



**Improve patient health by participating in the IOF Registry! Open to qualified and dedicated physicians to advance patient care.**

The IOF Patient Database Registry is a confidential and secure, web-based data collection system that tracks patient-reported outcomes and complications from patient surveys. It was established to provide physicians with valuable information to assist in making long-term decisions regarding interventional orthopedic treatments, such as platelet rich plasma (PRP) and bone marrow derived stem cells.

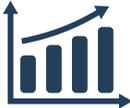
**7** **CURRENT STUDIES**  
Hip, Knee, Shoulder & Spine/Ankle/Foot,  
Elbow/Wrist/Hand, General Ortho

|                              |                         |                         |                            |
|------------------------------|-------------------------|-------------------------|----------------------------|
| <b>7,006</b>                 | <b>28,834</b>           | <b>40</b>               | <b>260</b>                 |
| <b>TOTAL ACTIVE PATIENTS</b> | <b>TOTAL PROCEDURES</b> | <b>TOTAL PHYSICIANS</b> | <b>DATA POINTS TRACKED</b> |

**BENEFITS TO PARTICIPATING PHYSICIANS**

- Learn the true benefit and possible complications of percutaneous orthopedic therapies
- Unlimited potential to present new data through presentations, publications (data results, insights and case studies) and posters
- Make informed decisions by identifying new areas for quality improvement



**DATA WE COLLECT** 

- Patient Candidacy Grade
- Pain Scale
- Body part specific, validated functional questionnaires such as LEFS, IKDC, Oxford, DASH, and FRI
- Procedure details including the following injectates: Bone Marrow, Platelet Rich Plasma, Platelet Lysate, Fat Graft, Prolotherapy & Others
- Patient reported complications/adverse events and illnesses
- Outcomes (i.e., Improvement %) and perspectives that are crucial to understanding how we can further ensure safety and improve the care delivered

 **PATIENT FOLLOW-UP METHOD**

Our program allows patients to participate via online surveys, which can be completed in the comfort and privacy of their home or office. Communicating through an email address the IOF sends follow-up surveys via an online portal. Follow-up survey intervals are emailed at 1, 3, 6, 12, 18, 24 months and every year thereafter up to 20 years. The IOF Registry allows participating patients the opportunity to report their health status relating to:

- Quality of Life
- Joint Function and Pain
- Activity Levels
- Long-Term Procedure Success

Through the voluntary participation of Registry members, questions about percutaneous orthopedic procedures can be answered, enabling a more thorough understanding of these non-invasive treatment options.