

**Characteristics of cohorts in the Childhood Cancer Research Group
at the Danish Cancer Society's Research Center**

Cohort name (acronym)	Adult Life after Childhood Cancer in Scandinavia - ALiCCS
Main focus areas	Somatic late effects in all organs and organ systems after treatment for childhood cancer
Source of identification of study population	Nordic cancer registries and central population registries
Study population	33,160 1-year survivors of childhood cancer from Denmark, Finland, Iceland, Norway and Sweden diagnosed at age 0-19 years between 1943-2008 and five as many randomly selected population comparisons individually matched
Design and data collection	
Study design	Cohort and case-cohort studies
Setting	Population-based
Data sources (yes/no)	
Registers	Yes
Medical records	Yes
Questionnaires	No
Interviews	No
Physical examinations	No
Bio samples (if yes, specify)	No
Other tests	No
Website and/or contact person and link to cohort design paper	www.aliccs.org Jeanette Falck Winther, jeanette@cancer.dk https://onlinelibrary.wiley.com/doi/full/10.1002/pbc.25661