Guidance for the use of Oral Nutritional Supplements for Adults in General Practice

This guidance is based on NICE Guideline 32, “Nutrition Support in Adults” published in 2006.

**Aim**

To promote the appropriate, rational and cost-effective prescribing of Oral Nutritional Supplements by primary care in Kirklees.

**Rationale**

Oral nutritional supplements are nutritional supplements that are used in patients who have been identified as being nutritionally compromised.

Use of oral nutritional supplements requires regular monitoring of the patients progress.

Alternative methods can be used to supplement dietary input without recourse to Oral Nutritional Supplements.

Oral Nutritional Supplements are relatively expensive for the NHS. NHS Kirklees is keen to encourage a “food first” strategy and reserve the use of oral nutritional supplements for patients who have not responded to dietary measures alone.

Audits in recent years by the Medicines Management team have shown that oral supplements are often initiated inappropriately or continued unnecessarily or without adequate review.

**Objectives**

To ensure that alternative non-prescribable nutritional support methods have been considered and given an adequate trial prior to prescription of supplements.

To promote the use of explicit targets and review criteria for patients receiving oral nutritional supplements.

To provide audit criteria based on the above targets, by which adoption of this guidance can be evaluated.

**Process**

For any individual patient the following steps should apply. Oral nutritional supplements should only be introduced after steps 1-4 have been completed and the nutritional intake is still inadequate.
Step 1: Identification

Screening should assess body mass index (BMI) and percentage unintentional weight loss and should also consider the time over which nutrient intake has been unintentionally reduced and/or the likelihood of future impaired nutrient intake. The Malnutrition Universal Screening Tool (MUST), for example, may be used to do this.

People with disease related malnutrition are defined by NICE as:

- BMI < 18.5
- Unplanned weight loss >10% in past 3-6 months or
- BMI <20 and unplanned weight loss > 5% in past 3-6 months.

Nutritional support should be considered in people at risk of malnutrition who are defined by the following criteria.

- Have eaten little or nothing for more than 5 days and/or are likely to eat little or nothing for the next 5 days or longer
- Have poor absorptive capacity and/or have high nutrient losses and/or have increased nutritional needs from causes such as catabolism.

Step 2: Global nutritional assessment

Consider availability of adequate diet and patient’s ability to feed using appropriate utensils. Identify any problems with:

- Food intake including chewing (consider dental assessment) and swallowing (consider speech therapy assessment) appropriate food texture and temperature. Patient’s with dysphagia may require a modified consistency diet and are likely to require supplements for longer periods.
- Medication – consider any that may suppress appetite.
- Physical symptoms e.g. pain, sore mouth, vomiting.
- Medical prognosis (excessive input may be inappropriate in terminal illness)

Step 3: Goal setting

Realistic and measurable goals should be established and documented for each patient in order to identify the end-point of treatment and a review period agreed (usually monthly). Suitable goals might consist of attaining a target BMI or preventing further weight loss, completion of wound healing, or end of risk period (e.g. recovery from precipitating illness).

Step 4: First Line Dietary Advice (Non-prescribable supplementation)

This can be by fortification of normal diet, and/or addition of ‘over the counter’ nutritional supplements. Written information should be provided to reinforce the advice. The NHS Kirklees ‘food first’ booklet is recommended.

- Dietary fortification: suggest increasing frequency of intake (little and often) with snacks between meals. Add or increase amounts of high-energy foods such as full fat milk, butter, cream etc.
- Enriched drinks such as Build Up and Complan can be used in addition to meals.
Step 5: Prescribing Oral Nutritional Supplements

Prescription of oral nutritional supplements should only be initiated if first line dietary measures have failed despite adequate duration (around 4 weeks). Prescribing of oral nutritional supplements should only be undertaken where there are clinical benefits to be realised and clear nutritional goals to work towards. Repeat prescriptions should only be issued if there is an explicit plan for continuation (step 3).

Goals should be regularly reviewed and prescribing should cease when goals are achieved (step 6). Supplements should only be prescribed for the conditions specified under ACBS. These conditions are:

- Short bowel syndrome
- Intractable malabsorption
- Pre-op preparation of undernourished patients
- Proven inflammatory bowel disease
- Following total gastrectomy
- Bowel fistulae
- Disease related malnutrition
- Dysphagia.

First line choice of supplement in Kirklees are Complan milkshakes which the patient makes up with milk. Shakers are available free of charge from community pharmacies on patient request to assist in measuring and making the milkshakes and are included in starter packs. Please note: Complan milkshakes will not be a suitable product for all patients e.g. intolerant of cow’s milk, lactose intolerant and fluid restricted patients. See product information for further information.

Please note Complan and Complan® Shake are different products. Complan is not available on NHS prescription, whereas Complan® Shake is (if patient qualifies under ACBS criteria).

Second line supplements for use in patients for whom Complan milkshakes are not suitable should be selected from the Nutricia range of products (ie Fortijuice, Fortisip ranges). See appendix 1 for relative cost, calorie and protein content and flavours of some Nutricia products.

Prescriptions should be clearly marked “ACBS” and give clear directions for use (e.g. One to be taken twice daily between meals). “As directed” should not be used.

To avoid wastage, unless supplement has already been tried and the patient’s preference established, the initial prescription should be no more than 1 - 2 weeks supply and marked “Mixed Flavours”. Patient should then inform the practice of their preferred flavour choices. Sip feed prescriptions should be issued as monthly acute prescriptions for the first 3 months.

Monthly patient monitoring should take place during this period to assess progress toward treatment goals. Monitoring information should be shared with the prescriber when the next prescription is requested. Practices should have a system of ensuring that all homes report patient’s weight change when requesting further supplies of Oral Nutritional Supplements.

Hospital dietitians requesting continuation of supplements post discharge will identify the aims of treatment and patient’s target weight or BMI in a letter to the patients GP to enable continued prescribing and appropriate monitoring in primary care.

Retrospective prescriptions should not be normally be issued at the request of home delivery companies.

Patients stabilised on nutritional supplements should be assessed every 3 -6 months. Refer to the British National Formulary for the most up to date information on available formulations.
Step 6: Review by designated health professional
Any patient receiving oral nutritional supplements should be reviewed regularly. It is the responsibility of the prescriber to ensure that patients are adequately monitored in accordance with nutritional goals and review periods (stage 3). Effective interprofessional communication is essential to providing appropriate continuing nutritional support.

Step 7: Termination of Oral Nutritional Supplements and Follow Up
Providing that an effective plan has been prepared at the outset it should be possible to readily identify the point at which prescribed supplements can be stopped. Oral nutritional supplements should be gradually reduced and patients should continue to be monitored for 3 months after supplements have been discontinued.

Referral for specialist dietetic input (Steps 4-7)
Dietetic referral may be appropriate in any of the following circumstances:
- To advise on nutritional supplementation strategies and the appropriateness or otherwise of initiating oral nutritional supplements.
- To assist in appropriate planning and goal setting for nutritional support for individual patients
- Deterioration in nutritional status despite supplementation after excluding other contributory pathology.
- Apparent requirement for supplementation longer then three months.
- The presence of co-existing medical conditions such as diabetes, renal failure, coeliac disease or high cardiovascular risk.
- Where swallowing difficulties or other indications for modified food texture exist.

Audit Criteria
1. Is there documented assessment of patient’s needs (ie BMI, % unintentional weight loss or MUST score) recorded in the medical records?
2. Is there a record of the condition for which oral nutritional supplement prescribed?
3. Has the oral nutritional supplement been prescribed for an approved condition? (ACBS categories)
4. Is there an explicit nutritional goal identified in the medical records?
5. Is there a specified review interval?
6. Is there evidence that review has been undertaken in relation to set goals?
7. Have first line dietary measures been tried for 4 weeks prior to prescribing oral nutritional supplements?
8. Do prescriptions contain clear instructions about the dose and quantity to be used?

Further Information
The MUST screening tool can be downloaded from www.bapen.org.uk

Acknowledgments
With thanks to South Staffordshire PCT who’s ‘Guidance for the use of oral nutritional supplements for Adults in General Practice’ was used as a basis for this guidance.

References
National Prescribing Centre 1998 Oral nutrition support part 1, MeReC Bulletin 9(7)
National Prescribing Centre 1998 Oral nutrition support part 2, MeReC Bulletin 9(9)
Appendix 1.
The contract for enteral feeding products in the Calderdale and Huddersfield Foundation Trust footprint is for Nutricia products. These should be prescribed first line unless advised otherwise by a dietitian.

<table>
<thead>
<tr>
<th>Product</th>
<th>Cost per Unit</th>
<th>Calorie Content</th>
<th>Style</th>
<th>Protein Content</th>
<th>Flavours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Complan Shake</td>
<td>£0.90</td>
<td>387 K/cal - when made with 200mls Whole Milk</td>
<td>Milkshake</td>
<td>15g</td>
<td>Chocolate, Banana, Vanilla, Strawberry, Original</td>
</tr>
<tr>
<td>237ml (made with 200mls Whole Milk)</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Fortisip Compact</td>
<td>£1.85</td>
<td>300</td>
<td>Milkshake</td>
<td>12g</td>
<td>Chocolate, Mocha, Forest Fruits, Apricot, Banana, Strawberry, Vanilla</td>
</tr>
<tr>
<td>125ml bottle</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Forticreme</td>
<td>£1.80</td>
<td>200</td>
<td>Pudding</td>
<td>12g</td>
<td>Forest Fruits, Vanilla, Banana, Chocolate</td>
</tr>
<tr>
<td>125g pot</td>
<td></td>
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<td></td>
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</tr>
<tr>
<td>Fortijuce</td>
<td>£1.89</td>
<td>300</td>
<td>Juice</td>
<td>8g</td>
<td>Orange, Lemon, Apple, Blackcurrant, Strawberry, Tropical, Forest Fruits</td>
</tr>
<tr>
<td>200ml bottle</td>
<td></td>
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