

Memorial Sloan Kettering Cancer Center

Abstract No: 521

Background

•Cytomegalovirus (CMV) infection is the most common viral infection in allogeneic hematopoietic stem cell transplantation (HCT) recipients • Preemptive therapy (PET) is effective in preventing CMV end-organ disease (EOD) but poses substantial healthcare resource use (HCRU) and cost •Established benchmarks for HCRU and cost in the era of PET are required to perform cost benefit analyses of novel interventions for CMV

Objectives

- •To compare direct all-cause HCRU and cost in CMV R⁺ by receipt of PET (PET versus no PET) through 180 days post HCT (D180)
- •To quantify CMV-related HCRU and cost among PET recipients through D180
- •To compare CMV-related with non CMV related HCRU and cost through D180

Methods

•Study Design: Retrospective cohort study

•**Study Population:** CMV R+ adult allogeneic HCT recipients of first peripheral blood or marrow allograft at a single center from 3/2013 to 12/2017 • Data Sources: Clinical data including PET reasons and dates of

hospitalizations were extracted from the electronic medical records. Inpatient hospital charges were obtained from the Vizient billing database and converted to costs using institutional cost-to-charge ratios, wage index and inflation rate to 2017 US Dollars

•CMV monitoring: CMV+ recipients were monitored weekly by quantitative PCR assay starting on day 14 through D180 and managed by PET per standards of care

•CMV risk: Patients were categorized in two mutually exclusive groups: Low risk (LR) recipients of unmodified graft from matched related or unrelated donors or High risk (HR): recipients of unmodified graft from mismatched donor or ex vivo T-cell depleted graft from any donor.

•**PET group**: Receipt of pre-emptive antiviral therapy for CMV viremia •Follow-up period: Date of Index HCT to the earliest of D180 or death/relapse/2nd transplant, whichever occurred first.

•Healthcare resource utilization:

- •Length of inpatient stay (LOS) for index admission, and subsequent readmissions by day 180 post HCT
- •Number of all-cause and CMV-related readmissions by 180 post HCT •CMV-related readmissions were defined as readmissions for initiation of PET, work up or management of CMV End Organ Disease and any readmission where PET was initiated

Baseline Characteristics

• Of 368 HCT recipients, 192 (52%) were HR and 176 (48%) LR

- Overall, 208 (56.5%) patients received PET
- HR comprised 72% of PET group but only 26% of No PET group. PET Group (n=208) NO PET (n=160)



Economic Burden of Cytomegalovirus Infection in CMV-Seropositive (CMV R⁺) Hematopoietic Stem Cell Transplant Recipients Managed With Pre-Emptive Therapy: A Single Center Experience

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