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Table 1. Description of the main clinical and epidemiological studies included

FIRST AUTHOR, YEAR OF PUBLICATION	TYPE OF ARTICLE	NUMBER OF SUBJECTS	AGE AT DIAGNOSIS	OBJECTIVES	SOURCE OF DATA	KEY RESULTS
REULEN, 2011 ⁸	Multicentric retrospectve cohort study	17 981 cancer survivors	< 15 years	(1) to investigate the long term risks of subsequent primary neoplasms in survivors of childhood cancer; (2) to identify subsequent primary neoplasm types that contribute most to the long-term excess risk; (3) to identify subgroups of survivors at substantially increased risk of particular subsequent primary neoplasms who may require specific interventions.	The British Childhood Cancer Survivor Study; 5 years cancer survivors	Cumulative incidence of colorectal cancer for survivors treated with direct abdominopelvic irradiation 1.4% by age 50 years, comparable with the 1.2% risk in individuals with at least 2 first-degree relatives affected by colorectal cancer. The absolute excess risks (AERs) in survivors over 40 years are highest for digestive subsequent primary neoplasms (AER= 5.9, 95% CI: 2.5-9.3 per 10 000 person-years) and genitourinary subsequent primary neoplasms (AER=6.0, 95% CI: 2.3-9.6 per 10 000 person-years); these patients may require specific interventions (prevention, screening).
BAULD, 2005 ¹¹	Comparison study	153 cancer survivors; sample of6 377 healthy peers to match the adolescent survivor age range	13 - 24 years	To investigate smoking, alcohol use, illicit drug use and sexual risk in adolescent survivors of childhood cancer.	Adolescent cancer survivors off treatment for 12 months or more, recruited through the Royal Children's Hospital, Melbourne, Australia, compared with age matched healthy adolescents	Among participants, the most prevalent cancer diagnosis was leukaemia (53%) bone cancer (14%), Wilms tumour (10%) non-Hodgkins lymphoma (7%). Compared to their healthy peers, younger survivors (13 to 17-years) were at a decreased risk of reporting alcohol use (OR=0.44, 95% CI: 0.39-0.49), binge drinking

CARSWELL, 2008 ¹²	Comparative national, multi-centre, population-based study	1263 cancer survivors; 1422 controls from provincial health insurance agencies.	< 20 years	(1) To describe the prevalence of smoking and binge drinking among survivors of childhood and adolescent cancer, comparison with age and gender matched healthy controls. (2) To identify factors associated with these behaviors.	Late Effects study, part of the Canadian Childhood Cancer Surveillance and Control Program. Patient who survived at least 5 years after diagnosis.	Survivors were less likely to be current smokers (OR adj=0.65; 95% CI: 0.54–0.77) and binge drinkers (OR adj=0.66, 95% CI: 0.55–0.78) than controls. Survivors' smoking (23%) and binge drinking (25%) did not vary according to clinical factors.
TERCYAK, 2006 ¹³	Randomized controlled trial of health promotion intervention	75 cancer survivors	11 – 21 years	To test the efficacy of health education and health behavior counseling (cigarette use, insufficient physical activity, and nonadherence to sun protection recommendations) among adolescent survivors of childhood cancer. Focus on the prevalence and co-occurrence of three behavioral risk factors.	Telephone-based assessment of health behaviors and stress; behavioral record of the past 7 days.	28% of the patients reported one of three risk factors, 12% reported two of three risk factors, and 7% reported all three risk factors. Non-adherence to sun protection was the single most common risk factor; physical inactivity and non-adherent sun protection were the most common cooccurring risk factors. Greater age and stress were significantly associated with the presence of more than one behavioral risk factors.
KAHALLEY, 2012 ¹⁴	Multicentric cohort study	307 cancer survivors; 97 healthy siblings	< 21 years ; 14 - 20 years (healthy siblings)	To estimate the rate of smoking and identify factors associated with smoking in adolescent survivors	The Childhood Cancer Survivor Study Cohort*.Participants completed a self-report survey of health, quality of life, and health behaviors	Survivors: 28% ever smokers; 10% recent smokers. Sibling groups: 33% ever smokers; 9% recent smokers.Ever smoking was significantly associated with purging (RR = 2.49, 95% CI:1.60-3.88, p < 0.001), binging (RR = 1.57, 95%

						behavior (RR = 2.12, 95% CI:1.47-3.07 p < 0.001), peer smoking (RR = 3.21, 95% CI:2.35-4.38 p < 0.001), having smokers in the household (RR = 1.73, 95% CI:1.22-2.46 p < 0.01), and having no history of cranial radiation therapy (RR = 1.92, 95% CI:1.05-3.45, p < 0.05). Recent smoking was significantly associated with purging (RR = 4.44, 95% CI:2.08-9.50 p < 0.001), binging (RR = 2.55, 95% CI:1.32-4.90 p < 0.01), suicidal behavior (RR = 3.61, 95% CI:1.89-6.89 p < 0.001), emotional discomfort (RR = 2.75, 95% CI:1.38-5.49 p < 0.01), peer smoking (RR = 7.18, 95% CI:3.85-13.40 p < 0.001), and having smokers in the household (RR = 3.03, 95% CI:1.58-5.82 p < 0.01).
TYC, 2009 ¹⁵	Multicentric case-control study	94 (patients undergoing cancer treatment); cancer	8 - 11 years	To compare preadolescents treated for cancer to their healthy peers on a number of tobacco-related risk factors.	The 94 preadolescents undergoing treatment for cancer and a matched comparison sample of 190 participants without cancer completed questionnaires about their smoking habits, intentions to smoke and tobacco-related psychosocial risk factors.	There were no current smokers in the cancer cohort and only 2 current (0.5%) in the original school sample. Healthy preadolescents were more likely to report future intentions to smoke (34.1%) relative to preadolescents with cancer (14.0%, p<0.001). Healthy preadolescents were more likely to have at least one close friend who smoked (16.8%) as compared to those with cancer (7.5%, p=0.030). Compared to those without

						cancer had higher tobaccorelated knowledge (p=0.001), perceived themselves to be more vulnerable to tobacco-related illness (p<0.001), were more optimistic (p<0.001), and attributed more value to overall health (p=0.001). They also perceived themselves to be more vulnerable to general health problems (p<0.001), and had lower levels of rebelliousness/risk taking (p=0.001) as well as perceived instrumental value of smoking (p=0.002). Multivariable analyses suggest that the ages of 8-11 years may be a critical period for a child's development of attitudes about smoking in that every year of age conferred approximately a 50% increase in the odds of intending to smoke.
EMMONS, 2002 ¹⁶	Multicentric, retrospective cohort study	9 709 cancer survivors	≥ 18 years	To examine smoking behaviors and to evaluate predictors of cigarette smoking initiation and cessation	The Childhood Cancer Survivor Study*	Ever smoking: 28 %; current smokers: 17%. Factors independently associated with a statistically significant relative risk of smoking initiation included older age at cancer diagnosis, lower household income, less education, not having had pulmonary-related cancer treatment, and not having had brain radiation. Blacks were less likely to start smoking. The frequency of smoking

						initiation was significantly lower among survivors.
GREEN, 2012 ¹⁷	Multicentric, retrospective cohort study	9 284 cancer survivors	≥ 18 years	To evaluate the potential contribution of demographic, lifestyle, treatment, and intrapersonal factors and self-reported pharmaceutical use to obesity	T The Childhood Cancer Survivor Study*.	The risk of obesity (BMI** > 30 kg/m²) was increased among those 5 to 9 years of age at diagnosis (RR=1.12; 95% CI: 1.01-1.24; p=0.03), those who received 20 to 30 Gy of hypothalamic/pituitary radiation dose (RR=1.17; 95% CI: 1.05-1.30; p=0.01), and those with abnormal SF-36 physical function (RR=1.19; 95% CI: 1.06-1.33; p=0.001). The risk of obesity was decreased among those who met the US Centers for Disease Control and Prevention guidelines for vigorous physical activity (RR=0.90; 95%CI: 0.82-0. 97; p=0.01) and among those with a medium amount of cancerrelated anxiety (RR=0.86; 95% CI: 0.75-0.99; p=0.04). Of the pharmaceuticals evaluated, only paroxetine (antidepressant) was independently associated with an increased risk for obesity (RR=1.29; 95% CI: 1.08-1.54; p=0.01).

ROBIEN,	Clinical study	72 cancer	≤ 20 years	(1) To evaluate the typical	The Childhood Cancer	Half the participants met minimal
2008 ¹⁹	Jgar staay	survivors		dietary intake of adult	Survivor Study Cohort*	goals for fruit and vegetable
				survivors of childhood		intake: this is significantly greater
				acute lymphocytic		than the population median
				leukemia (ALL) and to		percentage of adults nationwide
				compare these data with		(23.2%) and in the state of
				major dietary		Minnesota (24.5%); and half the
				recommendations related		participants met minimal goals
				to cancer and		for dietary fat restrictions.
				cardiovascular disease		Participants reported dietary
				prevention.		sodium and added sugar intake in
				(2) To evaluate whether		excess of recommendations, and
				adherence with dietary		suboptimal consumption of
				guidelines was associated		dietary fiber. 18% of participants
				with BMI and waist		met recommendations for 30
				circumference among		minutes of physical activity, 5
				adult survivors of		days/week.
				childhood ALL.		days, week.
GURNEY,	Retrospective	921	≤ 20 years	To compare final height	The Childhood Cancer	40 % of participants were very
2003 ²⁰	cohort study	childhood	20 years	and BMI** between adult	Survivor Study Cohort*	short of stature: below the 10th
2003	conorcatady	brain cancer		survivors of childhood	Salvivoi Stady Collect	percentile for height.
		survivors		brain cancer and age- sex-		The strongest risk factors for
		301111013		matched population		adult short stature: young age at
				norms.		diagnosis and radiation
				To quantify the effects of		treatment involving the
				treatment and cancer-		hypothalamic-pituitary axis.
				related factors on the risk		Obesity risk factor: younger age
				of final height below the		and cranial radiation in females.
				10th percentile (adult		and cramar radiation in Terriales.
				short stature) or having		
				BMI of 30kg/m2 or more		
				(obesity).		
		1	1	(Obesity).		

BUCHANAN, 2009 ²³	Retrospective cohort study	9 298 childhood cancer survivors; 2 950 sibling controls	≤ 20 years	To determine the current sun protection behaviors of childhood cancer survivors, and to compare these behaviors to a sibling population. Secondary objective: to determine whether survivors who are at higher risk of skin cancer because of previous radiation therapy have improved sun protection behavior.	The Childhood Cancer Survivor Study Cohort*	66% of siblings and 67% of cancer survivors practiced at least some sunscreen use in the past summer. Childhood cancer survivors were less likely to have sunbathed in the past year (none vs. any: RR=0,92, 95%CI: 0,89-0,95) or use artificial tanning methods in the past year (none vs. any: RR=0,76, 95%CI: 0,70-0,83). Factors associated with survivor's sunscreen use in the previous year: exposure to therapeutic radiation, being female, having lighter skin complexions, having previously examined for skin cancer, and higher predisposition to sunburn. Variables associated with increased sunbathing in the previous year among survivors: no previous radiation, being female, younger age (≤45 years), the number of years post diagnosis (≤30 years), darker skin color, and lower predisposition to sunburn. Subject with a prior skin cancer were more likely to use sunscreen (RR=1,12, 95%IC: 1,06-1,18) and less likely to sunbathe (RR=0,87, 95%IC: 0,79-0,96). Three most common diagnoses
2006 ²⁴	sectional survey	adolescent cancer survivors	years	cancer diagnosis on adolescent physical activity behaviors across the cancer experience (ie. prediagnosis, during	western Canada (15-20 years old at the time of initial study recruitment) completed a mailed, self-	included some form of lymphoma or Hodgkin disease (29,9%), leukemia (27,8%) and tumors of central nervous system (14,4%).

	treatment, and post-treatment).	administered questionnaire (measures of physical activity at 3 time points).	Based on the 27 MET (metabolic equivalent) criterion (selected as criterion for 'active'): 84.5% were active prediagnosis, 26.4% were active during treatment, 73.6% were active post-treatment. The decline in physical activity persisted following the completion of treatment. No systematic differences or relationships between the
			1 2

UPADHYAYA, 2004 ²⁶	Open-label pilot study	16 ado- lescents without history of cancer	12 - 19 years	To examine the feasibility and preliminary tolerability of bupropion SR in adolescents with nicotine dependence.	Adolescents who were titrated over 1 week to bupropion SR 150 mg b.i.d and maintained at this dosage for 6 weeks. Participants also received two 30-minutes individual smoking cessation counseling sessions.	Nine participants received at least 4 weeks of medication: significant decrease in the average number of cigarettes smoked and carbon monoxide levels. 31.25 % of the adolescents were abstinent after 4 weeks of taking bupropion SR. Bupropion SR along with counseling may be safe and potentially efficacious for this population. Lack of weight gain among participants during the smoking cessation effort. Possibility that bupropion might have a harm reduction effect due to reduction of the number of cigarettes smoked.
KILLEN, 2004 ²⁷	Smoking cessation randomized clinical trial	211 adolescents smokers, whitout history of cancer	15 - 18 years	To examine the efficacy of a treatment for adolescent smokers that combines nicotine patches with bupropion.	Randomization to 1 of 2 groups: i) nicotine patch + bupropion SR ii) nicotine patch + placebo. Participants met weekly in 45 minutes session (group skills training).	Percentage abstinent assessed by time and treatment group: - Nicotine patch + placebo: 28 % week 10, 7 % week 26 - Nicotine patch + bupropion SR: 23% week 10, 8% week 26 Bupropion might failed to improve abstinence rates because of the dosage used in this study (150 mg per day whereas the recommendation dosage for adult smokers is 300 mg per day). The medications appeared to be safe and were well tolerated. The large majority of adolescents in both treatment groups reduced their

HOLLEN,	Prospective	64 cancer	13 - 21	To test the hypothesis that	Intervention group with	consumption to a few cigarettes per day or less. Effect of the intervention for
1999 ²⁸	clinical trial	survivors	years	teen survivors who receive education to enhance decision-making skills will report increased quality decision making, maintained or lowered riskbehavior status (in smoking, alcohol use, or illicit drug use) 1-, 6-, and 12-months post-intervention more than the comparison group.	21 participants who attend a workshop, and comparison group with 43 survivors who did not attend the workshop. The intervention included three components: five one-hour educational units administered in one day, three short videocassettes (one on decision making, two on alcohol use), and four weekly home assignments given during the first post-intervention month.	improving decision-making knowledge, and decision making was significant at 1-month post-intervention, and highly significant at 12-months post-intervention. The intervention did not affect smoking risk motivation at any of the three time points. The intervention had a significant effect on motivation for engaging in alcohol use at 1-month post-intervention, and a marginally significant effect at 6 months, but no effect at 12-months. Home assignments for remediation with cancersurviving adolescents most likely need to be expanded over a longer follow-up period to obtain a lasting effect for risk behaviors.

TYC, 2003 ³⁰	Randomized controlled trial	103 cancer survivors	10 - 18 years	To evaluate the efficacy of a tobacco risk counseling intervention on knowledge, perceived vulnerability and future intentions to smoke among preadolescents and adolescents cancer survivors, compared with a standard care control condition.	Participants from St Jude Children's Research Hospital, who were randomized in 2 groups: standard care control group (standard advice about the risks of tobacco use), and tobacco intervention group (with more intensive late effects risk counselling in addition to an educational video, goal setting, written physician feedback, smoking literature, follow-up telephone counseling).	Compared with the standard care control group, intervention group's participants had significantly higher knowledge scores, higher perceived vulnerability scores, and lower intention scores at 12 months following the intervention. No significant differences between the two groups at 6 months, across all measures, were found. Participants whose parents used tobacco had significantly higher intentions scores at 12 months compared with scores obtained at 6 months and baseline. Participants whose parents were non-tobacco users and who received the intervention reported significantly lower intention scores than the control group patients. Informing adolescent cancer survivors about their personal susceptibility to negative health outcomes can play a role in promoting tobacco abstinence in this yulnerable population.
SMITH, 1996 ³³	Non- randomized open-label clinical trial	22 ado- lescents smokers	13 - 17 years	To evaluate the safety, tolerance, and efficacy of 24-hour nicotine patch therapy in adolescent	The intervention was a daily nicotine patch therapy for 8 weeks (22mg/d for 6 weeks	82% experienced at least one adverse event during the 8-week patch phase; 68% reported some kind of skin reaction; 55%

				smokers with current smoking rate of 20 or more cigarettes/day, who were trying to stop smoking.	followed by 11mg/d for 2 weeks). Weekly individual behavioral counseling and group support continued for 8 weeks with follow up visits at 3 & 6 months; and a mailed survey at 1 year.	reported erythema only; 32% reported no skin reactions (common adverse events also reported with patch use adult smokers in the literature). 14% of the participants reported not smoking during week 8. Patch is safe to use in this population, and well tolerated as an adjunct to a smoking cessation program.
MOOLCHAN, 2005 ³⁴	Double- blind, double dummy, randomized 3-arm trial	120 adolescents without history of cancer	13 - 17 years	To determine the safety and efficacy of the nicotine patch and gum for adolescents who smoked ≥ 10 cigarettes per day and who want to quit smoking.	Participants received twelve weeks of nicotine patch or gum therapy with cognitive-behavioral therapy, with a follow-up visit at 6 months. They were randomized in 3 groups: nicotine patch 21 mg (34 participants), nicotine gum 2 and 4 mg (46 participants), placebo patch and gum (40 participants).	Mean compliance across groups was higher for the patch than for the gum. Both patch and gum were well tolerated, and adverse events were similar to those reported in adult trials. No significant effect of patch versus gum placebo on cessation outcomes. Significant difference between the active patch and placebo arms with prolonged abstinence rates: 18% for the active patch group, 2.5% for the placebo group. The nicotine patch was significantly more effective than placebo in helping dependent adolescent smoking receiving cognitive-behavioral therapy quit smoking (prolonged abstinence).

HUDSON, 2002 ³⁵	Behavioral health promotion study	266 cancer survivors	12 - 18 years	To assess the impact of an intervention which is a multi-component behavioral intervention on changing health knowledge, health perceptions (perceived susceptibility, perceived seriousness, perceived benefits, perceived barriers), and health behavior practices	From a childhood cancer survivor cohort attenting the St. Jude Children's Research Hospital after completion of therapy clinic for annual evaluation, in remission 2 or more years after completion of cancer therapy; randomly assigned to a controlarm (standard care, n=135) or treatmentarm (standard care plus the multi-behavioral intervention, n=131). 251 participants were evaluable at the both time points for the calculation of change scores.	Baseline evaluation: no significant differences in health practices, perceptions or knowledge. 52% reported to be unaware of the potential risk of 2 nd cancer which was the most concerning potential treatment sequelae in 48% of participants. No significant differences in the change scores between the standard care group and the intervention group at 1 year follow-up. Perceptions of seriousness for risks factors secondary to the cancer treatment of adolescent cancer patients were increased as a result of the intervention. On selected patient subgroups: significant differences between males and females in change scores for health knowledge; females in the cohort demonstrating greater improvement in knowledge at 12-months among females
						improvement in knowledge at

^{*} The Childhood Cancer Survivor Study (CCSS) is a retrospective cohort study of 14359 survivors of childhood cancer diagnosed prior to 21 years of age between January 1, 1970 and December 31, 1986, with a longitudinal follow-up of 5-year survivors of childhood cancer treated in 26 institutions in the United-states and Canada. Eligible cancer diagnoses included leukemia, central nervous system malignancy, Hodgkin lymphoma, non-Hodgkin lymphoma, Wilms tumor, neuroblastoma, soft tissue sarcoma, and bone tumors. **Body Mass Index (BMI) is a measure of body fat based on height and weight that applies to men and women. AERs: Absolute Excess Risks; ALL: Acute Lymphoblastic Leukemia; CI: Confidence Interval; HR: Hazard Ratio; RR: Relative Risk; SIR: Standardized Incidence Ratio