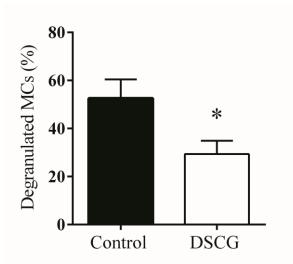
| | | Total MPO-ANCA patient population | <u>Subset for MC</u> <u>degranulation</u> <u>analysis</u> |
|-------------------------------------|--|--------------------------------------|---|
| <u>Demographic data</u> | Patient Number | 44 | 25 |
| | Sex (f/m) | 14/30 | 9/16 |
| | Age ¹ (years) | 67±2 | 66±3 |
| <u>Renal involvement</u> | | | |
| | Serum creatinine ¹ (µmo | al/L) 395±53 | 355±55 |
| | eGFR ¹ (mL/min/1.73m ² | ²) 21.6±2.4 | 23.8±3.5 |
| | Proteinuria ^{1,2} (g/day) | 1.8 ± 0.4 | 1.2±4.2 |
| | RBC ^{1,3} (urine cells/hpf) | 547±682 | 474±122 |
| <u>Extra renal</u> involvement | Lung or upper respirato tract or skin or arthralag presenting creatinine | | 10/25 |
| <u>Immunological</u> <u>data</u> | | | |
| | ANCA (MPO) titer ¹ (U | /ml) 134±16 | 120±21 |
| | CRP ⁴ (mmol/L) | 70±13.9 | 75±0.4 |

Supplementary Table 1. Comparisons of clinical, demographic and immunological data and indices of renal disease characteristics between total MPO-AAV patient population and subset population used for MC degranulation analysis.

¹Reported as mean \pm SEM, ²urinary total protein over 24 hr, ³Red Blood Cell excretion, ⁴C reactive protein on admission.

Supplementary Figure 1



Supplementary Figure 1: DSCG significantly reduces MC degranulation in a model of passive cutaneous anaphylaxis. C57BL/6 mice were sensitized subcutaneously (footpad) with 10ng α -DNP IgE and challenged 24hrs later with 100µg DNP-BSA intravenously. Two hours prior challenge (triggering MC degranulation), mice were injected intraperitoneally with either saline (control; n=10) or DSCG (n=10). Footpads were obtained and stained with toluidine blue to assess MC degranulation. Error bars represent mean ± SEM with statistical analysis by Mann-Whitney *t*-test **P*<0.05.