morphology. Severe degree of myelin loss was noted in four (22.2%) cases, and among them two cases had BT, and one case each had BB and TT morphology. Two (11.1%) cases (BL and BB morphology) had moderate degree of myelinated fiber loss. Among six cases of BT leprosy in the nerves, three (50%) cases exhibited well-defined endo-, peri-, and endoneurial epithelioid granulomas, whereas endo- and perineurial epithelioid granulomas were seen in two (33.33%) cases and only endoneurial granulomas in one (16.67%) case. Among five cases of BB leprosy in the nerves, three (60%) cases exhibited poorly defined granulomas involving only the endoneurial compartment, one (20%) case each involving all three and both endo- and perineurial compartments. The only case of TT showed necrotizing epithelioid

granulomas having Langhan's giant cells and rimmed by dense lymphoplasmacytic infiltrate, involving the endo-, peri-, and epineurium. Hence, epithelioid granulomas were detectable in 66.67% (12) cases. All three (100%) cases of BL did not show granulomas; rather, the fascicles were effaced by sheets of lymphohistiocytic infiltrate and interspersed foam cells.

All three (100%) cases of burnt-out HN were characterized by dense endoneurial collagenization or sclerosis, mild-to-moderate lymphohistiocytic inflammation in all the three compartments which was localized predominantly around blood vessels and thickened nutrient vessels. Foam cells were seen in eight (44.4%) cases, which included all cases of BL and BB, and two cases of BT. Plasma cells were present in all cases of BT and TT. Perineurial thickening was present in four (22.2%) cases of HN, which included two cases of BL, and one case each of BB and burnt-out HN. Out of 18 HN biopsies, epineurial perivascular inflammation, neovascularization, and vascular thickening were present in 17 (94.44%), 4 (22.22%), and 3 (16.67%) cases, respectively [Table 1]. There were not any microscopic features to differentiate between PNL and HN with skin lesions.

BI in HN cases ranged from 0 to 6 in five (83.33%) out of six cases of BT leprosy and three (100%) cases of burnt-out HN. One case each of TT (100%) and BL (33.3%) had a BI of zero. lepra bacilli were detected on Fite-Faraco stain in 8 (44.4%) out of 18 cases. BI was 1 and 4 in one case each of BT (16.7%) and BL (33.3%), respectively. Toward the lepromatous end of the spectrum, density of lepra bacilli in the tissue increased with a BI of 2 in 2 cases of BB, to BI-5 in 1 case each of BB and BL and BI-6 in 1 case each of BB and BL. Globi were encountered in 3 (16.67%) out of 18 cases, that is cases -5 (BL; BI-5), -11 (BL; BI-6), and -17 (BB; BI-6).

Out of 18 cases, 14 cases were classified as multibacillary (MB) and treated with multidrug

Figure 1: Case 1 mid-borderline – (a) Thickened right greater auricular nerve; (b) bilateral lower motor neuron facial palsy exhibiting lagophthalmos; (c and d) gross of thickened right greater auricular nerve; (e) whole-mount view of transverse section of thickened nerve; (f) enlarged fascicles ($\times 40$, H and E); (g) fascicles completely effaced by inflammatory cells ($\times 100$, H and E); (h) mononuclear cells, foam cells, and histiocytes ($\times 200$, H and E); (i and j) poorly formed granuloma ($\times 400$, H and E); (k) few remnant myelinated nerve fibers ($\times 400$, KPal); (l) lepra bacilli (OIF, Fite); (m) ($\times 40$, H and E), (n) ($\times 100$, H and E), (o) ($\times 400$, H and E); well-defined epithelioid granuloma in papillary and reticular dermis

therapy (MDT). In two cases, treatment history was not available and two cases of burnt-out HN did not receive any treatment. Complete recovery was seen in eight cases. There was resolution of sensory symptoms in two cases, persistence of motor weakness and foot drop in

Figure 2: Case 3 borderline tuberculoid – (a) Distinct inflammation in endo- and perineurium, marked endoneurial fibrosis (inset, Masson trichrome) and severe loss of myelinated nerve fibers ($\times 40$, H and E); (b) thickened perineurium and epineurial perivascular inflammation ($\times 100$, H and E); (c) thickened perineurium and endoneurial inflammation ($\times 100$, H and E); (d) epithelioid granuloma ($\times 400$, H and E)

one case each. One case is still undergoing MB-MDT and three cases were lost to follow-up. Leprea reactions were not reported in any of the cases.

DISCUSSION

In 1903, Albert Neisser described a “neural type of leprosy/lepra nervorum” for the first time and added the same to the already accepted “nodular” and “anesthetic” forms of leprosy.^[5] The Indian Association of Leprologists (IALs) included the distinct form of “neural leprosy” in their official six group classification in 1955 and named it “polyneuritic leprosy.”^[5] HN clinically manifests as thickened or tender nerves leading to sensory, motor, or autonomic disturbances and formation of trophic ulcers as well as loss of tissue. World Health Organization (WHO) in 1997 classified Hansen’s disease based on number skin lesions as paucibacillary (PB) (1 lesion), PB (2–5 lesions), and MB (>5 lesions).^[6] Later, WHO categorized LL, BL, and BB cases of the Ridley–Jopling classification, with a bacteriological index of ≥ 2 at any site in the initial skin smears as MB leprosy. On the other hand, PB leprosy included indeterminate, TT, and BT with a bacteriological index of < 2 . At least one of the cardinal signs is mandatory for the diagnosis of leprosy, which include definite loss of sensation in a pale (hypopigmented) or reddish skin patch, a thickened or enlarged peripheral nerve with a loss of sensation and/or weakness of the muscles supplied by that nerve, or the presence of acid-fast bacilli in a slit-skin smear. The cases can be classified as PB or MB based on number of skin lesions when slit-skin smear examination

Figure 3: Case 5 borderline lepromatous – (a) Inflammation in endo-, peri-, and epineurium ($\times 40$, H and E); (b) thickened perineurium ($\times 100$, H and E); (c) marked endoneurial fibrosis and severe loss of myelinated nerve fibres ($\times 100$, Masson trichrome); (d) endoneurial inflammation ($\times 200$, H and E); (e) dense lymphocytic infiltrate admixed with foam cells ($\times 400$, H and E); (f) numerous lepra bacilli and globi (OIF, Fite)

Figure 4: Case 8 burnt-out Hansen's – (a) ($\times 40$, H and E), (b) ($\times 100$, H and E): mild lymphocytic infiltrate in endo-, peri-, and epineurium; (c) neovascularization and mild perivascular lymphocytic infiltrate in epineurium ($\times 400$, H and E); (d) ($\times 40$, Masson trichrome), (d) inset, (e) ($\times 100$, Masson trichrome): Dense collagenization in the endoneurium; (f) complete loss of myelinated nerve fibers ($\times 100$, KPal)

is not available. It is recommended that the patient should be treated as a MB Hansen's case whenever the classification of an individual case is in doubt.^[7] There is the absence of any WHO guidelines for subclassification of PNL as PB or MB disease or its treatment.^[8] In 1982, IAL re-christened “polyneuritic leprosy” as “pure neuritic type of leprosy.”^[12] Single or multiple larger nerve trunks or their branches are enlarged in PNL with sensory loss in the corresponding dermatome, with neither skin lesions nor slit-skin smear positivity. Microscopy may reveal TT, borderline nonspecific, or even lepromatous morphology along with acid-fast bacilli.^[2] According to the present National Leprosy Eradication Programme guidelines in India for therapeutic purpose, involvement of one or more nerve trunks is considered as PB and MB, respectively.^[9]

In 2014, throughout the world, 213,899 people were newly diagnosed with Hansen's disease, which corresponds to a detection rate of 3.0/100,000 of population. Southeast Asian region registered the highest incidence of 8.12/100,000 of population. The overall registered prevalence in the world and Southeast Asia is 0.25 and 0.63/100,000 population, respectively. Of the newly diagnosed cases, 94% patients were reported in 13 countries: Bangladesh, Brazil, Democratic Republic of Congo, Ethiopia, India, Indonesia, Madagascar, Myanmar, Nepal, Nigeria, the Philippines, Sri Lanka, and the United Republic of Tanzania. MB cases constituted 61% of the patients, of which 74% were males.^[10]

India contributes to more than 50% of new cases detected globally every year. In the year 2011–2012,

Figure 5: Case 15 polar tuberculoid – (a) Enlarged fascicles with moderate endo-, peri-, and epineurial inflammation, and epithelioid granulomas ($\times 40$, H and E); (b) granuloma with Langhans giant cell (arrow) in epineurium ($\times 200$, H and E); (c) caseating (asterix) epithelioid granuloma ($\times 40$, H and E); (d) caseating (asterix) epithelioid granuloma and Langhans giant cell (arrow) ($\times 400$, H and E); (e) complete loss of myelinated nerve fibers ($\times 200$, KPal); (f) moderate degree loss of myelinated nerve fibres ($\times 200$, KPal)

127,000 new cases were diagnosed with an annual new case detection rate of 10.35/100,000 of population. As on April 1, 2012, a total of 83,000 cases are on record with a prevalence rate of 0.68/10,000 population.^[9] According to Indian data, PNL constituted about 4%–18% of leprosy patients.^[11] The incidence is reportedly higher in South India comprising up to 18% of new cases. PNL is more common in men,^[12] as in the present study. In the study by Mendiratta *et al.*, most number of cases occurred in the age group of 15–30 years,^[12] unlike the preponderance in 40–50 years revealed in the present study.

Among 46 cases of PNL, peripheral nerves were thickened in 32.6% cases, followed by trophic ulcers in 21.7% cases, muscle wasting, claw hand, and foot drop in 15.2%, 10.8%, and 8.7% cases, respectively.^[13] Mononeuritis multiplex was the most frequent clinical and electrophysiological pattern of nerve dysfunction, with sensory (89% of all cases) more than motor (81%) dysfunction and predominant axonal neuropathy^[14] comparable to the current study. Bilateral facial palsy is less common than unilateral facial palsy and the incidence of the same in leprosy varies from 3% to 24.59%.^[15] Reddy *et al.* described dichotomy between immunological grading of skin and nerve lesions in cases where both are involved simultaneously, with the nerve showing lower spectrum of the disease,^[16] and such a finding was noted only in case-4.

In the study by Hui *et al.* in 46 cases of PNL, there were 47.8% cases of BT, followed by 23.9%, 13%, and 10.8% cases of BL, TT, and BB, respectively.^[13] Endoneurial, perineurial, and epineurial inflammation was seen in

39.8%, 39.8%, and 27.7% cases, respectively. Epithelioid granulomas were described in 13.2% cases. Fibrosis was most common (34.7% cases) in perineurial compartment, followed by endoneurial and epineurial compartments in 31.6% and 26.3% cases, respectively. Reduction in myelinated fibres was noted in 55.6% cases. Mononuclear infiltrate and foamy macrophages were encountered in 42.3% and 19% cases. Endoneurial edema was described in 7.8% cases. In 19% cases, perineurial enlargement was noted. Microfascicles are nest-like structures composed of a variable quantity of small myelinated or nonmyelinated fibres and Schwann cells and surrounded by perineurial cells. These structures were discerned in 10.42% samples. In the same study, mononuclear cell infiltrate and fibrosis were seen in all three compartments in almost all the cases, epithelioid granulomas in 66.7% cases, foam cells in 44.4% cases, and perineurial thickening in 33.3% cases.^[17] Microfascicles were not seen in any of the cases in the present study. Another study reported epithelioid granulomas in 14% cases.^[13] Reduction in myelinated fibres was noted in 55.6% cases,^[16] unlike moderate-to-complete degree of myelinated fiber loss in all the cases in this study. Loss of myelinated fiber and Schwann cells is consistent with decrease in immunohistochemical staining of neurofilament in 100% HN biopsies, nerve growth factor receptor loss in 81.8%, PGP 9.5 staining loss in 100%, and loss of S-100 as well as myelin basic protein immunoreactivity in 90.9% cases each.^[18] Caseous necrosis was recorded in one case of TT (case-15; present study) as was reported by Hui *et al.*^[13]

Tests for Hansen's disease include slit-skin smears, skin and nerve biopsies, polymerase chain reaction (PCR),

Lepromin test, and serum assays for phenolic glycolipid 1 antibody (anti-PLG1).^[19] Anti-PLG1 titer correlates with bacterial load, being higher in lepromatous than TT cases.^[4] It can be useful for monitoring chemotherapy as titers correlate with the BI following treatment.^[20] Antunes *et al.* demonstrated lepra bacilli in 36.1% (52 out of 144) cases by Fite-Faraco stain. In the Lepra-negative cases, diagnosis of PNL was based on the PCR detection of *M. leprae* (28 out of 92 cases) and positivity of anti-PLG1 in the rest.^[17] In the present study, lepra bacilli were highlighted by Fite-Faraco stain in 44.4% cases, comparable to 47.8% in the study by Hui *et al.*^[13] Other ancillary tests were not done for the cases reviewed in our study.

CONCLUSION

HN is a debilitating but a completely treatable disease. Mononeuritis multiplex is the most common clinical presentation. Histological features such as mononuclear inflammation and foam cells in all compartments of the peripheral nerve, epithelioid granulomas, dense fibrosis, and less commonly perineurial thickening, with or without positivity for lepra bacilli are clues to diagnosis of HN. Knowing the morphological spectrum of HN is of cardinal importance because of the curable nature of the disease. Last but not the least, HN is one of the few neurological diseases with idiosyncratic clinical features and therefore a diligent neurological examination along with nerve biopsy examination is qualified for.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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Conflicts of interest

There are no conflicts of interest.

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