

Majority of patients were treated with combination therapy of pegylated interferon α 2a and ribavirin whilst a small proportion (28) have received (triple therapy) protease inhibitors. The total number of patients who achieved sustained virologic response (SVR) at the end of treatment were 264 (60.41%). 196 (74.42%) of those were genotype 3a, 57 (21.60%) were genotype 1a/1b and 11 (4.17%) were genotype 2b/2a. 43 (9.84%) were considered non-responders. 49 (11.21%) patients were unable to complete treatment due to critical physical or mental illness with 12 of those (24.50%) have achieved SVR. Patients' feedback for this nurse-led service has been very positive.

Conclusion Specialist nurse-led and clinicians supported hepatitis C service has delivered a high quality of care. Our dedicated specialist nurses working closely with clinicians have achieved high successful treatment rates in such a large cohort of patients.

Disclosure of Interest None Declared.

PTU-121 LLT1 IS UPREGULATED IN HEPATOCELLULAR CARCINOMA AND INHIBITS NK CELL CYTOTOXICITY

N Kumar*, A Mroz, M Aletrari, R Goldin, M Purbhoo. *Imperial College London, London, UK*

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Introduction Liver cancer is on the rise and prognosis is poor partly due to late diagnosis and a lack of systemic chemotherapy. Identification of novel markers allowing early diagnosis of hepatocellular carcinoma (HCC) and to act as targets for immunotherapies is essential to curb rising mortality. Lectin-like transcript 1 (LLT1) interacts with CD161¹ a receptor expressed on almost all NK cells. For the first time, we investigate the expression of LLT1 in the liver and HCC, both *in vivo* and *in vitro*, and determine the effect of LLT1 on NK cell function.

Methods LLT1 expression *in vivo* was determined by immunohistochemistry comparing 10 HCC specimens (resections and biopsies) to 15 normal liver controls. *In vitro* expression was demonstrated using flow cytometry to review HCC cell lines (Huh7, HepG2). Peripheral blood mononuclear cells (PBMCs), including NK cells, from healthy donors were incubated with target cells expressing different levels of LLT1: Huh7 cells, Huh7 cells expressing increased levels of LLT1 (achieved by transfection), and Jurkat cells lacking LLT1. The effect of these incubations on CD161⁺ and CD161-NK cell cytotoxicity was assessed by measuring CD107a expression using flow cytometry. This assay was performed in triplicate using 2 different donors on 3 separate occasions. Statistical analysis of variance (ANOVA) was performed to assess differences in NK cell cytotoxicity between each incubation condition.

Results Hepatocytes in normal and cirrhotic liver do not express LLT1. However, LLT1 is extensively upregulated in HCC *in vivo* with strong, diffuse staining in 9 out of 10 cases, and more focal staining seen in the remaining case. LLT1 is expressed *in vitro* on HCC cell lines (Huh7 and HepG2). CD161⁺NK cells show reduced cytotoxicity, compared to CD161-NK cells, when incubated with Huh7 cells expressing LLT1. Incubation with Huh7 cells correlates with significantly reduced activity by CD161⁺NK cells ($p \leq 0.001$) when compared to CD161⁺NK cell activity against non-LLT1 expressing Jurkat cells. CD161⁺NK cell cytotoxicity is further reduced when incubated with Huh7 cells expressing increased LLT1 levels (as achieved by transfection; $p \leq 0.05$ when compared to CD161⁺NK cell activity when incubated with non-transfected Huh7 cells). In contrast, the activity of CD161⁺ and CD161-NK cells is not

significantly different when the target cell does not express LLT1 (Jurkat cell).

Conclusion We demonstrate for the first time that LLT1 is not expressed by normal liver tissue, but upregulated in HCC. LLT1 inhibits NK cell cytotoxicity, representing a possible mechanism for HCC to evade the immune response to cancer. Therefore, not only may LLT1 be used as a diagnostic marker for HCC, it represents a novel immunotherapy target.

REFERENCE

1 Aldemir *et al.* Cutting edge: LLT1 is a ligand for the CD161 receptor. *J Immunol* 2005:7791

Disclosure of Interest None Declared.

PTU-122 A GMP TREG EXPANSION PROTOCOL RESTORES TREG SUPPRESSOR FUNCTION IN END-STAGE LIVER DISEASE; IMPLICATIONS FOR ADOPTIVE TRANSFER THERAPY

N Safinia*, T Vaikunthanathan, H Fraser, C Scotta, R Lechler, G Lombardi. *MRC Centre for Transplantation, King's College London, London, UK*

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Introduction Long-term survival in liver transplant recipients remains suboptimal because of the morbidity and mortality associated with the long-term use of immunosuppression (IS). However, IS weaning early post liver transplantation (LT) has largely been unsuccessful, supporting the need for active tolerance induction strategies. CD4⁺CD25⁺FOXP3⁺ (Tregs) play an important role in immunoregulation and have been shown in animal models to promote transplantation tolerance. Phase I trials in bone marrow transplantation have shown that *ex vivo* expanded Tregs have an excellent safety profile, which is encouraging for the broader application of these cells. The clinical trial, ThRIL, soon to be initiated at King's College London, aims to investigate the therapeutic potential of Tregs in the setting of LT.

We have devised a GMP compatible protocol that ensures the successful isolation and expansion of a functional and stable human Treg population in preparation for this trial.

Methods Tregs were isolated from 150ml of blood from patients with end-stage liver disease by a CliniMACS-based GMP isolation technique and expanded using anti-CD3/CD28 beads, IL-2 and rapamycin.

Results A 580-fold expansion of pure Tregs was achieved (97.4% CD4⁺CD25⁺ and 0.008% CD8⁺ cells) and the cells maintained FoxP3 expression (99.6% of the CD4⁺CD25⁺ cells express FoxP3). The populations of Tregs obtained were also stable and did not convert to Th17 cells when cultured in the presence of pro-inflammatory stimuli.

This protocol further proved to be ideal for the expansion of Tregs from patients with liver disease in view of restoring the Tregs' suppressive function (1:1 ratio – expanded Tregs 91.1% vs. freshly isolated Tregs 28.6% suppression, 1:10 ratio – 80.7% vs. 20.8% respectively). Based on these findings, we subsequently conducted an in-depth phenotypic characterisation of freshly isolated Tregs in order to delineate a population responsible for the apparent lack of suppressive function. An investigation into the possible mechanisms is currently ongoing.

Conclusion The feasibility of Treg based therapy is now widely accepted, provided that tailor-made clinical grade procedures for isolation and ex-vivo cell handling are available. Our rapamycin-based protocol is ideal in this setting as it not only satisfies the rigours of GMP manufacturing standards, but also