

Generation Healthy Kids

Main study

[Generation Sunde Børn - hovedstudie]

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STØTTET AF **novo nordisk fonden**

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3 Abbreviations and Definition of Terms

AE	Adverse Event
BMI	Body Mass Index
CAPL	Canadian Assessment of Physical Literacy
CCRP	Center for Clinical Research and Prevention
CRF	Case Report Form
DAPL	Danish Assessment of Physical Literacy
FM	Fat mass
GDPR	General Data Protection Regulation
LCPG	Local Community Partnership Group
MVPA	Moderate to Vigorous Physical Activity
NEXS	Department of Nutrition, Exercise and Sports
PA	Physical Activity
PE	Physical Education
PL	Physical Literacy
SDU	University of Southern Denmark
SES	Socioeconomic Status
SFO	After School Club
UCPH	University of Copenhagen
VPA	Vigorous Physical Activity

4 Background

4.1 The global problem of overweight and obesity

Overweight and obesity among children is a global problem affecting more than 41 million children under the age of 5 years. The World Health Organization (WHO) describes childhood obesity as one of the most serious public health challenges worldwide for the 21st century (1,2). The prevalence of overweight and obesity in Danish children is 12-13% in primary school and increases to 18-19% at the time of graduation (3), why early prevention is important. In the adult population 51% are estimated to be overweight or obese (4,5).

Overweight and obesity have many negative health consequences for children during their early years and in later life. For instance, children who have overweight or obesity are more likely to experience bullying, stigmatization and suffer from low self-esteem, anxiety, and lower quality of life (6). In addition, children with overweight and obesity have an increased risk of overweight and obesity as adults, which is well known to lead to poorer health and well-being and marked economic negative effects for the individual and for society.

There are stark inequalities in the prevalence of childhood overweight, both worldwide and in Denmark, contributing to the health inequality found in most countries (4,7). In Denmark, the overweight prevalence among children aged 6-7 years in high-income families is 11% compared to 26% in low-income families. Among children aged 14-15 years, the prevalence of overweight is almost 30% in families with short education compared to 10% among those from families with long education (8).

Many trials have been conducted to reduce childhood overweight and obesity, and several comprehensive systematic reviews have investigated the effect of different interventions in preventing childhood overweight and obesity (9–16). However, most of the earlier studies find no or only small effects on children's body mass index (BMI), and the evidence of long-term effects is limited.

Thus, there is a strong need to develop new innovative approaches and programs that effectively can prevent overweight among all children, starting early in life and embedding promising programs and integrated approaches into everyday practices.

4.2 Important characteristics to increase likelihood of success

Some of the earlier studies did show a significant effect on overweight measures. Across these studies, some characteristics and elements seem to increase the likelihood of success:

- A multi-component, multi-setting intervention, including well-planned involvement of relevant stakeholders, is likely to yield the highest level of effect in promoting healthy weight in children (17,18).

- Multi-component interventions targeting two or more health behaviours (i.e., physical activity, dietary outcomes, sedentary behaviour, sleep) seems to be more successful in improving adiposity outcomes compared to single-component interventions (12–14,19–24). Involving parents was found to be essential in the prevention of childhood overweight and obesity, especially among children younger than 12 years (9,11,13,23,25–29). However, less than half of the earlier studies involved parents (25). Among the studies that did involve parents, three out of four demonstrated positive outcomes in reducing BMI or weight status.

Other characteristics that seem promising are:

- Combining educational components (e.g. targeting the children's knowledge, competences, literacy levels, and motivation) with environmental or structural components (e.g. free healthy school meals, smaller portion sizes, easy access to fruit and vegetables, physical activity as a part of the school day, free and/or easy access to leisure time activities, healthy space management of supermarkets, healthy alternatives in restaurants, decreased availability of fast food and sugary beverages, and healthy school policies) (9,13,16,28,29). The environment around the child, both at school and at leisure time (home and local community) must support healthy living.
- Targeting a multitude of the settings where children spend their time (23,30) and including multiple intervention components (9–16). Several systematic reviews have shown limited effects of interventions that are purely school based (31).
- Long-term interventions (twelve months or longer) appear to be more effective compared to studies of shorter duration (9,13).
- Co-creation and involvement of relevant stakeholders was also found to be an important factor in earlier, successful studies. This means involving stakeholders in identifying, prioritizing, developing, and implementing interventions using co-creative approaches and capacity building processes (32–37). (16,38,39).

4.3 Key components in prevention - diet, physical activity, screen time and sleep

Healthy diets and balanced energy intake are key factors in prevention of overweight in childhood, which must be addressed simultaneously in different settings for maximal effect, in a manner that supports food literacy, practical competence and long-term behaviour change via environmental changes. Danish children consume too little vegetables, wholegrains, and fish, and too much sugar and fat, compared to the Danish food-based guidelines and the Nordic Nutrition Recommendations (40–42). This worsens with age, which means that preventive efforts are needed at an early age. There is considerable social inequality in dietary behaviours, as Danish children from households with short compared to long education consume less fruits, vegetables, dietary fibre, and fish and often skip breakfast

(43). School food programs can improve dietary quality (44), and in Sweden, provision of free school lunches during basic school has been associated with long term effect on different indicators for better health and well-being, including education attainment, higher salaries, and higher final height later in life (45).

The diet of Danish children has been found to be less healthy during weekends compared to weekdays (46), and studies have shown that children primarily learn about eating behaviour, dieting, cooking etc. by watching their parents. Further, parents define their children's home food environment, e.g., access to healthy and unhealthy food items and whether there are family rules limiting the intake of unhealthy foods (47,48).

Physical activity (PA) is an important part of a healthy lifestyle for children, as it improves physical fitness, motor competences and well-being, and reduces the risk of developing overweight and non-communicable diseases later in life (49). Current recommendations are that children and adolescents should participate in at least 60 minutes of moderate-to-vigorous physical activity (MVPA) per day and 3x30 minutes of vigorous physical activity (VPA) per week. However, in Scandinavia and the rest of Europe, many children do not reach these targets (50,51).

A high percentage (about 70 percent) of 7-9-year-old children in Denmark participate in one or more club-organized sports. This contributes significantly to these children's MVPA and increases their chance of being active at a later age, when physically active free play during e.g., recess decreases. The participation rates, however, are much lower for children whose parents have low sporting experience and in families with limited socioeconomic resources.

Excessive recreational screen media use and short- and poor-quality sleep are important risk factors for the development of childhood obesity (52,53). While high screen use may be obesogenic by displacing engagement in PA during leisure time, it may also lead to insufficient sleep among children (54). Sleep is not only critical for learning, memory, and school performance. Inadequate sleep also disrupts hormone levels that regulate appetite and food intake (53). A third of Danish 6-11-year-old children use screen media >4 hours/day during weekends, and 24% of children in this age group report screen use just before bedtime on all weekdays (7,55). Furthermore, significant socioeconomic disparity is seen in children's screen media use, with higher prevalence of excessive recreational screen use and screen use just before bedtime in children from socioeconomic disadvantaged families. Insufficient sleep is also more common among children from families of low socioeconomic status (56).

In addition, we observe that more healthy screen use practices appear to be promoted by well-established parental rules of screen media use hygiene (when to use screen media,

content- and amount of use), and we find a graded relationship of total recreational screen use, screen use just before bedtime and sleep time of children (57).

4.4 Generation Healthy Kids – the innovative approach

Generation Healthy Kids will build on the existing evidence, combining the characteristics and elements described above to improve the chance of success. The innovative aspects of the project are the combination of a randomized controlled trial targeting several important risk factors for excessive weight gain (dietary habits, physical activity, sleep, and screen time) with community capacity building and a systems approach (58). The unique combination is closely monitored for effect as well as process. The integrated intervention program will be unique in combining already tested effective intervention strategies with intervention components developed using co-creation and a systems mapping (58). Systems thinking will furthermore be used to ensure that a health equity lens is employed to understand the relationship between obesity and health inequalities locally (59). To reach children and families with low socio-economic status, focus will be on developing interventions that make healthy choices easy by creating healthy environments around the children, both at school and during leisure time.

4.5 The pilot study and the main trial

Generation Healthy Kids is a 4-year project including a pilot study which will be conducted in January-March 2023, and a 2-year cluster-randomized controlled main trial to be conducted in August 2023 – June 2025. The present protocol describes the main trial.

The pilot study has previously been approved by The Regional Committee on Health Research Ethics for Southern Denmark (Project ID 568021-22 S-20220059).

A thorough development and piloting is an essential prerequisite for ensuring feasibility and quality of the final intervention program to be assessed via the main trial. Based on the learnings and results from the pilot study, the final intervention program will be adjusted accordingly. Due to the timeline of the main trial, this protocol is submitted before the pilot study is conducted. Adjustments based on learnings and results from the pilot study will be included in amendment(s) to the Committee for change of the present protocol.

5 Study Objectives

5.1 Overall Objective

Generation Healthy Kids is a cluster-randomized community trial in which 24 local communities will be randomly allocated to intervention or control. The overall study objective is to investigate if a 2-school-year multi-setting, multi-component intervention focusing on healthy diets, physical activity, sleep and screen media habits in the local community can promote healthy weight and body composition in children who are in 1st and 2nd grade at inclusion (i.e., age 6–9 years at inclusion). We will also investigate the intervention's effects on dietary intake and nutritional status; food competences; physical literacy, activity levels, and fitness; sleep and screen media habits; growth; cardio metabolic health; cognitive and motor functions; school performance; and mental health and wellbeing. Furthermore, we will evaluate the implementation of the intervention.

In continuation of the above, we will investigate if the intervention can reduce social inequality in the outcomes, as well as explore potential effect modifiers such as sex, ethnicity, genetics, epigenetics etc. We will also explore associations between sociodemographic characteristics, health behaviours and outcomes cross-sectionally and longitudinally and use these data to validate and compare relevant measurement tools.

The primary outcome is fat mass (FM) in kg. We hypothesize that the combined intervention components will counter the development of increasing overweight and unhealthy weight with age and thus will cause less FM gain in the intervention group compared with the control group.

5.2 Specific Objectives

5.2.1 Main objective

The main objective is to investigate the impact of the 2-school-year intervention program on development of FM in the intervention group compared with the control group.

5.2.2 Secondary objectives

As secondary objectives, we will investigate the impact of the intervention program after 1 and 2 years on:

- Other body composition measures, including fat free mass (FFM), FFM index, FM index, FFM-to-FM ratio, % FM;
- Anthropometric measures and growth, including BMI z-score, prevalence of obesity, overweight, normal weight and underweight, waist circumference, linear growth and growth-related biomarkers;
- Cardio metabolic health, including blood pressure, blood lipid profile, insulin resistance markers;
- Other health related outcomes, including inflammation;
- Cognitive and motor functions, physical literacy, and school performance;
- Mental health and wellbeing;
- Dietary intake, biomarkers of food and nutrient intake and status and appetite and food competence;
- Physical activity, physical fitness, leisure-time activities, sleep- and screen media habits;
- Long-term (>2 years) body composition based on height and weight obtained from routine data in The National Child Health Register (Børnedatabasen).

Also, the collected data will be used for validations of associations between biological mechanisms and various measurements and outcomes (data analysed as observational data - i.e., based on baseline data, longitudinal data or using the data from the control group only), as well as the impact of background characteristics, such as socioeconomic position, family characteristics, sex, ethnicity, genetics, and epigenetics etc. on the effectiveness of the intervention.

A specific objective of Generation Healthy Kids is to involve relevant local and national partners in the development and implementation of the multi-component intervention program, and to create local engagement and ownership for the implementation of the intervention. The project will work with multiple public, private and non-governmental organizations in the local community, including leisure time- and sports organizations, supermarkets, restaurants, and fast-food chains, to create healthier environments, e.g., by increasing access to healthy foods in supermarkets. The effects of such initiatives will be monitored and evaluated by e.g., sales data from supermarkets.

In addition, an objective of the study is to understand and document implementation-related factors, causal mechanisms and contextual factors which shape the intervention's outcomes. Therefore, a thorough process evaluation will be conducted, see section 6.5.4.

A protocol résumé is found in Appendix 1.

6 Investigational Plan

6.1 Study Design

The study will be designed as a community-based open label cluster randomized superiority trial with two arms (intervention and control). Individual school districts will form the basis of the cluster. A comprehensive list of public schools (“folkeskoler”) will form the school district sampling frame, which is further explained in section 6.2.4. Teachers and pedagogues at each school will have a key role in implementing, coordinating, and anchoring the intervention.

6.1.1 Justification of study design

The study is designed as an open label cluster randomized trial as the primary objective of the study is to examine the effectiveness of the Generation Healthy Kids Program on the primary and secondary outcomes specified in sections 5.1 and 5.2.

6.1.2 Randomization

In this cluster-randomized trial, randomization will occur at school district level and will be performed prior to inclusion of participants (i.e., children) and collection of informed consent. A stratified block randomization will be carried out in a 1:1 ratio. To prevent baseline imbalance of cluster characteristics across the two study arms, we will use covariate-constrained randomization (60). The covariates will be at school- and school district level with tentative factors being number of classes per school year (at least two classes per school year required); the socioeconomic profile of the school based on parental educational level (proportion of parents with a shorter education); ethnicity of the children (proportion of pupils with non-Danish origin in the respective school); and degree of urbanization of school district. The randomization procedure will be carried out by a statistician not otherwise involved in the project (i.e., not involved in recruitment of school districts or participants), and the procedure will be concealed to the investigators.

6.1.3 Timeframe of the trial

The trial will run over two school years (2023-24 and 2024-25). Baseline data will be obtained before the start of the intervention, and follow-up data will be collected during the two school years period.

6.1.4 Sample size justification

The sample size calculations are based on the primary outcome for the study being FM (kg) at 22 months as measured by air displacement plethysmography (BODPOD). For the power calculation, a mean cluster size of $n=90$ children is assumed, which we expect based on our inclusion criteria of ≥ 2 classes per school year, an anticipated mean of 20 enrolled children per class, and a drop-out/missing data of 15%. Others and we have managed to limit missing data to approximately this amount in previous large-scale school-based cluster randomized trials in young people (61–63). Thus, we aim to enrol a mean of $n=105$ children from each school district. We assume a cluster size coefficient of variation of 0.24 based on the OPUS School Meal Study (64) PHASAR data (65) (1st to 4th grade children), a standard deviation of FM of 4.65 kg in both groups based on OPUS, and a conservative estimate of the intra-cluster correlation coefficient in FM (kg) of 0.03 based on PHASAR data (PI's of the OPUS and PHASAR study are part of the scientific team in Generation Healthy Kids). Finally, because we plan to adjust for baseline FM (kg) in our primary analysis, we also assume a correlation between baseline and follow-up FM (kg) of 0.8 in the power calculation. With an expected enrolment of 24 clusters in 1:1 ratio allocation to intervention and control, we have decided to additionally control for type-1 error due to a planned correction of small (cluster) sample using the Satterthwaite correction for mixed linear models (66) by lowering alpha to 0.025 in the power calculations. With a cluster size of 90 and 24 clusters (expected $n=2,160$ children in the analytical sample and 2,520 enrolled children at baseline), we have a power of 80% to detect a significant mean difference of -0.76 kg FM or greater at 22 months between intervention and control. In OPUS data, this difference corresponds to -0.15 BMI-z score (95% CI -0.144; -0.158), and a previous cohort study based on The Copenhagen School Health Records Register reported that a 1 kg increase in FM at 10 years of age was associated with a 12% and 15% increased risk of type 2 diabetes at 50 years of age for boys and girls, respectively (67), which corresponds to an 8% and 10% decrease in risk per -0.76 kg FM. Thus, the planned study is powered to detect a fairly small, but also clinically relevant size of mean difference in FM at 22 months between intervention and control.

6.1.5 Blinding

The study will be open label, as it will not be possible to blind participants due to the nature of the intervention. Also, it is not possible to blind personnel delivering the intervention (i.e., teachers). Outcome assessors will not be blinded, as we judge this to be impossible to implement with success. However, the primary outcome (FM) is objectively assessed not involving judgement by the assessor during the assessment and therefore unlikely to be influenced by knowledge of intervention allocation.

6.2 Study Participants

The study population will be approximately 2,500-2,600 children in 1st and 2nd grade (i.e., age 6–9 years at baseline) who attend public schools in 24 school districts from selected municipalities.

We have selected young school children as the target group for the intervention, because the prevalence of overweight and obesity increases from early childhood until adolescence, from approximately 12-13% at ages 6-7 years to approximately 18-19% at ages 14-15 years (68–72). Furthermore, early childhood is an important time period for formation of habits in relation to diet and physical activity which may persist into adulthood (73). Thus, early intervention is crucial to prevent unhealthy weight development.

6.2.1 Inclusion Criteria

All children attending 1st or 2nd grade in schools recruited into the study will be eligible to participate.

6.2.2 Exclusion Criteria

There are no exclusion criteria for the children. If a child has food allergies, plaster allergies, or chronic disease (Appendix 2a and 2b) or if parents, teachers, or research team judge that a child has physical or mental disabilities which cannot be accommodated in the dietary or physical interventions, the child can participate in the remaining parts of the program.

6.2.3 Withdrawal Criteria

The parents will be informed that they can withdraw their child from the study at any time and for whatever reason without any explanation or consequences for the child's daily life at school. If parents withdraw their child, they will be asked if they are willing to explain the reason, but the response is optional, i.e., entirely voluntary (see Appendix 3). In case of participant withdrawal, the data collected up to the point of withdrawal will be used in data analyses. Withdrawn participants will not be replaced.

The Primary Investigator and Clinical Responsible Physician (independently of each other) have the right and obligation to terminate participation of any child at any time if it is deemed to be in the best interest of the child, e.g.:

- If a child's safety or well-being is compromised by further participation

- If the investigator judges that the child does not want to continue his/her participation

Or in the interest of completion of the study, e.g.:

- If the child does not comply with the study procedures in a way that impacts the scientific integrity of the study.
- Reasons for withdrawal will be recorded, and parents will be informed about the decision.

Dropouts and children who do not participate in the study will not be excluded from participating in activities which are incorporated as a part of the school's regular schedule and activities, including physical activity during school time, school meals, and learning activities during school time regarding screen media habits, sleep habits and food competences. However, no data will be collected from these children.

6.2.4 Recruitment of schools

The sampling frame of the study will be constructed based on information on all public schools ("folkeskoler") in Denmark obtained from The Danish Ministry of Children and Education data-warehouse. Information obtained at school-level will include the number of children attending each school in 1st and 2nd grade. Special schools and schools in areas that already take part in similar research projects will not be eligible for participation. An initial sampling frame list will prioritize schools with at least two classes per year group and relative geographical proximity to the study centres. This sampling frame list will be expanded if the success of recruiting schools is lower than expected. To minimize contamination, eligible school districts are not allowed to lie geographically next to each other.

The schools selected to be part of the sampling frame will be contacted directly via the school leader and the head of the school board, and the school will be invited to participate in the study. Furthermore, municipalities will be contacted to assist with recruitment of schools and to ensure overall support for the study. Finally, in addition to this direct recruitment procedure, an open advertisement for the study will be placed on relevant platforms (e.g., Folkeskolen.dk) to recruit potentially interested schools for the study.

6.2.5 Recruitment and enrolment of study participants

6.2.5.1 Invitations

Families in the selected schools and classes will receive invitations to the study through Aula or E-boks and in printed form from teachers. Aula is an online communication platform used by all schools and approved for the processing of sensitive information.

The invitation message on Aula or E-boks will include:

- An invitation letter, including an invitation to information meetings about the study at the school (Appendix 4A and 4B);
- Written participant information about the study (Appendix 5A and 5B);
- Informed consent form (standard form S6) (Appendix 6)
- The pamphlet '*Forsøgspersonens rettigheder i et sundhedsvidenskabeligt forskningsprojekt*' reviewed and published by the ethical committee system, Sep 2019
- Information about processing of participants' personal data and biological material in accordance with the European General Data Protection Regulation (GDPR).
- Power of attorney for joint custody holders for one custody holder to give consent on behalf of the other custody holder (Appendix 7).

Parents (or custody holders) are asked to carefully read all the information material and to explain the content to their child/children. The information material will be translated into the languages judged by the school and research team to be most relevant for the included classes.

Parents sign up for information meetings at the school, either via a secure online system or by handing in a physical "paper-slip" to their teacher. Parents are informed that they must bring their child to the information meeting, and that they can also bring an assessor ("bisidder") if they wish.

Prior to the information meetings, members of the research team will visit the schools in the morning during the time slot where children are usually dropped off at school, and/or in the afternoon when children are usually picked up or at other times considered relevant by the school. The researchers will hand out one-page information leaflets about the study (Appendix 8), encourage parents to participate in the information meetings, and answer any questions from parents about the study. In addition, one-page information leaflets will be handed out to parents in the involved classes by staff in the afterschool clubs, and posters about the project will be posted at the schools (Appendix 9). Members of the research team may also visit the involved classes during school time and inform the children about the study activities

in a language and at a level appropriate for their understanding, cf. Committee Act § 5, stk. 2. The parents and children will always be informed that participation is entirely voluntary, and that the decision on whether to participate will in no way impact the child's schooling.

6.2.5.2 Information meetings for families at the school

The information meetings will be held at the school in groups of maximum 25 families per meeting. Parents will be informed that they must bring their child to the information meeting.

At the meetings, the background and objectives of the study will be presented by the Investigator, or by a qualified person who is delegated to give the information, and the parts of the study that involve the family and child will be explained in detail to both the parents and the children (see the content of the information meeting in "Table 1. Overview of study procedures and activities"). Also, the parents will be informed that in case of an incidental finding concerning their child, e.g. hyperglycemia, or hypertension, they will be contacted either by phone or e-mail if it is judged to be clinically relevant by the Clinically Responsible Physician.

In line with the Committee Act § 5, stk. 2, information to the children will be provided in a language and at a level that is appropriate and suited for the children's age and comprehension. The information to the children will be provided by project staff experienced in communicating to children or who have been trained specifically for this task. There will be time and room for asking questions, also in a separate room one-to-one. If the parents or the child are unable to participate in the information meetings, they can have the oral information for themselves (see below section 6.2.5.4). If the project staff judges that some parents need a second explanation of the study, this will be encouraged. It will be emphasized that the participants have the right to at least 24 hours consideration time, that participation is voluntary, that the participants can withdraw from the study at any time without giving a reason and without any consequences, and that all personal data will be kept strictly confidential. It will be ensured that parents with insufficient knowledge of Danish will be adequately supported to fully understand the information provided.

The research team will register which parents and children are present at the information meetings, and during information provided to children in the classroom, in order to make sure that the custody holders and the child have received thorough oral participant information and fully understand the study and its procedures.

6.2.5.3 Informed consent process

In case the parents/custody holders and the child want the child to participate and do not wish further consideration time, both parents/custody holders together with investigator or a

delegated and qualified person fill out the consent form (Appendix 6) at the information meeting. They will also be informed that by signing the informed consent they give permission for the controlling authorities to access their data, and that a person carrying out these functions is subject to the rules of confidentiality.

In the supplementary information about GDPR, parents and participants will receive additional information about the way data and biological material are collected, shared, and made available for other researchers and data processors in compliance with GDPR.

If parents/custody holders or children wish further time for consideration, the parents/custody holders will receive an informed consent form to take home. The informed consent form must be signed by the parents/custody holders and the investigator or a delegated and qualified person, and additional questions from the parents must be answered. No study procedures can be initiated before the informed consent is obtained. One parent or custody holder can empower the other parent or custody holder to decide whether the child should participate (Appendix 7). The parents will also be given the opportunity to sign the informed consent form by electronic signature, via a link which will be sent out on Aula or E-boks immediately after the information meeting only to those who participated in the meeting.

As the study participants are under the age of 18, they cannot give informed consent. However, the project staff will be aware of the child's response and reactions to the study (both verbal and non-verbal responses); if the project staff at any time judges that the child does not want to participate in the study, the child will not be included.

6.2.5.4. Parents who do not participate in information meetings

Follow-up information meetings, either in groups or as individual meetings, online or face-to-face, will be arranged for parents/custody holders and children who were not able to participate in the initial meetings.

School staff will be asked to encourage families to participate in information meetings to learn about the study, but it will always be stressed that participation in the study is completely voluntary, and that a decision not to participate will in no way impact their child. We will ask school staff (e.g., teachers and pedagogues) to offer parents the possibility to be contacted by telephone by a member of the research team, if the parents wish further information about the study before deciding whether to participate in one of the information meetings.

6.2.5.5. New students enrolled in the schools during the study period

If new students are enrolled in the participating school classes during the intervention period, they and their parents will be invited to participate in the study. Parents will initially be contacted by the schoolteacher via Aula or in person. If they are interested in hearing more about the study, the parents and children will receive oral and written participant information by the research team following the same procedures as described above in section 6.2.5.3. Written informed consent will be obtained from custody holders before enrolment into the study.

6.2.5.6 Recruitment of local stakeholders

In each local intervention community, a stakeholder analysis will be done together with representatives from the municipality and school. Potentially relevant stakeholders will initially be invited for a bilateral meeting. Relevant stakeholders will initially be defined as community leaders who have authority to initiate action within child health and well-being, including leaders from the municipality, the local school, local sports- and leisure time organizations, and private businesses such as local supermarkets and restaurants. These leaders will build a system map of drivers of obesity in the local community as further described in section 6.3.1. The system map captures drivers of childhood obesity in the local community. Subsequently, a larger group of community representatives will be engaged from organizations whose activities and agenda includes remit and capacity to influence children's food and activity environments, screen time and sleep. Furthermore, selected local leaders and stakeholders in each community will be invited into a Local Community Partnership Group (LCPG) that will function as a steering group during the project period and help ensure coordination and long-term local anchoring of the intervention program.

6.3 Intervention

6.3.1 Principles of the intervention

The development of the intervention in the intervention school districts will be guided by the Medical Research Council framework on designing complex interventions as well as a systems thinking approach (58,74,75). Furthermore, the super-setting approach will guide the process of creating synergy between intervention components (76).

The development of the intervention will take place on three levels:

- a) Core intervention components at school level based on existing evidence

- b) Intervention components developed in co-creation with national stakeholders
- c) Community capacity building in each intervention local community

The core intervention components at school level are pre-defined based on existing evidence and will be described in section 6.3.2 Intervention in schools and after school clubs. The intervention at school level will be refined based on experiences from the pilot study. Furthermore, the intervention components will be adapted to the local context of each school.

Other interventions will be developed primarily outside school, focusing especially on changing the environmental drivers of childhood obesity (social, cultural, physical, economic, and political). Inspired by the Public Health England's publication "Whole systems approach to obesity: A guide to support local approaches to promoting a healthy weight" (59), a co-creation process with national stakeholders took place prior to the pilot study to develop relevant interventions. The national stakeholders included representatives from national supermarket chains, fast food restaurants, Rådet for Sund Mad [Council for Healthy Food], national leisure time organisations, Local Government Denmark (KL), The Danish Healthy Cities Network, Danish Health Authority, The Danish Food Administration etc. After the initial stakeholder analysis, the process of co-creation consisted of a series of bilateral meetings and two workshops. In the first workshop, a systems map was developed using the software STICKE (77). In the second workshop, stakeholders came together to prioritise areas to intervene in the local system and propose collaborative and aligned actions to bridge interventions between settings. The project drew on the expertise and resources of The Intersectoral Prevention Laboratory (www.forebyggelseslaboratoriet.dk) who is responsible for the process of facilitating the creation of a systems map using STICKE.

Community capacity building will take place in each of the local intervention communities. Initially, a stakeholder analysis will be conducted to identify community leaders and stakeholders that have authority to initiate action and that would be relevant to engage in an intervention anchored in the local community promoting healthy weight among local children. Inspired by the cluster-randomized trial, WHO STOPS (78), the process will consist of the phases listed below:

- Collecting and communicating local data and scientific evidence to raise local awareness of childhood obesity and to engage and recruit community leaders.
- Identifying, inviting, and working with a smaller group of community leaders who have authority to initiate action, including leaders from the municipality, the local school, local leisure time organisations, and private businesses including local supermarkets and restaurants. These leaders will build a causal loop diagram of the causes of childhood obesity in their community (79) using STICKE (77).

- Engaging a larger group of community representatives from organizations whose activities and agenda includes remit and capacity to influence children's food and activity environments, screen media habits and sleep.
- Involving the larger group of community representatives in working together to design actions to prevent childhood obesity that they can carry out across the community, inspired by the causal loop diagram or systems map and informed by a prepared evidence brief on obesity prevention, including case studies from previous successful interventions.
- Ongoing data collection and updates of the systems map to enhance implementation and diffusion of the selected actions and stimulate new ideas in a constructive, capacity-building cycle.
- Selected local leaders and stakeholders in each local community will be invited into a LCPG that will function as a steering group.

To develop an overall coordinated intervention programme, the different intervention components developed at different levels and in different settings will be combined using the Super-setting Approach (76). To promote integration and synergy between interventions, common themes for all settings (e.g., mental well-being, taste and senses etc.) will run throughout the intervention period for 2-3 months each.

6.3.2 Interventions in schools and after-school clubs

Prior to initiation of the intervention and measurement schedule (cf. section 6.6.), project staff may visit the school classes and explain the study activities to the children during school time in their classroom. This will occur at a level appropriate for the children's age group by project staff specifically trained for this task. Children whose parents have not provided informed consent to participate in the study will be exempted from this information.

6.3.2.1 Food and nutrition

The dietary intervention will be based on the national Danish food-based dietary guidelines (40) and the Nordic Nutrition Recommendations (41) and seeks to promote a balanced energy intake, increased healthy food intakes, and improving children's healthy food competences. A key focus will be on increasing intakes of wholegrains, fruits, vegetables and legumes, and fish, reducing the intake of sugar, and on drinking water.

The dietary intervention in school will include pre-prepared do-it-yourself cold lunch and possibly a midmorning snack, consisting of for example rye bread sandwiches, pasta salad, raw vegetables with dip etc. The children will be involved in the table setting and cleaning after the meals. We expect that extra time will be needed to be scheduled for eating by the

schools. Lunch will be offered to all children regardless of their participation in the measurement schedule and will be free of charge. We will collect information about allergies and other dietary considerations from the parents of participating children at enrolment. Parents will be informed that the meals cannot accommodate special dietary needs and that the menu with full list of ingredients (including allergens) will be posted on a specific website one week before, for them to check which parts their child can eat. The teachers/pedagogues will help handling allergies in the eating situation as in a normal school day. We will ask the teachers to note any allergies in children who wish to have the meals, but who are not participating in the measurements. To encourage water consumption, cold-water dispensers will be made available if this is feasible and needed at each school, and each child will receive a reusable water bottle.

In addition to this, we will develop fun and engaging hands-on food competence exercises that the teachers can use in their teaching to support the children's knowledge, self-efficacy and liking of healthy foods as well as practical skills in food preparation. Examples are sensorics taste exercises, vegetable puzzles and following a recipe to cook a simple dish together in the school kitchen.

In the after-school clubs (SFOs), we will work with their existing offers of breakfast and afternoon snacks and provide tips and inspiration to encourage the staff to follow the national food-based guidelines. Other activities focusing on healthy diet may be arranged in collaboration with the SFO, such as afternoon café for parents where children serve healthy snacks and/or excursions to local farmers, greengrocers, or fishmongers.

6.3.2.2 Physical activity

The amount of moderate and vigorous PA during school time will be increased through three approaches: a) 3x45 minutes of organized vigorous PA, b) education of Physical Education (PE) teachers and c) improved facilities and structure for PA during recess and in after school clubs.

a) 3x45 minutes per week of organized vigorous physical activity during school time

To ensure that all children reach the recommended 3x30 minutes of vigorous activity per week, we will implement three weekly 45-minute sessions in the school curriculum. These sessions will be carried out throughout the study period and will encompass modified and varied sporting activities (80) and games, which promote motivation, active involvement of all children, and intensity. These 45-min sessions are developed to promote joy through play, provide appropriate challenges of the children's motor competences and physical capacity and, importantly, to achieve at least 30 minutes of varied, vigorous activity in relation to cardiopulmonary and musculoskeletal loading for all children, irrespectively of sex, fitness

levels, skills, and prior experience with sporting activities (80). The activities will thus be adapted to allow participation for all children.

b) Physical activity during school recess

In primary schools, recess time amounts to almost 5 hours weekly and is a setting where young school children on average have a high level of PA (81). This makes recess an important part of the total daily and weekly amount of MVPA for young school children. It has been shown that improved facilities, both in terms of permanent play and sport facilities (e.g., climbing frames, football goals) and loose play equipment (balls, skipping ropes, skateboards), increases mean daily amount of PA among young school children as well as the proportion reaching daily recommended levels of MVPA (82,83). This part of the integrated PA intervention centres on distributing packages of loose play and sports equipment to school classes and making agreements with schools and teachers about how these are distributed and collected at the beginning and end of recess. Previous projects such as Move@School have shown this to be feasible and a popular approach for improving children's opportunities to be active during recess. This will be supplemented by the concept of drawing lines in the schoolyards for 4-square and similar games and, if relevant, providing small goals for basketball, floorball and football.

c) Physical activity in after-school clubs

Danish school children at age 7-8 years spend about 10 hours a week in after-school clubs. Mean activity level is moderately high, but great variation is seen between children (81). More physical activity in this setting will be facilitated by implementing policies limiting computer gaming as an activity and facilitating outdoor activities as much as possible. In collaboration with the staff at the after-school clubs, activities suiting the institutional facilities and time structures will be developed and the equipment needed will be identified.

6.3.2.3 Sleep and screen media habits

During the study period, teachers/pedagogues will be asked to facilitate small assignments on screen media and sleep practices with the children during school time. The children will be asked to discuss and reflect upon screen media and sleep practices in their family, based on material from e.g., Medierådet (The Danish Media Council). The results of the children's assignments will be presented at later workshops for the parents (see below in section 6.3.3.2). This is done to integrate the children's perspective in the parent workshops, especially in relation to screen media habits.

6.3.2.4 Education of school staff

School staff (teachers and pedagogues) involved with the 1st and 2nd grade classes at the intervention schools will be asked to participate in three courses in the beginning of the study period:

Course on food and nutrition (approx. 4 hours): Teachers and pedagogues involved in preparation of the school food and having lunch with the children will receive education and material in e.g., the hygienic and pedagogical aspects of food and eating, nutrition, sensorics as well as tips and exercises to be used in the classroom.

Course on PA (6 hours): The aim of the educational program for PE teachers and other teachers with responsibility of PA in the class is to develop children's understanding, motivation, and abilities for being active in different sports and exercise activities throughout life. In many ways, the aim is to develop children's physical literacy (PL). We aim to help schools and PE teachers fulfil the ambitious aim of PE by offering a short educational program developed and tested in the PE component of the Y-path program (84). PE teachers will be offered this short education in combination with peer supervision and sparring on how to support children's PL development and how to create a motivating and inclusive climate for all.

Course on children's screen media and sleep habits (3 hours): Teachers and/or pedagogues with overall responsibility for the class will participate in an educational program related to children's sleep and screen media habits. The educational program will inform teachers about evidence and recommendations on sleep and screen media habits, and examples of class exercises and activities for the workshops will be presented.

6.3.3 Interventions focused on the child through the family

The interventions focusing on the child through the family will include fixed intervention components determined à priori by the research team, as well as co-created intervention components developed in collaboration with families.

During the intervention period, up to four parent or family workshops will be held at the schools each year. The workshops will be used to convey important scientific knowledge within each of the project's focus areas (diet, physical activity, screen media habits and sleep) to the parents. The workshops will also include practical exercises, such as inspiration for ways that parents can engage in physically active play with their children and/or tips and tricks for involving children in daily cooking. Furthermore, the workshops will be used to ask parents to provide their views on perceived individual-level and structural barriers for adopting healthier

habits, and in collaboration with parents, we will co-create interventions which could support behavioural changes within each focus area.

The form of each workshop will be decided in collaboration with the school. The workshops will be facilitated by members of the research team, teachers, pedagogues and/or external experts (such as Medierådet, Børns Vilkår, Hello Kitchen etc.), depending on the content of the specific workshop. Participants in the workshops may be parents alone and/or parents and children together.

We will pay special attention to avoid lecturing, parent bashing, stigma, and unhealthy focus on body weight. Focus will be on actively engaging and involving parents in promoting healthier lives in their family and in the local community. To reach all parents irrespective of ethnic and educational background, we will provide simple messages supported by visuals, and we will translate material into numerous languages if recommended by the school.

Written study material, such as invitations for workshops and inspirational material, will be sent to parents by e-mail via the software REDCap and/or via AULA and in print via the teachers. Inspirational material may include, for example, inspiration for birthday parties and playdates including fun physical activities that promote movement as well as suggestions for simple, fun and healthy food which we will develop together with the school and/or other stakeholders.

The planned intervention components which involve the families within diet and sleep/screen media habits are described in more detail in the following sections.

6.3.3.1 Diet

At least one of the parent workshops held at the school during the intervention period will be used to provide motivation, tips, and strategies for healthier eating in the family. If feasible, there may also be cooking workshops for the families at the school, where families will be invited to cook with their children following simple and hands-on recipes to gain practical healthy food competences in the families.

6.3.3.2 Screen media habits and sleep

Based on the current knowledge of determinants of screen media habits in children, as well as our experiences in the family-based 'SCREENS trial' (57) and the ongoing school-based 'Screen-Free Time with Friends', the intervention will mainly focus on the interpersonal- and institutional (school) level, i.e., the family and peer social networks (85,86).

The overall aim of the screen media habits and sleep intervention component is to promote healthy screen media habits, encourage healthy sleep practices in the participating children, and to reduce the gap between the parents' and child's understanding of digital literacy.

The intervention will contain fixed generic components such as facilitating class culture and providing parents with the latest knowledge and recommendations within sleep hygiene and screen media habits.

Inspiration sheets

All parents, (also those who have not provided consent on behalf of their child to participate in the study), will receive inspiration sheets introducing knowledge and recommendations about screen media habits and/or sleep, potential tools to address this knowledge as well as suggestions for child and parent activities to support dialogues about sleep and screen media habits between the children and parents. Through these inspiration sheets we seek to encourage parents to reflect and consider the sleep and screen media habits in their family. We will provide parents with tools they can choose to use to support the introduction and maintenance of e.g., screen media rules. Importantly, any specific screen media rules will be determined by the parents/family and not by the researchers.

Parent workshops on sleep and screen media habits

Several parent workshops will be held over the two-year intervention period. The workshops will include information and dialogue focusing on children's sleep and screen media practices. The workshops will focus on the assignments on sleep and screen media habits that children completed during school hours (see 6.3.2.3).

The goal of the workshops is to increase parent knowledge on their own and their children's screen media practices, strengthen the dialogue with other parents, their own child, and between children within the provided subject (e.g., evening screen media use).

The topic for the workshops is planned to be an introduction to the knowledgebase for the importance of better sleep and balanced screen habits. The meetings will also focus on the exchange of experiences (pros and cons) with sleep and screen media habits at home and during child playdates among parents and reflections on potential strategies to support better sleep and screen media practices among the children during leisure time periods, in the family, and to increase the awareness of alternative leisure time activities (e.g., participation in after-school clubs, sports clubs).

All parents will be invited to participate in the workshops even though they have not consented on behalf of their child to participate in the study.

6.3.4 Intervention focused on the local environment

6.3.4.1 Local supermarkets, restaurants, and fast-food outlets

To promote healthier eating, children and families need to be supported within their local community by food environments that make the healthy choice the easy choice and not a challenge in their everyday life (87–92).

Within the national partnership established in connection with the study (see section 6.3.1.), new solutions will be developed to promote healthier eating among children, e.g., decreasing portion sizes, improving the availability of fruit, vegetables, cold fresh water, and healthy snacks, and decreasing the availability of sweets and sugar-sweetened beverages in supermarkets, restaurants, cafés, fast-food outlets and local sports clubs. Priority will be given to increasing the availability of fruit and vegetables, including convenience products, to make cooking with vegetables easier. The local community partnership groups (LPCGs) established in each intervention school district (see section 6.3.1) will have an important role in adapting and anchoring the interventions in the local setting.

Furthermore, stakeholders from each of the 12 local intervention communities will be invited into a co-creation process and to join the LPCGs. Invited stakeholders will be representatives from local sports organizations and other leisure time organizations for children, the school, the municipality, local supermarkets, restaurants and other private companies, parents, and other relevant local actors. The LPCGs will be responsible for developing and implementing activities within the local community that can promote healthier environments for children and families, including healthier food environment, easier access to active living including participation in sporting activities and outdoor activities, community-building activities etc.

6.3.4.2 Sports organizations

To support children that are not in sports clubs to join and become members, we will establish collaborations between schools, municipalities, and local sports clubs. Such collaborative initiatives have been shown to benefit families with low sporting experience, as these families are more positive towards sporting activities initiated via the school.

6.3.5 Compliance

Adherence to the intervention components will be explored at different levels using multiple method e.g., interviews, questionnaires, observations, logs, accelerometry, dietary registrations and biomarkers. As part of the evaluation of the implementation, we will assess whether the different intervention components are implemented by intervention providers,

e.g., teachers and other pedagogical staff, as intended and to what extent (dose delivered). Moreover, we will explore to what degree children and parents are exposed to the intervention component (dose received and reach). We will also explore participation in general.

6.4 Control schools

The control schools will be offered education related to physical activity for the eldest classes in the school, and the participating children in the present study will be offered participation in an activity day by the end of the study period.

6.5 Overview of study procedures and activities

The table below summarizes the activities described above and provides an overview of the planned study procedures as well as measurements, which will be conducted in 4 test rounds i.e., at the beginning and end of each of the two school years. The measurements will be further detailed in section 6.6.

Table 1. Overview of study procedures and activities

May 2023	Initial contact <ul style="list-style-type: none"> • Invitation letters and participant information are distributed to parents / custody holders
May to September 2023	Information meetings at the school <ul style="list-style-type: none"> • An understandable presentation of the context and purpose of the study • An understandable explanation of the study interventions and measurements to parents and children. • Description of rights and considerations before decision • Description of foreseeable risks, adverse events, complications, and inconveniences • Unpredictable risks • Voluntary participation and right to withdraw at any time • Review of pamphlet “Before you decide” (11.NVK marts 2019)

	<ul style="list-style-type: none"> • Information regarding handling of personal data/biological material, confidentiality, and general compliance with The European General Data Protection Regulation/Databeskyttelsesforordningen. • That all data collected during the participation in the study will be used in the event of a participant chooses to withdraw from the study • The right to time for reflection • Questions • Explanation of informed consent and withdrawal • Collection of informed consent (including the possibility for 24 hours of consideration time) • Evaluation of inclusions/exclusions criteria including the allergy questionnaire <p>Staff education</p> <ul style="list-style-type: none"> • Courses at schools – an education program for teachers and related staff involved in the intervention (see section 6.3.2.4) <p>Preparation of measurements</p> <ul style="list-style-type: none"> • Instructions on dietary recording and accelerometer measurement
<p>Round 1: August-November 2023 (baseline)</p> <p>+</p> <p>Round 2: April-June 2024</p> <p>+</p> <p>Round 4: April-June 2025 (endpoint)</p>	<p>Measurements at home:</p> <ul style="list-style-type: none"> • Three-day dietary record, app. 30 min/day • Accelerometry • Screen/app time • Questionnaires <p>Measurements in school:</p> <ul style="list-style-type: none"> • Fasting blood sample and time for breakfast • Height, weight, and waist circumference • Body composition (BOD POD and bioimpedance) • Blood pressure and heart rate • Mounting of accelerometer device

	<ul style="list-style-type: none"> • Cognitive functions and school performance (math and reading) • Physical and motor function tests (handgrip strength, balance, vertical jump, sprint, agility, gross and fine motor function) • Yoyo running test • Adverse events • Questionnaires
Round 3: August-November 2024 (Subset of round 1,2 and 4)	Measurements at home: <ul style="list-style-type: none"> • Three-day dietary record, app. 30 min/day • Questionnaires Measurements in school: <ul style="list-style-type: none"> • Height, weight, and waist circumference • Body composition (BOD POD and bioimpedance) • Blood pressure and heart rate • Physical and motor function tests (handgrip strength, balance, vertical jump, sprint, agility, gross and fine motor function) • Cognitive functions • Yoyo running test • Questionnaires and school tests (math and reading)
October 2023 to June 2025	Activities in school and in the community <ul style="list-style-type: none"> • School lunch 4-5 times per week • Possibly a midmorning snack 4-5 times per week • 3x40 minutes of organized vigorous PA per week in school. • PA in breaks and after school clubs. • Inspiration sheets for families on screen media habits, sleep and dietary habits etc. • Tips and inspiration for staff in after-school clubs on how to follow the national food-based guidelines • Cold-water dispensers and reusable water bottles, if relevant • Parent workshops at the school • Assignments / exercises for children during school time about screen media habits, sleep, nutrition and food competences

	<p>Other</p> <ul style="list-style-type: none"> • School staff education • Packages of loose play and sports equipment for school classes • Community events • Fieldwork, interviews, logbooks, and short questionnaires for e.g., parents, school staff and local stakeholders • Observations
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6.6 Measurements and study procedures

As mentioned, measurements will be conducted in 4 test rounds i.e., at the beginning and end of each of the two school years. Each measurement will be conducted in all 4 rounds, unless stated otherwise below.

6.6.1 Questionnaires to be filled in at home

The questionnaires will be administered via text message and e-mail with a link to the electronic survey to parents.

The following questionnaires will be administered:

- Family background (round 1 or at round 2, 3, 4 or at new enrolments if not answered at previous data collection rounds): Data on children's SES will be collected from the parent(s) or legal guardian(s) using the Danish Occupational Social Class Measurement (93) (appendix 10).
- Sun habits (vitamin D) (in appendix 10)
- Child's leisure time sports participation (in appendix 10).
- Pubertal stage (round 1 and 4) (appendix 11-12).
- Food allergies, plaster allergies, or chronic disease (Appendix 2a and 2b (control schools))
- Neurodevelopmental disorders and medical conditions (in appendix 2a and 2b) (at round 1 or at round 2, 3, 4 or at new enrolments if not answered at previous data collection rounds).

- Breakfast consumption, intake of wholegrains, sugary foods, beverages, fish, and dietary supplements (appendix 13).
- Wellbeing & Quality of Life: Children's well-being will be measured by KIDSCREEN 27 (parental edition) (appendix 14) (round 1, 2, 3 and 4) (94). As no valid measure exists for children under age 11, the parent version will be used to supplement the child version.
- Child and parent screen time, type and current screen time practices (round 1, 2 and 4) will be collected using the modified parent reported SCREENS questionnaire (95) (appendix 15).
- Child quality of sleep (round 1, 2 and 4) will be measured using the Children's Sleep Habits Questionnaire translated to Danish (96) (appendix 16).
- Perception of the project and the activities (see section 6.6.4).
- Adverse events and concomitant medicine (round 2, 3 and 4) (appendix 17).
- Drop-out questionnaire (at dropout) (appendix 3).

6.6.2 Other procedures at home

6.6.2.1 Dietary records

Children's daily intake of food and drinks will be recorded by the parents for three consecutive days (two weekdays and one weekend day). We will use the validated, web-based tool, myfood24, which uses data from the national food composition tables. The tool is user-friendly, suitable for the target group and recording links can be sent via email.

6.6.2.2 Screen time and categorized app time measurement

At baseline, we kindly ask the parents to install a research-based app, Ethica (<https://ethicadata.com>), on the participating child's *own* smartphone (if they have one) at round 1, 2 and 4 over a period of 14 days collecting information on screen time and categorized app usage and screen time patterns. After installation, the app runs automatically in the background. The app has been developed to meet legal requirements for data security.

6.6.2.3 Physical activity and sleep assessments (Axivity)

At round 1, 2, and 4 detailed, objective information on the whole spectrum of behaviours will be assessed by continuous 24-hour/7-8-day Axivity® accelerometer measurement using thigh positioned monitors. Using the thigh measurement position the modified age-adjusted method of Skotte et al. (97) provides the ability to classify postural allocations like sitting, standing and activities like walking and running with high sensitivity and specificity in children

(98). Time lying in bed and sleep time will be objectively measured using an accelerometer classification algorithm that is currently under development.

The Axivity® AX3 monitor is a small (23 x 32.5 x 7.6 mm), lightweight (11 g), waterproof, 3-axis accelerometer data logger (Axivity, 2016). The Axivity® AX3 will be attached to the skin using thigh belts developed by a Danish company (Elas) or tape (Opsite Flexifix®, Fixumull®) directly on the body using compress (gaze) or artificial skin (DuoDerm) between the monitor and the skin. We have good experience with both methods attaching the Axivity® AX3 monitor to the body, but using plaster is more time consuming so we need to find out which method is most feasible in the study. The monitors will be attached at the school. All observers/staff in contact with the children will undergo pedagogical and professional training in the tests and in carrying out the measurements with the children and in adherence with Danish law have the necessary criminal records bureau check. It is important to ensure that the children also give their assent to participate in the study, if their parents have provided a written consent allowing them to participate. During the study period we will be aware of the children's reactions and whether they feel comfortable or not with the test situations. If the children do not want to wear accelerometers, we will respect this. Children with plaster allergy, childhood eczema or sensitive skin will always be offered to use belts. The children or parents will receive an information leaflet to take home or on the parent email including relevant information regarding the physical activity measurement.

6.6.3 Measurements in school

Two program centres (UCPH and SDU), with specially trained test teams, will be responsible for the measurements in the schools.

6.6.3.1 Measures of anthropometry including height, weight, and waist circumference

The children will be weighed once in underwear/light clothes on a digital scale. Standing height will be measured three times to the nearest millimetre with a stadiometer. Waist circumference will be measured three times to the nearest millimetre by a non-elastic measuring tape at the level of the umbilicus. The mean of three measurements will be used for height and waist circumference.

6.6.3.2 Body composition and blood pressure

Body composition (i.e., FM and fat free mass) will be measured by air-displacement plethysmography using a BOD POD and a bioimpedance weight (InBody 270). The children will be wearing underwear/light clothes and a bathing cap. The children must sit in a transparent capsule for approximately 5 minutes (BOD POD), the bioimpedance measures takes approximately 1 min.

Blood pressure and resting heart rate will be measured three times by an automated device while the child is lying down after 10 minutes of rest. The appropriate cuff size for the child's arm circumference will be used.

6.6.3.3 Blood samples

At test round 1, 2 and 4 blood samples will be collected in the overnight fasted state (except water) in the children. The children can have breakfast shortly after. We will instruct the parents to provide the children with a breakfast pack on the measurement day but will also have food available for children that do not have any. The children will be provided with local anaesthetic patches (EMLA), which the parents will be instructed to apply on the child's arms in the morning (appendix 18). After removal of the patches, max. 30 ml blood will be drawn from the child's forearm by trained laboratory technicians. The blood samples will be brought back to University of Copenhagen or University of Southern Denmark for processing and storage at -70°C until analysis.

6.6.3.4 Health related fitness and motor functions

Aerobic fitness and intermittent exercise performance will be evaluated by the Yo-Yo Intermittent Recovery Level 1 Children's test (80), where the children will have to run 2x16m at increasing speed with 10 s break between each run. The test continues until the children cannot reach the finish line in time and takes around 20 min. for up to 50 pupils, depending on the size of the indoor gym. Heart rate will be measured continuously during the test, to determine endurance capacity and maximal heart rate. The test is simple and validated for fitness and maximal heart rate assessment for 6-10-year-old children (99). Heart rate and locomotor activities will also be measured during selected physical activities throughout the study period to describe the intensity.

Muscular fitness, strength and sprint performance will be determined by the hand grip strength test and a 20-m sprint test. The hand grip strength test is conducted with a hand grip Dynamometer. The child squeezes the dynamometer with all their strength, two times with each hand. An average score is then calculated using the measurements from both hands. Sprint time is measured by photocells. The children will complete each test twice and will be instructed to use all their power.

Gross motor functions are assessed as static and dynamic balance ability, upper extremity gross-motor function, jump height and agility. Jump height and balance ability are measured on a force plate measuring the vertical ground reaction force. Jump height is measured during the vertical countermovement jump test. Balance ability is measured during one-leg maintained balance and as maximum reaching ability during maintained balance, Agility is

measured as time to complete an agility running exercise preferably as a T-drill. Time to completion is measured by photocells. The children will complete each test twice and will be instructed to use all their power. Upper extremity gross-motor function is tested as speed and accuracy in the ability to perform accurate, goal-directed reaching movements towards a number of visually displayed targets within 30s or 1 minute. Fine motor function is measured by completion of a pegboard test where the children use both the dominant and nondominant hand to manually place small rods in the pegboard within 30s or 1 minute (100).

6.6.3.5 Test of cognitive functions and school performance

At test round 1, 2 and 4 the children will complete a short battery of standardized, validated and age-appropriate neurocognitive tests (100,101) including tests of executive function, memory, and attention. The tests resemble computer games, and they are administered individually via a tablet or iPad. At these time points, school performance will also be assessed by use of standard and age-appropriate tests of Danish spelling, reading and mathematics proficiency (100,102). We have good experience administering these methods in a school setting and for children at this age.

6.6.3.6 Wellbeing & Quality of Life: KIDSCREEN 27

Children's well-being will be measured using KIDSCREEN 27 (appendix 14). This validated questionnaire consists of 27 items that measure five dimensions of children's quality of life and well-being. These are physical well-being, psychological well-being, relationship with parents, relations to peers & social support, and life quality in the school environment (94). Due to the respondents' low age and inability to read a video and speech assisted child version of these questionnaires will be developed. The children will fill in this questionnaire using tablets and headphones enabling all items of the questionnaire to be read aloud, while the text appears on the screen. The answer scale points are illustrated with smileys.

To assess pupils' psychological well-being and functioning during school-based intervention activities, basic psychological needs satisfaction and intrinsic motivation is measured acutely right after occurrences of the school intervention activities. Satisfaction of children's three basic psychological needs for experiencing relatedness; - competence and autonomy as well as intrinsic motivation - will be measured with 3 items each adapted from Basic Psychological Needs Scale (BPNS-R) for PE (103) and the Intrinsic Motivation Inventory (104,105) adapted to the age-group and context (Appendix 18). A smiley based 4-point answer scale will be used.

To assess the motivational climate during intervention activities, systematic observations and scoring of indicators of empowering (task-orientation, relatedness support and autonomy

support) dis-empowering climate (ego-orientation and controlling behaviour) will be collected during intervention activities.

6.6.3.7 Motivation, confidence, and knowledge in relation to physical activity

A Danish version of the Canadian Assessment of Physical Literacy (CAPL) questionnaire, validated and context adapted in the Danish Assessment of Physical Literacy (DAPL) study (106), will be used to measure children's intrinsic motivation and confidence for physical activity as well as their knowledge and understanding of physical activity. A Danish video and speech assisted child version of these questionnaires has been developed and validated in the DAPL study (school grade 1 to 6). The children will fill in this questionnaire using tablets and headphones enabling all items of the questionnaire to be read aloud, while the text appears on the screen, will fill out this questionnaire (see: <https://www.surveyxact.dk/LinkCollector?key=PHC22ZQNUJ9K> or Appendix 19).

The children will also be asked about their participation in leisure time sports club activities (Appendix 20).

6.6.3.8 Food competences and meal culture

Children's food competence and meal culture will be assessed using a self-constructed questionnaire developed for and validated in the age group (Appendix 21).

6.6.3.9 Registration of snack and lunch consumption

On selected days, portion sizes and food/beverage intake at breakfast, midmorning snack, lunch and afternoon snack in schools/after school clubs will be recorded at class level in a subsample of the schools. Also, teachers/pedagogical staff present in the classes during the children's consumption of midmorning snack (if provided by the project) and lunch will be asked to roughly assess and register the intake of each of these meals at class level.

6.6.4 Evaluation of measures related to implementation of the intervention and organizational aspects

To explore acceptability, perceived outcomes, and consequences of the intervention, as well as structural and organizational aspects related to implementation of the intervention, school management at each school, and/or other relevant personnel with coordinating/managerial responsibilities related to this project, will be asked to complete a questionnaire and to participate in interviews. Moreover, all teachers and/or other relevant pedagogical staff, affiliated with the various intervention components, will be asked to complete a survey and

to participate in group interviews. Field work during school hours, e.g., classes, PA sessions, recess, and lunch with focus on how the intervention unfolds in practice, reasons for adaptation, and participant responsiveness will be conducted via observation methods etc.

Parents will be asked to participate in a focus group interview and to complete a questionnaire regarding perception of the project and the activities. Moreover, PE teachers are asked to roughly assess and register attendance and active participation during the 3x45 minutes sessions of PA at class level. Local coordinators will be asked to register the number of parents who attend school events, to explore implementation and perceived outcomes of the intervention. Moreover, fieldwork will be conducted when possible.

To explore relevant structural and organizational aspects as well as initiatives focusing on diet, physical activity, screen time and/or sleep in the control schools, the school management at each control school will be asked to complete a (few) questionnaire(s) and participate in an interview.

6.6.5 Measurements at further school, community, and national level

During the study, we will track the implementation and stakeholder engagement. Implementation data will include a register of activities in each local community, a database of stakeholder engagement and key engagement activities. Data will be collected on a regular basis and use methods recently described by Maitland et al. (107).

Furthermore, we will collect structural data on food environments, including accessibility and spatial layout of stores, using digital maps, GIS data (e.g., Smiley register) and field studies (90).

We plan to conduct both a) interviews with the manager of the after-school club and facilitators of activities in the leisure time clubs and the active summer holiday camp as well as b) systematic observations on selected days to explore acceptability, perceived outcomes and consequences of these intervention components. Moreover, we expect that actors involved in the implementation of intervention activities, e.g., schoolboard members, managers of after-school club, volunteers in leisure time clubs, will be asked to fill out questionnaires and/or take part in interviews exploring structural and organizational factors influencing implementation. Several core theories, models and frameworks within implementation science will serve as reference points for the mentioned investigations.

We will measure local and national partnerships using the Danish translation of the VicHealth partnerships analysis tool (108). Furthermore, this tool will be adjusted and adapted to the current project inspired by the work done by Indig et al. (109).

6.6.6 Use of the school's recordings of children's absence

As an indicator-measure of intervention effects on the children's general health and school wellbeing, we will seek to obtain the schools' recordings of children's attendance and absence, by parental consent.

6.7 Adverse Events

Adverse events (AEs) will be registered during the intervention. At each round of measurements, parents/custody holders will be asked by questionnaire if their child has had any AEs or reactions and any new medicine or changes in dosage taken since the last measurement as well as the start and end date (Appendix 17). In addition, parents will be asked to contact us if any events or reactions occur. The medically responsible physician will evaluate and record the intensity, causality, actions needed and outcomes of reported events.

6.8 Handling of biological samples

6.8.1 Sample analysis

Biological samples will mainly be analysed at University of Copenhagen, Denmark. Samples could also be analysed at other laboratories within and outside the EU, when this is judged necessary for the quality of the analyses. Therefore, it is not possible for researchers to predict in advance where analyses will be carried out and by whom. If data is transferred to countries outside EU to a country which cannot ensure an adequate level of protection, the transfer will be in accordance with chapter V of the GDPR. Parents will be informed that their data may be sent to data processors in a country like these if this is necessary for the quality of the overall result of the study. They are also informed that their data will be sent as pseudonymised/coded data and that the transfer will comply with the GDPR.

Parents will be informed separately about the handling of the data and all data transferring will be in accordance with GDPR.

The processing of data and biological samples in the project and at external data processors will be recorded at the mandatory lists to the Danish Data Protection Agency, and all data processing will follow GDPR/Databeskyttelsesforordningen and Databeskyttelsesloven. Data set and samples will always only be sent to data processors in coded s without any accompanying data that could potentially identify the participants.

Table 2 gives an overview of the planned analyses of the biological samples.

Table 2. Biological sample analyses – blood

Biomarker field	Analyses
Compliance & nutrition biomarkers	<ul style="list-style-type: none"> • Fatty acids, lipids and other nutrient biomarkers • Alkylresorcinols (wholegrains) and other food biomarkers • 25-hydroxyvitamin D, vitamin D binding proteins and - metabolites • Haemoglobin, ferritin, and transferrin receptor (iron)
Growth & development	<ul style="list-style-type: none"> • IGF-I, IGFBP-3 and other growth factors • Osteocalcin, bone specific alkaline phosphatase and other bone markers • Parathyroid hormone • Sex hormones
Cardiometabolic	<ul style="list-style-type: none"> • Triacylglycerol and total, LDL and HDL cholesterol and others • Glucose, insulin, C-peptide, glycosylated hemoglobin (HbA1c) etc.
Appetite	<ul style="list-style-type: none"> • Appetite hormones
Inflammation	<ul style="list-style-type: none"> • C-reactive protein, cytokines, adipokines, and related markers • Immune cells and immune markers
Cognition	<ul style="list-style-type: none"> • Blood markers related to brain function such as brain-derived neurotrophic factor (BDNF) and serotonin
Mechanisms	<ul style="list-style-type: none"> • Metabolomics, proteomics and lipidomics • Short chain fatty acids and other metabolites
Genetics	<ul style="list-style-type: none"> • Genotypes and epigenetics related to the study outcomes and research questions (single nucleotides polymorphisms and genome wide association studies, NOT full genome sequencing, and epigenetics)

6.8.2 Genotypes and epigenetic analyses

Gene polymorphisms and epigenetic analyses will be used to investigate biological mechanisms and effect modifications, as described in section 5.2.2. We will only investigate genes related to the exposure variables and study outcomes, e.g. involved in nutrient status, growth and body weight, metabolism, cognitive function etc. Genotyping will not include any full genome or exome sequencing, but will either be performed by DNA microarray (such as the Illumina Infinium HumanCoreExome Beadchip), that gives information on common gene variants (>5%), or by targeted analysis of common single nucleotide polymorphisms. It is therefore considered highly unlikely that we will identify any genetic variants that have any direct relevance for illnesses.

6.8.3 Research Biobank

In relation to the planned analyses in this study a research biobank will be created at Department of Nutrition, Exercise and Sports, University of Copenhagen, at Department of Sports Science and Clinical Biomechanics, University of Southern Denmark and at the Odense University Hospital for storage of blood samples. The research biobank will include maximum 3 x 30 mL = 90 mL of blood from each child.

The research biobank is registered at University of Copenhagen and University of Southern Denmark, which have shared data responsibility in the project and record all data responsible and data handlers, as well as their handling of biological material and other personal data. Biological material will be stored at maximum -70 C in the research biobank until analysis is completed and will be destroyed no later than 10 years after last participant last visit (June 2035), unless custody holders have signed a separate consent form that eventual remains can be transferred to a biobank (see 6.7.3).

All personal data/biological material is handled with confidentiality and stored in accordance with applicable law, GDPR and Danish Data Protection Act. no. 502 of 23rd May 2018.

6.8.4 Biobank for future research purposes

Eventual remains of material of blood (maximum 3x 10 mL=30 mL) from the research biobank, will, if the custody holders of participants have signed a separate consent form for this, be transferred from the Research Biobanks at University of Southern Denmark and University of Copenhagen to a biobank at University of Copenhagen. The biobank material will be stored at maximum -70 C at the Department of Nutrition, Exercise and Sports, University of Copenhagen, Rolighedsvej 26, 1958 Frederiksberg C. The biological samples in this biobank

will be stored for a maximum of 15 years after last participant last visit (June 2040) and will only be used in new research projects after these have been approved by the Danish Regional Committee on Biomedical Research Ethics, unless the samples have been totally anonymized. This means that the ID-log that connects study participants and the ID on the biological material has been destroyed and it is no more possible to identify the participant. If the study participants or custody holders of participants choose to donate remains of biological material from the research biobank to the biobank and later regret it, they can contact NEXS and have the samples destroyed if the samples have not been anonymized.

PI knows that establishment of a biobank for future related research projects of the excess material from the research biobank is not covered by the committee by law, but by the General Data Protection Regulation (Databeskyttelsesloven/ Databeskyttelsesforordningen). As the procedure is considered relevant for the entire research protocol in Denmark and to be able to prove that the biobank is established on a legal basis, the information is included in the material for the ethical committee.

6.9 Data Management and Quality Control

6.9.1 Data Collection and Processing

All collected data will be stored in accordance with the General Data Protection Regulation/Databeskyttelsesforordningen for data handling.

6.9.1.1 Data storage in REDCap

Data from questionnaires and most outcome data collected during school measurement days will be collected and stored directly in REDCap, which is a secure platform for building and managing online surveys and databases. REDCap is managed by the researcher service organization OPEN in the Region of Southern Denmark.

Research personnel will collect and enter data following formulated standard operating procedures. Data quality will be ensured using tailored validation fields in REDCap.

6.9.1.2 Data storage on other platforms

Outcome data (smartphone use, cognitive tests and dietary record) will also be stored on online servers provided by external companies (EthicaData, Cambridge Cognition, MyFood24). We will establish data handling agreements with the companies to ensure that data is securely and not used for other purposes.

Data from heart rate monitors (collected during the fitness test) will be extracted and stored pseudonymised in safe folders on university servers at SDU or UCPH.

Accelerometers will be collected at the schools (after 7-10 days) and transported back to the University of Southern Denmark or University of Copenhagen where data will be extracted from the measurement devices and stored in safe folders on the University servers.

Collected blood samples will each day be transported back to the University of Southern Denmark or University of Copenhagen where they will be processed and stored at maximum –70 degrees Celsius until analysis.

Process data such as field notes, observations, logbooks, and interviews, which will be collected in paper form or recorded, will be entered and stored in safe folders on university and/or region servers at SDU and/or Center for Clinical Research and Prevention.

Aggregated datasets from all data storage platforms other than REDCap will be stored in the REDCap file repository. The aggregated datasets will be uploaded to the REDCap file repository and include the ID log variable to allow for combination of the datasets.

6.9.2 Confidentiality

To maintain confidentiality, participants will only be identified by their initials and an assigned participant number. Documents that identify the participant (such as the signed informed consent document and the ID log) will be maintained in strict confidence by the investigator, except to the extent necessary to allow auditing and/or monitoring by the appropriate regulatory authority. All information obtained during the study will be handled according to local regulations and the GDPR/Databeskyttelsesforordningen. The study is registered on a central list at the Institutions/UCPH and SDU. The universities conduct lists of all data controllers/data processors and their handling of the personal data, and the lists are available in case of inspections from the Danish Data Protection Agency.

6.9.3 Information/data from patient hospital records

Data and information from the participants' Hospital medical records will not be included in this project.

6.10 Use of data from Danish registers

We will obtain information on eligible children and their families from Danish registries with the purposes to 1) obtain contact information (name, address, and CPR-number) on eligible

children and their parents (for distribution of all participant materials related to the study), 2) compare socio-demographic characteristics (parent education and employment) of participants with eligible non-participants to better judge generalizability and explore this analytically, 3) investigate the long-term (i.e., a 7-year follow-up) effectiveness of the program on body composition, and 4) lower the burden of the parental survey with respect to sociodemographic information, and 5) limit missing information on sociodemographic information due to survey non-response. Registries that will be used for these purposes include at least the following: The Civil Registration System (Det Centrale Personregister), The National Child Health Register (Børnedatabasen), and the DREAM database.

In the participant information (Appendix 5a and 5b), we inform all parents that we plan to collect information on the above-mentioned factors from Danish registers on all eligible children regardless of whether consent to participation in this project has been obtained or not. The Legal authority for said collection of data will be Section 10 in Databeskyttelsesloven and Article 6 (1) (e) of General Data Protection Regulation

Furthermore, we will obtain information from Danish national health registers and clinical databases, e.g., the National Patient Register, the National Prescription Register, the National Health Services Register, the Danish Psychiatric Central Register, the Medical Birth Register, and the National Child Health Register, to study associations between lifestyle factors and biomarkers collected in the study and subsequent long-term health and morbidity among participants. Such registry linkages will be performed for up to 10 years after last participant last visit (June 2035) to be able to passively follow the cohort during adolescence and adulthood, and thereby study early childhood determinants of later health and morbidity. All registry linkages will be performed on secure servers at Statistics Denmark or the National Board of Health Data, and all data sets will be pseudonymized before analysis.

The establishment of a database containing data from this project as well as the subsequent transfer of data for related future research projects are not covered by the Danish Committee Act. Rather, for said actions, PI must comply with the General Data Protection Regulation (Databeskyttelsesforordningen) and Databeskyttelsesloven. Accordingly, the parents must be informed separately about this in the mandatory GDPR information sheet and a valid consent must be obtained.

6.11 Statistical methods

6.11.1 Primary analyses

All analyses of the predefined primary and secondary outcomes will be planned a priori and made publicly available in statistical analyses plans at clinicaltrials.org. Reporting of the trial will be done in accordance with the CONSORT guidelines for cluster randomized trials. The primary analysis will be conducted on an intention-to-treat basis using mixed regression analysis with school district as random effect using degrees-of-freedom correction (Satterthwaite) to control for type 1 error due to the moderate number of clusters. Because the school district is the unit of randomization (not municipality, region, or school-class) and the intervention is being applied at this level, no further cluster-level will need to be considered statistically. Analyses will be adjusted for baseline factors that are found to be related to individual school districts (clusters) to control for possible baseline imbalance of cluster- and participant level data. In analyses using more than two assessment points, random slope effects of time will be considered to examine the possible heterogeneity of effects according follow-up time. In all intention-to-treat analyses, data from all participating children at inclusion will be included. All hypotheses testing will be based on two-sided tests at the $\alpha=0.05$ level.

6.11.2 Secondary analyses

Analysis and reporting of predefined secondary outcomes will be mentioned explicitly as secondary outcome analyses in the papers.

In addition, several sub-group analyses will be predefined and outlined in the final statistical analysis plan before the start of the trial. This will include subgroup analyses by e.g., parent socio-economic status, school district characteristics, and baseline adiposity status (being normal weight at baseline vs. being overweight or obese at baseline). Sub-group analysis among normal weight children at baseline will also be used to examine the effectiveness of the program on incidence of overweight or obesity. Other outcomes or sub-group analysis that are not predefined in the final study statistical analyses plan published before the start of the trial, will be defined explicitly as exploratory post hoc analyses in the papers and the exploratory nature of these analyses will be mentioned as a major limitation. Per-protocol and complete case analyses of the primary and secondary outcomes will also be carried out. We will analyze the association between implementation fidelity and actual dose delivered of each intervention activity in each intervention setting and its association with intervention outcomes among children.

In the final statistical analyses plan, a detailed plan to adjust for multiplicity based on a priority of the secondary outcomes and follow-up time points will be outlined.

6.11.3 Exploratory analyses

All analyses that were not pre-planned will be labelled as exploratory analyses.

6.11.4 Safety analysis

A safety analysis will be carried out to summarize any reported adverse events associated with participation in the study.

6.11.5 Definition of datasets

Quantitative datasets will be generated from the REDCap databases for each specific research aim guided by a predefined analysis plan. When registry data are needed for analyses, and these cannot be stored locally in REDCap, data stored in REDCap will be uploaded to 'Forskermaskinen' at Statistics Denmark.

6.11.6 Handling of missing values

Missing values will be handled via the mixed regression modelling approach to analyses (110,111). If alternative analyses are carried out missing values will be imputed using multiple imputation or inverse-probability weighting.

6.12 Suspension or early termination of the study

The principal investigator may suspend or prematurely terminate parts of or the entire clinical study for significant and documented reasons. If a suspicion relating to an unacceptable risk to study participants arises during the study, the principal investigator will suspend the study while the risk is assessed. The principal investigator will terminate the study if an unacceptable risk is confirmed. The principal investigator must ensure that the premature termination is justified in writing (fill out the termination form) and that the parents of the participants are informed.

6.13 Serious adverse events and reporting

6.13.1 Serious adverse events

A serious adverse event is defined as any untoward medical occurrence in a study participant that:

- Results in death
- Is life-threatening at the time of the event

- Requires inpatient hospitalization
- Results in persistent or significant disability or incapacity.

6.13.2 Reporting of Serious adverse events

The principal investigator will immediately inform the ethics committees if a serious adverse event occurs during the project. Reporting will take place no later than 7 days after the principal investigator became aware of any such adverse event. The standard forms from the ethics committees will be used for this purpose. In case of serious adverse events resulting from the project, the investigator will make any information requested by the committee available.

When the study ends, the principal investigator will provide a list of all serious expected and unexpected adverse reactions and all serious adverse events that have occurred during the study period. The report will be accompanied by an assessment of the participants' safety. The report will be delivered using the forms prepared by the committee system.

6.14 Study Documentation

6.14.1 Publication of Results

Results from the study, whether positive, negative, or inconclusive, will be summarized and published in peer-reviewed scientific journals and/or as part of scientific conferences. Authorship of publications should in agreement with the publication approval procedure reflect the collaborative nature of the project.

The project is registered in www.clinicaltrials.gov

7 Ethical conducts of the study

The study will be conducted in accordance with the ethical principles set forth in the current version of the Declaration of Helsinki. Children in the participation classes that are not enrolled in the study will be invited to participate in all activities that are like what could happen during the school day anyway, but without any examinations and data registration.

It always, and especially for families with small and younger children, interferes with everyday life to participate in a clinical study as it requires time to be set aside to comply with the procedures of the study, and some procedures may be associated with a risk.

7.1 Ethical approval and study registration

Ethical approval is obtained by (Project ID xxxxxx-22 S-xxxxxxx) and the data processing of the participants' personal data and biological material and the study is registered on statutory lists before enrolment of any study participants.

7.2 Justification of the study

Investigation of the effectiveness of a health promotion intervention program in children on health outcomes is not possible to conduct with high validity in a non-randomized experiment or in an observational study. Thus, for the present study there is a need for a control group and to investigate the aims of the study using an experimental randomized study design. The study will provide important new knowledge about the effectiveness of the Generation Healthy Kids program, a novel multicomponent intervention developed using a systems approach that can benefit children in Denmark and elsewhere in the future.

7.3 Underage study participants - justification and precautions

Justification of inclusion of children in the clinical study is in line with the Committee act (Komitéloven §19, stk. 3) which concerns studies not involving medicine, as the project can only be conducted in the described age group to answer the objective stated in section 5. Further, it is justified to recruit children of this young age as the overall project and main trial is focused on early prevention of overweight at school start before the overweight prevalence starts to rise markedly. Also, as described below, the study gives minimal risk of harm for the participants, and it is our assessment that the benefits outweigh the potential harms.

7.4 Children of ethnic minority

All children in the participating classes are eligible for the study. Written information materials will be translated depending on what is needed in the local school district. Verbal information will as standard be in Danish, but interpreters will be present at meetings if needed.

7.5 Risks and safety in relation to conducting the study

The interventions will not include focus on weight, but instead promote well-being, healthy habits, sense of community, and play. All communication will be prepared carefully in terms of non-stigmatization of children or families and there will not be any focus on weight related issues.

Participation in the study is associated with minimal risks. We have thoroughly considered the ethical aspects of the discomfort of the biological sampling. Since the blood samples are crucial for investigating the outcomes of the study (including diet and health related biomarkers), the discomfort of the blood sample is considered to be outweighed by its usefulness. The discomfort of blood sampling will be minimized by use of local anaesthetic patches before blood sampling and blood will be sampled by skilled laboratory technicians or other trained and qualified staff. The only expected risk of the biological sampling is a mild pain at the site of the blood sampling on the arm and potentially a small blue bruise at the site of the prick, which will disappear within a few days. Risks for infections that are associated with blood sampling are extremely low.

The children will be asked to meet for the visits after an overnight fast to standardize the blood results. They have time to eat immediately after the blood sample, at 10:30 am the latest, and, as in our previous trials with blood sampling in children they are allowed to drink 100-200 ml during the fasting period and up to 1-2 hours before the blood sampling. A fasting period of this duration is relatively short, and we do not expect the children to experience considerable discomfort due to the fasting. However, if this should be the case, the examinations will be discontinued immediately, and the child will be offered drinks and food.

The BOD POD body composition measurement is crucial for investigating the primary outcome of the main trial (fat mass). The measurement uses air-displacement plethysmography, is safe and approved for children, and takes few minutes. The measurement is constantly monitored by the operator and if the child expresses discomfort sitting inside the BOD POD, the measurement will be stopped immediately.

There is a risk of more focus on weight related issues e.g., overweight when measuring height, weight, and waist circumference. Therefore, the personnel will be trained to talk carefully about the measurement and to handle comments on body weight related issues from peers. Moreover, the children will not be told the measurement numbers (numbers on digital scale is hidden) and they can wear light clothing during the examinations.

There are no considerable risks related to the school meals which are prepared using foods that are part of a normal diet. The parents will be asked about food allergies at project start-

up and will be informed to check the menu plans with ingredients allergens online the week before. The children will be instructed in hygiene e.g., hand washing. The risk of injury or discomfort of performing the muscular and fitness tests are comparable to the risk children are exposed to during daily life and play. In case of any discomfort or accident the activity will be discontinued.

The risk or side effect of participating in the measurement of physical activity based on our extensive experience in carrying out physical activity measurement in paediatric populations is minimal. However, attaching the monitor to the skin on the thigh, might be unpleasant for some children and might in some rare cases result in skin irritation or an allergic reaction to the adhesive tape. The parents and the child will receive written and oral instruction about the instruments, and how the instrument can be removed gently. Furthermore, the written information will contain information on which symptoms to be aware of to reduce any physical inconveniences in case of an allergic reaction, and whom to contact in case of an allergic reaction. If the children participating in this project are allergic to adhesive tape, have eczema or other skin problems (see appendix 56), they will be offered to wear a monitor in a thigh belt. The belts include skin friendly silicone to keep the belt in place during the child's play. Children/parents can switch the belt between the right and left leg if necessary and to avoid/reduce potential skin irritation.

7.6 Benefits of the study and risk-benefit evaluation

The overall aim of Generation Healthy Kids is to promote healthy weight and wellbeing in children and is expected to benefit future generations of children. Furthermore, it is expected that the study will generate new and important knowledge on how to promote sustainable changes and potentially how to scale-up.

The participation in a study can cause some inconvenience to daily life and can be time-consuming. During the study, the families in the intervention school districts will receive school meals for the children, inspiration, and support for healthier lifestyle habits, and the control schools will be invited to an activity day by the end of the study period. After completion of the study all children receive a diploma (Appendix 22) and after data analysis, and publication of the main scientific papers, the parents can read about the study's main conclusions on the project's website.

The schools and local communities are also expected to benefit from the study in terms of elevated knowledge and focus on a healthy lifestyle.

All things considered, the study is judged to be ethically justified and proper as the individual, scientific and societal benefits are believed to outweigh the limited discomfort and very low risk of harm associated with the study.

7.7 Participant Information and Informed Consent

Written, informed consent will be obtained from all parents/guardians to the subjects prior to entry into the study and execution of any study-related procedures. PI or delegated and qualified persons staff will explain verbally to parents/guards the objectives, nature, significance, risks, and implications of the study before inclusion. This information will also be included in the written subject information sheet. The information will be given at meetings at each school. If the parents/guards are not able to join the meeting, they can arrange a separate meeting with project staff.

The parents/guardians will be offered to bring with them an assessor to the information meeting and after the meeting they are given time to discuss any questions and decide on participation in the study. All parents/guardians will receive a copy of the subject information sheet. The children will be informed of the objectives and procedures of tests in a language they can understand. The procedures are explained in more detail in section 6.2.5 and in the participant information.

7.8 Protocol Changes

Substantial amendments to this protocol may be implemented only after a favourable opinion of the Ethics Committees has been obtained. Amendments to the protocol are regarded as substantial if they have a significant impact on:

- The safety, physical health, and mental integrity of the study subjects
- The scientific value of the study
- The conduct or the management of the study

7.9 Protocol Deviations

No systematic deviations from the protocol are allowed. All significant protocol deviations noted during the study (by the Investigator) will be recorded in the e-CRF (REDCap).

7.9 Insurance

The participants are insured in accordance with the Law: “Lov om arbejdsskadeforsikring” (cf. LBK nr 1186 af 19/08/2022) by the current insurance at University of Southern Denmark and University of Copenhagen. During the study the participants will be covered by law: “Lov om klage- og erstatningsadgang inden for sundhedsvæsenet” (cf. LBK no. 995 14/06/2018 www.retsinformation.dk).

8 Funding

The initial concept of the Generation Healthy Kids was developed by the Investigators of the study. Generation Healthy Kids is funded by the Novo Nordisk Foundation with 80 mio. DKK in total, covering both the pilot study and the main trial. The study funds have been transferred to a research account at University of Copenhagen, which is subject to public audit. The funding will primarily cover salary for project staff (PhD students, postdocs, medical laboratory technicians, project managers and communication staff) and expenses related to operation e.g., school food, buy out of teachers, measurement equipment, analysis of blood samples, car rent, measuring instrument (one Bod Pod) and communication materials. Funding will not be used for paying subjects.

The Investigators declare to have no financial interests in the results of the study or any economic affiliation with any of the project partners.

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10 List of Appendices (all in Danish)

Appendix 1: Protokol resumé

Appendix 2a: Spørgeskema om fødevareallergi, allergi over for plaster, kronisk sygdom og andre hensyn

Appendix 2b: Spørgeskema om allergi over for plaster, kronisk sygdom og andre hensyn (kontrolskoler)

Appendix 3: Drop-out skema

Appendix 4a og 4b: Invitationsbrev til interventionsskoler og kontrolskoler

Appendix 5a og 5b: Deltagerinformation til interventionsskoler og kontrolskoler

Appendix 6: Samtykkeerklæring

Appendix 7: Forældrefuldmagt

Appendix 8: Pjece om informationsmøde

Appendix 9: Plakat om informationsmøde

Appendix 10: Spørgeskema om familiebaggrund, idræt og solvaner

Appendix 11: Spørgeskema om pubertetsudvikling til forældre med drenge

Appendix 12: Spørgeskema om pubertetsudvikling til forældre med piger

Appendix 13: Spørgeskema om barnets kost

Appendix 14: Spørgeskemaet KIDSCREEN 27

Appendix 15: SCREENS modificeret spørgeskema

Appendix 16: Børns søvnvaner spørgeskemaet

Appendix 17: Uønskede hændelser og samtidig medicin

Appendix 18: Spørgeskema om dækning af basale psykologiske behov og IM under interventionsaktiviteter

Appendix 19: Spørgeskema til børn om physical literacy

Appendix 20: Spørgeskema til børn om deltagelse i fritidsidræt

Appendix 21: Spørgeskema om madkompetence og måltidskultur

Appendix 22: Diplom