Platelet-rich plasma in osteoarthritis treatment: review of current evidence

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Abstract: Platelet-rich plasma (PRP) is defined as a volume of plasma with a platelet concentration higher than the average in peripheral blood. Many basic, preclinical and even clinical case studies and trials report PRP’s ability to improve musculoskeletal conditions including osteoarthritis, but paradoxically, just as many conclude it has no effect. The purpose of this narrative review is to discuss the available relevant evidence that supports the clinical use of PRP in osteoarthritis, highlighting those variables we perceive as critical. Here, recent systematic reviews and meta-analyses were used to identify the latest randomized controlled trials (RCTs) testing a PRP product as an intra-articular treatment for knee osteoarthritis, compared with an intra-articular control (mostly hyaluronic acid). Conclusions in the identified RCTs are examined and compared. In total, five recent meta-analyses and systematic reviews were found meeting the above criteria. A total of 19 individual trials were identified in the five reviews but only 9 were level of evidence I RCTs, and many had moderate or high risks of bias. At present, results from these RCTs seem to favor PRP use over other intra-articular treatments to improve pain scales in the short and medium term (6–12 months), but the overall level of evidence is low. As a result, clinical effectiveness of PRP for knee osteoarthritis treatment is still under debate. This is, prominently, the result of a lack of standardization of PRP products, scarceness of high quality RCTs not showing high risks of bias, and poor patient stratification for inclusion in the RCTs.

Keywords: allogenic products, anti-inflammatory intra-articular therapies, clinical evidence, clinical trials, knee osteoarthritis, patient stratification, platelet-rich plasma

Introduction
Platelets, also known as thrombocytes, are small cytoplasmic fragments derived from bone marrow megakaryocytes. Most platelet functions are directly connected with platelet activation, a process that occurs naturally after an injury in the wall of a blood vessel. Platelets are then exposed to collagen and other extracellular matrix proteins that stimulate their activation, resulting in the release of the content of their cytoplasmic granules. Overall, platelets contain over 800 proteins and molecules, comprising cytokines, chemokines, membrane proteins, metabolites, messenger molecules, growth factors (GFs) and numerous soluble proteins. As a result, besides their role in coagulation and hemostasis, platelets are also involved in vasoconstriction, inflammation, immune response, angiogenesis and tissue regeneration and consequently, they participate in numerous physiologic signaling mechanisms and are related to multiple pathologies.

The therapeutic use of platelet concentrates was first described by Whitman in 1997, although blood-derived fibrin glues were already used 30 years earlier to seal wounds and stimulate their healing. In 1998, platelet concentrates started to be known as platelet-rich plasma (PRP), generally defined as a volume of autologous plasma containing a higher platelet count than peripheral blood (150,000–350,000 platelets/μl). Thereafter...
multiple systems have been developed to concentrate platelets and remove erythrocytes (red blood cells; RBCs) and, in some cases, also leukocytes (white blood cells; WBCs), as reviewed elsewhere.9 For the purpose of this review, PRP or PRP products refer to any product derived from a platelet concentrate (PC), optionally containing WBCs, from whole blood.

The rationale for its use is quite strong: it is considered well tolerated, it very rarely leads to complications, it is easy to prepare and administer and it is less aggressive than other therapeutic options that might be indicated for some patients, such as corticoid intra-articular injection or even surgery.10,11

The number of studies on PRP preparation methods and applications has grown exponentially since the late 1990s; however, there are still concerns regarding its clinical efficacy, mainly due to the heterogeneity of preparation methods and resulting products, the scarceness of high quality randomized controlled trials (RCTs), and the contradictory results that have been found so far.

**Custom and commercial preparation methods for human use**

Methodologies to prepare PRP and derived products vary widely. Briefly, they can rely on single centrifugation, double centrifugation, or blood selective filtration procedures, and on manual or automatic systems operated in open or closed circuits. *Ex vivo*, platelet activation can be triggered mechanically with freeze–thawing cycles, chemically with thrombin or calcium chloride, or endogenously. This last option implies the direct application of nonactivated PRP, to allow local tissue factors to elicit the process.9,12

The high number of variables involved has led to innumerable custom protocols developed in-house in the different research labs and also to many (proprietary) commercial systems for producing PRP products (see Table 1).

Due to their complex composition and to inter- and intra-individual differences,13 each PRP is unique and very difficult, if not impossible, to characterize. As a result, PRPs are usually defined in terms of a few critical variables and a detailed description of the preparation protocol. These variables usually include (1) the proportion of platelets in PRP to platelets in whole blood (platelet enrichment factor, PEF), (2) presence/absence of WBCs, and (3) method of activation. Yet, many authors believe this is not enough and different systems for a more complete and standardized description of platelet-derived product characteristics have been proposed (see Table 2).

Mishra and colleagues proposed to classify PRPs with two parameters: firstly, ‘type’ of PRP: (1) increased WBCs and no activation; (2) increased WBCs and activated; (3) minimal/no WBCs and no activation; (4) minimal/no WBCs and activated; and secondly, its platelet enrichment factor, A if the PRP contains a platelet concentration at or above five times the baseline, or B if platelet concentration is less than five times the baseline.14 On the other hand, the PAW (Platelets, Activation, White cells) classification system includes at least three variables: (1) the absolute platelet concentration (P); (2) the method of activation (A); (3) the presence or absence of WBCs and neutrophils (W) relative to the baseline. Platelets are categorized as P1 (baseline) to P4 (>1.2 million platelets/ul), activation as either exogenous (X) or not, and WBCs and neutrophils as either above or below baseline.15 Mautner and colleagues advocated for reporting at least platelet concentration (cells/μl, volume injected), leukocyte concentration, including the concentration of neutrophils (if >1%), RBC concentration and activation by exogenous agents.16 And more recently still, the DEPA (Dose of injected platelets, Efficiency of production, Purity of the PRP, Activation of the PRP) classification system was proposed.17 This system takes into account (1) the dose of injected platelets from A (>5 billion) to D (<1 billion); (2) platelet capture efficiency from blood from A (>90%) to D (<30%); (3) the % platelets compared with RBCs and leukocytes in the PRP from A (>90%, very pure) to D (<30%, whole blood PRP); and, (4) the activation process. Thus, an ‘AAA’ DEPA score refers to an injection of PRP with a very high dose of platelets (>5 billion) with little contamination from RBCs and with very high platelet recovery efficiency from blood. However, none of these methods has been formally adopted and PRP descriptions are still very heterogeneous.

**Current regulatory framework**

Across the European Union (EU) PRP is regulated under Directive 2001/83/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating
**Table 1.** Examples of commercial PRP preparation systems.

| Name (manufacturer, country)         | Approval | System | Circuit | Procedure | Activator      | Composition PEF/WBC/RBC | Indications                                      |
|--------------------------------------|----------|--------|---------|-----------|----------------|-------------------------|-------------------------------------------------|
| Accelerate PRP Sport (Exactech, UK)  | CE       | M      | SC      | SS        | Thrombin       | 3–8×                    | Orthopedics, sports medicine, pain, cosmetic    |
| ACP (Arthrex, USA)                   | FDA; CE  | M      | C       | SS        | -              | 2–3×/no WBCs/no RBCs    | Orthopedics                                     |
| Angel (Arthrex, USA)                 | FDA; CE  | A      | C       | SS        | -              | 2–18×/custom WBCs       | Orthopedics                                     |
| BioCUE (Biomet, USA)                 | FDA; CE  | A      | C       | SS        | -              | Equivalent to GPS II    | Mix with bone graft                             |
| Cascade (MTF/CONMED, USA)            | FDA      | M      | SC      | SS        | CaCl₂          | 5×                      | Sports medicine, wound healing                  |
| Centrepid (CellMedix, USA)           | FDA      | M      | SC      | DS        | -              | 5×/2–3× WBCs            | Mix with bone graft                             |
| Cyclone (Alliance Partners, USA)     | FDA      | NR     | NR      | NR        | -              | Equivalent to Pure PRP II | Mix with bone graft                             |
| Dr. PRP (Dr. PRP USA LCC, USA)       | FDA; CE  | M      | SC      | DS        | -              | 15×/high WBCs/low RBCs  | Ophthalmology, orthopedics, sports medicine, cosmetics, pain |
| GLD PRP (Glotech/Glofinn Oy, S Korea)| CE       | M      | SC      | DS        | -              | 4–8×                    | Orthopedics, wound healing, ophthalmology, cosmetic |
| GPS II and III (Biomet, USA)         | FDA; CE  | M      | SC      | SS        | -              | 3–9.3×/5× WBCs          | Orthopedics, surgeries, wound healing           |
| Kyocera Medical PRP Kit (Kyocera, Japan) | CE     | M      | SC      | DS        | CaCl₂          | 7.8×/high WBCs/high RBCs | Not specified                                   |
| Magellan (Isto Biologics, USA)       | FDA      | A      | C       | DS        | Thrombin CaCl₂ | 2–13×                  | Not specified                                   |
| Name (manufacturer, country) | Approval | System | Circuit | Procedure | Activator | Composition PEF/WBC/RBC | Indications |
|-----------------------------|----------|--------|---------|-----------|-----------|------------------------|-------------|
| Peak (Depuy, USA)           | FDA      | A      | C       | SS        | -         | 8×/4–6× WBCs/1/3 RBCs  | Mix with bone graft |
| PRGF-Endoret (BTI, Spain)   | FDA; CE  | M      | C       | SS        | CaCl₂     | 2–3×                   | Oral surgery, dermatology, cosmetic, wound healing, ophthalmology, OA, orthopedics |
| Pure PRP II (Emcyte, USA)   | FDA; CE  | M      | C       | DS        | -         | 6–16×/3–7× monocytes/no RBCs | Orthopedics, surgeries |
| Regenexx-SCP (Regenex, USA) | FDA      | M      | C       | SS        | -         | 20–40×/low WBCs/low RBCs | Orthopedics, sports med., pain |
| RegenKit (RegenLab, Switzerland) | FDA; CE | M      | SC      | SS        | -/Ca gluconate | 1–2×1.5×10³ WBCs/no RBCs | Orthopedics, OA |
| Res-Q 60 PRP (ThermoGenesis, USA) | FDA   | SA     | C       | SS        | -         | Equivalent to Pure PRP II | Mix with bone graft, cardiovascular |
| Selphyl (Cascade Medical, USA) | FDA; CE | M      | SC      | SS        | CaCl₂     | 1–2×                   | Direct injection; Mix with bone grafts |
| SmartPREP2 (Harvest/Terumo, Japan) | FDA; CE | SA     | SC      | DS        | -         | 4–8×/3–4× WBCs/1× RBCs | Orthopedics |
| TropoCells (Estar Medical, Israel) | FDA; CE | M      | O       | SS        | -         | 4–5×/no WBCs/no RBCs   | Orthopedics, wound healing, dentistry, ophthalmology. |

A, automatic; ACP, Autologous Conditioned Plasma; C, closed; CE, European Union CE mark; DS, double spin; F, filtration; FDA, United States Food and Drug Administration; GPS, Gravitational Platelet Separation System; M, manual; NR, not reported; O, open; OA, osteoarthritis; PEF, platelet enrichment factor ([platelets in PRP]/[platelets in whole blood]); PRP, platelet-rich plasma; RBC, red blood cell; SA, semi-automatic; SC, semi-closed; SS, single spin; WBC, white blood cell.
to medicinal products for human use. The Directive is then assumed and adapted in each Member State according to their own national regulatory frame. For instance, in Spain, the AEMPS (Agencia Española de Medicamentos y Productos Sanitarios, Spanish Agency of Medicines and Health Care Products) published in May 2013 a ‘Report on the Use of Platelet-Rich Plasma’ (INFORME/V1/23052013). As a result, its regulation is much stricter than when it was considered a blood-derived product. It was well received, but surprisingly, with the current European and national regulatory framework, only the preparation procedure, and not the product itself, is regulated, and it does not include any requirements about its composition or its effectiveness. In fact, the AEMPS itself points out that there exists some confusion in this type of autologous product between the medicament production procedures and the medicament itself. In commercial products, a CE marking is sought, to attest to the manufacturer’s declaration that the product complies with the essential requirements of the relevant European health, safety and environmental protection legislation, but this marking

Table 2. Most commonly used PRP classification systems.

| Classification system | Variables | Platelet enrichment |
|-----------------------|-----------|---------------------|
| Mishra and colleagues<sup>14</sup> | Type of PRP | 1: increased WBCs and no activation 2: increased WBCs and activated 3: minimal/no WBCs and no activation 4: minimal/no WBCs and activated |
| | | A: [platelet] above 5× baseline B: [platelet] below 5× baseline |
| PAW system<sup>15</sup> | P [platelets/µl] | A (method of activation) |
| | | P1: [platelet] ≤ baseline P2: baseline < [platelet] < 7.5 · 10⁵ P3: 7.5 · 10⁵ < [platelet] < 1.25 · 10⁶ P4 (> 1.25·10⁶ platelets/µl) |
| | | Exogenous Not exogenous |
| | | W (Presence of WBCs) Above baseline Below baseline |
| Mautner and colleagues<sup>16</sup> | Absolute [platelet] | [Leukocytes] [RBCs] |
| | | Cells/µl + volume injected Including neutrophils if > 1% % Yes/no |
| DEPA<sup>17</sup> | Dose injected | Platelet capture efficiency Purity of PRP [Platelet] with respect to WBC + RBC |
| | | A: [platelet] > 5·10⁹ A: > 90% A > 90% |
| | | B: 3·10⁹ < [platelet] < 5·10⁹ B: 70% to 90% |
| | | C: 10⁹ < [platelet] < 3·10⁹ C: 30% to 70% |
| | | D: [platelet] < 10⁹ D: < 30% |

PRP, platelet-rich plasma; RBC, red blood cell; WBC, white blood cell.
makes no reference to its clinical efficacy.\textsuperscript{21} The above-mentioned EU Directive has not been translated into national law all across Europe, and therefore PRP products are still considered as blood derivatives in some EU countries.

In the United States (US), on the other hand, PRP products are regulated according to the Food and Drug Administration (FDA)’s 21 CFR 1271 of the Code of Regulations, and do not follow the US FDA’s traditional regulatory pathway that includes animal studies and clinical trials.\textsuperscript{22} PRP preparation systems are generally brought to the market through a 510(k) application. This route implies that the device is ‘substantially equivalent’ to another previously cleared and limited to the same indications.\textsuperscript{23} However, it should be noted that, as is the case in EU with the CE label, the clearance applies only to the device and its intended use in an operative setting and makes no claim about its effectiveness for a particular indication.\textsuperscript{23} Evidently, all commercial systems included in Table 1 have achieved a EU CE mark or US FDA 501(k) approval.

Evidence of clinical efficacy of PRP-derived products in musculoskeletal injuries/diseases

The efficacy of PRP-derived products in its various forms (direct liquid/injection, gel, clot, release rate, etc.) either from commercial brands or in-house protocols, has been tested by means of \textit{in vitro} studies, preclinical and clinical trials for oral and maxillofacial surgery, treatment of chronic ulcers, ophthalmology, dermatology, and injuries and pathologies associated to tendon, muscle, cartilage and bone, among other fields.\textsuperscript{24–31} Ligament and tendon repair focus the attention of most studies addressing soft tissue regeneration. However, even in this extensively investigated field there is no consensus on PRP efficacy and most authors highlight the need for more rigorous RCTs. A recent review by the Cochrane collaboration evaluated the evidence supporting clinical efficacy in soft tissue injuries. Of the identified trials (19 randomized and quasi-randomized trials), 16 were judged at high or unclear risk of bias.\textsuperscript{32} It was also pointed out that PRP preparation methods lacked standardization and quantification of the PRP applied to the patient. The review concluded that there is currently insufficient evidence to support the use of platelet-rich therapy for treating musculoskeletal soft tissue injuries.

Application of PRP for cartilage regeneration and OA treatment, our field of interest, has been getting more and more attention over the last decade. Comprehensive review of \textit{in vitro} and preclinical validation studies is beyond the scope of this review but can be found elsewhere.\textsuperscript{33} Here we comment on available evidence compiled from five recent meta-analyses and systematic reviews.\textsuperscript{34–38} These included randomized or quasi-randomized clinical trials that evaluated intra-articular (IA) injection(s) of PRP-derived products (commercial or custom protocols) with other IA interventions [hyaluronic acid (HA), corticosteroids (CSs), saline or other] for osteoarthritis (OA) treatment. A total of 19 individual trials were identified in the five reviews but only 9 were level of evidence I RCTs. Some were evaluated in more than one or even in all five reviews. Trial details are summarized in Tables 3 and 4, including the characteristics of the PRP used, the interventions, controls, outcome measures and the main results. The main conclusions of the five reviews consistently favored the use of PRP products over other IA treatments (in most cases HA), in particular in terms of pain improvement up to 12 months, but the global levels of evidence assigned in the reviews varied. Scales to evaluate methodological quality or risk of bias differed among the reviews which might account for these differences. Laudy and colleagues, Anitua and colleagues and Shen and colleagues used some version of the Cochrane collaboration risk of bias tool.\textsuperscript{34–36} Meheux and colleagues used the modified Coleman methodology score and Chang and coworkers used the Jadad scale for RCTs and the Newcastle–Ottawa Scale for quasi-experimental studies.\textsuperscript{37,38}

As a result, for instance, the RCT by Cerza and colleagues, which was included in four out of the five reviews, received different assessments depending of the review selected. Risk of bias was considered high in two of them,\textsuperscript{34,36} while methodological quality was either considered low\textsuperscript{38} or ‘excellent or good’.\textsuperscript{37} In another example, the RCT by Patel and colleagues\textsuperscript{40} was considered to have a high risk of bias in two reviews,\textsuperscript{34,36} while its methodological quality was graded as excellent in the other two.\textsuperscript{37,38}
| Reference | PRP used | PRP characteristics (SS versus DS/Mishra classification/activation) | Intervention (n of injections/time interval, w/ volume, ml) | Control(s) (n injections/time interval, w/ volume, ml) |
|-----------|---------|-------------------------------------------------|-------------------------------------------------|-------------------------------------------------|
| Vaquerizo and colleagues<sup>39</sup> Level I evidence RCT | PRGF-Endoret | LP-PRP SS/4B/CaCl<sub>2</sub> | 3/1/8* | HA [Durolane]: 1/-/NR |
| Patel and colleagues<sup>40</sup> Level I evidence RCT | Custom | LP-PRP SS/4B/CaCl<sub>2</sub> | 1/-/8 | PRP: [2/3/8] Saline: [1/-/8] |
| Filardo and colleagues<sup>41</sup> Level I evidence RCT | Custom | LR-PRP DS/(NR)A/NR | 3/1/5 | HA [Hyalubrix >1500 kDa]: 3/1/NR |
| Cerza and colleagues<sup>42</sup> Level I evidence RCT | ACP | LP-PRP SS/3A/No | 4/1/5.5 | HA [Hyalgan]: 4/1/20 mg |
| Sanchez and colleagues<sup>43</sup> Level I evidence RCT | PRGF-Endoret | LP-PRP SS/4B/CaCl<sub>2</sub> | 3/1/8 | HA [Euflexxa]: 3/1/NR |
| Say and colleagues<sup>44</sup> Prospective comparative clinical study | Custom (PRGF-Endoret protocol) | LP-PRP SS/4B/CaCl<sub>2</sub> | 1/-/2.5 | HA [NR]: 3/1/2.5 |
| Spakova and colleagues<sup>45</sup> Prospective, cohort study with a control group | Custom | LR-PRP Triple S/1B/No | 3/1/3 | HA [Erectus]: 3/1/NR |
| Li and colleagues<sup>46</sup> Article in Chinese | Weigao kit | LP-PRP DS/NR/NR | 3/3/3.5 | HA [Sofast] |
| Filardo and colleagues<sup>47</sup> Level of evidence II Observational study | LP-PRP (PRGF-Endoret protocol) | LP-PRP SS/4B/CaCl<sub>2</sub> | 3/3/5 | LR-PRP [DS/2B/CaCl<sub>2</sub>]: 3/3/5 ml |
| Kon and colleagues<sup>48</sup> Level of Evidence II Prospective comparative study | Custom | LR-PRP DS/2A/CaCl<sub>2</sub> | 3/2/5 | HA [MW: 1000–2900 kDa]: 30 mg/2 ml HA [MW: 500–730 kDa]: 20 mg/2 ml |
| Sanchez and colleagues<sup>49</sup> | PRGF-Endoret | LP-PRP SS/4B/CaCl<sub>2</sub> | 3/1/8 | HA |
| Duymus and colleagues<sup>50</sup> Level of evidence I RCT | Ycellbio kit | LR-PRP SS/1A/No | 2/4/5 | HA [Ostenil Plus]: 1/-/40 mg ozone gas: 4/1/15 ml |

(Continued)
Table 3. (Continued)

| Reference                          | PRP used | PRP characteristics (SS versus DS/Mishra classification/activation) | Intervention (n of injections/time interval, w/volume, ml) | Control(s) (n injections/time interval, w/volume, ml) |
|------------------------------------|----------|---------------------------------------------------------------------|----------------------------------------------------------|------------------------------------------------------|
| Kon and colleagues                               | Custom | LR-PRP DS/2B/CaCl₂                                                       | 3/1/5                                                   | HA (Hyalubrix): 3/1/30 mg                             |
| Forogh and colleagues                 | TUBEX kit | LR-PRP DS/(NR)B/Ca gluconate                                          | 1/-/5                                                  | Depo Medrol (CS): 1/-/40 mg                           |
| Görmeli and colleagues               | Custom | LR-PRP DS/2A/CaCl₂                                                       | 3/1/5                                                   | PRP: 1/-/5 ml [single injection]                       |
|                                     |          |                                                                     |                                                          | HA [Orthovisc]: 3/1/30 mg                             |
|                                     |          |                                                                     |                                                          | Saline: 3/1/NR                                        |
| Montanez-Heredia and colleagues     | Custom | LP-PRP DS/(NR)A/NR                                                      | 3/2/NR                                                  | HA [Adant]: 3/15 d/NR                                 |
| Paterson and colleagues             | Custom | LR-PRP DS/NR/ultraviolet                                               | 3/1/3                                                   | HA [Hylan G-F 20]: 3/1/3ml                             |
| Raeissatat and colleagues           | Rooyagen kit | LR-PRP DS/1A*/No [5.2±1.5]×[4.8±1.8]× baseline values in 1st and 2nd injections | 2/4/4–6                                             | HA [Hyalgan]: 3/1/20 mg                               |
| Smith                               | ACP      | LP-PRP SS/NR/NR                                                         | 3/1/3–8                                                 | Saline: 3/1/3–8 ml                                    |

*i.e. 3/1/8 means three injections at a 1-week interval at 8 ml of PRP each.
CS, corticosteroid; DS, double spin; HA, hyaluronic acid; LP-PRP, leukocyte-poor PRP; LR-PRP, leukocyte-rich PRP; MW, molecular weight; NR, not reported; PRGF, plasma rich in growth factors; PRP, platelet-rich plasma; RCT, randomized clinical trial; SS, single spin.
# is connected to the values immediately below, [5.2±1.5]×[4.8±1.8]× baseline values in 1st and 2nd injections.
According to Mishra classification, PRP is labelled “A” if platelet concentration is “at or above 5 times the baseline” and B otherwise, and in this case, concentration is in the cut-off number.
### Table 4. Outcomes, main results and methodological quality of trials reviewed in various works.34–38

| Reference | Outcome | Main results | 'Risk of bias' according to reviews34–36 | 'Methodological quality' assessment according to reviews37,38 |
|-----------|---------|--------------|------------------------------------------|-----------------------------------------------------------|
| Vaquerizo and colleagues39 Level I evidence RCT | WOMAC score, Lequesne index, adverse events, 24, 48w | The rate of response to PRGF-Endoret was significantly higher for all scores including pain, stiffness, and physical function on the WOMAC, Lequesne index, and OMERACT-OARSI responders at 24 and 48 weeks. | High | Excellent or good |
| Patel and colleagues40 Level I evidence RCT | WOMAC, VAS, patient satisfaction, adverse event before, 6 w, 3m, 6m | There were significant improvements in all WOMAC parameters in both PRP groups up until the final follow up (6 m) | High | Excellent or good |
| Filardo and colleagues41 Level I evidence RCT | IKDC, EQ-VAS, Tegner and KOOS before treatment and at 2, 6 and 12 m | PRP injections offered a significant clinical improvement up to 1 year of follow up. | High | Excellent or good |
| Cerza and colleagues42 Level I evidence RCT | WOMAC score, before treatment and at 4, 12, 24 w | 'Treatment with ACP showed a significantly better clinical outcome than did treatment with HA, with sustained lower WOMAC scores. Treatment with HA did not seem to be effective in the patients with grade III gonarthrosis.' | High | Excellent or good |
| Sanchez and colleagues43 Level I evidence RCT | Normalized WOMAC score, Lequesne index, adverse effects, 6 m | Rate of response [50% decrease in knee pain from baseline to week 24] to PRGF was significantly higher compared with HA. | Low | Excellent or good |
| Say and colleagues44 Prospective comparative clinical study | KOOS, VAS 3, 6 m | KOOS score and visual pain scale were significantly better in the PRP group compared with HA group at 3 and 6 m follow up. | High | - |
| Spakova and colleagues45 prospective, cohort study with a control group | WOMAC score, 11-point numeric rating scale, adverse events, at 3 and 6 m | Comparison of PRP with HA showed significantly better results in WOMAC Index and NRS scores in the former, in both follow-up periods. | High | - |
| Li and colleagues46 [article in Chinese] | IKDC, WOMAC total, Lequesne score, adverse events, at 6 m | There were significant differences in IKDC score, WOMAC score and Lequesne index between pre- and post-injection in both groups. The effectiveness of the PRP group was significantly better than HA at 6 m. | High | - |

(Continued)
| Reference                        | Outcome                                                                 | Main results                                                                                                                                                                                                 | 'Risk of bias' according to reviews[^34-36] | 'Methodological quality' assessment according to reviews[^37,38] |
|--------------------------------|-------------------------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------|---------------------------------------------------------------|
| Filardo and colleagues[^47]    | IKDC, EQ-VAS, Tegner at 2, 6, 12 m                                      | Both groups showed a statistically significant improvement of all clinical scores from preoperative to final follow up. Comparing the two groups, no differences were found in the subjective IKDC, EQ-VAS, or Tegner scores at 2, 6, and 12 m follow up. | High                                         | 5*                                                            |
| Kon and colleagues[^48]         | IKDC and EQ-VAS, adverse events 2 and 6 m                              | There was a statistically significant improvement in all clinical scores from basal to 2 and 6 m follow up in all treatment groups. At 6 m follow up, PRP group showed significantly better IKDC and EQ-VAS results compared with both HA groups. | High                                         | -                                                             |
| Sanchez and colleagues[^49]     | Patients with 40% decrease in WOMAC pain score (5 w)                   | Success rate was 33.3% in the PRGF group and 10% in the control. PRGF showed higher percent reductions in physical function subscale and overall WOMAC at 5 w compared with HA. | -                                            | High                                                         |
| Duymus and colleagues[^50]      | VAS, WOMAC scores and total at 1, 3, 6, 12 m                           | Effective clinical results were achieved in all groups at 1 m. At 6 m, clinical effects continued in the PRP and HA groups. At 12 m, PRP was significantly superior to HA. | -                                            | High                                                         |
| Kon and colleagues[^48]         | IKDC subjective, KOOS, EQ-VAS, Tegner score, ROM, transpatellar circumference, patient satisfaction, adverse events at 2, 6, 12 m | IKDC was significantly better at 12 m compared with baseline for both PRP and HA. No significant differences were found between both groups. | -                                            | Moderate                                                     |
| Forogh and colleagues[^10]      | KOOS, EQ-VAS, ROM, 20-meter walk test, patient satisfaction at 2, 6 m   | Pain relief in the group treated with PRP was significantly higher than in the CS group. PRP was more effective than CSs in improving ADL and QoL. | -                                            | High                                                         |
| Reference                          | Outcome                                | Main results                                                                                           | 'Risk of bias' according to reviews 34–36 | 'Methodological quality' assessment according to reviews 37,38 |
|-----------------------------------|----------------------------------------|--------------------------------------------------------------------------------------------------------|------------------------------------------|-----------------------------------------------------------|
| Görmeli and colleagues51          | EQ-VAS, IKDC                          | Scores of patients treated with one PRP injection or HA were not significantly different. Scores after three PRP injections resulted in significantly better scores than those of the other groups. | -                                        | High                                                     |
| Montanez and colleagues52         | VAS, KOOS, EuroQoL, adverse events     | Both PRP and HA treatments improved pain in knee OA patients. There were no significant differences among groups. | -                                        | High                                                     |
| Paterson and colleagues53         | VAS, KOOS, knee-QoL, functional tests, | No significant difference with the HA group.                                                          | -                                        | Moderate                                                 |
| Raeissadat and colleagues54       | WOMAC total score, SF-36 at 52 w       | WOMAC mean pain was significantly reduced in both groups at 52w and PRP group was significantly better than the HA group*. Physical function, stiffness, and total WOMAC were only significantly improved in the PRP group. *Note that baseline WOMAC pain, function and total scores were already significantly different at baseline. | -                                        | High                                                     |
| Smith55                           | WOMAC score, adverse events at 1, 2 w, | In the ACP group WOMAC scores were significantly decreased starting at 1 w. The decrease remained significant throughout the study. Scores in the ACP group were statistically better than those of the placebo group starting at 2 weeks throughout. | -                                        | Moderate                                                 |

51 Jadad scale was used to assess the quality of RCT. The aggregate scores ranged from 0 to 5 points. Trials with scores <3 were assumed to have a lower methodological quality.
52 Newcastle–Ottawa Scale was used to assess the quality of selection, comparability, exposure, and outcome of prospective follow up and quasi-experimental studies. The maximum scores observed were 9 points, and total scores <4 points were considered low in quality.

ACP, Autologous Conditioned Plasma; ADL, Activities of Daily Living; CS, corticosteroid; EQ-VAS, EuroQol-visual analogue scale; HA, hyaluronic acid; IKDC, International Knee Documentation Committee; KOOS, knee injury and osteoarthritis outcome score; NR, Not reported; NRS, Numeric Rating Scale; OA, osteoarthritis; PRGF, plasma rich in growth factors; PRP, platelet-rich plasma; QoL, quality of life; ROM, Range of Movement; WOMAC, The Western Ontario and McMaster Universities Osteoarthritis Index.

Table 4. (Continued)
The clinical trial performed by Sanchez and colleagues was generally well considered by all reviews, with low or moderate risk of bias, and with 'excellent or good' methodological quality and with the best score for RCTs.

Additional relevant points consistently raised in the reviews were that more high quality RCTs are needed and that PRP formulations need to be standardized to allow comparison across studies.

At present, more clinical trials are underway. At least 60 clinical trials can be found in the EU (clinicaltrialsregister.eu) and US (clinicaltrials.gov) trial registries with the search terms ‘platelet-rich plasma’ and ‘osteoarthritis’ (access date: March 4, 2018) from which further results will hopefully soon be available.

Proposal for a new approach

As mentioned, in light of the above results, current evidence seems to favor PRP over other IA treatments (HA or CSs) for the treatment of OA. However, the clinical efficacy of PRP therapy remains an open discussion. Here we advocate for a change in the focus of the field to consider PRP (and derived products) as conventional drugs, in the sense that compositions (not only platelet concentrations or preparation methods), doses, and indications with demonstrated clinical efficacy, are established, and all regardless of the patient basal platelet count. We believe this approach would facilitate the demonstration of its clinical efficacy, and therefore is more likely to result in clinically effective products.

This would involve defining additional variables for each application, which could be classified into those related to product characterization and dosage, and those related to PRP administration procedure:

[A]. Product characterization and dosage [what, how much]

Product description should include volume administered and platelet absolute concentration, concentration of WBCs (including neutrophils) and RBCs, and (optimally) GFs. The concentrations of platelets, WBCs and RBCs are fairly easy to determine, but routine quantification of GF concentration is more troublesome. Further, due to the highly complex nature of PRP, it is not fully understood which of the many factors/proteins in its composition are responsible for its effects. The most abundant GFs [platelet-derived growth factor (PDGF)-BB, transforming growth factor (TGF)-β1, vascular endothelial growth factor (VEGF), and epidermal growth factor (EGF)] are the most probable candidates, but there are not many mechanistic studies that have looked at this. As an example, Bendinelli and colleagues showed that PRP anti-inflammatory effects on human chondrocytes are the result of an inhibition of nuclear factor (NF)-κB transactivation mediated by hepatocyte growth factor (HGF) using a specific competitive inhibitor of HGF (an HGF-antagonist/angiogenesis inhibitor, NK4). Some research supports that platelet concentration is correlated with the concentration of GFs. Positive correlations have been found between platelet concentration and PDGF-AB and TGF-β1; VEGF; and EGF and PDGF-BB. But some other studies found poor or no correlations. Therefore, standardizing the platelet dose administered might not warrant equivalent efficacy.

In this context, as with cellular therapy products and biologicals, the application of potency assays may play a key role in defining the quality of PRP-derived products and assuring its efficacy for the desired indication. Potency is the specific ability or capacity of a product, as indicated by appropriate laboratory tests, to effect a given result. In the case of PRP-derived products, given their high complexity, a combination of multiple biological or analytical assays (i.e. an assay matrix) should be considered. Ideally, these assays should be rapid, sensitive, reproducible and cost-effective. This approach has already been used, for example, to evaluate the potential benefit of PRP treatment in patients with long bone nonunion, and to establish a relevant angiogenic potency assay for a commercial product. Normalizing the use of potency assays in commercial PRP products may be helpful to fully characterize available PRP products and evaluate their different biological effects (anti-inflammatory, chondroprotective, proliferative effects, etc.) to select the most appropriate product for each indication.

[B]. Procedure variables [when, where, how, how often]

Variables such as proper timing, treatment periodicity, location and technique for injection would need to be selected to establish efficacy in each
indication. Clinical trials included in Tables 3 and 4 are very dissimilar with respect to these details.

Interventions vary from a single injection,\textsuperscript{10,40,44} two monthly injections,\textsuperscript{54} three injections at 15-day intervals\textsuperscript{48,52} or 21-day intervals;\textsuperscript{41,46} however, the most frequent treatment strategy is to apply 3-weekly PRP injections.\textsuperscript{39,43,45,47,49,51,53,55} With respect to indications, inclusion criteria were also diverse across studies. It is relevant that some trials discriminated between patients with early and consolidated OA using the Kellgren-Lawrence (K–L) grading system,\textsuperscript{48,52} the Ahlbäck classification\textsuperscript{40} or cartilage degeneration by magnetic resonance imaging.\textsuperscript{41} These trials have consistently found that PRP, compared with HA, showed better performance in younger patients affected by cartilage lesions or early OA than in older patients or with more advanced pathology.\textsuperscript{40,41,48,52}

But, in addition, it should be taken into account that OA is a very heterogeneous and multifactorial pathology. It is now known that in OA patients who present the same symptoms (disease), the underlying mechanisms causing them (illness) might be different.\textsuperscript{70} Different OA phenotypes are emerging that will most likely require different treatment approaches. Notably, mechanical, inflammatory and metabolic phenotypes have been proposed.\textsuperscript{71} Therefore, PRP might be effective for some patients but not others. In particular, treatment with PRP-derived products might only be effective in patients with ‘inflammatory’ or ‘mechanical’ OA due to PRP’s anti-inflammatory and regenerative potential suggested by existing preclinical and clinical trials. Clinical efficacy might be easier to demonstrate if OA patients are stratified.

Location and technique for the injection, as well as post-injection recommendations are, also very diverse. As for location and technique, various approaches have been used including lateral,\textsuperscript{41,45,48} supero-lateral,\textsuperscript{10,42} para-patellar,\textsuperscript{55} and lateral mid-patellar,\textsuperscript{54} among others. After-injection recommendations also vary, including rest (10 or 20 min of immobilization\textsuperscript{40,51,54}); movement (flexion and extension, 5 min\textsuperscript{39}); passive flexion and extension 10 times, followed by 10 min rest in the supine position\textsuperscript{10} or 15–20 min of rest, followed by active knee flexing and extending so PRP could spread evenly.\textsuperscript{54} But this variability is not likely to affect efficacy.

**Novel approaches: allogenic PRPs**

At least three recent works have described the use of allogenic PRP clinically. Smrke and colleagues performed a case study in which a 50-year-old male with type 2 diabetes suffering from a comminuted fracture of the tibia and delayed union was treated with a graft composed of allogenic platelet gel mixed with autologous cancellous bone.\textsuperscript{72} To avoid immunogenic reactions, the allogeneic PC was ABO- and RhD-matched, leukocyte-depleted, irradiated and activated by human thrombin. At week 14 after the procedure, no platelet or human leukocyte antigen (HLA) class I antibodies were detected. At 12 months after the procedure, the non-union was resolved, and full load-bearing was achieved.

As a follow up, a prospective clinical study was conducted by the same group to treat long bone nonunions using the same type of allogenic product (allogenic platelet gel and autologous cancellous bone) in nine patients.\textsuperscript{73} They used random single-donor allogenic PCs [ABO and RhD-matched, serologically HIV, hepatitis B virus, hepatitis C virus and lues-negative, leukocyte-depleted, and irradiated] from standard blood bank stocks. As in the case study, screening for HLA antibodies class I, and human platelet antibodies was performed before implantation and after 3 months, also without detecting any sign of immunologic reactions. At 1 year after surgery, seven out of the nine patients treated achieved complete healing.

More recently, Bottegoni and colleagues performed a prospective open-label, uncontrolled, single-center, pilot study with 60 patients.\textsuperscript{74} Participating patients (aged 65–86 years) suffered symptomatic early or moderate knee OA (Ahlbäck grade I–III) and were affected by hematologic disorders, preventing autologous PRP treatment. Effectiveness, as measured with the International Knee Documentation Committee (IKDC), knee injury and osteoarthritis outcome score (KOOS) and EuroQol-visual analogue scales (EQ-VAS), was varied. As noted in other trials, younger patients with lower degree of degeneration showed a better response. In addition, they did
not report any severe complications related to the allogenic nature of the PRP.

Interestingly, a multicentric study was recently performed to standardize a clinical grade procedure for the preparation of allogeneic PCs from umbilical cord blood. As mentioned by the authors, many umbilical cord blood units donated for hematopoietic transplant are deemed unsuitable due to low stem cell content. This prompted some banks from the Italian Cord Blood Network to perform a cooperation program among delivery rooms, cord blood banks and blood transfusion services to develop this standardized protocol, scalable to other national centers, or for translation to other countries.

Other experts have also highlighted that this avenue ought to be explored in future research as a way to deliver PRP therapy to the highest possible number of patients, including those with hematologic disorders or elderly patients.

Still, important issues need to be resolved before the use of this type of product could be generalized. Most notably, safety issues, including what measures need to be taken to eliminate the allergenic potential and to inactivate all pathogens; and secondly, regulatory issues, such as the global consideration of PRP as a drug/medicament for human use.

Final remarks/conclusions
In this work, we have reviewed current available evidence supporting the use of PRP derivatives to treat OA. At present, results from randomized clinical trials seem to favor PRP used over other IA treatments such as HA injections, to improve pain scales in the short and medium term (6–12 months). However, poor methodological quality of available trials and variability in PRP preparations, confound definite demonstration of clinical efficacy. Here we have proposed an alternative approach in which PRPs are considered as conventional drugs for human use. That is, compositions (not platelet concentrations or preparation methods), doses and indications for which clinical efficacy can be demonstrated are identified, regardless of the individual patient basal platelet count.

Additionally, RCTs in which OA patients were stratified suggest that PRP is more effective in those with lower degree of cartilage degeneration or OA grade. Therefore, careful patient stratification, even with respect to OA phenotype, should be encouraged for all trials since it might facilitate proving clinical efficacy for each indication.

The use of allogenic PRP should also be explored, provided safety and regulatory issues can be resolved satisfactorily in order to make PRP available for the highest number of patients possible.

Concluding, at present the therapeutic potential of PRP products in OA remains unfulfilled and without further standardization its clinical efficacy will remain an open debate.

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Conflict of interest statement
The authors declare that there is no conflict of interest.

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References
1. Dovlatova N. Current status and future prospects for platelet function testing in the diagnosis of inherited bleeding disorders. Br J Haematol 2015; 170: 150–161.
2. Di Michele M, Van Geet C and Freson K. Recent advances in platelet proteomics. *Expert Rev Proteomics* 2012; 9: 451–466.

3. Idzko M, Pitchford S and Page C. Role of platelets in allergic airway inflammation. *J Allergy Clin Immunol* 2015; 135: 1416–1423.

4. Speth C, Rambach G, Wuerzner R, et al. Complement and platelets: mutual interference in the immune network. *Mol Immunol* 2015; 67: 108–118.

5. Weyrich AS. Platelets: more than a sack of glue. *Hematology Am Soc Hematol Educ Program* 2014: 400–403.

6. Whitman DH, Berry RL and Green DM. Platelet gel: an autologous alternative to fibrin glue with applications in oral and maxillofacial surgery. *J Oral Maxillofac Surg* 1997; 55: 1294–1299.

7. Matras H. Effect of various fibrin preparations on reimplantations in the rat skin. *Osterr Z Stomatol* 1970; 67: 338–359.

8. Marx RE, Carlson ER, Eichstaedt RM, et al. Platelet-rich plasma - Growth factor enhancement for bone grafts. *Oral Surg Oral Med Oral Pathol Oral Radiol Endod* 1998; 85: 638–646.

9. Dhurat R and Sukesh M. Principles and methods of preparation of platelet-rich plasma: a review and author’s perspective. *J Cutan Aesthet Surg* 2014; 7: 189–197.

10. Forogh B, Mianehsaz E, Shoaee S, et al. Effect of single injection of platelet-rich plasma in comparison with corticosteroid on knee osteoarthritis: a double-blind randomized clinical trial. *J Sports Med Phys Fitness* 2016; 56: 901–908.

11. Ayhan E, Kesmezacar H and Akgun I. Intraarticular injections (corticosteroid, hyaluronic acid, platelet rich plasma) for the knee osteoarthritis. *World J Orthop* 2014; 5: 351–361.

12. Fioravanti C, Frustaci I, Armellin E, et al. Autologous blood preparations rich in platelets, fibrin and growth factors. *Oral Implantol* 2015; 8: 96–113.

13. Mazzocca AD, McCarthy MBR, Chowaniec DM, et al. Platelet-rich plasma differs according to preparation method and human variability. *J Bone Joint Surg Am* 2012; 94A: 308–316.

14. Mishra A, Harmon K, Woodall J, et al. Sports medicine applications of platelet rich plasma. *Curr Pharm Biotechnol* 2012; 13: 1185–1195.

15. DeLong JM, Russell RP and Mazzocca AD. Platelet-rich plasma: the PAW classification system. *Arthroscopy* 2012; 28: 998–1009.

16. Mautner K, Malanga GA, Smith J, et al. A call for a standard classification system for future biologic research: the rationale for new PRP nomenclature. *PM&R* 2015; 7: S53–S59.

17. Magalon J, Chateau AL, Bertrand B, et al. DEPA classification: a proposal for standardising PRP use and a retrospective application of available devices. *BMJ Open Sport Exerc Med* 2016; 2: e000060.

18. Directive 2001/83/EC of the European Parliament and of the Council of 6 November 2001 on the Community code relating to medicinal products for human use. (Official Journal L 311, 28.11.2001).

19. INFORME/V1/23052013. Informe de la Agencia Española de Medicamentos y Productos Sanitarios sobre el uso de plasma rico en plaquetas. Agencia Española de Medicamentos y Productos Sanitarios, https://www.aemps.gob.es/medicamentosUsoHumano/medSitucionesEspeciales/docs/PRP-AEMPSDLDEF-mayo13.pdf (2013).

20. Anitua E, Prado R and Orive G. Closing regulatory gaps: new ground rules for platelet-rich plasma. *Trends Biotechnol* 2015; 33: 492–495.

21. Council Directive 93/42/EEC of 14 June 1993 concerning medical devices. (Official Journal L 169, 12.07.1993).

22. 21 CFR 1271 – Human cells, tissues, and cellular and tissue-based products. (Code of Federal Regulations, 01.04.2011).

23. Beitzel K, Allen D, Apostolakos J, et al. US definitions, current use, and FDA stance on use of platelet-rich plasma in sports medicine. *J Knee Surg* 2015; 28: 29–33.

24. He Y, Chen J, Huang Y, et al. Local application of platelet-rich fibrin during lower third molar extraction improves treatment outcomes. *J Oral Maxillofac Surg* 2017; 75: 2497–2506.

25. Sriram S, Sankaralingam R, Mani M, et al. Autologous platelet rich plasma in the management of non-healing vasculitic ulcers. *Int J Rheum Dis* 2016; 19: 1331–1336.

26. Alio JL, Rodriguez AE, Ferreira-Oliveira R, et al. Treatment of dry eye disease with autologous platelet-rich plasma: a prospective, interventional, non-randomized study. *Ophthalmol Ther* 2017; 6: 285–293.

27. Ibrahim ZA, El-Ashmawy AA and Shora OA. Therapeutic effect of microneedling and autologous platelet-rich plasma in the treatment of atrophic scars: a randomized study. *J Cosmet Dermatol* 2017; 16: 388–399.
28. Yan R, Gu Y, Ran J, Hu Y, et al. Intra-tendon delivery of leukocyte-poor platelet-rich plasma improves healing compared with leukocyte-rich platelet-rich plasma in a rabbit Achilles tendinopathy model. *Am J Sports Med* 2017; 45: 1909–1920.

29. Reurink G, Goudsward GJ, Moen MH, et al. Rationale, secondary outcome scores and 1-year follow up of a randomised trial of platelet-rich plasma injections in acute hamstring muscle injury: the Dutch Hamstring Injection Therapy study. *Br J Sports Med* 2015; 49: 1206–1212.

30. Sengul AT, Buyukkarabacak YB, Altunkaynak BZ, et al. Effects of platelet-rich plasma on cartilage regeneration after costal cartilage resection: a stereological and histopathological study. *Acta Chir Belg* 2017; 117: 21–28.

31. Namazi H and Mehbudi A. Investigating the effect of intra-articular PRP injection on pain and function improvement in patients with distal radius fracture. *Orthop Traumatol Surg Res* 2016; 102: 47–52.

32. Moraes VY, Lenza M, Tamaoki MJ, et al. Platelet-rich therapies for musculoskeletal soft tissue injuries. *Cochrane Database Syst Rev* 2014, Issue 4. Art. No.: CD010071. DOI: 10.1002/14651858.CD010071.pub3

33. Xie X, Zhang C and Tuan RS. Biology of platelet-rich plasma and its clinical application in cartilage repair. *Arthritis Res Ther* 2014; 16: 204.

34. Laudy AB, Bakker EW, Rekers M, et al. Efficacy of platelet-rich plasma injections in osteoarthritis of the knee: a systematic review and meta-analysis. *Br J Sports Med* 2015; 49: 657–672.

35. Anitua E, Sanchez M, Javier Aguirre J, et al. Efficacy and safety of plasma rich in growth factors intra-articular infiltrations in the treatment of knee osteoarthritis. *Arthroscopy* 2014; 30: 1006–1017.

36. Shen L, Yuan T, Chen S, et al. The temporal effect of platelet-rich plasma on pain and physical function in the treatment of knee osteoarthritis: systematic review and meta-analysis of randomized controlled trials. *J Orthop Surg Res* 2017; 12.

37. Meheux CJ, McCulloch PC, Lintner DM, et al. Efficacy of intra-articular platelet-rich plasma injections in knee osteoarthritis: a systematic review. *Arthroscopy* 2016;32:495–505

38. Chang KV, Hung CY, Aliwarga F, et al. Comparative effectiveness of platelet-rich plasma injections for treating knee joint cartilage degenerative pathology: a systematic review and meta-analysis. *Arch Phys Med Rehabil* 2014; 95: 562–575.

39. Vaquerizo V, Angel Plasencia M, et al. Comparison of intra-articular injections of plasma rich in growth factors (PRGF-Endoret) versus durolane hyaluronic acid in the treatment of patients with symptomatic osteoarthritis: a randomized controlled trial. *Arthroscopy* 2013; 29: 1635–1643.

40. Patel S, Dhillon MS, Aggarwal S, et al. Treatment with platelet-rich plasma is more effective than placebo for knee osteoarthritis a prospective, double-blind, randomized trial. *Am J Sports Med* 2013; 41: 356–364.

41. Filardo G, Kon E, Ruiz MTP, et al. Platelet-rich plasma intra-articular injections for cartilage degeneration and osteoarthritis: single- versus double-spinning approach. *Knee Surg Sports Traumatol Arthrosc* 2012; 20: 2078–2087.

42. Cerza F, Carni S, Carcangiu A, et al. Comparison between hyaluronic acid and platelet-rich plasma, intra-articular infiltration in the treatment of gonarthrosis. *Am J Sports Med* 2012; 40: 2822–2827.

43. Sanchez M, Fiz N, Azofra J, et al. A randomized clinical trial evaluating plasma rich in growth factors (PRGF-Endoret) versus hyaluronic acid in the short-term treatment of symptomatic knee osteoarthritis. *Arthroscopy* 2012; 28: 1070–1078.

44. Say F, Guler D, Yener K, et al. Platelet-rich plasma injection is more effective than hyaluronic acid in the treatment of knee osteoarthritis. *Acta Chir Orthop Traumatol Cech* 2013; 80: 278–283.

45. Spakova T, Rosocha J, Lacko M, et al. Treatment of knee joint osteoarthritis with autologous platelet-rich plasma in comparison with hyaluronic acid. *Am J Phys Med Rehabil* 2012; 91: 411–417.

46. Li M, Zhang C, Ai Z, et al. Therapeutic effectiveness of intra-knee-articular injection of platelet-rich plasma on knee articular cartilage degeneration. *Zhongguo Xiu Fu Chong Jian Wai Ke Za Zhi* 2011; 25: 1192–1196.

47. Filardo G, Kon E, Di Martino A, et al. Platelet-rich plasma vs hyaluronic acid to treat knee degenerative pathology: study design and preliminary results of a randomized controlled trial. *BMC Musculoskel Disord* 2012; 13: 229.

48. Kon E, Mandelbaum B, Buda R, et al. Platelet-rich plasma intra-articular injection versus hyaluronic acid viscosupplementation as treatments for cartilage pathology: from early
49. Sanchez M, Anitua E, Azofra J, et al. Intra-articular injection of an autologous preparation rich in growth factors for the treatment of knee OA: a retrospective cohort study. *Clin Exp Rheumatol* 2008; 26: 910–913.

50. Duymus TM, Mutlu S, Dernek B, et al. Choice of intra-articular injection in treatment of knee osteoarthritis: platelet-rich plasma, hyaluronic acid or ozone options. *Knee Surg Sports Traumatol Arthrosc* 2017; 25: 485–492.

51. Görmeli G, Görmeli CA, Ataoglu B, et al. Multiple PRP injections are more effective than single injections and hyaluronic acid in knees with early osteoarthritis: a randomized, double-blind, placebo-controlled trial. *Knee Surg Sports Traumatol Arthrosc* 2017; 25: 958–965.

52. Montanez-Heredia E, Irizar S, Huertas PJ, et al. Intra-articular injections of platelet-rich plasma versus hyaluronic acid in the treatment of osteoarthritic knee pain: a randomized clinical trial in the context of the Spanish national health care system. *Int J Mol Sci* 2016, 17, 1064; doi:10.3390/ijms17071064

53. Paterson KL, Nicholls M, Bennell KL, et al. Intra-articular injection of photo-activated platelet-rich plasma in patients with knee osteoarthritis: a double-blind, randomized controlled pilot study. *BMC Musculoskel Disord* 2016; 17: 67.

54. Raeissadat SA, Rayegani SM, Hassanabadi H, et al. Knee Osteoarthritis injection choices: platelet-rich plasma (PRP) versus hyaluronic acid (a one-year randomized clinical trial). *Clin Med Insights Arthritis Musculoskelet Disord* 2015; 8: 1–8.

55. Smith PA. Intra-articular autologous conditioned plasma injections provide safe and efficacious treatment for knee osteoarthritis: an FDA-sanctioned, randomized, double-blind, placebo-controlled clinical trial. *Am J Sports Med* 2016; 44: 884–891.

56. Bendinelli P, Matteucci E, Dogliotti G, et al. Molecular basis of anti-inflammatory action of platelet-rich plasma on human chondrocytes: mechanisms of NF-kappa B inhibition via HGF. *J Cell Physiol* 2010; 225: 757–766.

57. Magalon J, Bausset O, Serratrice N, et al. Characterization and comparison of 5 platelet-rich plasma preparations in a single-donor model. *Arthroscopy* 2014; 30: 629–638.

58. Castillo TN, Pouliot MA, Kim HJ, et al. Comparison of growth factor and platelet concentration from commercial platelet-rich plasma separation systems. *Am J Sports Med* 2011; 39: 266–271.

59. Sundman EA, Cole BJ and Fortier LA. Growth factor and catabolic cytokine concentrations are influenced by the cellular composition of platelet-rich plasma. *Am J Sports Med* 2011; 39: 2135–2140.

60. Eppley BL, Woodell JE and Higgins J. Platelet quantification and growth factor analysis from platelet-rich plasma: implications for wound healing. *Plast Reconstr Surg* 2004; 114: 1502–1508.

61. Weibrich G, Kleis WKG, Hafner G, et al. Growth factor levels in platelet-rich plasma and correlations with donor age, sex, and platelet count. *J Craniofaciof Surg* 2002; 30: 97–102.

62. Andrea Gomez L, Escobar M and Penuela O. Standardization of a protocol for obtaining platelet rich plasma from blood donors; a tool for tissue regeneration procedures. *Clin Lab* 2015; 61: 973–980.

63. Bravery CA, Carmen J, Fong T, et al. Potency assay development for cellular therapy products: an ISCT review of the requirements and experiences in the industry. *Cytotherapy* 2013; 15: 9–19.

64. Hematti P. Characterization of mesenchymal stromal cells: potency assay development. *Transfusion* 2016; 56: 32S–33S.

65. Basu J and Ludlow JW. Cell-based therapeutic products: potency assay development and application. *Regen Med* 2014; 9: 497–512.

66. Guthrie K, Bruce A, Sangha N, et al. Potency evaluation of tissue engineered and regenerative medicine products. *Trends Biotechnol* 2013; 31: 505–514.

67. Anitua E, Prado R and Orive G. PRP therapies – Is it time for potency assays? Letter to the Editor. *Am J Sports Med* 2016; 44: NP63–NP64.

68. Perut F, Dallari D, Rani N, et al. Cell-based assay system for predicting bone regeneration in patient affected by aseptic nonunion and treated with platelet rich fibrin. *Curr Pharm Biotechnol* 2016; 17: 1079–1088.

69. Bloom DD, Centanni JM, Bhatia N, et al. A reproducible immunopotency assay to measure mesenchymal stromal cell-mediated T-cell suppression. *Cytotherapy* 2015; 17: 140–151.

70. Kraus VB, Blanco FJ, Englund M, et al. Call for standardized definitions of osteoarthritis and risk stratification for clinical trials and clinical use. *Osteoarthritis Cartilage* 2015; 23: 1233–1241.
71. Berenbaum F. Osteoarthritis as an inflammatory disease (osteoarthritis is not osteoarthrosis!). *Osteoarthritis Cartilage* 2013; 21: 16–21.

72. Smrke D, Gubina B, Domanovic D, *et al.* Allogeneic platelet gel with autologous cancellous bone graft for the treatment of a large bone defect. *Eur Surg Res* 2007; 39: 170–174.

73. Gubina B, Rožman P, Bišcević M, *et al.* The influence of allogeneic platelet gel on the morphology of human long bones. *Coll Antropol* 2014; 38: 865–870.

74. Bottegoni C, Dei Giudici L, Salvemini S, *et al.* Homologous platelet-rich plasma for the treatment of knee osteoarthritis in selected elderly patients: an open-label, uncontrolled, pilot study. *Ther Adv Musculoskelet Dis* 2016; 8: 35–41.

75. Rebulla P, Pupella S, Santodirocco M, *et al.* Multicentre standardisation of a clinical grade procedure for the preparation of allogeneic platelet concentrates from umbilical cord blood. *Blood Transfusion* 2016; 14: 73–79.

76. Anitua E, Prado R and Orive G. Allogeneic platelet-rich plasma: at the Dawn of an off-the-shelf therapy? *Trends Biotechnol* 2017; 35: 91–93.

77. Andia I and Abate M. Platelet-rich plasma in the treatment of skeletal muscle injuries. *Expert Opin Biol Ther* 2015; 15: 987–999.