

# AN ALTERNATIVE APPROACH TO CALCULATING PEAK BONE MINERAL DENSITIES AND PEAK BONE MASSES

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Abstract— Polynomial regression analyses, commonly used for curve fitting in Bone Mineral Density (BMD) versus Age or Bone Mineral Content (BMC) versus Age graphs, do not seem to be entirely satisfactory. Instead, LOESS (Locally Weighted Smoothing) regression works better in depicting the trend in experimental results. Bootstrapping method has been employed to generate 100 samples from each set of data obtained from DEXA machines located in different parts of Bangladesh and these have been analyzed using Loess regression to calculate the peak bone mineral density and peak bone mass along with the ages at which they appear at 95% confidence intervals. Results have been discussed in the light of previous ones.

*Keywords*—**BMD**, **BMC**, **Osteoporosis**, **Loess**, **Bootstrapping**, **T-score** 

# I. INTRODUCTION

Peak Bone Mineral Density (pBMD) and Peak Bone Mineral Content/Mass (PBM) are important parameters, indicative of the bone health of a population. The greater are the values of these parameters; the better is the bone health of that population. pBMD is important from another point of view. It is used to calculate the T-score defined by the relation,

T-score = (measured BMD – mean BMD of young healthy reference group) / standard deviation [1].

According to WHO, a person with a T-score of -1.0 and above is normal, with a T-score within the range, -1.0 and -2.5, has low bone density or osteopenia and with a T-score of -2.5 and below is suffering from osteoporosis, a bone disease that makes the bone brittle [2]. The problem arises as to how to choose this "mean BMD of young healthy reference group", because there is no universal value. This 'mean BMD' which is nothing but pBMD has been found to depend on sex and ethnicities [3, 4]. Thus to calculate a person's T-score, his ethnicity and sex has to be kept under consideration [5, 6]. Ho-Pham et al [7] have shown for Vietnamese men and women that improper reference values can yield results that can lead to wrong diagnosis of osteoporosis. As far as we know, no such reference value exists in Bangladesh. In our previous paper [8], we calculated pBMD using the traditional method of polynomial regression. In the present paper we have used the LOESS (Locally Weighted Smoothing) curve fitting method.

Although pBMD (g/cm<sup>2</sup>) and PBM (g) are closely related, their relationship may not be exactly linear. BMC depends on both the size and density of the skeletal bone and differs because of either bone size or bone density. Thus it has been found that at the spine 86.2 % of BMD variation is due to BMC and 12.6 % to bone areal size, whereas at the hip it is 98 % and 1.1 % [9]. An increase of PBM by one standard deviation is supposed to reduce the fracture risk by 50%. We have therefore decided to estimate PBM as well by the new method.

# II. METHODS

Details of the experimental procedure for measuring BMD and BMC have been discussed elsewhere [8]. R software was used for statistical analysis.

#### III. RESULTS AND DISCUSSION

A graph of BMD versus age for females has been shown in Fig. 1 where polynomials of degrees, 1-4 have been tried. None of the polynomials seems able to satisfactorily explain the pattern of variation, although most authors have used polynomial of degree 3 and a few have used degree 4 for curve fitting. It is common knowledge to anyone familiar with bone health that bone density increases from childhood to youth, remains steady for sometime and then gradually decreases in old age. The rate of increase in initial years is much higher than the rate of decrease in later years. Linear equation gives no maximum whereas the quadratic one gives a maximum followed by a much sharper fall than in real life. Both third and fourth degree polynomials predict an additional rise in BMD in contrast to reality. Loess curve which is given in Fig. 2 depicts the expected behavior.



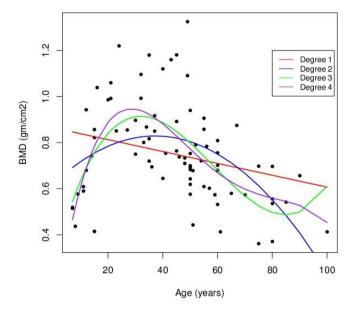


Fig. 1. BMD versus Age graph for female lumbar spine showing curve fitting by polynomials

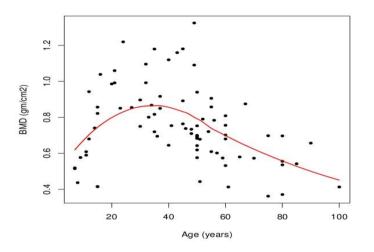


Fig. 2. Loess graph for BMD female

It is found on statistical analysis that although correlation gets better on going from first to fourth degree polynomial, p values suggest no significant improvement. All these considerations lead us to conclude that Loess is the best alternative.

Fig. 3 shows the polynomial fitting of male lumbar spine results.

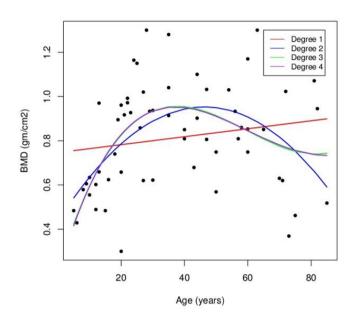


Fig. 3. BMD versus Age graph for male lumbar spine showing polynomial fitting

This figure is almost similar to Fig. 1 except that polynomials of degree 3 and 4 completely overlap each other.

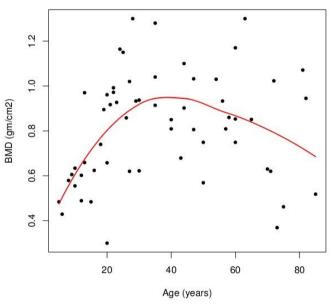


Fig. 4. Loess graph for BMD male

This graph is very similar to that in Fig. 2. Graphs on Bone Mineral Content (BMC) have been shown in Figs. 5-8.



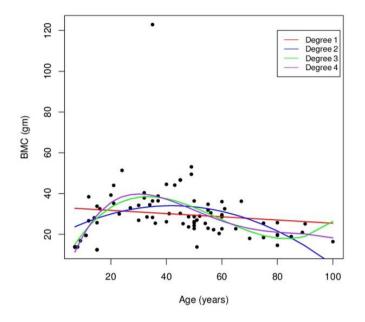


Fig. 5. BMC versus Age graph with polynomial fittings for female lumbar spine

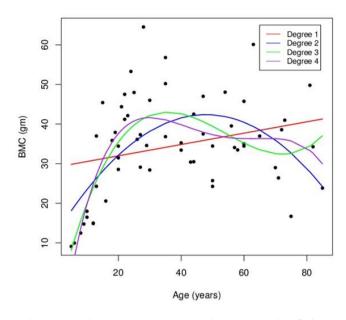


Fig. 7. BMC versus Age graph with polynomial fittings for male lumbar spine

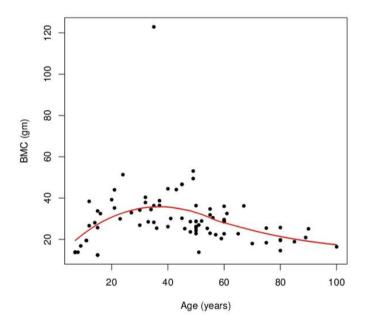


Fig. 6. Loess graph for BMC female

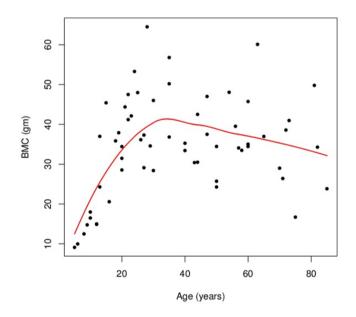


Fig. 8. Loess graph for BMC male

It would be seen that BMC graphs are similar to the corresponding female and male BMD graphs. Calculated values of pBMD and PBM along with their standard deviations and 95% confidence intervals are shown in Tables 1 and 2.



	Original sample		After bootstrapping						
Sex	pBMD	Age	pBMD	Standard	95% conf.	Age	Standard	95% conf.	
	$(g/cm^2)$	(years)	$(g/cm^2)$	deviation	interval	(years)	deviation	interval	
Female	0.865	35.58	0.868	0.0201	0.8653 -	34.71	1.479	34.50 -	
					0.8709			34.92	
Male	0.949	38.72	0.951	0.0226	0.9478 -	39.76	2.729	39.38 -	
					0.9541			40.14	

Table 1. pBMD and Age of occurrence for female and male lumbar spine

Table 2. PBM and Age of occurrence for female and male lumbar spine

Sex	Original sample		After bootstrapping						
	PBM (g)	Age (years)	PBM (g)	Standard deviation	95% conf. interval	Age (years)	Standard deviation	95% conf. interval	
Female	35.80	37.72	36.02	1.729	35.77 – 36.26	36.99	1.141	36.83 – 37.15	
Male	35.80	34.26	41.50	1.136	41.35 – 41.66	35.26	1.724	35.02 – 35.50	

Our results look different from the literature values [8]. Strictly speaking they are not comparable. If we look at the polynomial fittings carefully, we find that the peak values and the corresponding ages differ with the degree chosen. As the degree goes from 2 to 4, the ages shift toward the left and peak values tend to increase ( degree 1 does not give a maximum as it is a straight line). Most of the literature values are based on curve fitting by polynomials of degree 3 and 4, which are often very close and overlapping. Another very important point is that to calculate the peak value, a particular age group is selected, usually 20-29, which makes the age limited to twenties. Unless BMD or BMC values remain steady in this age group, the results are likely to be in error. Besides peak vaues have been obtained in the age group of thirties as well, especially for females [10-13]. By using bootstapping method we have created 100 samples from each data set and calculated the individual age at which the peak appears. That bootstrapping works well is proven by the fact that the results for original sample and the bootstrapped ones are quite close to one another.

From the graphs presented here it is evident through both experimental observation and statistical analysis that Loess is the best alternative for polynomial curve fitting. Bootstrapping is now an established method of resampling and is being widely used in different areas of research including environmental science where data collection may be difficult and expensive. The only precondition for the successful application of bootstrapping is that the original data sample must be reliable. Unfortunately we are not too happy with our data set as they were obtained from hospital sources, which included patients with and without bone weakness and some volunteers. We have not excluded any outliers for plotting curves.

The method proposed here seems to be logical but its robustness has to tested against reliable data sets of different countries.

# IV. CONCLUSION

Important as they are in predicting and diagnosing osteoporosis, the reference values for BMD should be consistent, reliable and uniform all throughout the world. In order to achieve that objective, the international experts on bone health should come to consensus on the following points:

1) Given that the pattern of variation of BMD/BMC against Age is the same for all peoples, decision should made as to whether curve fitting should be done by polynomials of degree 3, 4 or by Loess method;

2) For calculating peak values, it has to be decided whether particular age groups should be chosen or individual values should be obtained on the basis of bootstap resampling;

3) Pressure should be brought to bear upon the manufacturers of different brand of instruments to make sure that all brands give the same value on measurement as is the case with other diagnostic and analytical equipments.



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