INFORMED CONSENT

Adipose Derived Stem Cell & Platelet Rich Plasma

l,	have been advised and consulted about the injection technique of adipose
derived stem cells and of platelet rich plasma for	or the treatment of orthopedic conditions.
I understand and voluntarily consent an	nd authorize the following procedure: re-injection of my own adipose derived
stem cells to treat joint, tendon, or ligament pa	ain. I understand the procedure may require follow up treatments.
I have been informed that even though t	this is not yet a FDA approved procedure, this procedure has been used safely
and successfully on other patients.	
I have been advised that stem cell injecti	ion treatments are used to tighten and strengthen weak and damaged ligaments
and tendons which are believed to cause pain a	and instability. It is also used to decrease pain and improve function in some
forms of arthritis. The technique requires the in	injection of stem cells derived from my own adipose tissue according to standard
fat harvesting and injection techniques. The sit	te of injection is where the ligament or tendon attaches to the bone, at the joint
capsule, or inside the joint.	
I have been advised that the procedure r	may initially increase the painful area or reproduce symptoms for one to three
days (and occasionally, as long as ten days), and	d then may decrease in intensity, but may not completely eradicate my
symptoms.	
I have been advised that platelet rich pla	asma is an established treatment technique used to tighten and strengthen weak
and damaged ligaments and tendons which are	e believed to cause pain and instability. It is also used to decrease pain and
improve function in some forms of arthritis. Th	ne technique requires the injection of platelet rich plasma derived from my own
blood according to standard blood collection ar	nd injection techniques. The site of the injection is where the ligament or tendor
attaches to the bone, at the joint capsule, or in:	side the joint.
I have been informed that the procedure	e has been used on many patients and has been proven safe. The procedure
may initially increase the painful area or reprod	duce symptoms fro one to three days (and occasionally, as long as ten days), and
then may decrease in intensity, but may not co	mpletely eradicate my symptoms.
I understand the possible benefits of the	e procedure are to improve or resolve pain and improve function. I acknowledge
that no guarantee has been given by anyone as	s to the results that I may have.
I have been informed that the alternativ	ves to stem cell injections are:
*Surgical Intervention may be a possi	ibility *Injection with steroids
*Manipulation may provide temporar	ry relief *Acupuncture
I have been informed that the risks and	complications of stem cell injections are:
*Immediate pain at injection site	*Stiffness in the injected point
*Bruising	*Allergic reaction
*Infection	*Nerve or muscle injury
*Nausea/Vomiting	*Dizziness or fainting
*Swelling after joint injections	*Bleeding
*Temporary blood sugar increase	*Itching at injection site
I have been informed that the risks of not	t having the treatment are:
*No pain relief	
*Continued instability of the damaged	d joint or ligament and probable worsening of pain
I understand that this procedure is usua	ally not covered by insurance and I am responsible for the total charges.
I certify that I understand all the information	ation above in its entirety, have had my questions answered, and the potential
side effects explained.	
Patients Signature/Date	Consultants Signature/Date
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