Correlation of Age and Bone Marrow Derived CD 34+ Cells and Leucocytes in 873 Patients.

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Research

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Abstract

Background:

The use of regenerative medicine, such as autologous chondrocyte implantation (ACI), matrix associated stem cell therapy (MAST) and bone marrow derived stem cell therapy against arthritis is the gold standard for certain indications. However, the clinical improvement of patients using these novel therapies remains heterogeneous and the reasons for this are not fully understood.

The impact of age is always a concern for patients and doctors and elderly patients can only be mobilized with lower total collected CD34+ cells, older age correlates with inferior results, fatty degeneration of the bone marrow, delayed fracture-healing and osteoporosis, but solid data are missing.

Purpose:

This is the first study to determine the average quantity of leukocytes and CD 34+ cells and their relationship in human bone marrow.

Study design: Descriptive Laboratory Study

Methods:

We evaluated the laboratory results of 873 patients (aged 1-90 years), who underwent stem cell transplantation for non malignant diseases.

Results:

We found no age-related decrease regarding the number and the vitality of leukocytes and CD 34+ stem cells. The number of bone marrow derived leucocytes and CD 34+ cells showed a strong and significant correlation.

Conclusion:

The amount of bone marrow derived stem cells can be predicted by leukocytes. This study makes further research possible in order to link clinical outcome to the absolute number of stem cells and leukocytes.

An upper age-limit for stem cell therapy can therefore not be defined from the donor-site perspective.

Clinical Relevance:

The number of leucocytes might be used to predict the amount of stem cells in order to select the ideal patient.

What Is Known About The Subject
Cartilage undergoes remarkable alterations in composition, organization, and mechanical properties with aging. Therefore several cartilage restoring methods have been established in the past (ACI, MAST, etc.). Regardless of the purpose of a cell therapy, successful cell transplantations require the use of a sufficient number of specific cells and their engraftment. Specific qualitative age-dependent findings were published up to now, but the quantity of stem cells according to age has not yet been studied sufficiently. To our knowledge this is the first study, assessing viability and quantitative number of stem cells over age in approximately 1000 patients.

What this study adds to existing knowledge:

Based on our findings, no upper age limit can be derived from a quantitative point of view. Furthermore we found a strong correlation between leukocytes and bone marrow derived CD+ stem cells. This is the first study to determine the average quantity of leukocytes and CD 34+ cells and their relationship in human bone marrow. We also found an inter-individual variation of cell number. This study makes further research possible in order to link clinical outcome to the absolute number of stem cells and leukocytes.

Introduction

The use of cell therapy, such as autologous chondrocyte implantation (ACI) and matrix associated stem cell therapy (MAST) for the repair of damaged cartilage is well established, demonstrating good short to medium term outcomes.\textsuperscript{9, 24} However, the clinical improvement of patients using these novel therapies remains heterogenous and the effects are not fully understood. Recently bone marrow derived stem cells gained further attraction in orthopedic diagnoses, where they are also used as bone marrow aspirate (BMA) or bone marrow aspirate concentration (BMAC) for stem cell therapy.

(1) Arthritis is treated in this way, aiming to generate additional cartilage, and stop the inflammatory process.\textsuperscript{8, 11, 12, 18, 27, 31, 33, 35}

(2) In patients with spinal cord injuries, bone marrow derived stem cells can improve some neural function.\textsuperscript{6, 7, 13, 15, 17, 23}

Regardless of the purpose of a cell therapy, successful cell transplantations requires the use of a sufficient number of specific cells and their engraftment.\textsuperscript{20} Better clinical results can be achieved with a higher donor-site stem cell count\textsuperscript{19, 30} and a lower age.\textsuperscript{14} The impact of age is not fully understood for stem cell procedures and controversial:

Specific qualitative age-dependent findings were published up to now (lower proliferation and extra cellular matrix forming potential\textsuperscript{16}, decreasing growth rate and telomere length,\textsuperscript{4} lower mobilization rate,\textsuperscript{1, 28} fatty degeneration of the bone marrow, delayed fracture-healing and osteoporosis,\textsuperscript{21} acquired mitochondrial DNA mutations\textsuperscript{34}) but the quantity of stem cells according to age has not yet been studied sufficiently. In hemato-oncology is is known, that grafts from older donors do not adversely affect
outcomes of allogeneic hematopoietic cell transplantation as compared to grafts from younger donors in.\textsuperscript{10,28}

Since an age-dependent range of the physiological leukocyte- and stem-cell-numbers and their vitality in the human bone marrow has not yet been reported and most clinical studies about CD 34+ stem cells refer to patients with malignant diseases, we wanted to establish a normal range of bone marrow derived leukocytes and stem cells of patients without malignancies according to age, harvested by bone marrow aspiration (BMA).

Therefore the aim of this study is to

- evaluate the number and the vitality of bone marrow derived leukocytes and CD 34+ cells in a large number of patients undergoing autologous stem cell transplantation for nonmalignant diseases
- study the connection to age in order to find out, if a possible age limit is present, after which cell number and vitality decreases.
- predictor exists for the amount of CD 34+ cells in the bone marrow.

**Material And Methods**

**Level IIb: Retrospective cohort study**

In a retrospective study the laboratory results all patients, who underwent stem cell transplantation for nonmalignant diseases were evaluated. All bone marrow punctures were done by the same surgeon (K.G.). The stem cells were harvested with a Yamshidi Needle (15 ga x 2.688 in MAX Bone Marrow Aspiration Needle, ARGON Medical devices, Athens, USA, [www.argonmedical.com](http://www.argonmedical.com)) under sedoanalgesia and in compliance with all applicable laws and regulations. 90 ml of bone marrow aspirate was retrieved using the Technique of Kristin Oliver\textsuperscript{26} (using 10 ml syringes and changing direction repeatedly). One milliliter of this sample was immediately transferred to a laboratory and analyzed with FACS (Fluorescence Activated Cell sorter) using a stem Cell Kit from Beckman Coulter and the ISHAGE protocol ([https://www.bc-cytometry.com/PDF/DataSheet/IM3630.pdf](https://www.bc-cytometry.com/PDF/DataSheet/IM3630.pdf)).

**Inclusion criteria were:**

- Age between 1 and 90 years
- Bone marrow puncture

**Exclusion criteria were:**

- Malignant disease
- Known autoimmune disease
- Inflammatory disease
- Prior bone marrow stimulation or treatment with GCSF (granulocyte colony stimulating factor)
- Prior treatment with hormones or corticoids in the last 6 months

**Statistical Analysis was done using:**

- Pearson's chi-squared tests
- One-way analyses of variance (with post-hoc Bonferroni-adjusted pairwise comparisons of estimated marginal means)
- Pearson's Correlation

**Results**

873 datasets were found in the laboratory patient database. Age ranged from 1-90 years (mean 28, median 25) and patients were clustered into age groups of 10 years. Gender distribution was 29% female and 71% male. There was no gender-difference. The anonymous laboratory data was statistically evaluated by a blinded observer. (Table 1)

<table>
<thead>
<tr>
<th>Age</th>
<th>n</th>
<th>gender</th>
<th>Leukocyte count</th>
<th>Leukocyte vitality</th>
<th>CD 34+ cell count</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;10</td>
<td>296</td>
<td>male</td>
<td>19 (±16)</td>
<td>87 (±9)</td>
<td>298 (±299)</td>
</tr>
<tr>
<td>11-20</td>
<td>102</td>
<td>74%</td>
<td>14 (±8)</td>
<td>86 (±8)</td>
<td>147 (±134)</td>
</tr>
<tr>
<td>21-30</td>
<td>89</td>
<td>72%</td>
<td>14 (±7)</td>
<td>84 (±16)</td>
<td>125 (±106)</td>
</tr>
<tr>
<td>31-40</td>
<td>92</td>
<td>80%</td>
<td>11 (±6)</td>
<td>86 (±9)</td>
<td>97 (±69)</td>
</tr>
<tr>
<td>41-50</td>
<td>109</td>
<td>75%</td>
<td>14 (±8)</td>
<td>91 (±3)</td>
<td>90 (±73)</td>
</tr>
<tr>
<td>51-60</td>
<td>98</td>
<td>72%</td>
<td>8 (±4)</td>
<td>85 (±13)</td>
<td>94 (±149)</td>
</tr>
<tr>
<td>61-90</td>
<td>86</td>
<td>51%</td>
<td>7 (±2)</td>
<td>86 (±7)</td>
<td>79 (±89)</td>
</tr>
<tr>
<td>mean</td>
<td>28</td>
<td>71%</td>
<td>16 (±13)</td>
<td>87 (±9)</td>
<td>183 (±226)</td>
</tr>
</tbody>
</table>

**Bone marrow derived leucocyte cell count**

No significant differences between age and gender and no interaction. Even in a pairwise comparison of patients below the age of 20 versus patients at the age of 20 years and over, no significant differences
were found \((p=.9)\). The correlation between age and bone marrow derived leucocyte cell count is negligible \((r=-.255, p<.001, \text{Figure 1})\).

**Bone marrow derived leucocyte vitality**

No significant differences between age and gender and no interaction in all age groups, no significant correlation. (Table 1)

**Bone marrow derived CD34+ cell count**

No significant differences between age and gender and no interaction. Even in a pairwise comparison of patients below the age of 20 versus patients at the age of 20 years and over, no significant differences were found \((p=.8)\). The correlation between age and bone marrow derived CD 34+ cell count is negligible and comes from patients below the age of 10 years \((r=-.361, p<.001, \text{Figure 2})\).

**Correlation / Predictor**

The number of bone marrow derived leucocytes and CD 34+ cells (as a subset of leucocytes, respectively) had a great variation between individual patients, but both cell types correlated strong and significant \((p<.001, r^2=822, \text{Figure 3})\) within the respective patients. Bone marrow derived leukocytes are therefore a viable predictor for the amount of stem cells. The number of stem cells can be calculated as follows:

\[
\text{Stem cells (CD 34+ cells / Microliter) = 13.5 \times Leukocytes (per Nanoliter bone marrow aspirate)}.
\]

Bone marrow derived leucocytes and and CD 34+ cells had a negligible tendency \((r=-.255, p<.001 \text{ and } r=-.361, p<.001)\) to decrease over life time. The weak correlation was only due to the group of children below the age of 10 years. Thereafter there was a variation between individual patients, but no decrease over time.

**Discussion**

Arthritis and musculoskeletal disorders constitute a major cause of disability and the burden of musculoskeletal diseases will increase with an increasing ageing population. Stem cells remain at the forefront of efforts in Regenerative Medicine, based on a conviction that this technology can provide an effective treatment paradigm for major diseases where there is still an unmet need.

In 2017 the first prospective, single blind, placebo-controlled trial of bone marrow aspirate concentrate for knee osteoarthritis described a positive clinical outcome. A recent review of 1500 papers on stem cell therapy in orthopaedics revealed, that studies reported information on only 42\% (range, 25%-60\%) of the variables included within established minimum reporting standards, leaving it unclear, which amount of
stem cells was really harvested and injected.\textsuperscript{25,29} A higher donor-site cell count correlates with a better outcome.\textsuperscript{8,14,18,19,27,35} In hemato-oncology, where bone marrow transplantations are performed routinely since decades, elderly patients show inferior mobilization rates and inferior outcome in some studies.\textsuperscript{1,28,14} Likewise, in a mice model an age-related fatty degeneration of the bone marrow was described.\textsuperscript{21}

The current study looked at the donor site of 873 healthy patients (without bone marrow diseases), applicable for orthopaedic interventions, to understand the vitality and the quality of the bone marrow derived (stem) cells.

**Age and cell counts**

Due to its large sample size this study is the first to establish a normal range of leukocytes and CD34+ stem cells in bone marrow aspirate (BMA) in the average population. The amount and vitality of bone marrow derived leucocytes and the number of mononuclear cells (stem cells) do not deteriorate over age. The formerly described lower mobilization rate of bone marrow derived (stem) cells in the elderly in some studies\textsuperscript{1,28} conflicts with a recent study, where little of the parameter variability could be explained by age.\textsuperscript{5} Our finding, that the number of bone marrow derived leucocytes remains stable in adults was confirmed in a prior study with 24 goats, but we did not find further papers referring to humans.\textsuperscript{2}

**Vitality**

The vitality of bone marrow derived cells was always high (87-91\%) in all age groups using FACS (fluorescence-activated cell sorting) analysis. Regarding dental pulp stem, the proliferation rate decreases in elderly patients.\textsuperscript{36} We could not confirm this finding in bone marrow derived cells. It is well known, that jawbones have an different bone metabolism.\textsuperscript{32}

**Correlation / Predictor**

Stem cells are mononuclear cells and therefore a 7.4\% fraction (1 / 13.5) of the bone marrow derived leucocytes, as we demonstrated.

To our knowledge no prior study described the positive and strong correlation (p<.001, $r^2=822$) between both parameters. Since stem cells can only be identified using specific CD antigen sets (CD 34, CD 90, CD 45, CD 107,...), which is costly and laborious, this correlation can be utilized, to predict the amount of stem cells based on the number of bone marrow derived leucocytes alone, which is much easier. As a matter of fact, only a negligible share of publications reports the absolute number of stem cells used per patient.\textsuperscript{29} Using the new described correlation, a much cheaper possibility exists, to assess, if a specific
patient has a high or low stem cell number, since counting leukocytes can be done in any operating room, but counting stem cells requires at least a FACS analysis.

**Limitations:**

Since this was a retrospective study, we were not able to assess any social data (body weight, sport habits, smoking status,...). We did not perform colonization and differentiation experiments, since the lack of a specific MSC marker and the low frequency of MSCs in bone marrow necessitate their isolation by in vitro expansion.

Further research is needed to link clinical outcome with the absolute number of bone marrow derived leukocytes and stem cells since it is not yet clear, if the heterogenous clinical results of individual patients are linked to the heterogenous bone marrow derived stem cell counts. It remains speculative if patients with higher cell counts will have better outcomes and might therefore be better suitable for stem cell operations.

**Conclusion**

We established a normal range of bone marrow derived leukocytes and stem cells in 873 patients. No age-related decrease regarding the number and the vitality of leukocytes and stem cells was found. Furthermore the number of bone marrow derived leucocytes might be used to predict the amount of stem cells (usually a 7.4% share) on an individual basis in order to focus on the “ideal” patients.

**Declarations**

- **Ethics approval and consent to participate**

Use of anonymous secondary data, no trace to individual patients is possible. According to the declaration of Helsinky and the Ethics Commission of our University no informed consent necessary.

- **Consent for publication**

Approved by all authors.

- **Availability of data and material**

Free use of our data upon request.

- **Competing interests**

Nothing to declare.
• **Funding**

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• **Authors' contributions**

GK and CP contributed to the drafting and CP to writing and each author contributed to substantial review of this paper. HL did the statistic analysis. CP, HL, GS, MK produced the figures and tables. All authors have read and approved the final submitted manuscript.

• **Acknowledgements**

Nothing to declare.

**References**


**Figures**
Figure 1

There is no relevant decrease of bone marrow derived leucocyte cell count with increasing age.
Even in elderly patients (up to 90 years) there is no relevant decrease of stem cells.
A strong correlation between bone marrow derived leucocytes and bone marrow derived CD34+ cells was found.