Abstract

Tuberculosis (TB) of the head and neck can be contained in the lymph nodes, larynx, oropharynx, salivary glands, nose and paranasal sinuses, ear; skin and skull. Head and neck TB presentations are varied in nature and thus difficult to diagnose. The clinical features, radiological findings, microbiological diagnostic modalities, surgical and medical management and outcomes of nine cases of head and neck TB are discussed in detail here, together with a thorough review of the literature. Patients presented with atypical symptoms such as discharging sinus, ear lobule swelling, otitis media, vision loss and facial weakness, long refractory otorhoea and granulation tissue in the ear canal. We diagnosed tubercular skull base osteomyelitis (one case) and laryngeal tuberculosis (two cases), mastoid tuberculosis (one case) and non-tubercular mycobacterial infection involving the temporal bone (two cases), sino-nasal region (one case), maxilla (one cases) and ear lobule (one case) over a period of 8 months. All patients were managed successfully with a combination of surgery and a well-planned treatment regimen for non-tuberculous mycobacteria (NTM) or anti-tubercular drugs for TB. All had successful outcomes except one patient with tubercular skull base osteomyelitis who expired before the initiation of anti-tubercular therapy (ATT). High clinical suspicion followed by thorough diagnostic work-up for both TB and NTM would enable early diagnosis and complete treatment.

INTRODUCTION

According to the World Health Organization (WHO) Global TB report, approximately 10 million people developed tuberculosis (TB) in 2020 and 1.4 million died in 2019 [1]. There were an estimated 1.2 million TB deaths among HIV-negative people in 2019 and an additional 208 000 deaths among HIV-positive people. South-East Asia accounts for 44% of cases globally and India accounts for 26% of the burden of TB cases. Two million four hundred thousand TB cases were incident TB cases (new and relapse/recurrent) from India. Tuberculosis affects the pulmonary system (80% of cases) and extrapulmonary sites in 20% cases. Head and neck TB constitutes 10–15% of all EPTB cases, with the majority of these cases being cervical lymph node TB, and <1% extra nodal head and neck TB cases. Head and neck TB is caused by Mycobacterium tuberculosis (MTB; most common) and non-tubercular mycobacteria (NTM; rare). Head and neck TB cases have been reported to involve the cervical lymph nodes, middle ear, larynx, pharynx, paranasal sinus, tonsils and salivary glands [2–4]. The commonest presentation of head and neck TB is tubercular cervical lymphadenopathy. With the lungs being the primary focus, the lymph nodes are involved and a subsequent cough results in the secondary involvement of the larynx, oropharynx and para nasal sinuses [2, 3]. Otomastoiditis due to NTM is extremely rare and has only been described in 56 well-documented cases in the literature in English [2, 5]. Predisposing factors for head and neck TB have been described as poor oral hygiene, local trauma, pre-existing oral leukoplakia, periapical granuloma, periodontitis and nutritional deficiency due to inadequate intake [3]. Clinical presentations of head and neck TB strongly mimic common ENT diseases, with occasional subtle atypical symptoms and signs, which, if missed, lead to misdiagnosis and treatment failure apart from disease progression. Unless strongly suspected, diagnosis of these cases is challenging. We intend to present nine cases of head and neck involvement, including only one case secondary to primary pulmonary TB. Clinical presentation, microbiological diagnostic modalities and treatment outcomes are discussed in detail, together with a thorough review of the literature.
SUBJECTS AND METHODS
This case series includes patients presenting to the Department of Otorhinolaryngology who were suspected to have tubercular involvement of the head and neck, which was confirmed by the mycobacteriology laboratory in the Department of Microbiology at the All India Institute of Medical Sciences. Detailed demographic findings, unusual symptoms/signs, radiological and histopathological investigations and microbiological tests used for confirmation of diagnosis, presence of primary pulmonary focus, treatment received and treatment response were noted.

Radiological investigations
All patients underwent thin-section computed tomography (CT) or magnetic resonance imaging (MRI) to evaluate the middle ear and mastoid and temporal bones. Imaging features, including the presence of soft tissue in the middle ear, the integrity of the ossicles and mastoid bone destruction, were analysed and compared with the surgical findings.

Microbiological investigations
Various samples, such as sputum, pus aspirates and tissue biopsy were collected, depending on the site and the lesion affected in suspected head and neck TB cases. Samples for microscopy and culture were processed using the N-acetyl-L-cysteine-sodium hydroxide method. Decontaminated samples were stained using the Ziehl–Neelsen (ZN) technique and microscopy was performed and graded according to RNTCP guidelines, i.e. scanty (1–9/100 fields), 1+ (10–99/100 fields), 2+ (1–10/field) and 3+ (10/field). Culture was performed using the automated liquid MGIT 960 culture system (BD). Positive cultures were subjected to differentiation into MTB complex and NTM (MGIT TBc Identification test; BD). Positive liquid culture was subjected to MGIT 960 liquid culture drug susceptibility testing. Samples were subjected to molecular detection of TB and rifampicin resistance using the GeneXpert MTB/RIF (Cepheid, Inc.) system and TrueNat MTB-RIF (MolBio, India) in accordance with the manufacturers’ instructions.

The Truenat MTB/NTM assay is a chip-based real-time polymerase chain reaction test for the semiquantitative detection and differentiation of MTB from NTM in human pulmonary and EPTB specimens. Truenat MTB/NTM runs on Truelab Real Time micro-PCR Analyzers. The target genes selected in this assay are the rpoB and nrdz genes. The rpoB gene is a common gene that is present in both MTB and NTM, and its will indicate the presence of total mycobacteria in a given sample. The nrdz only specific for MTB and is not present in NTM. The Truenat MTB/NTM test can detect a diverse range of NTM strains. The following NTM strains can be differentiated by the Truenat MTB/NTM assay: *M. avium*, *M. malmoense*, *M. scrofulaceum*, *M. ulcerans*, *M. abscessus*, *M. fortuitum*, *M. gordonae*, *M. szulgai*, *M. kansasi*, *M. asiaticum*, *M. celatum*, *M. simiae*, *M. triviale*, *M. terrae*, *M. flavescens*, *M. haemophilum*, *M. thermoresistibile*, *M. marinum*, *M. xenopi*, *M. vaccae*, *M. chelonae*, *M. smegmatis* and *M. intracellulare*.

RESULTS
A total of nine cases were included over a period of 8 months. Their demographic and clinical details are described in Table 1. Eight patients were adults with an age range from 25 to 85 years and one patient was a 6-year-old child. Seven patients were male and two were female. Two patients had co-morbid conditions; one was on anti-retroviral treatment and the other had diabetes mellitus with hypertension. Patients presented with atypical symptoms such as discharging sinus, ear lobe swelling, otitis media, vision loss and facial weakness, long refractory otorrhoea and granulation tissue in the ear canal. One patient had complained of right vision loss, proptosis and headache for 6 months. One of these nine patients had coexistent pulmonary tuberculosis. None of the patients had a significant past or family history. All patients were suffering for a long span of time and were non-responsive to broad-spectrum antibiotics. External auditory canal, laryngoscopy and audiometry examination showed atypical features that were consistent with tuberculosis or neoplasms. Radiological investigations showed typical features of osteomyelitis of the temporal bone and skull base, and others, such as destruction of bone, increased bone uptake and densities, which needed further evaluation (Table 2, Fig. 1). Histopathological evaluation revealed granuloma in few cases and atypical features in some, suggesting neoplasm. We were able to diagnose involvement of the temporal bone (two cases), tubercular skull base osteomyelitis (one case), mastoid bone (one cases), larynx (two case), sino-nasal (one case), maxillary sinus (one cases) and ear lobe (one case). Microbiological examination of samples confirmed the cases (Table 2). All samples were acid-fast bacilli-positive (ZN staining), while GeneXpert was positive in four cases for MTB and TrueNat was positive in all nine cases and detected MTB (four cases) and NTM (five cases) (Table 2). All patients were managed successfully with a combination of surgical and combination regimens including antibiotics (NTM cases) and anti-tubercular drugs, except one tubercular skull base osteomyelitis patient, who expired before the initiation of anti-tubercular therapy (ATT) (Table 3).

DISCUSSION
Ample published literature has discussed and described head and neck TB [2]. Non-tuberculous mycobacteria (NTM) have also been reported to cause head and neck involvement. NTM are ubiquitous micro-organisms in natural and constructed environments. NTM are known to cause a broad spectrum of diseases, including lymphadenitis and skin, soft tissue, ocular, abdominal, pulmonary, wound and disseminated infections. Tubercular cervical lymphadenopathy is the commonest presentation and is characterized by sub-acute to chronic or recurrent lymphadenopathy, usually bilateral, painless, with or without intermittently discharging sinuses leaving scars [2]. Common presented symptoms include low-grade fever episodes, loss of appetite, weight loss and occasionally history of close contact with patients of TB. Most of the head and neck TB cases have evidence of pulmonary disease and secondary
infection in the larynx and nose as well as paranasal sinuses through coughing and sneezing infected material through [2–4]. Secondary aural infection occurs through seeding of the middle ear by nasopharyngeal spread or by the passage of infected material up the auditory tube through coughing. Haematogenous implantation can lead to the spread of mycobacterial infection, potentially to any site of the body, including the head and neck. Pulmonary or systemic TB must be investigated to exclude it from head and neck TB patients. In our case series, only one laryngeal tuberculosis patient had co-existing active pulmonary tuberculosis, a direct inoculation of mycobacterium from the air inhaled through the eustachian tube or haematogenous inoculation could be some of the possible routes of infection. A healed primary pulmonary focus (with so many commonly prescribed antibiotics being effective against TB) but persistent bony involvement in temporal bone or a reactivation of an old healed focus in the airways and localizing in the head and neck region are a few other possibilities. The site of localization of the disease and hence the disease presentation could vary. Although there is more awareness of pulmonary TB, the importance of suspecting the disease in the head and neck airways needs to be understood.

In a retrospective analysis of 117 patients with primary TB of head and neck, Bhat et al., reported 95 % patients with cervical lymphadenopathy, two cases with laryngeal TB, and one each with TB of cervical spine, oropharynx, ear and retropharyngeal abscess [2].

### Table 1. Demographic and clinical details of all the patients

| Case | Age/sex | Comorbidity | Past H/O of PTB | Complaints | Clinical diagnosis | Clinical features/Atypical if any | Final diagnosis |
|------|---------|-------------|----------------|------------|-------------------|---------------------------------|----------------|
| 1    | 46/M    | AIDS (on ART) | No             | Left cheek discharging sinus since 3 years | Left maxillary sinus osteomyelitis | Absence of history of trauma/dental extraction, no H/O fever | Maxillary tuberculosis |
| 2    | 42/M    | No          | No             | Left ear lobe swelling for 6 months | Benign cyst | Cystic swelling with no discharge | Ear lobe tuberculosis |
| 3    | 82/M    | DM and HTN  | No             | Left ear ache, ipsilateral headache for 3 months, left facial weakness | Skull base osteomyelitis | Congested but intact tympanic membrane, grade 3 left facial weakness | Tubercular skull base osteomyelitis |
| 4    | 63/M    | No          | No             | Right vision loss and proptosis × 6 months; headache × 8 months | Right sphenoid sinus mucocele | Right visual acuity 6/60 with proptosis | Sinonasal tuberculosis |
| 5    | 29/F    | No          | No             | Right ear discharge × 3 months, conductive hearing loss (50/20), facial weakness × 2 months, H/O on-off immutus and vertigo | Complicated supplicative otitis media | Right tympanic membrane reveals oedematous external auditory canal and mucopurulent discharge with bulge in posterosuperior quadrant; pure tone audiometry revealed right side severe conducting hearing loss with normal left ear; facial weakness grade 5 laryngoscopy WNL | Temporal bone tuberculosis |
| 6    | 40/M    | No          | No             | Hoarseness × 3 months associated with H/O productive cough | Laryngeal growth with pulmonary tuberculosis | Bilateral true vocal cord and arytenoid irregularity; no H/O smoking / fever/weight loss/fuss of appetite | Laryngeal and pulmonary tuberculosis |
| 7    | 48/F    | No          | No             | Left ear discharge for 3 months (H/O MRM done 1 month back), diplopia for 20 days | Left complicated chronic otitis media with Gradenigo’s syndrome | Florid pale granulations in post-operative mastoid cavity, left mixed hearing loss (55/25), left sixth nerve paresis | Temporal bone tuberculosis |
| 8    | 42/M    | DM          | No             | Hoarseness of voice × 1 year associated with dry cough | Laryngeal TB | Mouse nibbled appearance and ulceration in post-glottis | Laryngeal tuberculosis |
| 9    | 6/M     | No          | No             | Left ear discharge for 3 months | B/l mastoiditis | Granulation tissue on post-EAC wall | Mastoid tuberculosis |

AIDS, acquired immunodeficiency syndrome; ART, anti retroviral therapy; DM, diabetes mellitus; EAC, external auditory canal; H/O, history of; HTN, hypertension; MRM, modified radical mastoidectomy; WNL, within normal limits.
| Case     | Clinical diagnosis                              | Radiological investigations                          | Sample                      | Histopathological investigations                                                                 | Microbiological investigations                                                                 | Organism isolated |
|----------|------------------------------------------------|-----------------------------------------------------|-----------------------------|---------------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------|-------------------|
| Case 1   | Left maxillary sinus tubercular osteomyelitis  | NCCT PNS – destruction of left maxilla; PETCT – infiltrative lesion of left maxillary bone (Fig. 1 a) | Biopsy from maxillary sinus | Markedly hyperplastic stratified squamous epithelium with marked papillomatous and superficial ulcerations; epithelium shows focal mild nuclear atypia | ZN smear – scanty 5–6 bacilli/100 oif; Gene Xpert – negative; TRUNAT- NTM MGIT culture – negative | NTM               |
| Case 2   | Benign cyst of ear lobule                       | None; chest X-ray – WNL                              | Excision biopsy             | Nodular lesion shows acanthotic epidermis; mild chronic inflammatory infiltrate is noted          | ZN smear – scanty 4 bacilli/100 oif; Gene Xpert- – negative; TRUNAT-NTM MGIT culture – negative | NTM               |
| Case 3   | Skull base osteomyelitis                        | HRCT temporal bone – soft tissue density – middle ear, mastoid with areas of lysis and sclerosis on mandibular condyle. (Fig. 1 b) | Myringotomy done – biopsy from hypertrophic middle ear mucosa | None                                                                                             | ZN smear – 1+; Gene Xpert – not done; TRUNAT-MTB MGIT culture – positive after 3 weeks | MTB               |
| Case 4   | Right sphenoid ethmoid mucocoele               | CECT PNS – expansile capsulated lesion involving right posterior ethmoid and sphenoid causing displacement of globe | Endoscopic drainage and tissue biopsy | Inflammatory aetiology                                                                         | ZN smear – scanty 4 bacilli/100 oif; Gene Xpert – negative; TRUNAT-NTM MGIT culture – negative | NTM               |
| Case 5   | Complicated suppurative otitis media            | HRCT temporal bone – STD filling middle ear/ mastoid; ((Fig. 1 c) bone scan – uptake in right temporal bone; laryngoscopy – WNL | Biopsy sample               | Inflammatory aetiology                                                                         | ZN smear – scanty 3 bacilli/100 oif Gene Xpert – negative; TRUNAT-NTM MGIT culture – negative | NTM               |
| Case 6   | Laryngeal and pulmonary tuberculosis           | X-ray chest – left upper lobe consolidation; laryngoscopic examination reveals bilateral irregular cords with granular lesion | Laryngeal biopsy and sputum sample | Direct laryngoscopic biopsy showed epithelioid cell granuloma with langhan type giant cell consistent with tuberculosis | ZN smear – scanty 2–3 bacilli/100 oif; Gene Xpert – positive; TRUNAT- MTB MGIT culture – negative | MTB               |
| Case 7   | Left gradenigo syndrome                        | HRCT temporal bone and CEMRI shows post-op cavity with destruction reaching petrous apex with enhancement along Meckel's cave and clival region; (Fig. 1 d) X-ray chest – WNL | Biopsy from cavity granulation | Inflammatory granulation tissue and occasional bone fragments, no necrosis or dysplasia seen | ZN smear – scanty 3 bacilli/100 oif; Gene Xpert – negative; TRUNAT- NTM MGIT culture – negative | NTM               |
| Case 8   | Laryngeal TB                                   | CECT neck shows diffuse mucosal thickening in B/L maxillary sinus and ethmoid sinus | Biopsy from larynx          | Poorly differentiated malignant tumour S/O non-Hodgkin lymphoma                                  | ZN smear – scanty; Gene Xpert – positive; TRUNAT – MTB; MGIT culture – negative               | MTB               |

Continued
cervical lymphadenopathy. Instead, we found temporal bone TB (two cases), tubercular skull base osteomyelitis (one case), laryngeal TB (one case), sino-nasal TB (two cases) and ear lobe TB (one case).

Tuberculosis of the middle ear and mastoid presents with persistent painless otorrhoea, otalgia, hearing loss and, in extreme cases, facial palsy. TOM (tuberculous otitis media) has a typical presentation of multiple tympanic membrane perforations [4, 6]. Thin mucoid or sanguineous otorrhoea and abundant pale polypoid tissue have been seen during otological examination [4, 6]. Radiology shows a background of pneumatized mastoid and features of bony involvement and sequestrum formation. Tubercular otitis media/mastoiditis should be suspected when signs suggestive of extensive disease as compared to minimal symptoms. Biopsies from the middle ear or mastoid granulations are supportive for confirming the diagnosis.

Tuberculosis of the nasal and the paranasal sinuses presents with non-specific symptoms such as nasal obstruction and blood-stained rhinorrhoea and headache, with mimicking as sinusitis and neoplastic lesions. Tubercular suspicion becomes stronger when signs of recurrent discharging sinuses over the cheek or zygomatic area are evident [5, 7]. Detailed clinical endoscopic examination and biopsies from the pale granular

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**Table 2.** Continued

| Case | Clinical diagnosis             | Radiological investigations | Sample          | Histopathological investigations | Microbiological investigations | Organism isolated |
|------|--------------------------------|----------------------------|-----------------|---------------------------------|-------------------------------|-------------------|
| Case 9 | Mastoid tuberculosis | No | Pus from ear | No | ZN smear – scanty 2–3 bacilli/100 oif; Gene Xpert – positive; TRUNAT – MTB; MGIT culture – negative | MTB |

B/L, bilateral; CEMRI, contrast enhanced magnetic resonance imaging; HRCT, high resonance computed tomography; MGIT, mycobacterium growth indicator tube; MTB, *Mycobacterium tuberculosis*; NTM, non-tubercular mycobacteria; S/O, suggestive of; STD, soft tissue densities; WNL, within limits; ZN, Ziehl–Neelsen.

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**Fig. 1.** (a) A CT scan of case 1 shows infiltrative lesion of the left maxillary bone. (b) A CT scan of case 3 shows temporal bone soft tissue density – middle ear, mastoid with area of lysis and sclerosis on mandibular condyle. (c) A CT scan of case 5 shows temporal bone soft tissue density filling middle ear and mastoid. (d) A CT scan of case 7 shows post-operative cavity with destruction reaching petrous apex with enhancement along Meckel’s cave and cival region.
tissue are the steps towards making a diagnosis. One case of maxillary tuberculosis presented with persistent discharging sinus for 3 years. The patient was known to have AIDS and was on antiretroviral therapy. One case of sino-nasal tuberculosis presented with atypical features, such as right-sided vision loss (visual acuity, 6/60) with proptosis for 6 months and headache for 8 months.

The case of ear lobule tuberculosis had very unusual features, and presented with a benign cyst of the ear lobule without

### Table 3. Medical and surgical management and outcome for all patients

| Serial no. | Age/sex | Final diagnosis                      | Medical treatment                          | Surgical treatment                          | Response to treatment                                      |
|------------|---------|--------------------------------------|--------------------------------------------|--------------------------------------------|----------------------------------------------------------|
| 1          | 46/M    | Maxillary tuberculosis               | ATT CATEGORY 1 9 months                    | Surgical debridement from maxillary sinus | Cured; Healed scar present over left cheek; Follow-up done for 18 months; no discharge |
| 2          | 42/M    | Ear lobule tuberculosis              | ATT CATEGORY 1 6 months                    | Excision biopsy with primary suturing      | Cured; Follow-up done for 7 months; Well-healed local site |
| 3          | 82/M    | Tubercular skull base osteomyelitis  | Patient expired before start of ATT due to diabetic nephropathy | Myringotomy done under LA                  | Expired                                                  |
| 4          | 63/M    | Sinonasal tuberculosis               | ATT CATEGORY 1 6 months                    | Endoscopic drainage and tissue debridement | Cured; Right eye vision improved (6/18); No proptosis; Well-healed sinonasal cavity |
| 5          | 29/F    | Temporal bone tuberculosis           | IV ceftazidime and amikacin ×4 days        | Cortical mastoidectomy with tympanoplasty placement | Cured; No ear discharge for 9 months; Facial nerve function improved – grade 3, intact Tympanic membrane |
| 6          | 40/M    | Laryngeal and pulmonary tuberculosis| ATT CATEGORY 1 6 months                    | Direct laryngoscopic biopsy.              | Cured; Follow-up for 1 year; Voice quality improved markedly; Laryngeal evaluation reveals no lesions over cords; only mild congestion seen |
| 7          | 48/F    | Temporal bone tuberculosis           | ATT CATEGORY 1 with tab clarithromycin 500 mg BD 12 months | Cortical mastoidectomy with tympanoplasty placement | Cured; Lateral rectus function improvement after 2 weeks of ATT; Well-healed mastoid cavity with no granulations at present |
| 8          | 42/M    | Laryngeal TB                         | ATT CATEGORY 1 6 months                    | Direct laryngoscopic biopsy              | Cured; Follow-up for 1 year; Voice quality improved markedly |
| 9          | 6/M     | Mastoid tuberculosis                 | Tab rifacept – kid 3 od Tab ethambutol 400 mg od | No                                        | Cured; follow-up is ongoing                              |

ATT, anti tubercular treatment; BD, twice a day; I/V, intravenous; OD, once a day.
discharge for 6 months or any history of trauma or ear piercing. A case report involving a 62-year-old woman with right pinna tuberculosis, diagnosed using histopathology and confirmed using microbiological methods, was treated successfully with ATT [7]. Another case report of Tuberculosis verrucosa cutis is recorded for a young woman with keloid over ear lobule (the keloid increased in size over 4 months) [8]. We diagnosed an unusual causative organism in our patient, an NTM identified as acid-fast bacilli microscopically and confirmed by Truernat. This patient was treated with a combination regimen including several antibiotics.

Laryngeal tuberculosis has been reported to be the second most commonly reported TB in the head and neck region followed by cervical lymphadenitis. Co-infection of pulmonary tuberculosis is found in ~1% of laryngeal tuberculosis cases. Laryngeal TB is commonly associated with an undiagnosed, inadequately treated or recurrent pulmonary tuberculosis. Our laryngeal TB case also had a co-existing active primary pulmonary tuberculosis. Presenting symptoms include dysphonia, cough, odynophagia, dyspnoea, dysphagia, hoarseness of voice and weight loss [9]. Patients undergoing direct laryngoscopy and biopsy have laryngeal TB detected in 6.25% of cases. Indirect laryngoscopy/fibroptic laryngoscopic examination commonly reveals diffuse pale and polypoidal thickening of the epiglottis, vocal cords and arytenoids, closely mimicking neoplastic lesions and requires pathological as well as microbiological evaluation to confirm the diagnosis [10]. The laryngeal tuberculosis case in our series presented with hoarseness of voice in the last 3 months, associated with a productive cough.

Skull base osteomyelitis (SBO) has been reported as a complication of otitis externa due to Pseudomonas aeruginosa in older diabetic patients and most commonly involves the temporal bone. Atypical or central skull base osteomyelitis arises from the sphenoid or occipital bones rather than the temporal bone and is not usually associated with otitis externa [11, 12]. Incidence of atypical SBO has been reported much less frequently and initially presents with headache as the only symptom, with cranial neuropathies occurring in the later course of disease. The published literature reveals that small-vessel disease of the external auditory canal (EAC) and an increase in the cerumen pH in diabetic persons can predispose patients to a rapid and at times devastating disease course [13]. When infection traverses the fissures of Santorini and the tympanomastoid suture to the Haversian system of the compact bone, and finally to the skull base, then SBO can lead to cranial nerve (CN) palsies. CN VII is most commonly affected, due to the proximity to the EAC on exiting the skull base, followed by CNs IX, X, XI and XII [14, 15]. Our case was a classic case of tubercular skull base osteomyelitis in a 82-year-old diabetic and hypertensive who presented with otorrhea, ear ache and headache with grade 5 left facial weakness and succumbed to the disease before the initiation of antitubercular therapy.

Computed tomography is helpful in making a diagnosis and to evaluate the extent and severity of extratemporal soft tissue involvement and skull base osteomyelitis. Soft tissue inflammation, usually seen/limited in the early stages of disease and advanced disease, involves skull base bone destruction and abscess formation. The HRCT findings in our case were consistent with those reported in advanced disease [14].

Temporal bone osteomyelitis (TBO) occurs as a complication from necrotizing external otitis media (NEO) and middle ear infection. TBO is a rare but very aggressive disease with different aetiologies [16]. TBO predisposes patients to clinical conditions such as trauma, bone surgery, or other diseases that can affect the vascularity of bone (osteoporosis, osteo- petrosis, Paget's disease, radiation and malignancy) [17]. Facial neuropathy is most commonly reported and facial paralysis occurs in 25% of patients due to the involvement of the stylomastoid foramen [18, 19]. A case of local temporal bone osteomyelitis caused by granulomatous tuberculous infection has been reported [20]. Two of our patients were suffering from complicated suppurative chronic otitis media and presented with otorrhea, conductive hearing loss, facial weakness and diplopia (in one case). The radiological findings in both the cases confirmed them as temporal bone osteomyelitis and microbiological examination detected NTM in both cases, guiding treatment decisions (as per the ATS guidelines, 2007) [23] and securing a favourable outcome. Yeh et al. discussed the fact that NTM otomastoiditis with TBO should be suspected in cases of chronic otorrhea, otalgia and granulation tissue in the ear [21].

Few cases of NTM otomastoiditis have been reported in the published literature. A retrospective case series has reported the rates of granulation tissue (90.9%), otalgia (31.8%), facial palsy (9.1%) and lung involvement (5.4%). Cases of temporal bone osteomyelitis have been reported, most commonly with the NTM M. abscessus (52.7% of cases) and M. avium-intracellulare, followed by M. fortuitum [21–23]. Three distinct routes have been suggested for the entry of NTM into the middle ear: direct inoculation through perforation of the tympanic membrane or a tympanostomy tube (most common), through the eustachian tube, and direct spread via improperly sterilized surgical instruments [22, 23]. We detected five patients with NTM. Two patients had NTM temporal bone osteomyelitis, and had presented with otalgia, otorrhea, facial palsy and granulation tissue and previous history of tympanic membrane perforation. Gardenigo syndrome caused by NTM has also been reported [24]. Our case fitted the definition of Gardenigo syndrome, diagnosed as temporal bone osteomyelitis caused by NTM.

Head and neck TB (MTB and NTM) remains a diagnostic and therapeutic challenge for clinicians due to varied presentation. Multiple diagnostic modalities aided to the accurate diagnosis and treatment. The published literature and ATS guidelines suggested a macrolide drug (clarithromycin) to be the most appropriate initial regimen with a combination of parenteral amikacin and/or imipenem for treatment of NTM [23]. The duration of antibiotic use depends on response to treatment for each case, but is at least 6 months with careful monitoring, and longer if required. Long-term multiple antibiotic therapy
is advised for at least 1 to 3 months after the symptoms have resolved. NTM have a tendency to reproduce and exist in biofilms, which makes them difficult to destroy [21, 23]. Surgical debridement or even repeated surgery with antibiotic therapy to remove the NTM and the surrounding biofilms is required to accomplish infection control and efficiently manage such infections.

**CONCLUSION**

Clinicians should maintain a high suspicion for head and neck TB. Unusual presentations such as refractory otorrhea, granulation tissue in the middle ear cavity and mastoid air cells, or cortical bone or ossicular chain destruction on imaging without evidence of a cholesteatoma suggests a need for head and neck TB work-up. Whenever accurate diagnosis has been confirmed, surgical and medical intervention (well-designed regimen) should be implemented and a symptom-free course should be ensured before antibiotics are stopped.

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**Author contributions**

Study design and manuscript writing: U.B.S. and K.B. Data acquisition and data analysis: S.K. and R.M. Data interpretation and literature search: P.S. and K.B.T. Figure design and composition: K.B. and S.K. Critical analysis and manuscript drafting and verification of data: A.T., S.C.S., U.B.S.

**Conflicts of interest**

The authors declare that there are no conflicts of interest.

**Ethical statement**

Ethical approval has been received (ref. no. IEC – 248/09.04.2021, RP-06/2021) from the Institute Ethics Committee.

**References**

1. World Health Organization. Global tuberculosis report. 2020. https://www.who.int/publications/i/item/9789240013131
2. Nalini B, Vinayak S. Tuberculosis in ear, nose, and throat practice: its presentation and diagnosis. American J Otolaryngology. 2006;27:39–45.
3. Lmekki S, Lecanu JB. Tuberculosis of the middle ear and nasal passage: A case report. Inter J Mycobact. 2013;2:51–53.
4. Cho Y-S, Lee H-S, Kim S-W, Chung K-H, Lee D-K, et al. Tuberculous otitis media: a clinical and radiologic analysis of 52 patients. Laryngoscope. 2006;116:921–927.
5. Linman S, Stokroos RJ, Linssen CFM. Mycobacterium abscessus, an uncommon cause of chronic otitis media: a case report and literature review. Arch Otolaryngol Head Neck Surg. 2008;134:1004–1006.
6. Mohamad Yusof AN, Mohamad I, Haron A, Salahuddin Z. Unusual presentations of tuberculosis – A case series. Egyptian J Ear, Nose, Throat and Allied Sciences. 2016;17:155–158.
7. Basal Y, Ermishe B, Eryilmaz A, Erçuğur B. Two rare cases of head and neck tuberculosis. BMJ Case Rep 2015;2015:bcr2015211897.
8. Halim MS, Ahmed SQ, Junaid M, Bashir MR. Tuberculosis verrucosa cutis in a patient with keloid over ear lobe. BMJ Case Rep 2013;2013:bcr2013010294.
9. Felemban T, Ashi A, Sindi A, Rajab M, Jehani ZA. Hoarseness of voice as a rare presentation of tuberculosis: a case report study. Open Access Maced J Med Sci. 2019;7:3262–3264.
10. Matimba A, Moncho M, Musoke J, Seedat RY. Diagnosis of laryngeal tuberculosis in a high TB burden area. Eur Arch Otorhinolaryngol. 2020;277:2137–2140.
11. Nawas MT, Daruwalla VJ, Spirer D, Micco AG. Nemeth complicated necrotizing otitis externa. American J Otolaryngology—Head and Neck Medicine and Surgery. 2013;34:70a–709.
12. Carfrae MJ, Kesser BW. Malignant otitis externa, Otolaryngologic. Clinics of North America. 2008;4:537–549.
13. Sreepada GS, Kwartler JA. Skull base osteomyelitis secondary to malignant otitis externa. Curr Opin Otolaryngol Head Neck Surg. 2003;11:316–323.
14. Kornilenko L, Arcehvo I. Aggressive bilateral temporal bone tuberculosis. a case report. Theory and Practice in Medicine. 2016;22:437–440.
15. Illing E, Zolotar M, Ross E, Oalaye O, Molony N. Malignant otitis externa with skull base osteomyelitis. J Surg Case Rep 2011;2011:6.
16. Patmore H, Jebreel A, Uppal S, McNinney,”Skull base infection presenting with multiple lower cranial nerve palsies. American Journal of Otolaryngology—Head and Neck Medicine and Surgery. 2010;31:376–380.
17. Rubin Grandis J, Branstetter BF 4th. Yu VL. The changing face of malignant (necrotising) external otitis: clinical, radiological, and anatomic correlations. Lancel Infect Dis. 2004;4:36–39.
18. Mani N, Sudhoff H, Rajagopai S, Moffat D, Axon PR. Cranial nerve involvement in malignant external otitis: implications for clinical outcome. Laryngoscope 2007;117:907–910.
19. Castro R, Robinson N, Klein J, Geimeier W. Malignant external otitis and mastoiditis associated with an IgG4 subclass deficiency in a child. Del Med J 1990;62:1417–1421.
20. Cassidy PM, Hedberg K, Saulson A, McNelly E, Winthrop KL. Nontuberculous mycobacterial disease prevalence and risk factors: a changing epidemiology. Clin Infect Dis 2009;49:124–9.
21. Yeh CF, Yang Tu T, Wang MC, Chu CH, Huang CY. Emergence of refractory otomastoiditis due to nontuberculous mycobacteria. Clin Infect Dis 2016;62:739–745.
22. KCW, Shu MT, Chen BN. Ototomastoiditis with acute left facial nerve paralysis caused by mycobacterium chelonea. Ear Nose Throat J 2011;90:18–22.
23. Griffith DE, Aksamit T, Brown-Elliot BA, Catanzaro A, Daley C, et al. An official ATS/IDSA statement: diagnosis, treatment, and prevention of nontuberculous mycobacterial diseases. Am J Respir Crit Care Med 2007;175:374–416.
24. Chen P-Y, Wu C-C, Yang T-L, Hsu C-J, Lin Y-T, et al. Gradenigo syndrome caused by nontuberculous mycobacteria. Audiol Neurootol. 2014;19:275–282.