Osteoinductivity of Biomet InterGro® in the Athymic Mouse Model

Michael G. Dunn, Ph.D., Director, Orthopaedic Research Laboratory
UMDNJ – Robert Wood Johnson Medical School, 1 Robert Wood Johnson Pl., New Brunswick, NJ 08903

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SUMMARY

The objective of this study was to characterize the osteoinductive properties of two commercially available products containing demineralized bone matrix: InterGro DBM Putty and InterGro DBM Plus (both Biomet). Osteoinductivity (OI), the ability to produce *de novo* heterotopic bone, was assessed histologically (OI ranked on a scale of 0-4) following intramuscular implantation of multiple samples for each test group in an athymic mouse model. Results of this study suggest that:

- **InterGro DBM Putty was not consistently osteoinductive** in this model; only 61% of the samples were osteoinductive, with an average osteoinduction score of 0.85 ± 0.79.

- **InterGro DBM Plus was not consistently osteoinductive** in this model: only 63% of the samples were osteoinductive, with an average osteoinduction score of 0.63 ± 0.49.

INTRODUCTION AND BACKGROUND

Demineralized bone matrix (DBM) is used for treating bony defects as a bone void filler. The purpose of this study was to characterize the osteoinductivity of InterGro DBM Putty and InterGro DBM Plus, which are commercially-available products containing DBM, both from Biomet.

When implanted into normal animals, human DBM is xenogeneic, and is expected to provoke an immune response that may compromise the analysis of osteoinduction. To avoid this, the athymic mouse model was used. The athymic mouse lacks a thymus gland and therefore cannot mount a humoral immune response to the human DBM implants. Precedence of the use of
an athymic mouse (Nu/Nu) model for studying the osteoinductive potential of
demineralized bone allograft was noted in Schwartz et al.\textsuperscript{1}

Samples of the test groups were implanted bilaterally into the mouse hamstring
muscle. Intramuscular implantation of active DBM is expected to induce
cartilage and then bone formation within the implants, a process termed
osteoinduction. The hamstring muscle group (biceps femoris muscle) is a large,
easily accessible muscle, which is commonly used as an implant site to evaluate
heterotopic bone formation. Histological evaluation of the test articles was
conducted 28 days after implantation to assess osteoinduction.

METHODS AND MATERIALS

This study utilized two test groups: \textbf{InterGro DBM Putty} (3 lots; \textit{Table 2}) and
\textbf{InterGro DBM Plus} (1 lot; \textit{Table 2}). For comparisons, this study references
osteoinductivity data on Enhance\textsuperscript{TM} Demineralized Cortical Fibers collected
by the same investigator using techniques identical to those described in this
study.\textsuperscript{2} In some cases, the reference data was obtained contemporaneously with
test samples in this study.

Eight samples (weighing 25 mg each) from each lot of material were prepared
for implantation. The samples were randomized and implanted bilaterally in the
hamstring muscles of athymic nude mice. Animals were sacrificed at 4 weeks
post-implantation. Decalcified histology was then performed on the explanted
samples; 5 histological slides with 3 sections per slide were prepared for each
sample (15 sections total per sample). Slides were stained with hematoxylin
and eosin, and samples were evaluated for osteoinductivity. A semi-quantitative
scoring system was utilized to assess osteoinduction.
The relative amount of osteoinduction was evaluated semi-quantitatively by the study investigator using the scoring system described below; the observer was blinded to the identification of the implant. Osteoinductive scores were based on the degree to which new bone, bone cells, osteoid, calcified cartilage remnants, and marrow elements were present. To be consistent with proposed standards in the industry, the scoring system in Table 1 was utilized.

<table>
<thead>
<tr>
<th>Score</th>
<th>Criteria</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>No evidence of new bone formation</td>
</tr>
<tr>
<td>1</td>
<td>1-25% of the section is covered by new bone</td>
</tr>
<tr>
<td>2</td>
<td>26%-50% of the section is covered by new bone</td>
</tr>
<tr>
<td>3</td>
<td>51%-75% of the section is covered by new bone</td>
</tr>
<tr>
<td>4</td>
<td>&gt;75% of the section is covered by new bone</td>
</tr>
</tbody>
</table>

Table 1: Osteoinductivity Scoring Scale and Criteria

The overall score for each sample was obtained by averaging the highest 5 scores from the histological slides; scores for each experimental group were determined by pooling the overall scores of the individual samples. The results of semi-quantitative scoring are presented as a mean ± standard deviation.

Images of histological slides from each test group were also captured and stored using a digital camera and computer system (Image-Pro Plus™ imaging software).
RESULTS & CONCLUSIONS

InterGro DBM Putty was not consistently osteoinductive in this model; only 61% of the samples were osteoinductive, with an average osteoinduction score of 0.85 ± 0.79 (*Tables 2 & 3*).

InterGro DBM Plus was not consistently osteoinductive in this model as well: 63% of the samples were osteoinductive, with an average osteoinduction score of 0.63 ± 0.52 (*Tables 2 & 3*).

Figures 1 and 2 show representative histological images demonstrating the osteoinductive response to the InterGro DBM Putty and Plus implants, respectively, with new bone formation.

The osteoinductivity scores for InterGro DBM (both Putty and Plus compositions) are significantly lower than the osteoinductivity scores for Enhance™ Demineralized Cortical Fibers. In all cases, 100% of Enhance™ Demineralized Cortical Fibers samples are osteoinductive when assessed using this model.²

In conclusion, these results suggest that under the conditions of this study, and for the batches (donors) tested, the osteoinductivity for both the Putty and Plus compositions of Biomet InterGro DBM is significantly less than that of Enhance™ Demineralized Cortical Fibers.

It is unknown how the osteoinductive potential, measured in the athymic mouse model, will correlate with clinical performance in humans.
Table 2: InterGro osteoinduction scores

<table>
<thead>
<tr>
<th>Article</th>
<th>Lot</th>
<th>Average Osteoinductive Score</th>
<th>Group Std Dev</th>
</tr>
</thead>
<tbody>
<tr>
<td>InterGro DBM Putty</td>
<td>004920</td>
<td>0.83</td>
<td>0.75</td>
</tr>
<tr>
<td>InterGro DBM Putty</td>
<td>090040</td>
<td>1.25</td>
<td>0.84</td>
</tr>
<tr>
<td>InterGro DBM Putty</td>
<td>744040</td>
<td>0.43</td>
<td>0.50</td>
</tr>
<tr>
<td>InterGro DBM Plus</td>
<td>747310</td>
<td>0.63</td>
<td>0.49</td>
</tr>
</tbody>
</table>

Table 3: Summary statistics, number of samples that could be histologically evaluated, and number of osteoinductive samples for each group. Number of osteoinductive samples is divided by the number of evaluated samples to give the % of osteoinductive samples for each group.
REFERENCES:


   Musculoskeletal Transplant Foundation (MKTG -810).

3. Draft Standard: Standard Guide for the Assessment of Bone Inductive Materials, ASTM F04.4 Division, Draft by Barbara Boyan,
   Univ. of Texas Health Science Center at San Antonio, downloaded from ASTM website 5-8-2000.

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