Abstract

Leprosy is the most ancient bacterial disease in the history of mankind. It remains still public health problem in the countries where it is endemic. Leprosy (Hansen’s disease) is a chronic infectious disease which is diagnosable and curable if recognized early and treated adequately. The infectious agent of leprosy is Mycobacterium leprae which is only bacteria to infect peripheral nerves. Leprosy is characterized by a variety of abnormal immune response. It depends on the host’s specific CMI response to M. leprae and it may be genetically determined. Antileprotic multidrugs therapy (MDT) as recommended by WHO is now the standard and accepted method for leprosy control. Leprosy can not be completely root out with physicians, control offices, leprosaria and propaganda; it will disappear when the economic and cultural factors change, because leprosy is the thermometer of civilization.

Key words: Leprosy, granulomatous diseases.

Introduction

Leprosy (Hansen’s disease) is a chronic infectious disease that primarily affects the peripheral nerves, skin, upper respiratory tract mucosa, eyes and certain other tissues. The causative agent is mycobacterium leprae which is first identified in 1873 by the Norwegian bacteriologist, Gerhard Henrik Armauer Hansen. It occurs at all ages but more common in between 10-20 and in 30-60 years. The ratio of male and female is 2:1 and common in India, Brazil, Indonesia, Myanmar, Madagascar, Nepal, Bangladesh, Thailand, Sub-Saharan Africa and Central America. The prevalence in Bangladesh is 0.29/10000 populations and endemic zones are: Nilphamari-1.61 per 10000 populations, Bandarban-1.58 per 10000 populations, Gaibandha-1.56 per 10000 populations and Khagrachari-1.32 per 10000 populations. It is not a hereditary disease. Infection may be genetically determined and monozygotic twins have 60-85% chance of leprosy. The disease is more common in low socio-economic status, overcrowding, poor nutrition and sanitation. The infectious agent of leprosy is Mycobacterium leprae which is obligate intracellular bacteria, gram positive, acid and alcohol fast, straight or slightly curved, rod shaped with parallel sides and rounded end, occur in clumps or globi or bundle of cigars in macrophages, only bacteria to infect peripheral nerves, best grow at below human core body temp 320c -350c. It can not grow in artificial culture media but may be cultivated in–nine banded armadillo, mouse foot pad, mangabey monkey. The mode of transmission is droplets infection, close contact with the patient for prolong period, ulcerated/abraded skin.

The incubation period is average 2-5 years. In tuberculoid leprosy it is up to 5 yrs and for lepromatous leprosy may be 20 yrs or longer.

Immunopathogenesis of leprosy

Leprosy is a spectrum disease characterized by a variety of abnormal immune response. It depends on the integrity of the host’s specific CMI response to M. leprae and it may be genetically determined. Whatever may the route of transmission, bacteremia occurs in all forms of leprosy and disseminated throughout the body via blood and lymphatics. M. leprae enter the target tissue, selectively schwann cells of peripheral nerves where it multiply and liberate from the infected schwann cells then infect neighbouring schwann cells and thus intraneural infection spreads. The infected nerve is then invaded by histocytes (macrophage) and lymphocytes with subsequent formation of granuloma leading to nerve damage resulting neurological manifestations. If the bodies CMI is capable of anchoring the infection within the nerves without skin involvement this will be limited to pure neural leprosy. But if bacilli or antigens escape from the nerve into surrounding tissue, skin lesions of TT develop at that site. Subsequently borderline leprosy and lepromatous leprosy may develop depending upon the immunity of the individual. Tuberculoid leprosy patient has strong T-cell and macrophage activation with release of type-1 cytokine (IL-2, IFN-Y and IL-12) which result in CMI response and to localize the infection. In lepromatous leprosy, release of type-2 cytokine (IL-4, IL-5, IL-10) cause strong antibody response and concomitantly inhibit T-cell and macrophage resulting in progression of the infection.

Figure 1: The Spectrum of Immunity in Leprosy.
Classification of leprosy

a. Ridlay & Jopling classification (on the basis of clinical, bacteriological, immunological & histopathological features):

1) Tuberculoid Leprosy (TT), (2)Borderline Tuberculoid (BT),(3) Borderline Bordline(BB), (4)Borderline Lepromatous (BL), (5)Lepromatous Leprosy (LL)

b. WHO classification (on the basis of BI/Mt):

(I) Paucibacillary leprosy: (a) Tuberculoid leprosy (TT), (b) Borderline tuberculoid leprosy(BT) (c)Indeterminate(I) (d)Pure neural leprosy (PN), where the skin smear is negative and/or the number of Skin lesion(s) are 1-5.

(II).Multibacillary leprosy (a)Borderline leprosy (BB), (b)Borderlinelepromatous(BL)

(c) Lepromatous leprosy (LL), Where the skin smear is positive and/or the number of skin lesion(s) are 6 or more. If a patient is clinically diagnosed as PB, but shows skin smear positivity he/she is classified as MB and given MB treatment7.

Evolution of leprosy lesion

Negative Contact (M.Leprae-)

Positive contact (M. leprae +)

Indeterminate pha s 2 8

Primary Neuritic (PN)

Tuberculoid (TT)

Borderline Lepromatous (LL)

BT

BB

BL

Spontaneous regression

* Usually an unstable muscle

Figure II: Evaluation of leprosy lesion

Clinical features

a. Indeterminate leprosy (I): It presents as single or multiple asymmetrical slightly hypo-pigmented or erythematous and usually ill-defined macules on skin, sensation on the affected area is normal or impaired while sweating and hair growth are usually unaffected. The peripheral nerves are not enlarged. Lepromin test may be either positive or negative. It is usually self healing but it may progress to other forms of leprosy7.

b. Tuberculoid leprosy (TT): It is usually single or multiple (Less than 5) asymmetrical hypo-pigmented or erythematous, oval or round macules, patches and plaques with a sharply defined elevated border that slopes down to a flattened atrophic center having the appearance of “a saucer right side up”. Sensation is markedly impaired or lost, sweating and hair growths are also impaired. Nerve involvement is early and prominent. Peripheral nerves are usually cord like, hard, enlarged and tender. Nerve abscess may occur. Damage to nerves may result in loss of sensation, pain, tingling and muscle weakness or paralysis resulting wrist drop, foot drop, and lagophthalmos. Lepromin test is strongly positive8.

c. Borderline tuberculoid leprosy (BT): The lesions are similar to tuberculoid leprosy except these are smaller and more numerous. Satellite lesions around large macules or plaques are characteristic3.

d. Borderline leprosy (BB): It is unstable. The lesions are generalized, asymmetrical numerous, hypopigmented or erythematous, irregular shaped plaques. Sensation, sweating and hair growth over the lesions are impaired1.

e. Borderline lepromatous leprosy (BL): The lesions are roughly symmetrical, numerous hypopigmented or erythematous macules, papules, plaques or nodules. Sensation and sweating over the lesion–normal. Peripheral nerve involvements appear later, mucosal and systemic involvement usually absent5.

f. Lepromatous leprosy ( LL)

(I) skin: The lesions are very numerous symmetrically distributed erythematous or coppery shiny macules, papules and nodules. The patient may have leonine face, loss of eyebrow and eyelashes (Madarosis). Lepromin test is negative.

(II) Nerve involvement: Involvement is bilaterally symmetrical, usually in a glove-stocking pattern. In advanced cases nerves become thin and hard due to fibrosis and resulting extensive anesthesia. Nerve changes are-

- Sensory - Anaesthesia
- Motor - Paralysis and wasting of muscles
- Autonomic - Loss of sweating and vasomotor dysfunction

(IV) Changes in Eye: Corneal erosion, exposure keratitis, ulceration, corneal opacity, vascular keratitis, interstitial keratitis, pannus formation, acute corneal leproma, iritis and iridocyclitis.

(V) Visceral / systemic involvement: Except for the GIT, lungs & brain virtually every organ may be involved. Lymph nodes, liver, bone marrow, spleen, testicles are heavily infected. Bone marrow of the phalanges is destroyed resulting shortening, osteoporosis and the bone fractures easily. Testes are most commonly & severely affected resulting testicular atrophy, gynecostasia, azoospermia and sterility. Glomerulonephritis, secondary amyloidosis may occur. Both smooth & striated muscles are invaded.

g. Special forms of leprosy

(I) Histoid leprosy: Uncommon form of multibacillary leprosy. May found in patient who have discontinued the treatment or leprosy bacilli have become drug-resistant. Erythematous, round or oval, shiny, large papules or plaques and glistening nodules on skin and subcutaneous tissue. Site- Buttocks, lower back, face, and bony prominence. Biopsy shows- elongated or spindle shaped histocytes containing bacilli and showing a whorled arrangement.

(II) Pure neuritic leprosy: Typical nerve damage occur without skin lesion. Ulnar nerve mostly affected. Histology most commonly shows the feature of TT or BL but commonly BT. Sensory change earlier than motor.

(III) Lucio leprosy: This is the special form of lepromatous leprosy found in Western Mexico and certain Latin American areas. It is characterized by a diffuse widespread infiltration of skin (without formation of nodules), loss of body hair, eye brow, eye lashes and wide spread sensory loss. It may smooth the facial wrinkle of older patient resulting youthful appearance, some times called “lepra Bonita” or pretty leprosy.

(IV) Drug resistance leprosy: It is characterized by progressive deterioration in the clinical and bacteriological status of patients inspite of treatment. Two types: Primary resistance occurs in persons who are infected by drug resistant M-leprae. May occurs in all types of leprosy. Secondary resistance-develops as a result of long term monotherapy and it is usually confined to LL.

Lepra reactions

It is a tissue destructive, immunologically driven, inflammatory process that may occur in patient of leprosy before, during or after institution of therapy. The triggering factors are institution of multidrug therapy (50% patient.), intercurrent infections (viral, malaria), anaemia, vaccination, pregnancy and parturition, puberty, drugs-vitamin A, iodides, and bromide, surgical intervention, mental and/ or physical stress. Types of lepra reaction are Type- I Lepra reaction (reversal or downgrading reactions) and Type-II Lepra reaction (Erythema Nodosum Leprosum).

Type I Lepra reaction (Reversal and Downgrading reaction) :
It is associated with alternation of cell-mediated immune reaction. When cell mediated immunity increase called reversal reaction and when decrease called downgrading reaction. It is a type-IV hypersensitivity reaction usually occurs within 06 months of treatment. It occurs in a borderline spectrum i.e. BT, BB and BL. Some or all the existing leprosy lesions develop pain or tenderness, erythema, oedema, look like erysipela. Necrosis and ulceration may occur. Affected nerve become rapidly swollen, extremely painful and tender. Nerve abscess may develop. Loss of motor function ie, claw hand, foot drop, facial palsy may occur suddenly. Untreated cases develop permanent paralysis.

Type II Lepra reaction (Erythema Nodosum Leprosum): It is type III hypersensitivity reaction. It occurs almost exclusively in LL, occasionally in BL cases. Existing lesions do not show clinical aggravation. Sudden appearance of crops of evanescent, pink coloured tender nodules or plaques. May become vesicular, purulent, bullous, gangrenous and undergo breakdown. Constitutional symptoms like fever, headache, malaise may develop. Patient may be toxic. It may be associated with- iritis, iridocyclitis, epistaxis, muscle, bone and nerve pain. Epididymo-orchitis, lymphadenitis and proteinuria may develop.

Investigations of leprosy

a. Blood for complete picture – Anaemia, lymphopenia and increased ESR.

b. Slit skin smear for acid fast bacillus(AFB): Stained films revealed that there were large number of acid and alcohol fast bacilli arranged in straight and curved parallel bundles with globular masses (cigar-bundle appearance), morphologically resembling Mycobacterium leprae.

C. Skin biopsy for histopathological examination and Fite-Faraco stain: Section from skin revealed extensive infiltration of macrophage in the dermis, separated from epidermis by narrow Grenz zone, with destruction of skin adnexa. Few foci of poorly defined granuloma in dermis is also noted. Large number of lepra bacilli were seen in Wade- Fite stained sections.
**Complications of leprosy**

Due to nerve damages-claw hand, wrist drop, foot drop, lagophthalmos and due to massive invasion of tissues by M. leprae-iritis, iridocyclitis, testicular atrophy, gynaecomastia, epididymoorchitis, leonine facies and saddle nose. Other complication is malum performance pedis.

**Prevention & control**


b. Immunoprophylaxis by BCG vaccination and a heat killed M. leprae plus BCG vaccination.

c. Improvement of living condition.

d. Health education and propaganda about leprosy.

**Rehabilitation**

To reintegration of the leprosy patient into the community, taking part in all its activities as a contributing member of the society.

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**References**


