Publication summary		
Title	Intermittent bolus versus continuous feeding in children receiving an	
	enteral formula with food derived ingredients: A national multicentre	
	retrospective study	
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Type of study	Observational / retrospective multi-centered	
Objective /	To evaluate the tolerance of different feeding	
hypothesis	modes (intermittent bolus/continuous/combination) in children	
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Results	who are fed with an enteral formula with food derived ingredients. Population: 43 children were included aged 1 to 17 years old (median: 6 years old). - 47% had a neurological or neuro-disability (20 of 43 children) - Median time children received an enteral formula before switching was 52 week (IQR, 24 – 120) - >50% was on intermittent bolus feeding - 81% were on a gastrostomy feeding tube (35 of 43 children) - 11% was fed in to the jejunum (5 of 43 children), of which 2 were fed as boluses over 2h at each feeding episode - Median feed volume in children receiving intermittent boluses: 150 mL (IQR 75-190mL) - Vomiting was the most reported feed intolerance prior to the switch Study outcomes: - Primary outcome: feed tolerance per feeding mode (intermittent bolus, continuous, and combination) - Bolus: - >80% improvement in vomiting and loose stools - >70% improvement in retching and constipation - >66% improvement in abdominal pain - Continuous: - 100% improvement in vomiting, retching, abdominal pain and loose stools - 75% improvement in constipation - Combination: - 100% improvement in vomiting, retching and abdominal pain Reported change in gastrointestinal symptoms after switching to an enteral formula with food ingredients in relation to mode of feeding. Gastrointestinal Symptoms after switching to an enteral formula with food ingredients in relation to mode of feeding. Gastrointestinal Symptoms after switching to an enteral formula with food ingredients in relation to mode of feeding.	
	Vomiting 12 (91.67%) 7 (85.71%) 3 (100%) 2 (100%) Retching 20 (85%) 11 (72.73%) 7 (100%) 2 (100%) Abdominal Pain 6 (83.33%) 3 (66.67%) 2 (100%) 1 (100%) Loose stool 11 (90.91%) 6 (83.33%) 5 (100%) 0 (0%) Constipation 13 (69.23%) 8 (75%) 4 (75%) 1 (0%)	



	 Secondary outcomes: weight-for-age and height-for-age Children who were fed intermittent bolus reported the greatest increase in weight (p-value: 0.003), children who were fed continuously or a combination also saw clinically significant weight gain (p-value of 0.0052 and 0.068 respectively) No significant differences in feed volume, total fluid or total daily calorie intake after switching or within different feeding modes >90% of dieticians reported that the nutritional goals were met after formula was changed. Children who were feeding continuously reported the highest achievement to meet dieticians' nutritional goals. Reason for parents to switch to a real-food formula: 1) previously on blended diet / unable to start blended diet and felt this formula was an appropriate compromise, 2) due to poor feeding tolerance to previous formula.
Conclusion	Children who were continuously fed reported the greatest improvements in feed tolerance symptoms. Conversely, children who were bolus fed reported the greatest weight gain. An enteral formula with food derived ingredients (e.g. Compleat Paediatric) is well tolerated whether delivered continuously, or as a bolus feed in achieving feed tolerance, weight gain and dietetic goals.
Short description of	Study characteristics:
the methods used (target group, duration intervention etc.)	 Inclusion criteria: children between 1 and 17 years who had switched to the new enteral formula (e.g. Compleat Paediatric) for at least one month and accounted for at least 80% of their total energy requirements Study sites: 4 National Health Service Trusts around England Duration intervention: 1 month Type of intervention: switch to an enteral formula with food derived ingredients (Compleat Paediatric) Data collection: by paediatric dieticians, via Microsoft Forms (anthropometric and gastrointestional outcomes over a 1 month period)
	Primary outcome: feed tolerance per feeding mode (intermittent bolus, continuous, and combination) Secondary outcomes: weight-for-age and height-for-age
Limitations	 Small sample size (43 participants) Short trial period (1 month) Retrospective design (less accurate)

