heart disease in 63% and 23% had diabetes mellitus. Patients were optimally treated (84% on β-blockers, 88% on ACE inhibitors, and 46% on spironolactone). The mean daily dose of furosemide was 60 (3) mg. Very few patients (5%) were sufficient in vitamin D. Patients with worse symptoms as measured by NYHA status had lower vitamin D levels than non-diabetics (p<0.001) and there was a negative correlation between vitamin D and fasting glucose levels (r=−0.13; p=0.02). There was no relationship between vitamin D levels and age, calcium, creatinine or CRP, and no differences between those patients taking and those not taking β-blockers and ACE inhibitors. In 8 unselected patients we found a negative relationship between tumour necrosis factor-alpha (TNF-α) levels and vitamin D (r=0.62; p=0.05). Although there was no relationship between vitamin D levels and baseline LVEF, in a subgroup of 150 patients followed up one year after titration to optimal CHF therapy, there was a significant positive relationship between change in LV dimensions and vitamin D levels at the time of the baseline scan (p<0.05).

Conclusions The vitamin D-PTH axis is abnormal in CHF, related to the severity of the condition. Our data suggest that reverse remodelling in response to optimal drug titration is greater in those with higher vitamin D levels. Whether vitamin D deficiency is causally related to CHF remains unknown and requires a long-term, randomised, placebo-controlled study in CHF patients with efficacy and mechanistic outcomes, using a dose of vitamin D capable of normalising both vitamin D and PTH levels.

**Background**

Red cell distribution width (RDW) is a surrogate of many aberrations (inflammation, malnutrition, iron deficiency (ID)) that may drive chronic heart failure (CHF) progression. While an elevated RDW and iron deficiency at baseline predict mortality in CHF, little is known about the prognostic implications of their temporal trends.

**Methods**

We analysed the relation of red cell indices on first consultation and over time with mortality in 274 outpatients with CHF (mean (±SD) age 70±14 years, LVEF 28±8%, NYHA class 2±1, 54% ischaemic). The combination of a rising RDW and a falling mean cell volume (MCV) identified evolving ID.

**Results**

On initial consultation, an RDW >15%, Hb<12.5 g/dl, and MCV <80 fl were evident in 41%, 46%, and 8% of patients. Over a median (±IQR) follow-up of 15±17 months, 60 (22%) patients died. On Cox proportional hazards analyses, a higher RDW independently predicted increased mortality (HR 1.21, p<0.0001). Over time, 51%, 58%, 40%, and 23% of patients had a rise in RDW, a fall in Hb, a fall in MCV, and evolving ID, respectively. A rising RDW predicted death (HR 1.18, p=0.002) independently of baseline RDWs and changes in Hb, with an absolute increase >1% conferring a twofold escalated risk of mortality (Abstract 107 figure 1A). Evolving ID was also associated with poorer survival (HR 2.89, p<0.0001, Abstract 107 figure 1B).
Conclusions An expanding RDW and evolving iron deficiency over time predict an amplified risk of death in CHF and could be utilised for risk stratification or therapeutically targeted to improve outcomes.

Results Significant vortical flow in any segment (defined as flow disturbance occupying more than one half of the aortic lumen) was present in all patients with MFS, but in only 7/18 controls (p<0.0005). The severity of flow disturbance was greater in MFS patients than controls (median severity score 3 for Marfan patients, 1 for controls, p<0.0005). There was marked regional variation in the prevalence of major flow disturbance (Abstract 108 figure 2), with the sinuses of Valsalva and proximal descending aorta being most frequently affected. Prior repaired aortic dissection was associated with marked abnormalities of blood flow (Abstract 108 figure 1C), with corresponding increases in axial WSS within the true lumen of the dissected aorta (typical axial WSS in the dissected ascending aorta was +0.9 N/m², compared to +0.54 N/m² in healthy controls). Aortic flow disturbance in MFS was of one of three types: Type A: flow disturbance confined to the sinuses of Valsalva, Type B: flow disturbance confined to the proximal descending aorta, Type C: flow disturbance in both the sinuses of Valsalva and the proximal descending aorta.

Abstract 108 Figure 2

Prevalence of vortical flow disturbance by region for Marfan patients and controls.