Level of serum beta carotene and its association with lumbar osteophyte formation

N. Selvam¹*, Raju Babu Rapeta²

¹Professor, ²Assistant Professor, Dept. of Orthopaedics, Sri Venkateswaraa Medical College and Research Centre, Ariyur – Pondicherry, India

*Corresponding Author: N. Selvam
Email: drselvam@rediffmail.com

Abstract

Introduction: Cause of chronic low back pain in persons over the age of 50 years include degenerative diseases like osteoarthritis of spine and spinal canal stenosis. Osteoarthritis of spine is a degenerative process¹⁵-⁶⁰ identified radiologically by joint space reduction, marginal osteophytes and sub-chondral sclerosis. Nathan et al³ classified significantly osteophyte positive if more than SIX osteophytes are found from L1 / L2 to L5 / S1 vertebrae in the lumbar spine X-rays. Serum beta carotene is a carotenoid, which has anti-oxidant effect¹⁵-⁵⁷ by increasing Glutathione levels in cells. Normal serum beta carotene level ranges from 0.74 nmol/ml to 3.72 nmols/ml. Based on literature, higher serum beta carotene levels are shown to halt the progression of many diseases including cardiac diseases³⁸,³⁹ and degenerative changes in knee joint. But so far only a single study is available to co-relate the levels of serum beta carotene and lumbar spine osteophytes. Hence this study was done.

Materials and Methods: This study was conducted at JIPMER, Pondicherry, from August 2015 to March 2017. It was conducted as OP procedure. 153 patients of over 50 years of age who complained of low back pain of more than 3 months duration were selected. Their personal history regarding alcohol drinking and smoking noted. Radiographs of lumbar spine A.P. and lateral views were taken. They were divided into osteophyte positive or negative as defined by Nathan.³ These patients’ serum beta carotene levels were assessed. The level of serum beta carotene and its relationship with lumbar spine osteophyte formation was analysed.

Results: Females predominated study 67% of patients were osteophyte negative and 33% were positive. 74% of osteophyte positive patients had serum beta carotene less than 0.7 nmol/ml and 26% had more than that. But, nearly all 98% of osteophyte negative had more than 0.7 nmol/ml of serum beta carotene, showing that the lumbar osteophyte formation is inversely proportional to the level of serum beta carotene levels. Our statistics also showed that there is a strong association between alcohol consumptions and smoking with lumbar osteophyte formation.

Conclusion: Our study showed that low serum beta carotene levels predisposed and increased lumbar osteophyte formation. It also showed strong association between Alcohol consumption and Tobacco use with lumbar osteophyte formation.

Keywords: Lumbar osteophyte, Serum beta carotene, Degenerative osteoarthritis, Alcohol consumption, Tobacco use, Chronic low back pain.

Introduction

Chronic low back¹⁵-¹² affects 60 to 85% of adults in their life time. Definition of chronic low back pain is pain persisting for more than three consecutive months without pain free period - Van Geen et al.¹ Causes of chronic low back pain in persons of more than 50 years of age include osteoarthritis of spine and spinal canal stenosis. Osteoarthritis of spine is defined radiologically by joint narrowing, osteophytes and sub-chondral sclerosis. (O’Neill TW et al.²)

Osteophyte formation is a physiological response to load bearing, alteration in oxygen tension and fluid dynamic pressure.¹³-¹⁷

There are two types of osteophytes: 1. Spondylosis deforms and 2. Inter-vertebral osteochondrosis.

Spondylosis deforms is a bony outgrowth that arises from anterior and lateral perimeters of the vertebral end plate apophyses. They occur due to stress of annular ligament and seen at lumbar 3 level and has no effect on vertebral height and is usually asymptomatic.¹¹,¹³-¹⁷

Keywords: Lumbar osteophyte, Serum beta carotene, Degenerative osteoarthritis, Alcohol consumption, Tobacco use, Chronic low back pain.

Fig. 1
The second category-inter-vertebral osteochondrosis are more aggressive with end plate osteophytes and disc space narrowing. If they protrude into spinal canal they may cause radiculopathy or spinal canal stenosis.

Radiologically osteophyte formation can be assessed by Nathan’s classification. If the total number of osteophytes from L1/L2 to L5/S1 is more than six, that patient is considered osteophyte positive. If it is less than six, the patient is considered as osteophyte negative.

Suprapaneni et al. stated reactive oxygen species are involved in age related degenerative diseases’ pathophysiology.

Cells within joint produce reactive oxygen species which are involved in oxidative damage to joint and osteophyte formation.

Serum beta carotene is a carotenoid which has antioxidant effect by increasing Glutathione levels in cells. Beta carotene had an effect halting the progression of many diseases cardiovascular and reducing lung cancer risk. It also halts the progression of degenerative changes in Knee joint. As an antioxidant it removes singlet oxygen, scavenges per-oxyl and hydroxyl radicals as well as inactivating anionic super oxide compounds.

The normal serum beta carotene level ranges from 0.74 nmol/ml to 3.72 nmol/ml. Human ELISA Kit can access serum beta carotene level.

Many literature studies showed that there is relation between serum beta carotene levels and the formation of osteophytes in degenerative joint disease. Mc Alindon et al. a co-hort study, Rose et al. showed that there is the association between degenerative osteophytes formation of the Knee joint and serum beta carotene levels.

Only Imgama et al. studied the association of the level of serum beta carotene and lumbar osteophyte
formation. He found that low serum beta carotene levels might cause lumbar osteophyte formation.

There is no other study co-relating the level of serum beta carotene and lumbar degenerative osteophyte formation. Hence this study undertaken.

 Imagama also studied the co-ratian between the alcohol intake and tobacco consumption. That study showed that there is a strong co-relation of degenerative joint changes with alcohol consumption and tobacco usage.

Here, we studied this too.

Materials and Methods

153 patients, aged above 50 years who came to the OPD of JIPMER Orthopaedic Department from August 2015 to March 2017 were studied. All of them all complained of chronic low back pain of more than 3 months duration.

Along with the routine demographic data of patients (Name, Age, Sex, Occupation, etc...) history of Tobacco use (10 cigarettes per day x 5 days) and Alcohol consumption (3 - 4 large per day x 5 years) were noted.

X-Rays of Lumbar Spine: A.P. and Lateral views were taken. Patients with structural and anatomical defects in the Lumbo-sacral spine like spondylolisthesis, scoliosis were excluded from the study.

Once the patient was included in the study 2ml of blood was taken for routine investigation of TC, DC, ESR, HB. The blood was centrifuged and the serum was collected, labelled and stored in the Bio-Chemistry department for analysis of serum beta carotene.

After completion of sample size of 153 patients, the serum beta carotene levels of these 153 patient were analysed by two Human ELISA 96 strip well kit (Bioassay co.)

All the 153 patients’ X-Rays of lumbar spine were reviewed by a single radiologist. If more than 6 osteophytes were observed from L1 to L5 that patient was considered to be osteophyte positive. Patients with less than 6 osteophytes from L1 to L5 were considered as osteophyte negative (Nathan, 1962)3

The expected mean and standard deviation of the level of serum beta carotene in Low back pain patients is 0.55 with standard deviation of 0.38 and the sample size is estimated at 5% of significance and 11% relative precision.

Inferential and descriptive statistical analyses were carried out in the present study. Results on continuous measurements were shown on Mean SD (Min – Max) and the results of categorical measurements were presented in numbers. The significance was assessed at 5% level of significance.

Student ‘t’ test (two tailed, independent) was used to find the significance of study parameters, on continuous scale.

Chi-square / Fischer Exact test was used to find the significance of study parameters on categorical scale.

P value: 0.05 < P < 0.10 significant

Results

All the 153 are from South India. It was conducted from August 2015 to March 2017.

Majority of patients were in the 50–60 years age group (64.7%) followed by 61–70 years (30.1%), 71–80 years (4.6%) and above 80 years (0.7%).

Table 1

<table>
<thead>
<tr>
<th>Age in years</th>
<th>Number of patients</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>50-60</td>
<td>99</td>
<td>64.7%</td>
</tr>
<tr>
<td>60-70</td>
<td>46</td>
<td>30.1%</td>
</tr>
<tr>
<td>70-80</td>
<td>7</td>
<td>4.6%</td>
</tr>
<tr>
<td>&gt;80</td>
<td>1</td>
<td>0.7%</td>
</tr>
<tr>
<td>Total</td>
<td>153</td>
<td>100%</td>
</tr>
</tbody>
</table>

A little more than two thirds (68%) were females and rest (32%) males.

50 patients in the study group (33%) were osteophyte positive, of which 22 were (44%) males and 28 were (44%) males and 28 were (56%) females.

Of the 49 males (100%) 22 were osteophyte positive (45%). Of the 104 females (100%), 28 were osteophyte positive (27%) suggested male predominance in osteophyte formation.

Table 2: Osteophyte association with gender

<table>
<thead>
<tr>
<th></th>
<th>Osteophyte Positive</th>
<th>Osteophyte Negative</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>22 (44.89%)</td>
<td>27 (55.11%)</td>
<td>49 (100%)</td>
</tr>
<tr>
<td>Female</td>
<td>28 (26.93%)</td>
<td>76 (73.07%)</td>
<td>104 (100%)</td>
</tr>
</tbody>
</table>

Serum Beta Carotene Levels: The normal serum beta carotene values are from 0.7 nmol/mL to 3.72 nmol/mL.

Out of the 104 female patients 22% had less than 0.7 nmol/mL serum beta carotene level and 78% had more than 0.7 nmol/mL.

Among the 49 male patients 32% had less than 0.7 nmol/mL of serum beta carotene level and 68% had more than 0.7 nmol/mL.

Table 3: Gender distribution serum beta carotene

<table>
<thead>
<tr>
<th>Serum Beta (NMOL/ML)</th>
<th>Female</th>
<th>Male</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;0.7(Low)</td>
<td>23(59%)</td>
<td>16(41%)</td>
<td>39</td>
</tr>
<tr>
<td>&gt;0.7(Normal)</td>
<td>81(71%)</td>
<td>33(29%)</td>
<td>114</td>
</tr>
</tbody>
</table>

The association of serum beta carotene and lumbar osteophyte formation was studied. Among the 50 osteophyte positive patients, 39 patients (74%) had serum beta carotene less than 0.7 nmol/mL and 13
patients (26%) had serum beta carotene more than 0.7 nmol/mL.
Out of 103 patients of osteophyte negative group only 2 patients (02%) had less than 0.7 nmol/ml and 101 patients (98%) had more than 0.7 nmol/mL of serum beta carotene levels. This is statistically significant with P value of 0.0001.

Table 4: Association of beta carotene and lumbar osteophyte

<table>
<thead>
<tr>
<th>Beta carotene</th>
<th>Osteophyte</th>
<th>Osteophyte</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMOL/ML</td>
<td>Negative &lt;6 number</td>
<td>Positive &gt;6 number</td>
</tr>
<tr>
<td>&lt;0.7</td>
<td>2(2%)</td>
<td>37(74%)</td>
</tr>
<tr>
<td>0.7-2.8</td>
<td>101(98%)</td>
<td>13(26%)</td>
</tr>
<tr>
<td>Total</td>
<td>103</td>
<td>50</td>
</tr>
</tbody>
</table>

The correlation between serum beta carotene and lumbar osteophyte was studied. There is moderate correlation between lumbar osteophytes formation with serum beta carotene level (r=-0.5822) which is statistically significant (p<0.0001).

The relationship between Tobacco consumption and Lumbar osteophyte formation was studied. Of the 50 osteophytes positive patients 44% were Tobacco consumers. Of the 103 osteophyte negative patients majority (84%) were non Tobacco consumers. These statistics show that there is a strong association between Tobacco consumption and lumbar osteophyte formation which is reflected in the significant P value of 0.001.

Table 5: Tobacco consumption and osteophyte relation

<table>
<thead>
<tr>
<th></th>
<th>Osteophyte negative</th>
<th>Osteophyte positive</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non tobacco consumers</td>
<td>87(84%)</td>
<td>28(56%)</td>
</tr>
<tr>
<td>Tobacco consumers</td>
<td>16(16%)</td>
<td>22(44%)</td>
</tr>
<tr>
<td>Total</td>
<td>103</td>
<td>50</td>
</tr>
</tbody>
</table>

The relationship between alcohol consumption and lumbar osteophyte was also studied. Of the 103 osteophyte negative patients 84% were non-alcoholic. Of the 50 osteophyte positive patients 62% were non alcoholics. With a P value of 0.003, relatively less significant compared to other variables.

Table 6: Association of alcohol and osteophyte formation

<table>
<thead>
<tr>
<th></th>
<th>Osteophyte negative</th>
<th>Osteophyte positive</th>
</tr>
</thead>
<tbody>
<tr>
<td>Non alcoholics</td>
<td>86(84%)</td>
<td>31(62%)</td>
</tr>
<tr>
<td>Alcoholics</td>
<td>17(16%)</td>
<td>19(38%)</td>
</tr>
<tr>
<td>Total</td>
<td>103</td>
<td>50</td>
</tr>
</tbody>
</table>
Discussion

This study was undertaken to identify the relationship between the level of serum beta carotene and the formation of lumbar osteophytes in patients older than 50 years.

Literature showed only one study by Imagma et al (2005) who studied the relationship of serum beta carotene and lumbar osteophytes in 286 patients of over 50 years of age. With a mean age of 68 years.

Our study had 153 patients with the mean age of 59 years.

Imagma et al observed that men had significantly more lumbar osteophyte positivity than women. In our study we found men had more osteophyte positivity with 44% of all male patients had osteophyte positivity whereas only 27% of all female patients had osteophyte positivity.

Out of 153 patients we studied 34% had serum beta carotene less than 0.7 nmol/mL. Among the 50 osteophyte positive patients 74% had less than 0.7 nmol/mL of serum beta carotene.

In our study the average serum beta carotene levels in osteophyte positive patients is 0.516 +/- 0.327 nmol/mL and among osteophyte negative patients is 1.454 +/- 0.521 nmol/mL with P value < 0.001 which is statistically significant.

The study of Imagma et al indicated that the serum beta carotene level and lumbar osteophyte formation are inversely correlated. Our results are in agreement with his study.

In spite of the geographical remoteness between the locations of these two studies, the average serum beta carotene levels had almost equal values in osteophyte positive patients. This may indicate a common reference range of this particular parameter which may have the potential to be used as a criterion for pre-emptive diagnosis of impending degenerative lumbar Spine disease. This requires more studies as to whether usage of beta carotenoid will stop the progression of lumbar degenerative disease.

The results of our study showed that association of Tobacco consumption and lumbar osteophyte formation is more significant with a P value of 0.001 compared to the association of alcohol and lumbar osteophyte formation with P value of 0.033.

But Imagma et al study showed 80% of smokers had no osteophytes and only 20% had osteophyte positivity with a P value of 0.247 which is not significant.

Our results are in accordance with Imagma et al study with respect to alcohol and lumbar osteophyte association. But our study found that the association is stronger with Tobacco consumption than alcohol intake.

Conclusion

Our study was limited as we did not study other anti - oxidants other than beta carotene. Further our study did not include other factors of lumbar spine degeneration like Body Mass Index, Lordosis, etc.

From the results of our study we conclude that low serum beta carotene levels is associated with lumbar osteophyte formation.

Tobacco consumption to a more level and Alcohol intake to a lesser level predisposes to lumbar osteophyte formation.

From our study suggest that dietary supplements of beta carotene may present the progression of lumbar osteophyte formation.

References

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