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3 **Spondylodiscitis presenting as pleural effusion in a geriatric**
4 **female: a case report**

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11 **Abstract**

12 Pleural effusion is a frequently seen medical problem caused by pulmonary and
13 non-pulmonary diseases. Spondylodiscitis is a very rare cause of pleural
14 effusion and is typically diagnosed based on clinical, laboratory,
15 microbiological and radiological findings. The low incidence and different
16 clinical presentations of Spondylodiscitis make its diagnosis and treatment
17 challenging. We present the case of a 78-year-old female who was initially
18 admitted due to chest pain and, upon chest radiography, was found to have
19 pleural effusion; and eventually diagnosed with spondylodiscitis.

20 **Keywords:** Spondylodiscitis, exudative pleural effusion, geriatrics, vertebra,
21 infection

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23 **Introduction**

24 Pleural effusion is a widespread medical problem and has more than 50 known
25 causes, including pulmonary and non-pulmonary diseases. The most common
26 causes of exudative pleural effusion are malignancies, pneumonia and
27 tuberculosis.¹ Spondylodiscitis is a very rare cause of pleural effusion.² In such
28 a case, the diagnosis of spondylodiscitis is often delayed, since the initial

29 diagnostic tests for pleural effusion typically focus on pleuropulmonary
30 diseases. Early diagnosis and treatment are critical in the management of
31 spondylodiscitis and its consequent critical, fatal complications.³⁻⁵ Therefore,
32 spondylodiscitis must be considered in the differential diagnosis of pleural
33 effusion. Here, we present the case of a geriatric female who came with
34 complaint of chest pain, was noted to have pleural effusion upon chest
35 radiography, and was eventually diagnosed with spondylodiscitis.

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37 **Case Report**

38 A 78-year-old woman presented with right-sided chest pain and dyspnoea for
39 five days to the emergency service of **Yedikule Chest Diseases and Thoracic**
40 **Surgery Training and Research Hospital** in May 2017. She had a two-month
41 history of back pain and was admitted for physical therapy owing to these
42 complaints. She was referred to our hospital two days later because of right-
43 sided pleural effusion, as revealed by chest radiography (**Figure 1**). She also
44 had a history of ischaemic heart disease, hypertension and type 2 diabetes
45 mellitus. She had smoked for 50 packs/years and consumed 200 ml alcohol per
46 day four days a week. Her body temperature was 37°C, breathing sounds were
47 absent in the right basal hemithorax, her leukocyte count was normal, C-
48 reactive protein (CRP) level was 412.4 mg/dL and erythrocyte sedimentation
49 rate (ESR) was 122 mm/h. Her arterial blood gas pH was 7.47, partial oxygen
50 pressure was 65 mmHg and partial carbon dioxide pressure was 29 mmHg. Her
51 D-dimer level was increased at 6.78 mg/L (0–0.060), while blood and urine
52 cultures were negative. Electrocardiography and echocardiography revealed
53 normal findings. Levofloxacin (500 mg/day, intravenously) and Enoxaparin 0.6
54 mL 2 × 1 (subcutaneously) were started empirically. Chest computed
55 tomography (CT) angiography ruled out pulmonary embolism; however, right-
56 sided pleural effusion and degenerative changes in bone structures were noted.
57 Thoracentesis yielded a fluid exudate (pleural fluid pH 7.26, the pleural fluid-

58 to-serum albumin ratio >0.5 , the pleural fluid-to-serum LDH ratio >0.6) with an
59 adenosine deaminase level of 10.4 IU/L, a lymphocyte percentage of 40% and a
60 neutrophil percentage of 60%. No acid-fast bacilli and malignant cells were
61 detected in the pleural fluid. The pleural fluid cultures grew Methicillin-
62 sensitive *Staphylococcus aureus*. Treatment was modified to Levofloxacin (500
63 mg/day, intravenously) and Piperacillin-Tazobactam (3×4.5 g/day,
64 intravenously) for 14 days. On partial regression of pleural effusion the patient
65 was discharged. One week later, the patient was readmitted to our clinic due to
66 persistent back pain and right-sided pain. Physical examination revealed slightly
67 diminished breathing sounds in the right basal hemithorax and localised
68 tenderness in the right paravertebral area of the lower thoracic spine. Her ESR
69 was 114 mm/h and CRP level was 182 mg/dL, and thoracic ultrasonography
70 revealed a 2.5-cm-thick pleural effusion in the right pleural space. The blood
71 and urine cultures were still negative. Thoracentesis yielded haemorrhagic
72 pleural fluid with a lymphocyte percentage of 80% and a neutrophil percentage
73 of 20%. The pleural fluid cultures were negative for bacteria, acid-fast bacilli
74 and malignant cells; however, malignancy was suspected because of severe pain
75 and haemorrhagic pleural effusion. 18F-fluoro-D-deoxyglucose (FDG) positron
76 emission tomography (PET)/CT revealed high FDG uptake in the right
77 paravertebral region accompanied by heterogeneous density in the vertebral
78 endplates at the T9-T10 vertebral level (maximum standardised uptake value is
79 10.3) indicating spondylodiscitis (**Figure 2**). The diagnosis was confirmed
80 using magnetic resonance imaging (MRI) (**Figure 3**). The patient was treated
81 with Fusidate Sodium (3×500 mg/day, orally) in combination with Teicoplanin
82 (400 mg/day, orally) for six weeks, Hyperbaric oxygen therapy and spinal
83 immobilisation using a thoracolumbar corset. A month later, the pleural effusion
84 had resolved completely (**Figure 4**), CRP was 36 mg/dL and ESR was 57
85 mm/h. After the resolution of pleural effusion, the patient was followed by a
86 neurosurgeon for one year for the possibility of recurrence. Control MRI was

87 unnecessary because of improvement of inflammation markers as well as
88 clinical symptoms.

89

90 **Discussion**

91 Spondylodiscitis is an infrequent and serious infection of the intervertebral disc
92 and the adjacent vertebrae, usually seen in adults in their fifth to seventh
93 decades of life.⁶ Spondylodiscitis affects more men than women,^{4, 5} and has a
94 prevalence of 4.8/100,000 individuals per year.⁴ Advanced age, diabetes
95 mellitus, immunosuppressive medication, intravenous drug use, surgical
96 interventions, urinary tract infections, infective endocarditis, human
97 immunodeficiency virus infection, malnutrition, malignancy, heart diseases,
98 alcoholism, chronic hepatitis and renal failure are the principal risk factors.^{3, 6}
99 The prevalence is constantly increasing due to an increase in intravenous drug
100 users, haemodialysis patients, immunocompromised hosts^{3, 4} and geriatric
101 population.^{5, 6} This patient is an atypical case of spondylodiscitis and was
102 initially misdiagnosed. Advanced age, diabetes mellitus and chronic alcoholism
103 were risk factors for the development of spondylodiscitis in the present case.
104 Pathogens affect the spinal column through haematogenous, external
105 inoculation or contiguity. Haematogenous spread from a distant focus (urinary
106 tract infection, infective endocarditis, skin and soft tissue infection) is the most
107 common route of infection.^{3, 6} *S. aureus* is the most frequent causative
108 pathogen.^{3, 5, 6} Back pain, fever, paravertebral muscle tenderness and spasms are
109 the predominant symptoms and physical findings.³ Based on the causative
110 pathogen and the affected spine areas, spondylodiscitis may clinically manifest
111 in various forms, such as pleural effusion, empyema,² confusion, meningitis and
112 tetraparesis;⁷ furthermore, it may mimic other diseases, such as lymphoma,
113 hepatic or biliary diseases.⁸ This patient initially presented with chest pain and
114 pleural effusion and spondylodiscitis was masqueraded. Leucocyte count may
115 be normal or elevated, but ESR and CRP levels are typically elevated.³

116 Microbiological tests are essential for choosing the appropriate therapeutic
117 agent. In approximately half of the patients, blood culture is the simplest
118 method of identifying the causative pathogens.⁶ If blood culture is negative and
119 no clinical response to treatment is noted, CT-guided biopsy of the
120 intervertebral disc is recommended;³ however, this strategy may delay the
121 treatment. According to the clinical presentation, the causative pathogen can be
122 determined in the urine or pleural fluid,^{2, 9} and intervertebral disc biopsy may
123 not be necessary. In the present case, *S. aureus* was detected in the pleural fluid
124 and treatment was initiated for this pathogen. The mean diagnosis time is 2–4
125 months.³ In this patient, the initial investigation was focussed on the lungs
126 because of pleural effusion which delayed the actual diagnosis by 41 days. MRI
127 is the most sensitive and specific radiological technique for diagnosis and CT
128 plays only a minor role. When there is uncertainty in diagnosis, positron
129 emission tomography/CT can provide additional information.⁶ Treatment is
130 based on antibiotic therapy but surgery may be necessary for some patients.^{3, 5}
131 The optimal duration of antibiotic therapy is debatable. Six weeks of parenteral
132 or highly bioavailable oral antimicrobial therapy seems adequate for most
133 patients; however, the duration should never be less than that.³ Advanced age,
134 infections with Methicillin-resistant *S. aureus*, diabetes mellitus, renal failure or
135 paravertebral abscesses are the risk factors for recurrence.¹⁰ In this patient,
136 although the initial antibiotic of choice was appropriate, the short treatment
137 duration, advanced age and diabetes mellitus may have led to treatment failure.
138 It may be questionable whether pleural effusion or spondylodiscitis has an
139 initiation role. Probably the primary source of pleural effusion is
140 spondylodiscitis in this patient. This is supported by clinical course and follow-
141 up period of the patient — initially, she had back pain two months before right-
142 sided pain, then pleural effusion was exudate from the beginning and empyema
143 was never observed. And lastly, since the focus was on pleuropulmonary
144 diseases, routine CT scans of the thorax, which was only one centimetre slice

145 thickness, were initially performed and no skeletal windows or level
146 adjustments were used at the first admission of the patient. Therefore, at that
147 time, changes in vertebral bodies could not be clearly evaluated and
148 spondylodiscitis was masqueraded.

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150 **Conclusion**

151 The diagnosis of spondylodiscitis is typically difficult. Infrequency of disease,
152 nonspecific symptoms and widespread back pain in geriatric population often
153 lead to delayed or incorrect diagnosis. Furthermore, the manifestation of the
154 disease with atypical clinical symptoms and findings makes the diagnostic
155 process extremely complicated, as it happened in this case. Although
156 spondylodiscitis is rare, it should be considered in the differential diagnosis of
157 pleural effusions, especially in elderly patients with risk factors and suffering
158 from back pain.

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160 **Disclaimer:** None to declare

161 **Conflict of interest:** None to declare

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163 **Informed Consent:** Written informed consent was obtained from the patient for
164 publication of this case in the text.

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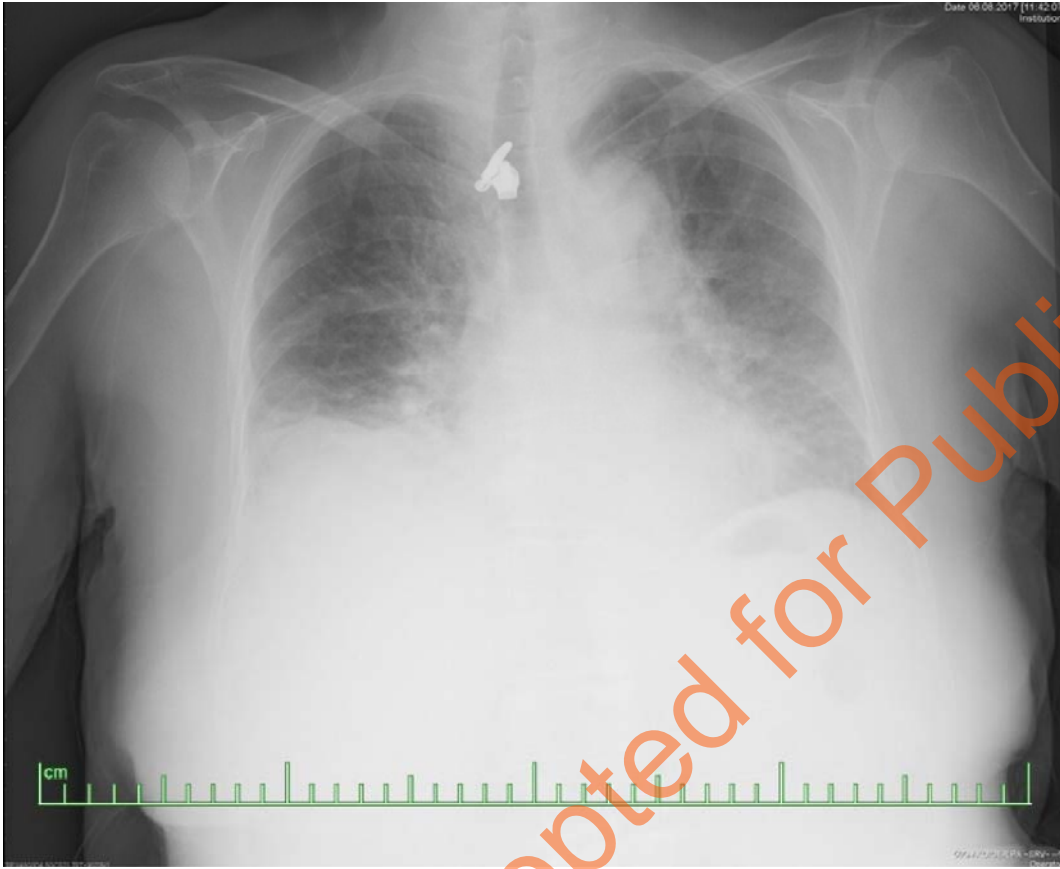
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204 **Figure 1: The initial chest radiograph showing pleural effusion on the right**
205 **side.**

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220 **Figure 2: PET/CT showing high FDG uptake in the right paravertebral**
 221 **region accompanied by heterogeneous density in the vertebral endplates at**
 222 **the T9-T10 vertebral level.**

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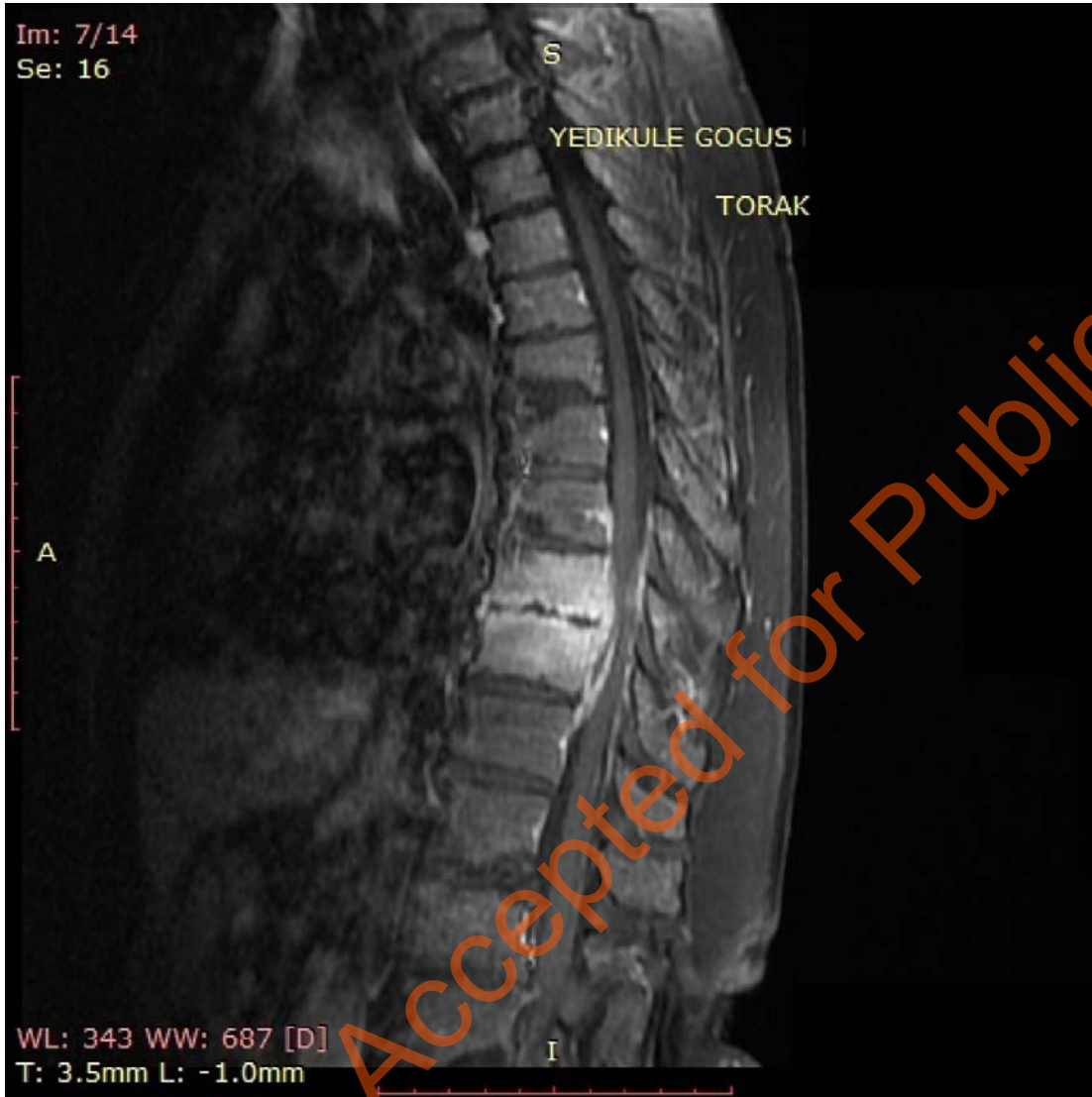
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238 **Figure 3: Post-contrast MRI of the spine showing contrast enhanced at the**
239 **level of T9-T10 and loss of disc space height.**

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Figure 4: Chest radiography showing no pleural effusion after 1 month of treatment.

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