



Recruiting PV Trials			
<p><b>KRT-232-102</b> (Interventional) <b>Polycythaemia Vera</b>  PI: Prof Adam Mead</p>	<p>A Two-Part, Randomized, Open-label, Multicenter, Phase 2a/2b Study of the Efficacy, Safety, and Pharmacokinetics of KRT-232 Compared to Ruxolitinib in Patients with Phlebotomy-Dependent Polycythemia Vera</p> <p><b>Eligibility Criteria:</b> Age 18 or over, Diagnosis of PV, Phlebotomy dependent Part A: either Splenomegaly <math>\geq 450 \text{ cm}^3</math> or absence of splenomegaly by CT/MRI; Part B: Splenomegaly by CT/MRI, Prior treatment with hydroxyurea (Part A and B) or interferon (Part A only) with resistance/intolerance to HU or IFN, ECOG performance status <math>\leq 2</math></p>	<p><b>MEASURES</b> (Non-interventional) <b>All MPN</b>  PI: Prof Adam Mead</p>	<p>The MPN Experimental Assessment of Symptoms by Utilizing Repetitive Evaluation (MEASURES) Trial: Serial Assessment of Symptomatic Response to Non Experimental Medical Therapies and/or Phlebotomy in Patients with Myeloproliferative Neoplasms</p> <p><b>Eligibility Criteria:</b> All Ph- MPN patients starting a new, therapeutic intervention</p>
<p><b>TAMARIN</b> (Interventional) <b>All MPN</b>  PI: Prof Adam Mead</p>	<p>Effects of TAMoxifen on the Mutant Allele Burden and Disease Course in Patients with Myeloproliferative Neoplasms</p> <p><b>Eligibility Criteria:</b> PV,ET,MF, Stable disease on hydroxyurea, interferon or ruxolitinib, No prior thrombosis, Age over 60, men aged between 50-59 may be considered, women must be post-menopausal</p>	<p><b>INForMeD</b> (Non-interventional) <b>All MPN</b>  PI: Prof Adam Mead</p>	<p>An observational and biological research study to investigate the genetic and cellular basis of sporadic and familial myeloid disorders</p> <p><b>Eligibility Criteria:</b> Age 2 or over, Patients under investigation for or diagnosed with a myeloid or related disorder, Patient willing to give consent to the study</p>
<p><b>PHAZAR</b> (Interventional) <b>Accelerated or blast phase MPN</b>  PI: Prof Adam Mead</p>	<p>A Phase 1b Study to assess the safety and tolerability of oral Ruxolitinib in combination with Azacytidine in patients with Advanced Phase MPN, including MDS or AML arising from MPN</p> <p><b>Eligibility Criteria:</b> Age 16 and over, Diagnosis of ET,PV or MF with <math>&gt; 10\%</math> bone marrow blasts (with or without dysplastic changes), ECOG performance status <math>\leq 3</math></p>	<b>Haematology Research contact details</b>	
		Prof Adam Mead	Email: adam.mead@imm.ox.ac.uk Tel: 01865 222324
		Research Nurse Team	Email: LP.haematology.RN@oxnet.nhs.uk