Diffuse Idiopathic Skeletal Hyperostosis in the Third Millennium: Is There (Yet) Cause for Concern?

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Diffuse Idiopathic Skeletal Hyperostosis in the Third Millennium: Is There (Yet) Cause for Concern?

To the Editor:

I read with great interest the recent editorial on diffuse idiopathic skeletal hyperostosis (DISH) by Dr. Mader1. The author affirms the clinical relevance of this skeletal disorder because of its various complications, some of which are very important in clinical practice (e.g., dysphagia, spinal fracture, spinal lumbar stenosis, neurologic complications, postsurgical heterotopic ossifications, etc.). He also emphasizes the need of updating diagnostic criteria (at least 3 sets are used) to obtain a more accurate and timely diagnosis of this condition in its early phases. Dr. Mader is to be congratulated for his interesting paper; however, while defining the still insufficient current diagnostic criteria, he omits comment on possible explanations for this critical diagnostic phase. I would like to express some considerations on the causes of the limited availability of diagnostic tools for DISH.

Forestier’s disease, a common name for DISH, particularly in European countries of Latin origin, is an ancient skeletal disorder whose pathologic alterations were described in human skeletons going back to 5000 years ago2. Many names have been proposed for this condition in the last century (Table 1) and this historical aspect may account for the difficulty in defining the disorder, and its nosographic placement among the various rheumatic diseases! The term diffuse idiopathic skeletal hyperostosis (and its famous acronym DISH), proposed by Resnick, et al in 19753, is largely accepted by the international scientific community because it more exactly expresses the widespread nature of the disorder and, unfortunately, also our ignorance about its etiology. DISH is a mysterious and fascinating condition; however, it remains a little more neglected in clinical practice and research for several reasons.

First, although common in middle-aged and elderly patients, especially those with metabolic disorders, DISH is still underdiagnosed. In their magisterial description Forestier and Rotes-Querol considered DISH “as a not rare disease but most often either goes unnoticed or is mistaken for some other form of vertebral ossification”4. More than 30 years ago Resnick included DISH among the disorders that are “less known, poorly recognized or misunderstood”5. Is it possible that these statements are still relevant today? In my clinical practice I continue to see many patients with DISH misdiagnosed as having osteoarthritis (OA), and I am regularly asked to differentiate DISH from ankylosing spondylitis (AS). Resnick’s criteria are useful for separating DISH from spondylosis deformans and AS, but they don’t help us to evaluate DISH in its early phases. DR. Mader is to be congratulated for his interesting paper; however, while defining the still insufficient current diagnostic criteria, he omits comment on possible explanations for this critical diagnostic phase. I would like to express some considerations on the causes of the limited availability of diagnostic tools for DISH.

<table>
<thead>
<tr>
<th>Period of Publication</th>
<th>Osteoarthritis</th>
<th>DISH</th>
</tr>
</thead>
<tbody>
<tr>
<td>01/01/1979 to 01/01/1989</td>
<td>4849</td>
<td>139</td>
</tr>
<tr>
<td>01/01/1989 to 01/01/1999</td>
<td>8699</td>
<td>233</td>
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<tr>
<td>01/01/1999 to 01/01/2009</td>
<td>18487</td>
<td>218</td>
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</tbody>
</table>

Table 2. Number of articles published on osteoarthritis versus DISH, as cited on PubMed.

Second, since Forestier’s and Rotes-Querol’s description in 19505, DISH has been considered an entity of little clinical relevance, and diagnosis is often made by radiographic examination prescribed for another clinical condition. Because of its mild clinical manifestations for which drug use is rarely required6, no clinical trial has ever been done, and no drug has shown to be efficacious in preventing or curing this condition7.

Third, as is well known, a disease is regarded by research with particular attention because of its clinical and epidemiological relevance and emotional impact on public opinion. DISH, much more than other rheumatic diseases, has not seemed to fit these characteristics.

Lastly, being considered a variant form of OA8,9, this degenerative disorder has come to be of less research interest than in the past. While in recent decades OA obtained much more attention from researchers, that was not the case for DISH (Table 2).

Is there still concern for this ancient disorder in the third millennium? Would it be too much to hope for more accurate diagnostic criteria in the future as invoked in Dr. Mader’s editorial? I think the answer to both questions will be affirmative. In the meantime it has become clear that this condition is a distinct clinical entity10-12 with many serious manifestations. This emerging aspect should stimulate much more interest among researchers than in the past. Every effort should be made for close collaboration among investigators from every field (endocrinology, orthopedics, pathology, radiology, rheumatology, etc.) interested in this disorder. This common effort should also translate into expanded knowledge and recognition of the disorder outside the borders of rheumatology.

In the digital age, with availability of ever more innovative and complex technologies, it should not be difficult to arrive at defining new criteria for a timely clinical-radiologic evaluation leading to diagnosis of DISH in its early phases.

REFERENCES
7. Yagan R, Khan MA. Confusion of roengenographic differential diagnosis between ankylosing hyperostosis (Forestier’s disease)


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