

Ankylosing spondylitis of the cervical spine - a case study

Grzegorz Wróbel, Jakub Spalek

Department of Anatomy, Faculty of Medicine and Health Sciences, Jan Kochanowski University, Kielce, Poland

Correspondence: Department of Anatomy, Faculty of Medicine and Health Sciences, Jan Kochanowski University, Al. IX Wieków Kielc 19A, 25-317 Kielce, Poland.

Tel: 413496965. E-mail: grzegorz.wrobel@ujk.edu.pl

Abstract

Ankylosing spondylitis (AS), (synonyms: Bechterew's disease, Marie Strumpell disease, rheumatoid spondylitis) is a chronic inflammatory disease that attacks the spine, causing its pain and stiffness. The severity of this disease ranges from mild to very heavy. Clinical manifestations of this disease are focused on changes in body posture, for example, inclining the body figure forward or may be a condition for severe disability. The analyzed case concerns the bone material obtained from archaeological excavations conducted in Kielce. During the osteological analysis of the female skeleton at the age of about 50 years, two vertebrae from the cervical segment were joined together. It was established that the fusion occurred between C6 and C7. In exact terms, the association occurred as a result of fusion of the vertebral body by marginal syndesmophytes and fusion of the posterior vertebral elements as well (facet joints). The exact cause of this disease is unknown. Probably for the disease to develop, several factors need to be imposed: genetic, immune (immune system disorders) and environmental factors, such as bacterial infections (especially the gastrointestinal tract and genitourinary tract).

Keywords: ankylosing spondylitis, cervical spine, osteology

1. Introduction

Ankylosing is a term meaning rigid or stiff. *Spondyl* refers to the spine, while *itis* means inflammation. Ankylosing spondylitis (AS), (synonyms: Bechterew's disease, Marie Strumpell disease, rheumatoid spondylitis) is a chronic inflammatory disease that attacks the spine, causing its pain and stiffness. The severity of this disease ranges from mild to very heavy. With regard to the advanced form of this disease, bony anastomoses form at the site of elastic vertebral joints, resulting in a progressive limitation of the mobility of the spine. Clinical manifestations of this disease are focused on changes in body posture, for example, inclining the body figure forward or may be a condition for severe disability. Pathologic changes in the course of this disease may undergo other joints (eg hip, shoulder) and various organs (including eyes, heart, and lungs). The basic changes in this disease are inflammation of the sacroiliac joints connecting the base of the spine and the pelvis. As a rule, the first pain usually appears in the cross, the pain of the cervical spine may appear. At the early stage of the disease may also occur a small fever, loss of appetite, a sense of general breakdown and fatigue. They are so-called general symptoms resulting from the body's defensive response to chronic inflammation. Replacement of ligaments or tendons by bone leads to develop of "Bamboo spine" and causes movement limitation [1-6].

2. Case presentation

The analyzed case concerns the bone material obtained from archaeological excavations conducted in Kielce. During the osteological analysis of the female skeleton at the age of about 50 years, two vertebrae from the cervical segment were joined together. It was established that the fusion occurred between C6 and C7. In exact terms, the association occurred as a result of fusion of the vertebral body by marginal syndesmophytes and fusion of the posterior vertebral elements as well (facet joints) (Figure 1 A , B, C) The range of cervical vertebrae fusion covered the whole circumference of the body of the vertebrae, whereas in the anterior projection of ankylosis it was observed in the range up to 13 mm (Figure 1 D).

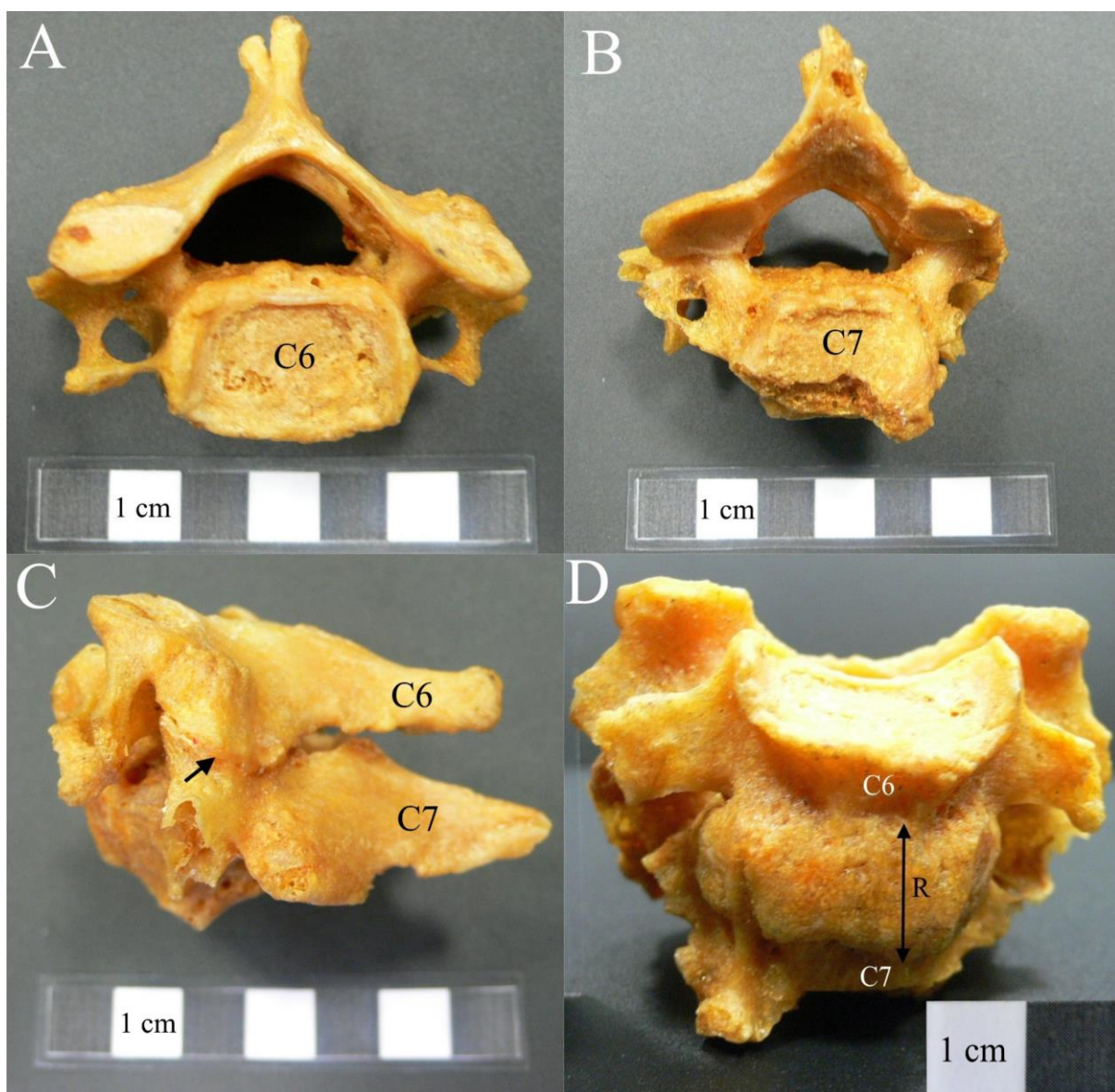


Figure 1. The fusion occurred between C6 and C7: A - view of the upper surface, B - view of the bottom surface, C - side view (the black arrowhead indicates the fusion in the area facet joints), D - front view (R - fusion range).

3. Discussion

The exact cause of this disease is unknown. Probably for the disease to develop, several factors need to be imposed: genetic, immune (immune system disorders) and environmental factors, such as bacterial infections (especially the gastrointestinal tract and genitourinary tract) [7-8]. An important role is played by genetic factors, which confirms the fact that the risk of getting sick is greater if a close relative has AS. In more than 90% of Caucasian patients, so-called HLA B27 antigen, which is the genetic marker of this disease. It should be noted, however, that the majority of people with this antigen do not have AS, and that they are sick with AS who do not have the current HLA B27 antigen. Several other genetic factors are also suspected [9-10]. The prevalence of AS varies considerably across regions of the world. In Central Europe, about 0.3-0.5% of the population suffer. In the United States, the prevalence of AS varies between 1.0% and 1.5%. The disease is more common in men than in women. It usually starts at the end of adolescence or young adults (between 17 and 35 years

of age), but it can also occur in children and older people [11-12].

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