

Supplementary Table 2: World Health Organization Trial Registration Data Set.

Data category	Information³²
Primary registry and trial identifying number	ClinicalTrials.gov NCT05734586
Date of registration in primary registry	8 February, 2023
Secondary identifying numbers	69HCL22_0474
Source(s) of monetary or material support	Hospices Civils de Lyon, France
Primary sponsor	Hospices Civils de Lyon, France
Secondary sponsor(s)	N/A
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Public title	Screening for Sarcopenic Dysphagia and the Implementation of Measures to Prevent Its Complications in Geriatric Patients [DYSPHAGING-PILOT]
Scientific title	Feasibility Study of Screening for Sarcopenic Dysphagia and the Implementation of Measures to Prevent Its Complications in Geriatric or Institutionalized Patients Aged ≥ 70 Years.
Countries of recruitment	France

Data category	Information ³²
Health condition(s) or problem(s) studied	Swallowing Disorder, Sarcopenic Dysphagia
Intervention(s)	<p>Other: EAT-10 (Eating assessment Tool) screening questionnaire After inclusion, issuance of the EAT-10 screening questionnaire for swallowing disorders by the healthcare team</p> <p>Procedure: Protective measures for the upper airways</p> <p>In the event of an EAT ≥ 2 score, immediate implementation or within three days by the healthcare team of protective measures for the upper airways in 3 sectors: 1: Postural maneuvers; 2: Hygienodietetic rules; 3: Food textures</p>
Key inclusion and exclusion criteria	<p>Inclusion Criteria:</p> <ul style="list-style-type: none"> • Patient aged ≥ 70 years, • Patient affiliated to a social security system, • Patient hospitalized in the health sector or in a medico-social institute, • Patient informed of the study (information leaflet provided) and having orally signified their consent to inclusion in the study. <p>Exclusion Criteria:</p> <ul style="list-style-type: none"> • Patient unable to feed orally, • Patient with an active pathology responsible for acute swallowing disorders (< 3 months) (neurodegenerative pathology with predominant motor impairment such as Charcot's disease, stroke, ENT disease). • Patient unable to answer the questionnaire.
Study type	<p>Interventional Allocation: N/A Intervention model: parallel assignment Masking: None (Open Label) Primary purpose: Other Phase II</p>
Date of first enrolment	June 1 st ,2023
Target sample size	102
Recruitment status	Recruiting
Primary outcome(s)	<p>Proportion of complete achievement of steps 1 and 2 [Time Frame: Three days] The judgment criterion is validated if</p> <ol style="list-style-type: none"> 1. Stage 1 is performed and the EAT-10 < 2 or if 2. Stage 1 is performed with an EAT-10 ≥ 2 and stage 2 is performed within 3 days after stage 1.

Data category	Information ³²
Key secondary outcomes	<ul style="list-style-type: none"> • Percentage of eligible patients refusing to participate in the study [Time Frame: 18 months] <ul style="list-style-type: none"> ○ Number of eligible patients who refused to participate in the study • Age, gender, comorbidities (CIRS-G), autonomy (ADL, IADL), co-medications [Time Frame: 19 months] <ul style="list-style-type: none"> ○ Patient characteristics will be collected at each site at the end of the study by a clinical research assistant based on their medical records. • Rate of partial completion of the protocol [Time Frame: 19 months] <ul style="list-style-type: none"> ○ Proportion of non-performance of step 1 and/or step 2 within the time limit. Proportion of steps 2 carried out incompletely), description of the reasons • Diagnosis of undernutrition and/or neurocognitive disorders and/or patent lung infection and/or COPD described in the patient's medical file, nutritional risk situation assessed by the Mini Nutritional Assessment® (MNA) [Time Frame: 19 months] <ul style="list-style-type: none"> ○ Patient characteristics will be collected at each site at the end of the study by a clinical research assistant based on their medical records. • Composition and disciplines of the care team [Time Frame: 19 months] <ul style="list-style-type: none"> ○ At the end of the study, all data on the each allied health professionals will be collected on the dysphaging sheet • Satisfaction of the care team (Likert scale). [Time Frame: 19 months] <ul style="list-style-type: none"> ○ At the end of the study, each allied health professionals who has been involved in the care of at least one patient will fill out a satisfaction questionnaire.