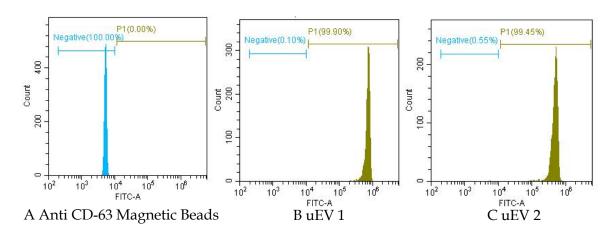
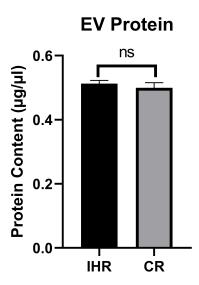


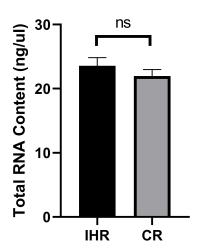
Supplementary Figure 1 Urinary extracellular vesicles (uEVs) Characterisation (*n* **= 3/group)** Representative Nanoparticle tracking analysis (NTA) showing exosome concentration (particles/mL)/size in pellet.A: uEVs isolated using in-house reagent (IHR); B: uEV isolated using commercial Invitrogen Reagent (CR).



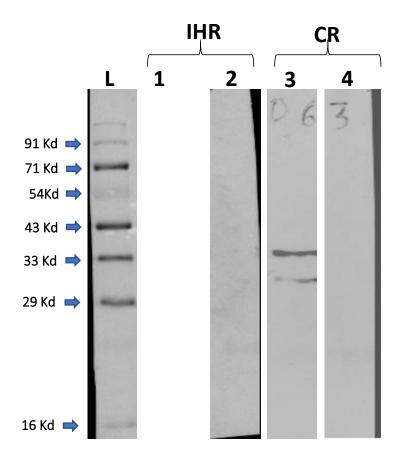
Supplementary Figure 2: Urinary Extracellular Vesicles (uEVs) Characterisation using Flow Cytometry (n=3/group) Representative histogram depicting the expression of CD63 on the surface of uEVs samples (uEV1-using in-house reagent (IHR) and uEV2- using commercial Invitrogen Reagent (CR)) captured on antihuman CD63 magnetic beads using Flow cytometry analysis. A: Unstained antihuman CD63 magnetic beads; B andC: Samples uEV1 and uEV2, captured on antihuman CD63 magnetic beads followed by anti-CD63 primary antibody and stained with detection antibody. Green colour on histogram represents CD63 positive uEV population.



Supplementary Figure 3: Total Urinary Extracellular Vesicles (uEVs) protein content using Pierce TM **BCA Protein assay Kit** Histogram depicting the uEV Total RNA concentration using in-house reagent (IHR) and uEV isolated using commercial Invitrogen Reagent (CR) (n=3/group). Statistical significance calculated *t*-test. ^a*P* value < 0.05, ^b*P* value < 0.01, ^c*P* value < 0.001.



Supplementary Figure 4 Total Urinary Extracellular Vesicles (uEVs) RNA Concentration Histogram depicting the uEV Total RNA concentration using inhouse reagent (IHR) and commercial Invitrogen Reagent (CR) (n=3/group). Statistical significance calculated *t*-test. ^aP value <0.05, ^bP value< 0.01, ^cP value < 0.001.



Supplementary Figure 5 Representative immunoblots showing the expression of CD63 (lane 1 and 3) and Calnexin (lane 2 and 4) in urinary extracellular vesicles (uEVs) isolated using in-house reagent (IHR) and commercial Invitrogen Reagent (CR). L: Protein ladder.

Supplementary Table 1 Shows the average protein concentration of the extracellular vesicle isolated using in-house reagent and commercial invitrogen reagent

| Number | Method employed | | Total | Urine | protein | uEV supernatant total protein | uEV | total | protein |
|--------|------------------------|--------|-----------------------|-------|---------|-------------------------------|-----------------------|-------|---------|
| | | | concentration (µg/µl) | | | concentration (μg/μl) | concentration (μg/μl) | | |
| 1 | In-house reagent (IHR) | | 8.268 ± 0.432 | | | 7.780 ± 0.141 | 0.205 ± 0.012 | | |
| 2 | Commercial Invit | trogen | 8.028 ± 0.064 | | | 7.531 ± 0.131 | 0.226 ± 0.013 | | |
| | Reagent (CR) | | | | | | | | |

Protein concentrations were detected using Pierce TM BCA Protein assay in total urine, urinary EVs supernatant (non-EV fraction) and uEVs. Values reported as mean \pm SD (n = 3/group).