The role of nutrition in aggressive haematological malignancies: A single-centre review of patients receiving intensive chemotherapy NHS Graham McIlroy¹, Deepa Muthukrishnan², Monika Widlak³, Nicola Burch³, Francesca Jones¹ **University Hospitals** ¹Department of Haematology; ²Department of Dietetics; ³Department of Nutrition and Gastroenterology **Coventry and Warwickshire** University Hospitals Coventry and Warwickshire, Coventry, UK. Graham.Mcllroy@doctors.org.uk

Introduction

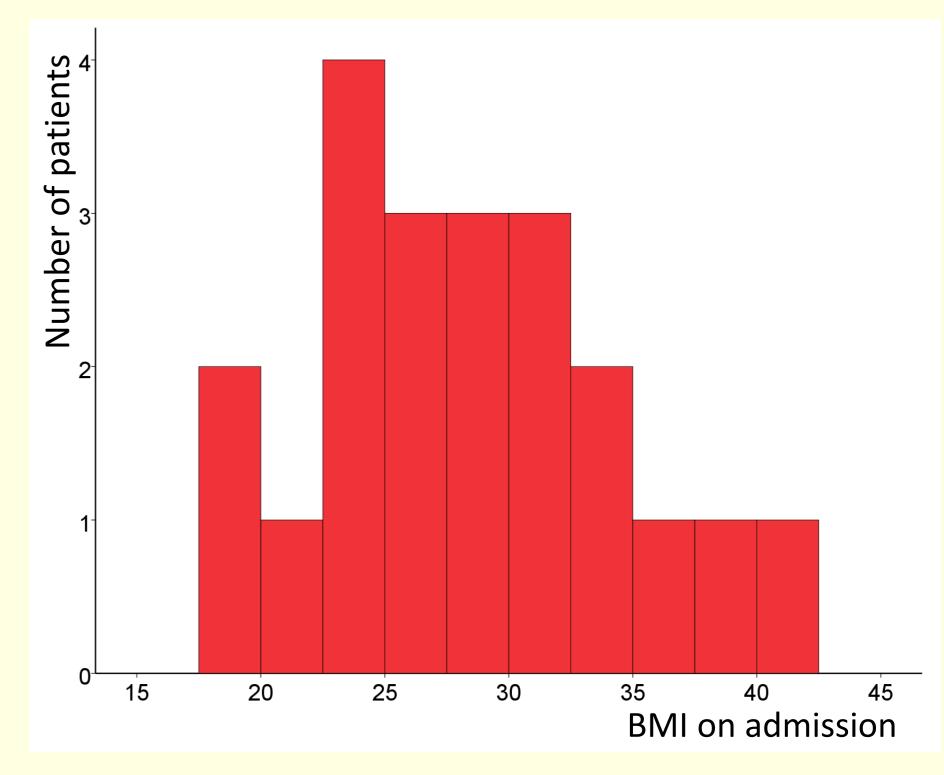
- Aggressive haematological malignancies need aggressive treatments. Tolerability of treatment depends on the level and maintenance of pre-morbid fitness, which require adequate nutrition.
- Malnutrition is common, reflecting patient, disease and treatment factors. It is associated with higher risk disease, poorer prognosis,

Malnutrition Universal Screening Tool (MUST)⁷

BMI		Unplanned v	veight loss		
>20	(0)	<5%	(0)		
18.5-20	(1)	5-10%	(1)		
<18.5	(2)	>10%	(2)		
Acutely ill and					
no nutrition fo	(2)				

Body mass index

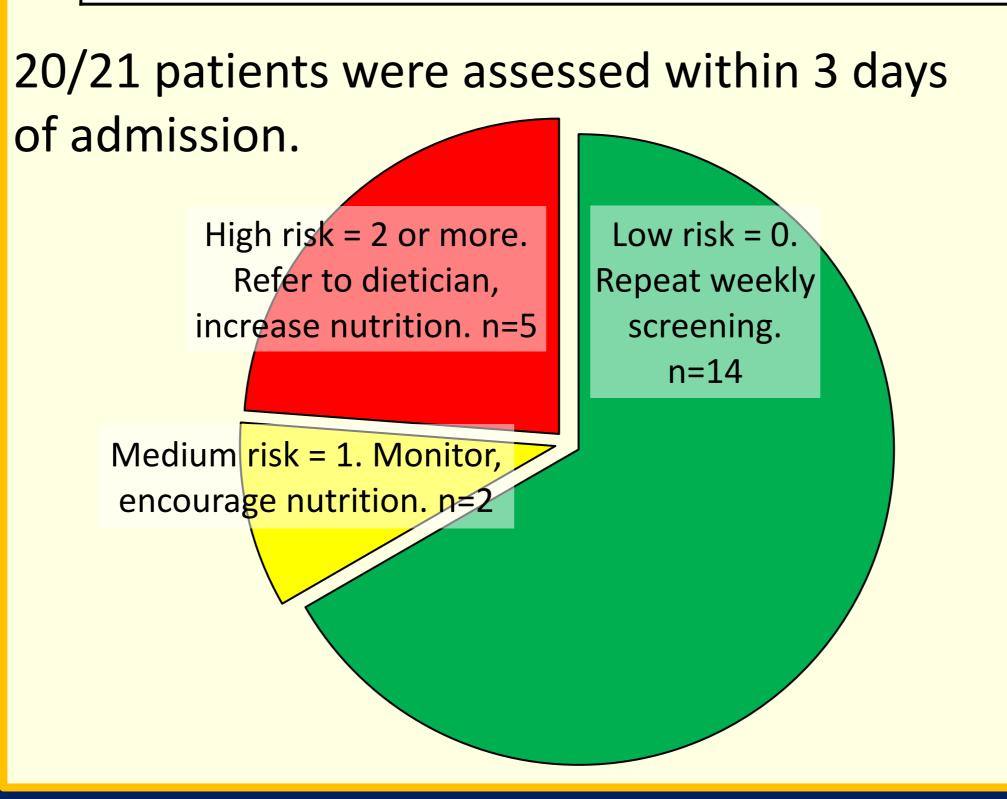
On admission, no patients were underweight (BMI <18.5). 14 patients (67%) were at least overweight.



and worse quality of life.¹⁻⁶

There is no consensus on how to identify and manage the risks of malnutrition in patients with haematological malignancies.

The aim of this service evaluation was to investigate the usefulness of existing nutritional assessments, measure weight loss during treatment, and explore the role of dietetic support for patients receiving their first cycle of intensive chemotherapy.



Weight change during treatment 20 patients (95%) lost weight during their admission for induction chemotherapy. 6 patients lost ≥10% of their body weight. Mean maximal weigh loss 5.6kg (range 0 to 12.1).

17 patients (81%) lost weight at discharge, compared with admission. Mean weight change over the admission -4.4kg (range -10.9 to +3kg).

Patients

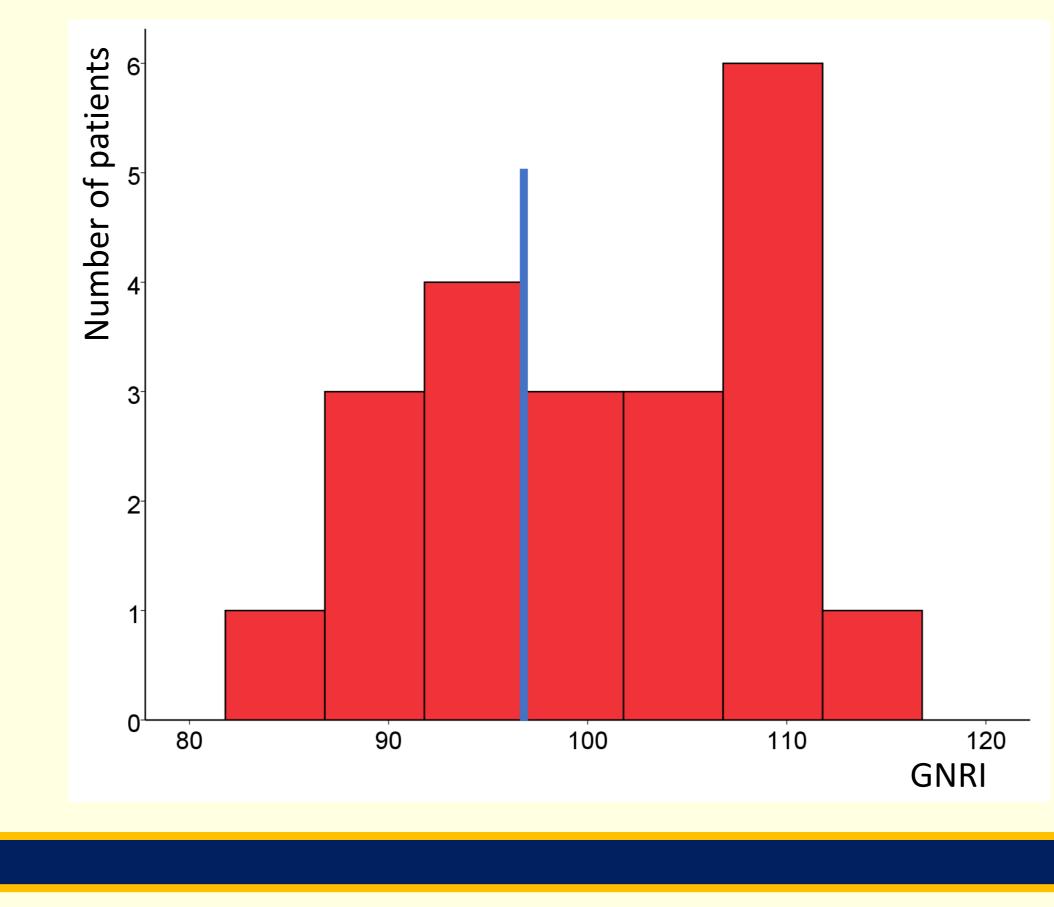
21 consecutive patients, admitted between June 2018 and May 2019 and treated with intensive chemotherapy for newly diagnosed AML or high-grade NHL, were included in this study. Patient characteristics shown in the table.

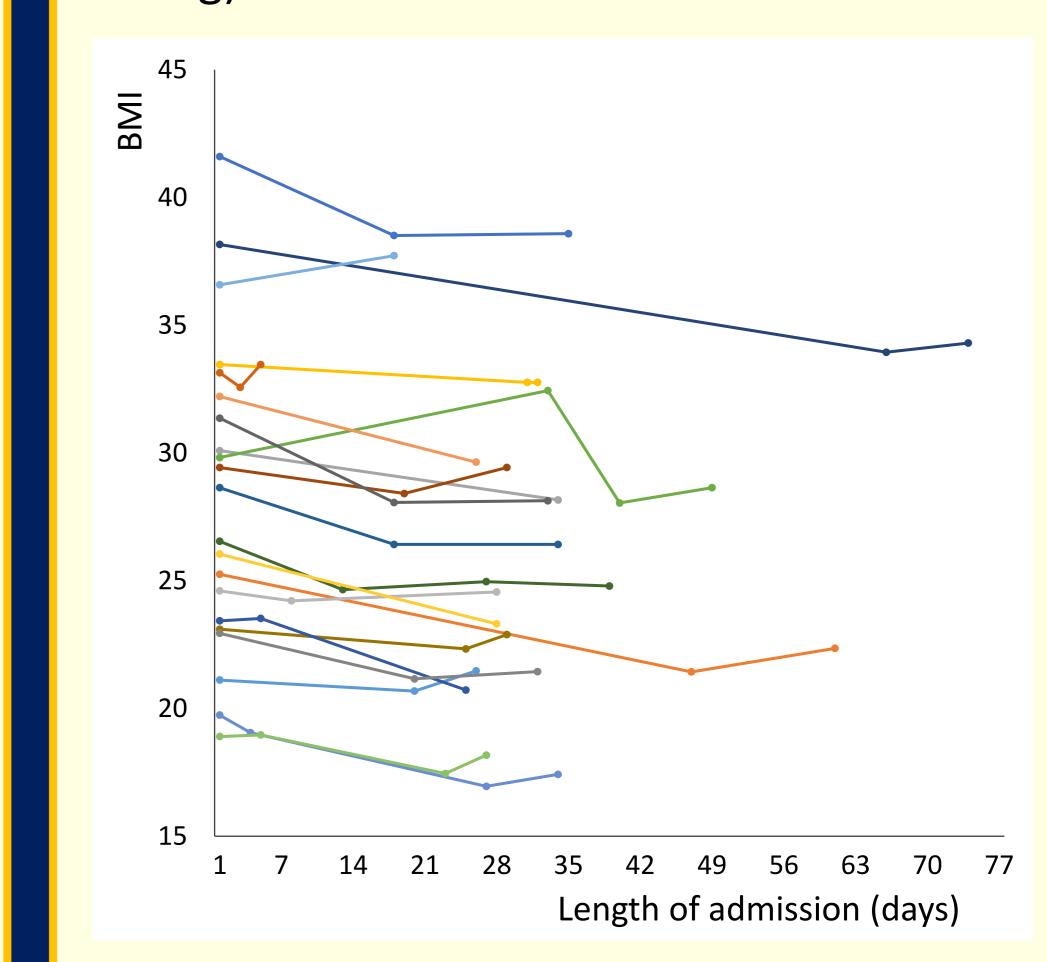
Geriatric Nutritional Risk Index⁶ GNRI =

{1.489 x albumin (g/L)} + {41.7 x [weight / IBW]} [Weight / ideal body weight] capped at 1 Lower score for higher nutritional risk. Score <96.8 shown to be associated with disease risk and poorer survival in DLBCL.⁶ This cohort:

Age	50 years	(27 to 69)
Men, number (%)	12	(57%)
Diagnosis and treatment		
AML, number (%)	15	(71%)
DA 3+10	11	
DA 3+10 + GO	2	
FLAG-IDA	1	
FLAG	1	
High-grade NHL	6	(29%)
R-CODOX-M	6	
Height	1.72 m	(1.58 to 1.9)
Admission weight	83.3 kg	(54.4 to 118.8)
Admission BMI	28.4	(18.9 to 41.6)
Admission albumin	40 g/L	(27 to 49)
Admission CRP	29 mg/L	(2 to 229)
Length of stay	33 days	(5 to 74)
Number of infective	2.1	(1 to 5)
episodes		

Mean score = 100.1 (range 81.9 to 112.1) 3 patients had BMI <22 *or* weight < IBW





Patients with the longest admission duration lost the most weight (r=0.68, p=0.001). Weight loss correlated with admission CRP (ρ=0.43, p=0.051).

Proportion of admission	64%	(27 to 100)
on IV antibiotics		

Average (range) shown, unless otherwise stated

Nutritional support

15 patients (71%) were referred to the dedicated Macmillan dietitian. Review was associated with complete nutrition records and use of NG feeding.

Weight loss not predicted by age, BMI on admission, albumin, or MUST or GNRI nutritional risk scores.

Conclusions

Existing measures of nutritional status translate poorly to patients receiving intensive chemotherapy for aggressive haematological malignancy, due to:

- Lack of sub-/acute weight loss
- Weight changes affected by fluid accumulation
- BMI often high at presentation
- Subjective element to MUST tool
- Confounding of biochemical parameters by severe acute illness

Early and proactive input from a specialist dietitian can support patients through treatment.

Better tools (e.g. mid-upper arm circumference, handgrip strength, bioelectrical impedance) are needed, to further target nutritional intervention.

References: ¹Hebuterne *et al.* (2014) J Parenter Enteral Nutr 38:196, ²Li *et al.* (2018) Medicine 97:3(e9663), ³Deluche *et al.* (2017) Nutrition 41:120, ⁴Malihi *et al.* (2013) J Hum Nutr Diet 26:123, ⁵Go et al. (2019) Ann Hematol 98:401, ⁶Kanemasa et al. (2018) Ann Hematol 97:999, ⁷BAPEN www.bapen.org.uk/screening-and-must/must-calculator



