## LYMPHOCYTE RECEPTOR SEQUENCING TO MONITOR ALLOIMMUNE RESPONSE: A SYSTEMATIC REVIEW

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## INTRODUCTION AND RATIONALE:

- We have few reliable ways of monitoring the T-cell response to the kidney donor after transplantation, so are unable to respond pro-actively to developing rejection
- Each of the millions of lymphocyte clones in the recipient recognizes just a single donor antigen target, and these clones expands dramatically as rejection occurs
- Monitoring the expansion of these clones offers a novel approach and a potential new tool to identify rejection at the earliest stage when it can be treated and abrogated
- We have conducted a systematic review to examine current research in lymphocyte receptor sequencing to do explore its potential use for transplant monitoring



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2010

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## CONCLUSIONS AND DEVELOPMENT OF A NEW BIOMARKER FOR KIDNEY TRANSPLANTATION

- Sequencing of the T-cell receptor is now a viable technology to monitor the expansion of T-cell clones and to define their response to highly specific antigen targets
- A very specific set of genes know as the CDR3 region of the T-cell receptor beta chain is the preferred sequencing target due to their relevance in HLA-antigen recognition
- Transplant patients show lower repertoire diversity (making identification potentially easier) and specific expansion of T-cell and B-cell clones which persist after rejection
- Patients who develop tolerance to their donor after transplantation show deletion of these important T-cell clones so do not attack the donor graft
- These data suggest that T-cell receptor monitoring may be an exciting new biomarker for monitoring rejection or tolerance, and a key guide to adjusting immune suppression
- BC has launched detailed studies to confirm this assay and to introduce this new test into our provincial transplant monitoring program, the first such biomarker in Canada

