



Please place collection kit  
barcode here.

**BLUE FIELDS ARE REQUIRED**

**PATIENT INFORMATION**

<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="checkbox"/> F <input type="checkbox"/> M
Patient Last Name	Patient First Name	Date of Birth (MM/DD/YY)	Biological Sex
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Patient Email	Telephone	Patient Weight (lbs.)	
<input type="text"/>	<input type="text"/>	<input type="text"/>	
Address	City	Country	

**ORDERING CLINICIAN**

<input type="text"/>	<input type="text"/>		
Clinic or Organization	Address		
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
Ordering Clinician Name	Natera® Clinic ID	City	Country
<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
E-mail	Contact Person	Telephone	Fax

**PROSPERA™ TEST ORDERING**

**PROSPERA:** Transplant Assessment

Kidney  SPK  Other \_\_\_\_\_

Date of Sample Collection (MM/DD/YY)

**Sample Requirements:** Two 10mL Tiger-top Streck Cell-Free DNA BCT® blood tubes

Prospera is not indicated in patients who are pregnant, <14 days post-transplant, or have active somatic disease such as cancer that involves large chromosomal events. Prospera is also not indicated for recipients of an allograft from an identical twin, multi-organ allografts with exception of SPK, or an allogeneic stem cell transplant.

**PATIENT HISTORY**

<input type="text"/>	<input type="checkbox"/> Living <input type="checkbox"/> Deceased	<input checked="" type="checkbox"/> Yes <input type="checkbox"/> No	<input type="text"/>
Date of Transplant (MM/DD/YY)	Donor Type	Donor Biologically Related	If related, define relationship

**DISPOSITION OR RETENTION OF SAMPLES**

Laboratory (Reseller) represents and confirms that the patient has given informed consent in compliance with applicable law to Natera's following sample disposition or retention policy: **PATIENT UNDERSTANDS AND CONSENTS THAT:** (i) her/his sample will be sent to the United States for performance of the test; (ii) Natera may retain the patient's leftover, de-identified samples to use for medical and technology advancement, research & development, product validation and quality assurance, independently or in collaboration with third-party partners, either in or outside the United States; and (iii) patient and patients heirs will not receive any payments, benefits, or rights to any resulting products or discoveries.

Check this box if the patient does not consent to (ii).