Oral eating and drinking procedure

This is a mandated procedure under the operational policy framework. Any edits to this procedure must follow the process outlined on the creating, updating and deleting operational policies page.

Overview

This procedure is a practical direction for all staff working in education and care to plan and manage children and young people who require oral eating and drinking support including oral sensitivities or fixation, pica, swallowing disorders, feeding, food phobia and choking.

This procedure must be read along with the Department for Education health support planning processes.

Scope

This procedure applies to educators, early childhood development specialists, principals, directors and education support staff working in education and care.

It also applies to all individuals and entities ("other parties") engaged or contracted by the department to perform work and/or to provide support/services, which involve interactions with children and young people or require close proximity to or access to children and young people (ie volunteers, contractors, employees from external agencies, third party service providers, etc).

This document describes:

- how education and care staff manage the health support needs both proactively and reactively in education and care for children and young people who require oral eating and drinking support, including:
 - o oral sensitivities or fixation (chewing, sucking, mouthing)
 - o pica (non-food items); swallowing disorders (dysphagia)
 - o feeding (oral issues and gastrostomy tubes)
 - o food phobias/sensory aversion
 - o choking
- signs and symptoms of swallowing and feeding disorders including choking and dysphagia
- the emergency response for any person experiencing a choking incident while at an education or care service
- education and training for managing oral eating and drinking
- risk minimisation strategies where a child or young person has feeding difficulties, dysphagia, oral sensitivities, and food or choking phobias.

This procedure applies from the time a child or young person is enrolled until they leave the education or care service.



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Detail

Treatment and management for oral eating and drinking

Managing oral eating and drinking difficulties and disorders in education and care services makes sure children and young people are:

- safe while eating, including strategies to minimise the risks of choking and aspiration
- adequately nourished and hydrated to enable attendance and full accessibility of the curriculum
- healthy (free from aspiration, pneumonia and other illnesses related to malnutrition or dehydration) to maximise attendance.

Most children and young people in education and care services manage their own eating and drinking. However, some may require supervision, coaching, guidance and support for their safety and wellbeing.

Where a child or young person has a risk of choking or aspiration, requires their food or fluid consistency to be modified or they need to be fed, an <u>oral eating and drinking care plan (DOC 254KB)</u> is needed to make sure they are supported and safe during mealtimes. Children and young people generally do not require an oral eating and drinking care plan for food phobias, eating disorders or oral sensitivities.

Where there is not a care plan in place and the education and care staff are concerned about the safety of a child or young person's oral eating and drinking this must be communicated to parent/guardians (an <u>oral eating and drinking observation log (DOC 66KB)</u> can be used to document and communicate). <u>Choking first aid (PDF 140KB)</u> must be instigated if a child or young person is choking.

Where a child or young person has invasive or complex healthcare needs, uncertain health or changing health, they may be eligible for and supported by the <u>Access Assistant program</u> or <u>Encompass</u>.

Choking

Alert! Choking is a medical emergency. Call 000 (ambulance) immediately.

When a child or young person is choking, it means that an object (food or non-food item) is stuck in the larynx (voice box) or trachea (airway) restricting the air flowing normally in and out of the lungs, so they cannot breathe properly. Most of the time the object only partially blocks the trachea and can be coughed up, and breathing returns to normal quickly.

Children under the age of 3 are most at risk for choking on food and small objects. However, children with oral fixation or pica, and children with a disability or chronic illness may be at a higher risk of choking.

Choking can be a **life-threatening emergency** when the object is completely blocking the air flow. When the brain does not receive oxygen for more than four minutes permanent brain damage or death can result.

Signs and symptoms of a choking emergency include:

- difficulty breathing, speaking, swallowing
- grabbing at throat or waving arms
- coughing, wheezing, gagging

- making a whistling or crowing sound, or no sound at all
- being unable to talk, cry or make noise
- turning blue blue lips, face, earlobes, fingernails
- appearing panicked
- becoming limp or unconscious
- becoming pale and sweaty.

First aid for choking emergency

Education and care staff must give first aid measures following any relevant care plan or first aid plan. Where this does not specify an individualised first aid response, standard first aid is to be followed.

- Choking First Aid for adult or child over 1 year (PDF 140KB)
- 1. Encourage the person to relax and ask to cough to remove the object.
- 2. If coughing does not remove the blockage CALL 000 (ambulance).
- 3. Bend the person well forward and give 5 sharp blows on the back between the shoulder blades with the heel of one hand.
- 4. Check if the blockage has been removed after each blow by looking in the mouth.
- 5. If the blockage has not cleared after step 3, give (up to) 5 chest thrusts.
- Give chest thrusts by placing one hand in the middle of the back for support and the heel of the other on the lower half of the sternum. Thrusts should be slower and sharper than CPR compressions.
- 7. Check if the blockage has been removed after each thrust.
- 8. If the blockage has not cleared after 5 thrusts, continue alternating 5 back blows with 5 chest thrusts until medical aid arrives; checking if the blockage has been removed after each action.
- Choking first aid for infant under 1 year (PDF 162KB).

Pulmonary aspiration

ALERT! Pulmonary aspiration is a medical emergency and can result in death from asphyxiation within minutes. Phone 000 (ambulance).

Pulmonary aspiration is the entry of materials (ie food, drink, stomach contents) from the gastrointestinal tract into the lower respiratory tract. When pulmonary aspiration occurs during eating or drinking, this is often referred to as 'going down the wrong pipe/hole'. Consequences of pulmonary aspiration range from no injury to pneumonia or death from asphyxiation (choking). Consequences are often dependent on the underlying health status of the person.

While there may be observable signs that accompany aspiration events, such as coughing, choking, eye reddening during eating, or a gurgly vocal quality after swallowing, children and young people can also aspirate 'silently', or with no observable symptoms.

Treatment and management are dependent on the medical consequence of pulmonary aspiration. Some treatments may include:

- removal of an object (if able by coughing up, or by bronchoscopy)
- endotracheal suctioning
- medication (if a bacterial infection)
- supportive therapies (fluid management, bronchodilators, physiotherapy)
- referral to speech pathologist.

Where a child or young person is at risk of aspiration, this must be documented in an <u>oral eating and</u> <u>drinking care plan (DOC 254KB)</u> and <u>health support agreement (DOC 131KB)</u>.

The health support agreement must include signs and symptoms of aspiration and emergency response actions. The <u>safety and risk management plan (DOCX 140KB)</u> must include specific risk minimisation strategies for the child or young person in the context of the education and care setting.

Feeding and swallowing difficulties

Speech pathologists specialise in treating children and young people with feeding and swallowing difficulties. The speech pathologist may work as part of a team including occupational therapists, nurses, dietitians and other specialists.

The team will evaluate posture, self-feeding abilities, medical status and nutritional intake and develop an <u>oral eating and drinking care plan (DOC 254KB)</u> that will include recommendations on positioning, equipment, diet/food preparation, feeding plan techniques and precautions.

Treatment for feeding and swallowing difficulties varies depending on the cause, symptoms and the problem, but may include:

- medical intervention (ie reflux medication)
- modified equipment (ie flexi-cup, built up handle spoon)
- nutritional change (different foods, more calories)
- food temperature and texture changes
- postural or positioning changes
- behaviour management or special techniques to help eating drinking and swallowing
- preparation of food and drinks (ie opening or unwrapping, appropriate texture).

Tube feeding

Education and care staff don't need to manage the feeding of children and young people with feeding tubes.

Children and young people with a nasal or gastric feeding tube will be supported by the <u>Access Assistant program</u>. A nursing care plan will be developed in consultation with the parent/guardian and nursing staff will give support and help for the child or young person in the education or care service.

The gastrostomy care instruction (PDF 327KB) and nasogastric care instruction (PDF 290KB) have been

developed to support staff and give first aid strategies for children and young people with a gastrostomy device or nasogastric tube.

Care plans and support agreements

Adequate nutrition is essential to support children and young people's growth and development and to allow them to participate in and benefit from their educational experiences.

A child or young person may have difficulties with eating and drinking for a variety of reasons and the nature of support will be dependent on the individual needs and requirements. Parent/guardians should be encouraged to consult a health or allied health professional (specifically a speech pathologist) where oral eating and drinking difficulties are identified.

Where a child or young person has a risk of choking or aspiration, requires their food or fluid consistency to be modified or they need to be fed, an <u>oral eating and drinking care plan (DOC 254KB)</u> is needed to make sure the child or young person is supported and safe during mealtimes. Children and young people generally do not require a care plan for food phobias, eating disorders or oral sensitivities.

The matrix below can be used as a guide for the requirement for a care and support plans. It is a guide only. It does not include every scenario for health care plan or support agreement requirements.

	OE&D Care Plan	Non-specific Health Care Plan	Health Support Agreement	Medication Agreement
Feeding and swallowing difficulties; food or choking phobias; eating disorder				
Food or fluid selectivity by type, texture, brand, shape or colour			✓	
Food or fluid consistency is modified	✓		~	
Food or fluids are fed to the child or young person	✓		~	
Behaviour management (where related only to food intake)			✓	
Behaviour management (with an underlying medical/health condition)		✓	✓	
Modified equipment			✓	
Medication management			✓	✓
Tube feeding				
Exclusively tube fed		✓		
Tube fed, with oral intake of food and/or drink	V			
Aspiration; Choking				
Where there is a known risk	V		✓	
Where there is a known risk with an underlying medical/health condition	V	✓	✓	
Oral sensitivities				
Where oral sensitivities are <u>diagnosed</u> and sensory activities required at education or care service			✓	
Pica where treatment involves medication or supplements only			✓	✓
Pica where medical or allied health have prescribed treatment or activities			✓	
Pica where there is an underlying medical/health condition		✓	✓	
Pica where parent or legal guardian has notified of behaviour therapy or activities			✓	

Oral eating and drinking care plan

An <u>oral eating and drinking care plan (DOC 254KB)</u> is completed and regularly reviewed by the speech pathologist or treating health professional, in consultation with the parent/guardian, where the child or young person requires feeding or modification to their food or drink.

The oral eating and drinking care plan should outline (where relevant) the:

- level of support needed (degree of supervision or help)
- type of support needed:
 - o equipment
 - o environment
 - o positioning
 - o mealtime processes and after mealtime care
 - o more time needed for mealtimes
- preparation and presentation of food and drink (the <u>International Dysphagia Diet Standardisation</u> <u>Initiative (IDDSI) Framework</u> gives descriptors for levels of food and fluid):
 - o food texture or consistency
 - o fluid thickness (refer to the <u>IDDSI Flow Test</u> for measuring liquid thickness)
 - o rate and order of intake
 - o strategies for spoon and finger food, and drinking
- communication strategies and needs (including education and care staff, parent/guardian, child or young person, medical or allied health professional)
- emergency response plan and emergency contacts
- risk-minimisation plan to make sure risks are assessed and minimised
- learning targets:
 - o increasing independence
 - o behaviour management
 - o increasing or decreasing food/fluid intake
- any requests for documentation and/or observations by education and care staff
- more information and recommended training for education and care staff to support the child or young person.

The speech pathologist or health professional completing the <u>oral eating and drinking care plan (DOC 254KB)</u> can give training, more information, clarification or demonstration as needed. Education and care staff will be accommodated to attend the hospital for education and training where health professionals can't attend the education or care service.

Non-specific health care plan

In some circumstances a <u>non-specific health care plan (DOCX 124KB)</u> may be developed by the treating health professional in consultation with the parent/guardians where a child or young person has oral eating and drinking difficulties.

Where a child or young person has invasive or complex health care needs, uncertain health or changing health they may be eligible for and supported by the Access Assistant program.

Individual first aid plan

In some circumstances an <u>individual first aid plan (DOCX 119KB)</u> may be developed by the treating health professional in consultation with the parent/guardian where a child or young person requires a first aid response that is **not** the standard first aid response for their health condition.

Health support agreement

Where a parent/guardian has identified a child or young person requires eating and/or drinking support, education and care staff should encourage them to consult with a speech pathologist and/or health professional for an appropriate diagnosis and treatment plan. Until a formal diagnosis and care plan is in place a health support agreement (DOC 131KB) must be developed.

In all circumstances where a child or young person has an <u>oral eating and drinking care plan (DOC 254KB)</u>, the education or care service must complete a health support agreement in consultation with parent/guardian. This should include completing a <u>safety and risk management plan (DOCX 140KB)</u> to identify risk minimisation strategies, management and treatment of the child or young person in the context of the education or care service.

The health support agreement should clearly identify cultural, spiritual and language needs and where a child or young person's oral eating and drinking difficulties are changeable.

Where the health support agreement is being completed for an Aboriginal child or young person this should be developed in consultation with the primary caregiver, who is not always the parent. Extended family members and Aboriginal community education officers or Aboriginal health workers may also help in the development of the health support agreement to make sure they are culturally appropriate and meaningful.

The development of health support agreements must be completed with an assurance that the content is understood and culturally valid. Children and families from culturally and linguistically diverse backgrounds may require support persons and access to interpreter services and community liaison officers.

A health support agreement and safety and risk management plan should be reviewed in consultation with the parent/guardian in each of the following circumstances:

- when the oral eating and drinking care plan has been reviewed and updated
- as soon as practicable after a medical incident at the education or care service
- before the child or young person participating in an offsite activity (ie camps or excursions) or at
 onsite special events (ie class parties, cultural days, fetes, incursions). Where the safety and risk
 management plan has not addressed risk minimisation strategies for offsite activities an offsite
 safety and risk management plan (DOC 134KB) may be developed.

Oral eating and drinking observation log

An <u>oral eating and drinking observation log (DOC 66KB)</u> is recommended to be completed where education and care staff notice a change in the child or young person's eating and drinking abilities or behaviours during or after mealtimes ie:

- a change in the child or young person's ability to chew or swallow or manage food and drink
- tires, is increasingly fatigued or loses consciousness during or after meals
- frequent and repeated coughing, spluttering, gagging, gurgling, near choking, or distress during or after meals
- rapid weight loss.

The oral eating and drinking observation log can be used as a communication tool between the education or care service, the parent/guardian and speech pathologist or health professional to:

- communicate with parents and guardians (where changes in mealtime or eating and drinking abilities are noted this may include both positive and negative changes)
- initiate a review of the oral eating and drinking care plan (DOC 254KB)
- give information and/or evidence for the review of the oral eating and drinking care plan (may be used to document eating and drinking all observations over a specific timeframe ie 1 day, 5 days).

Eating and drinking management

Supply of food and equipment

It is the responsibility of the parent/guardian to give appropriate food and drink for the child or young person during attendance at the education or care service. Some education and care services may give food and drink to the child or young person.

If the food or drink provided does not meet the requirements of the <u>health care plan</u> or <u>health support</u> <u>agreement (DOC 131KB)</u>, the following strategies can be implemented:

- contact the parent/guardian to request food or drink is brought in
- contact the parent/guardian to negotiate accessing food or drink from canteen or other appropriate source
- ask the parent/guardian to give appropriate food or drink that can be stored at the education or care service for use in emergencies.

Where the <u>oral eating and drinking care plan (DOC 254KB)</u> identifies specialised food, thickeners or equipment for oral eating and drinking management it is the parent/guardian's responsibility to give this to the education or care service.

Where equipment is needed for the child or young person there must be arrangements in place for the equipment to be cleaned and maintained. The arrangement must be clearly documented in the health support agreement (DOC 131KB). This could be the responsibility of either the:

education or care service – with a staff member nominated to be responsible for cleaning the

equipment

• parent/guardian – the equipment is sent home with the child or young person and returned to the education or care service. Where this option is selected, contingency arrangements should be identified if the equipment is not returned to the education or care service.

Mealtime routines

Predictable or regular mealtime routines can help children and young people with oral eating and drinking difficulties to become comfortable and familiar with what is expected of them and what to expect from education and care staff helping them.

The <u>oral eating and drinking care plan (DOC 254KB)</u> will give guidance around mealtime routines that should be followed for the child or young person. Specific requirements and instructions for education and care staff to follow for each child or young person will be documented in the <u>health support agreement (DOC 131KB)</u>.

General processes can be implemented to make sure successful mealtimes for children and young people with eating and drinking difficulties; combined with the specific requirements documented in the care plan and support agreement.

Before the meal

- communicate to the child or young person what is going to happen
- use language they will understand and/or visual cues to help them prepare for eating and drinking
- encourage the child or young person to be involved in meal preparation where possible
- be aware of the total environment in which meals occur
- comfortable, calm, enjoyable, distraction-free environments
- loose clothing and hair may drape over the child or young person or food tie back where possible
- encourage hand washing.

During the meal

- child or young person to be sitting down on a chair or sitting up straight with legs crossed
- allow the child or young person to pace their meal as much as possible
- children and young people with swallowing difficulties require more time to chew, swallow and fully clear their mouth before their next mouthful
- encourage further swallows for the child or young person to clear their mouth before presenting the next bite or sip
- if the health care or support plan includes the introduction of new foods do this when the child or young person is calm, alert and hungry
- keep foods at serving temperature (especially for slow eaters)
- keep mealtimes enjoyable.

After the meal

- finish the meal with a signal to clearly indicate mealtime is over, particularly for children and young people who have difficulty understanding or those with visual, hearing or sensory impairments
- encourage hand washing
- remove uneaten food and packaging from mealtime area and ensure it is appropriately disposed of.

Food and drink preparation

When preparing food or drink for children and young people with oral eating and drinking difficulties, education and care staff must make sure any specific requirements documented in the <u>oral eating and drinking care plan (DOC 254KB)</u> or <u>health support agreement (DOC 131KB)</u> are strictly adhered to.

The <u>IDDSI Framework</u> gives descriptors of food and fluid on a continuum of 8 levels (0-8) and the <u>IDDSI Flow</u> <u>Test</u> instructs how to test for the correct liquid consistency.

Safe food handling principles and standard infection prevention precautions must be adhered to (including not preparing food or drink if you are ill (diarrhoea or vomiting), wash and dry hands thoroughly, make sure clothes and equipment are clean).

When heating food and drinks, microwave in 10 - 20 second bursts, stir and test temperature after each burst. Test the temperature of food or drink by touching the spoon on the back of the hand.

Refer to the <u>eat well SA schools and preschools healthy eating guidelines (PDF 2.3MB)</u> (staff login required) for further information.

Food and drink storage

Food and drink should be safely and appropriately stored according to packaging and parent instructions. Storage facilities (ie refrigerator, cupboard) and waste bins should be out of reach of children and students and only accessed by staff. If necessary, to minimise risk to children and students, the storage facility should be always kept secure.

Positioning

Good positioning is essential for safe eating, drinking and swallowing. The child or young person's <u>oral eating and drinking care plan (DOC 254KB)</u> will describe the specific requirements on positioning, or a <u>transfer and positioning care plan (DOCX 175KB)</u> may be needed.

Work with the health professional where transfer and positioning needs are identified to make sure care plan directions are met.

A stable, upright and well-aligned position allows more energy and effort to be spent on the process of eating, drinking and swallowing. Awkward or inappropriate positioning can lead to food or fluid being inhaled (aspiration) and choking.

Some children and young people with eating and drinking difficulties may have physical impairments that interfere with their ability to maintain a stable position; ie muscle tone or weakness, difficulty controlling and coordinating movements.

<u>Guidelines for optimal positioning at mealtimes (PDF 173KB)</u> for children and young people with eating and drinking difficulties include:

- upright, with shoulders over hips, level shoulders and slightly forward
- bottom and hips well back into the chair to give a stable base for the trunk (eg use of pelvic strap)
- knees at around 90 degrees and comfortably apart (eg a pommel or padded wedge or block may be needed to help separate the knees)
- feet comfortably apart and resting on the floor or a support where possible (eg footstool or footplates on a wheelchair or highchair)
- head centred over the shoulders with an elongated neck, in a relaxed, upright and slightly chin-tuck position (remember head back will open the airway and swallowing is extremely difficult and unsafe)
- shoulder straps, chest harness give added support where advised
- head and neck positioning can be helped by a well-positioned head rest or even the use of a rolled towel, small cushion or even a neck brace (where advised)
- generally arms forward on the table or tray
- where possible the child or young person should be encouraged to move slightly forward towards the food, spoon, or drink as this is part of the normal active process for eating or drinking
- often a small wedge cushion or angled seat base can help for children and young people who have better trunk control and more active movement
- for children and young people who can't do this, make sure positioning is good and head on neck alignment is suitable for swallowing (eg elongated neck and slight chin tuck).

The position of the support person helping the child or young person during mealtimes is also important. They should be seated at head level with the person they are helping and their chair directly in front, or facing them across the table. This aids with communication and social interaction and allows the support person to effectively monitor the feeding and respond quickly if the child or young person is experiencing difficulty.

Adequate time must be taken to make sure the child or young person is positioned correctly for mealtimes, and recommended positioning equipment is used correctly.

Equipment

Adapted mealtime equipment may be prescribed for a child or young person with eating and drinking difficulties by the speech pathologist or health professional. The child or young person's <u>oral eating and drinking care plan (DOC 254KB)</u> will describe the specific equipment requirements.

Types of adapted mealtime equipment are endless and may include:

- drinking equipment:
 - o spout cups; help deliver liquid to the middle of the tongue
 - o cut-out cups; designed for cut out space for the nose when cup is tipped helping keep the head and neck in a safe position when drinking

- o regulating cups; allow only a small amount of liquid per sip
- o spill resistant cups; help prevent spillage
- o weighted cups; to help with minimising hand tremors
- o straws with non-returnable valves; liquid stays in the straw rather than returning to cup or bottle where the child or young person has difficulty with sucking
- o hands free drinking systems; can be placed on a table or mounted to personal equipment (wheelchair)

cutlery:

- o plastic-coated spoons; reduce damage to teeth and gums and generally have a shallower bowl than a regular spoon
- o maroon spoons; recommended where there is difficulty using lips to get the food from a regular spoon
- o cutlery with large grip; built-up handle, angled neck; increases independence to feed self
- plates, bowls, trays:
 - o plates and bowls with built up edges; increases independence to scoop out food
 - o plate guards; prevent spillage and allow scooping of food
 - o insulated plates and bowls; retains heat to keep the food warm over a longer period of time
- clothes protectors; to help with spillage and excessive loss of saliva.

Medication administration

Administration of medication in the education and care setting must be done following the <u>medication</u> <u>management procedure (PDF 933KB)</u>. Where medication is prescribed for a child or young person as a management strategy for their eating and drinking difficulty, a <u>medication agreement (DOC 172KB)</u> must be completed by the treating health professional. An <u>oral eating and drinking care plan (DOC 254KB)</u> cannot be used as a medication agreement.

Mental health and oral eating and drinking

Difficulty swallowing is a common symptom of anxiety, or having oral eating and drinking difficulties may contribute to increased anxiety for children and young people, as well as their parent/guardians.

In some cases anxiety may become debilitating, preventing the child or young person from engaging in daily activities at home, school, or socially.

Some symptoms of stress and anxiety may be mistaken for eating and drinking difficulties in children and young people. Education is important to make sure staff can identify anxiety symptoms from those of eating and drinking disorders, ensuring appropriate treatment (refer to <u>training and education</u> for further information).

High levels of anxiety are often seen in parent/guardians of children and young people with oral eating and drinking disorders. Regular and ongoing communication with parent/guardian is important to reassure them

of the strategies in place to manage the child or young person with an emphasis on the ability of the education or care service to provide a safe environment.

Training and education

All education and care settings must have at least one designated first aider who is trained in

Emergency First Aid Response in an Education and Care Setting">Education and Care Setting in attendance at all times who is immediately available to administer first aid.

The speech pathologist or health professional responsible for developing the <u>oral eating and drinking care</u> <u>plan (DOC 254KB)</u> may offer advice on implementation of the plan and may give training and education on techniques and equipment.

Education and care staff providing support and help to children and young people with eating and drinking difficulties, are encouraged to undertake the <u>Novita Children's Services Oral Eating and Drinking Support</u> (OEDS) workshop. The workshop includes practical, relevant information about safe practices and worksite risk management for children and young people requiring assistance and support with oral eating and drinking. Training and education can be arranged by contacting Central Intake Novita, training@novita.org.au or 1300 668 482; and is funded by education and care services.

The <u>Ellyn Satter Institute</u> has resources and links for education and care services that may help education and care staff that care for a child or young person requiring oral eating and drinking support.

<u>Feeding Matters</u> have a range of webinars and virtual workshops that can be viewed on demand and used as learning opportunities to support and manage children and young people requiring help with oral eating and drinking.

Communication and risk minimisation

Communication

Communication strategies for education and care services where a child or young person requiring support and help with eating and drinking should include:

- staff awareness and understanding of management strategies and emergency response for individuals
- regular communication with parent/guardians to give assurance that appropriate management, risk minimisation and emergency response strategies are in place
- education or care service complete <u>oral eating and drinking observation log (DOC 66KB)</u> to communicate changes in mealtime behaviours and/or eating and drinking abilities
- parent/guardian to communicate any changes to the child or young person's care to make sure education and care staff are up-to-date
- where age appropriate, communication to peers to identify early signs of a medical emergency and risk minimisation strategies.

Risk minimisation strategies

Individual risk minimisation strategies should be documented in the <u>safety and risk management plan (DOCX 140KB)</u> where a child or young person has feeding difficulties, dysphagia, oral sensitivities, food or choking phobias in consultation with the parent/guardian.

Risk minimisation and prevention strategies should be considered for all in-school and out-of-school settings.

General prevention strategies for education and care services to minimise the risk of:

- a choking incident may include:
 - o avoiding food that pose a choking risk to small airways (under 3s most vulnerable), any food that is small and firm is a choking hazard. ie popcorn, hotdogs, grapes, stone fruits, raw carrots, sausages, nuts, pieces of apple, corn chips, lollies. Further information available on International Dysphagia Diet Standardisation Initiative (IDDSI) Framework webpage.
 - encouraging children and young people to sit down for meals and snacks and not talking with food in mouths
 - o beware of toys and items that may be a choking hazard, ie deflated balloons, beads, small toy parts, batteries, coins, bottle caps, pebbles/stones in gardens
 - making sure safe, age-appropriate toys are available by following manufacturers age recommendations
 - o making sure all education and care staff are trained in first aid, including choking first aid and understand the emergency response process for a choking emergency
 - o review of hazard alerts (staff login required) and implementation of any actions
- an adverse event for a child or young person who has feeding difficulties, dysphagia, oral sensitivities, food or choking phobias may include:
 - o (all of the above listed under choking)
 - o making sure appropriate supervision of children and young people at risk
 - o identification and documentation of specific habits or behaviours in the health care or support plan
 - o advice received from the child or young people's parents or primary caregivers
 - recommendations made by the speech pathologist or health professional about mealtime management requirements are clearly documented, available, and strictly adhered to, including food preparation, positioning, environment, equipment etc.

Oral eating and drinking background

Syndromes and disorders

Certain syndromes and disorders may affect eating behaviour, including (but not limited to):

Prader-Willi Syndrome – a complex genetic disorder with a symptom including a chronic feeling of

hunger, leading to excessive eating and obesity

- Down Syndrome may have a small oral cavity and delayed development of teeth, resulting in difficulties chewing
- people with cerebral palsy, epilepsy or degenerative disorders may have difficulties eating and swallowing
- <u>hyperphagia or polyphagia</u> eating excessively without feeling full can be characteristics of other syndromes; can result in food grabbing, eating inedible objects, obesity and behavioural problems
- Autism Spectrum Disorder autistic children and young people commonly show eating difficulties such as food refusal, selective eating, over-eating and behavioural problems at mealtimes.

Eating disorders

An eating disorder is a serious mental illness with significant complications. The mortality rate for people with eating disorders is the highest of all psychiatric illnesses with the peak period of onset being during adolescence.

The <u>Diagnostic and Statistical Manual of Mental Disorders</u> (DSM-5) recognised 4 eating disorders:

- other specified feeding and eating disorders (OSFED)
- bulimia nervosa
- binge eating disorder
- anorexia nervosa.

For information about eating disorders, refer to National Eating Disorders Collaboration (NEDC).

Feeding

<u>Undernourishment</u> can lead to poor brain and linear growth and may affect the child or young person's mental health. Feeding disorders are not limited to children and young people with disabilities, however between <u>80% and 90%</u> of children with disabilities have some level of eating or drinking difficulty. As the level of disability increases, so does the severity of the oral feeding difficulty.

A feeding disorder is a child or young person's refusal or inability to eat certain food groups, textures, solids or liquids for a period of time (generally at least one month) causing weight loss, inability to grow naturally, and/or developmental delays. A feeding disorder can be caused by structural or functional abnormalities that affect physiology; or behavioural as a result of psychosocial or emotional difficulties. Most children and young people have multiple components to their feeding disorder.

Feeding disorders in children and young people may include (but are not limited to):

- extreme food selectivity (beyond pickiness) by type (excludes more than 1 food group), texture (only eats smooth or crunchy foods), or by brand, shape or colour
- gags when food is brought near the mouth
- has trouble paying attention long enough to eat a meal
- taking more than 30 minutes to eat a meal.

Medical interventions, such as nasogastric or gastronomy tubes, are used when children and young people can't take adequate nutrition by mouth (also be referred to as enteral nutrition or feeding, tube feeding, PEG feeding, gavage).

Some of the most common conditions that require enteral nutrition include (but are not limited to):

- congenital (present from birth) abnormalities of the mouth (cleft lip or palate), oesophagus, stomach, intestines, or heart
- sucking and swallowing disorders, which are often related to prematurity, brain injury, developmental delay, or certain neuromuscular conditions such as cerebral palsy
- failure to thrive, which is a general diagnosis that refers to a child's inability to gain weight and grow
 appropriately. Poor growth can be the result of an underlying medical condition such as cystic
 fibrosis, certain heart defects, cancer, intestinal problems, severe food allergies, or metabolic
 disorders, among other things.
- extreme difficulty taking medicines
- inability to burp to reduce reflux (stomach contents and acid moving backward from the stomach into the oesophagus).

Swallowing (dysphagia)

A swallowing disorder is called <u>dysphagia</u> and includes any problem with sucking, swallowing, drinking, chewing, eating, controlling saliva, taking medication or protecting the lungs from food and drink 'going down the wrong way'.

There are 3 phases of swallowing:

- oral (preparation and transit)
- pharyngeal
- oesophageal.

Dysphagia can occur in 1 or more of these phases.

Oral

May include:

- problems closing lips to keep food in mouth
- trouble chewing or moving food and liquid from the mouth to the throat
- unable to control food or liquid so it goes into the throat too quickly
- food stuck in cheek (pocketing)
- having residue left in mouth after swallowing.

Pharyngeal

May include:

- difficulty coordinating a swallow
- food or liquid entering into the airway (pulmonary aspiration)
- food or liquid remaining in the throat after swallowing (residue).

Oesophageal

May include:

- food getting stuck as it moves from the throat to the oesophagus
- poor movement of the oesophagus so food doesn't pass easily into the stomach
- food coming back up (reflux); food getting stuck in the oesophagus.

Signs and symptoms

Signs and symptoms of swallowing and feeding disorders vary based on the medical condition and age of the child or young person, but may include:

- back arching
- gagging
- breathing difficulties when feeding (may be signalled by increased respiratory rate)
- skin colour change (turning blue)
- coughing and/or choking during or after swallowing
- dehydration
- difficulty chewing foods or spitting out partially chewed foods
- difficulty initiating swallowing
- difficulty managing secretions (drooling)
- disengagement cues, such as facial grimacing, finger splaying, or head turning away from food source
- congestion, particularly after meals (including noisy or wet vocal quality)
- frequent respiratory illnesses
- food or fluids coming out of the nose
- refusing foods of certain textures or types
- taking only small volumes, over-packing the mouth, and/or pocketing foods
- vomiting
- weight loss or lack of appropriate weight gain.

Oral sensitivities (chewing, sucking, mouthing, pica)

The oral system is responsible for sending signals to our brain, organising information and receiving input. All

children and young people require this awareness and input for development; oral fixation can occur when a child or young person is either over responsive (hypersensitive) or under responsive (hyposensitive) to receiving information from the oral system. This is also sometimes referred to by the term 'oral sensory seeker'.

Signs and symptoms of a child or young person with oral defensiveness can include:

HYPERsensitivity

- don't like brushing their teeth, eating or washing their face
- difficulty sucking, chewing or swallowing
- may have frequent choking incidents
- picky eaters
- don't like eating food with certain temperatures or textures (food aversions), or strong flavours
- gagging when eating
- require liquid for food to go down
- may use teeth to remove food from fork or spoon
- may have other tactile sensitivities throughout the body

HYPOsensitivity

- pica licking, biting and chewing on nonfood objects
- biting (including themselves, or other children)
- little awareness of what is going on in mouth
- messy eater (often leaves food in or around the mouth after eating)
- can overfill their mouth resulting in choking
- drooling
- prefers strong flavours
- difficulty distinguishing between different tasting foods

Pica is a persistent eating disorder that involves eating items that are not typically thought of as food and do not contain significant nutritional value such as hair, dirt and paint chips. The eating of these substances must be developmentally inappropriate, children under 2 years of age should not be diagnosed with pica. The compulsive eating of material can sometimes be due to a nutritional deficiency but is most common in children and young people with autism, intellectual disabilities, or mental health disorders.

Chewing and sucking can form an important part of a child or young person's self-regulation of anxiety and attention levels. Chewing gives proprioceptive (body awareness) feedback to satisfy the sensory input that children and young people may seek orally; and indirectly gives calming and attention regulation through the trigeminal nerve pathways (transmitting sensations from the face to the brain).

Before beginning a treatment plan for oral sensitivities or pica it is recommended the parent/guardian consult a health professional (doctor and/or dentist) to rule out any medical issues that may be the underlying cause.

Oral sensitives

A speech pathologist and/or other health professionals can complete a sensory evaluation to determine individual needs for a child or young person that may have oral sensitivities.

A <u>non-specific health care plan (DOCX 124KB)</u> may be developed that includes recommendations, interventions, techniques and precautions; this may include oral sensory activities (chewing gum, drinking through a straw), textured foods (crunchy snacks that require a lot of chewing) or equipment (fidgets or non-food items to chew on).

Some oral sensory activities can be included in classroom activities ie blowing bubbles or whistles.

Pica

Given the risk of medical complications associated with pica (ie poisoning, tearing of the oesophagus or intestine) close monitoring of the child/young person is required to prevent and monitor for effects of ingestion.

If a potentially poisonous non-food substance has been consumed phone the **Poisons Information Centre 13 11 26** to speak with the, **for life-threatening or urgent medical attention, phone 000 (ambulance).**

The diagnosis and treatment for each individual will be different. Appropriate interventions and precautions should be included in a <u>non-specific health care plan (DOCX 124KB)</u> and/or <u>health support agreement (DOC 131KB)</u>.

Treatment techniques may include:

- behaviour therapy
- discrimination training between edible and inedible items
- self-protection devices that prohibit placement of objects in the mouth
- oral sensory activities
- environmental risk assessments or scans.

Food aversion or choking phobias

Food aversion or choking phobia is known as cibophobia.

Cibophobia and the fear of eating in public or eating disorders, eg anorexia, are not the same. The fear of eating in public is a social anxiety disorder. People with eating disorders fear the effects of food on the body image, whereas people with cibophobia have a fear of the food itself, or of swallowing the food. This often stems from a negative or traumatic episode (choking, vomiting, allergic reaction) after eating or drinking. In some cases, cibophobia may be associated with or result in anorexia, bulimia or other eating disorders.

Cibophobia might be about a particular food, brand or whole group of food such as meat, fruit or vegetables.

Treatment for food or choking phobias and/or eating disorders generally involves cognitive-behavioural therapy in combination with other treatments ie hypnotherapy. These therapies aim to change the attitude and beliefs relating to the phobia. Other treatment options include relaxation training and/or desensitisation.

A <u>non-specific health care plan (DOCX 124KB)</u> should be developed and any recommendations, interventions, techniques and precautions for the child or young person should be documented in the <u>health support agreement (DOC 131KB)</u> and <u>safety and risk management plan (DOCX 140KB)</u>.

Some treatment options may include:

- pureeing foods
- supervision while eating or prefers to eat alone
- eating smaller amounts over a longer period

- relaxation techniques before eating
- liquid forms of medications (if needed).

Definitions

Access Assistant program (Women's and Children's Health Network)

Supports children and young people with a disability and/or who have complex health support needs, so they can participate in the pre-school or school curriculum.

Provides professional health advice to support departmental health related policies.

allied health professional

Trained professionals (other than doctors, nurses and dental professionals) that can manage physical or mental health through diagnosis, treatment or rehabilitation. Allied health professionals include (but are not limited to) counsellors, dietitians, occupational therapists, physiotherapists, psychologists, social workers, speech pathologists.

aspiration

Also: pulmonary aspiration. Aspiration is the entry of material (food, drink, stomach contents) from the throat into the larynx (voice box) and lower respiratory tract (the portions of the respiratory system from the trachea—ie windpipe—to the lungs).

cibophobia

Also: food phobia, swallowing phobia. The fear and avoidance of food, chewing or swallowing fluids usually as the result of a negative or traumatic episode such as choking or vomiting after eating or drinking.

dysphagia

Difficulty or discomfort in swallowing solids or liquids.

education and care services

Includes children's centres, preschools, schools, family day care, home based childcare, respite care programs.

oral sensitivities

Also: oral sensory processing, mouthing, chewing. Oral sensitivity is defined as an atypical response to oral

stimulation. Atypical responses include hyporeactive responses, hyperreactive responses, or sensory defensiveness.

pica

The persistent craving and compulsive eating of non-food substances. Classified as a feeding and eating disorder of childhood.

tube feeding

Also: nasogastric tube, gastronomy tube, enteral nutrition, enteral feeding, PEG feeding, gavage. A medical device used to give nutrition to patients who cannot get nutrition by mouth, can't swallow safely, or need nutritional supplementation.

Supporting information

Choking first aid for adult or child over 1 year (PDF 140KB)

Choking first aid for infant under 1 year (PDF 162KB)

HSP120 Health support agreement (DOC 131KB)

HSP121 Safety and risk management plan (DOCX 140KB)

HSP124 Individual first aid plan (DOCX 119KB)

HSP125 Guide to planning health support (DOC 87KB)

HSP210 Oral eating and drinking care plan (DOC 254KB)

HSP211 Oral eating and drinking observation log (DOC 142KB)

HSP220 Transfer and positioning care plan (DOCX 175KB)

International Dysphagia Diet Standardisation Initiative (IDDSI) Framework

Kids Health Information: Safety: Choking, suffocation and strangulation prevention

Related legislation

Code of Practice First Aid in the Workplace 2012

Work Health and Safety Act 2012 (SA)

State Records Act 1997 (SA)

Disability Discrimination Act 1992 (Cth)

Disability Standards for Education 2005

National Disability Insurance Scheme Act 2013 (Cth)

Education and Early Childhood Services (Registration and Standards) Act 2011 (SA)

Education and Care Services National Law 2010

Education and Care Services National Regulations and within those regulations in particular: Regulation 12(b)(i); Regulation 77; Regulation 78; Regulation 79; Regulation 90; Regulation 91; Regulation 95; Regulation 96; Regulation 136; Regulation 153(1)(j); Regulation 162(c), (d) and (e); Regulation 168(2); Regulation 177(1)(c); and Regulation 183(2)(a), (b) and (c)

Related policies

Duty of care to children and young people policy (PDF 182KB) (staff login required)

First aid and infection control standard (PDF 258KB)

Medication management procedure (PDF 933KB)

Record history

Published date: February 2024

Approvals

OP number: 162

File number: 18/07533

Status: approved Version: 2.0

Policy officer: Senior Advisor Health Support, Disability Policy and Programs

Policy sponsor: Director, Inclusive Teaching and Learning

Responsible Executive Director: Executive Director, Support and Inclusion

Approved by: Director, Inclusive Teaching and Learning

Approved date: 19 February 2024 Next review date: 19 February 2027

Revision record

Version: 2.0

Approved by: Director, Inclusive Teaching and Learning

Approved date: 19 February 2024 Review date: 19 February 2027

Amendment(s): Major edit following Coronial inquest. Updated first aid guidelines for a choking emergency (to reflect those of the Australian Resuscitation Council). Updated to provide advice about the safe storage

and consumption of food and drinks on departmental sites.

Version: 1.2

Approved by: Director, Disability Policy and Programs

Approved date: 27 October 2021 Review date: 27 October 2024 Amendment(s): Updated hyperlinks.

Version: 1.1

Approved by: Director, Disability Policy and Programs

Approved date: 10 March 2020 Review date: 10 March 2023

Amendment(s): Reviewed and updated in new template.

Version: 1.0

Approved by: Senior Executive Group Approved date: 10 August 2018 Review date: 10 August 2019

Amendment(s): New procedure developed incorporating information from the former 'Oral eating and

drinking in education and children's services' FAQs revised edition December 2015.

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