SAS coding examples for case-cohort designs

*see "Case-cohort run code 042621.R" to do identical analyses in R https://github.com/obrienkm2/case-cohort/blob/main/Case-cohort%20run%20code%20042621.R

"Simple" scenario (Table, row 1) All cases selected, selection probability of sub-cohort = x% Example: O'Brien et al. (2017) Serum Vitamin D and Risk of Breast Cancer within Five Years. *Environ Health Persp*

```
cc1 = a data set containing the case-cohort data, including the following variables subcohort = 1 if in subcohort; 0 if not case = 1 if a case; 0 if not age_enrollment = age at enrollment age_eof = age at end of follow-up (e.g. event time or censoring time) exp = exposure of interest covar1, covar2, covar3 = covariates of interest (coded as categories) ID = identification variable
```

sampling rate= number of particants in sub-cohort / number of participants in full eligible cohort

```
*test code;
%LET epsilon=0.01; *or any number smaller than your smallest time
unit;
%LET sampling rate=0.05; *for the example data set cc1;
*restructure data set so that cases in sub-cohort weighted differently
according to time (will appear as two entries);
DATA ccnew1;
     SET wcc.cc1;
     *cases within subcohort - contribute fully until just before
     diagnosis;
     IF subcohort=1 AND case=1 THEN DO;
           start = age enrollment;
           stop= age eof - ε
           event = 0; *considered a censored observation;
     wt= 1/&sampling rate; *inverse probability of sampling weight;
     OUTPUT;
     END;
     *all cases contribute person-time right before event, count as
     event;
     IF case=1 THEN DO;
           start = age eof - ε
           stop = age eof;
          event = 1;
          wt=1;
     OUTPUT:
     END;
     *non-cases within subcohort - contribute full person time,
     censored;
```

```
ELSE IF subcohort=1 AND case=0 THEN DO;
           start = age enrollment;
           stop = age_eof;
           event = 0;
           wt= 1/&sampling rate;
           *inverse probability of sampling weight;
     OUTPUT;
     END;
RUN;
PROC PHREG DATA=ccnew1 covs(aggregate);
     CLASS covar1 covar2 covar3;
     MODEL (start, stop) *event(0) = exp covar1 covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN;
```

	Full-cohort,	Case-cohort,
	n=2,983 cases	n=2,983 cases
HR (95% CI)	1.14 (1.04, 1.19)	1.15 (1.00, 1.33)

Covariate-stratified case-cohort (Table, row 2) All cases selected, Sub-cohort selection probabilities of x_A % (Group A) and x_B % (Group B)

Example: Niehoff et al. (*in review*) Metals and breast cancer risk: a prospective study using toenail biomarkers

```
cc2 = a data set containing the case-cohort data, including the following variables subcohort = 1 if in subcohort; 0 if not case = 1 if a case; 0 if not age_enrollment = age at enrollment age_eof = age at end of follow-up (e.g. event time or censoring time) exp = exposure of interest covar2, covar3 = covariates of interest (coded as categories) ID = identification variable groupA=1 if in group A; 0 if in group B
```

sampling_rateA= number in sub-cohort from group A / number in full cohort from group A sampling_rateB= number in sub-cohort from group B / number in full cohort from group B

```
%LET sampling rateA=0.08; *for the example data set cc2;
%LET sampling rateB=0.15; *for the example data set cc2;
*restructure data set so that cases in sub-cohort weighted differently
according to time (will appear as two entries);
DATA ccnew2;
     SET wcc.cc2;
     *cases within subcohort - contribute fully until just before
     diagnosis;
     IF subcohort=1 AND case=1 THEN DO;
           start = age enrollment;
          stop= age eof - ε
          event = 0; *considered a censored observation;
     IF groupA=1 THEN wt= 1/&sampling rateA;
     ELSE IF groupA=0 THEN wt= 1/&sampling rateB;
     *inverse probability of sampling weights;
     OUTPUT:
     END;
     *all cases contribute person-time right before event, count as
     event;
     IF case=1 THEN DO;
          start = age eof - ε
          stop = age eof;
          event = 1;
          wt=1;
     OUTPUT;
     END:
     *non-cases within subcohort - contribute full person time, c
     ensored;
     ELSE IF subcohort=1 AND case=0 THEN DO;
```

```
start = age enrollment;
           stop = age eof;
           event = 0;
     IF groupA=1 THEN wt= 1/&sampling rateA;
     ELSE IF groupA=0 THEN wt= 1/&sampling rateB;
     *inverse probability of sampling weights;
     OUTPUT;
     END;
RUN;
PROC PHREG DATA=ccnew2 covs(aggregate);
     CLASS covar2 covar3;
     MODEL (start, stop) *event(0) = exp groupA covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN:
*group-stratified;
PROC PHREG DATA=ccnew2 covs(aggregate);
     WHERE groupA=1;
     CLASS covar2 covar3;
     MODEL (start, stop) *event(0) = exp covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN;
PROC PHREG DATA=ccnew2 covs(aggregate);
     WHERE groupA=0;
     CLASS covar2 covar3;
     MODEL (start, stop) *event(0) = exp covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN;
```

	Full-cohort,	Case-cohort
	n=2,983 cases	n=2,983 cases
	(2,573 Group A,	(2,573 Group A,
	214 Group B)	214 Group B)
Overall, HR (95% CI)	1.14 (1.04, 1.25)	
Groups A and B only	1.12 (1.02, 1.23)	1.12 (0.99, 1.27)
Group A, HR (95% CI)	1.12 (1.01, 1.23)	1.12 (0.98, 1.28)
Group B, HR (95% CI)	1.11 (0.81, 1.50)	1.10 (0.76, 1.59)

Outcome-stratified case-cohort (Table, row 3) 100% of type I cases and y% of type 2 cases selected; sub-cohort selection probability x% for all

Example: Sampling 100% of estrogen receptor-negative breast cancers and 50% of estrogen receptor-positive breast cancers, with the desire to look at subtype-specific and overall exposure-disease associations

```
cc3 = a data set containing the case-cohort data, including the following variables subcohort = 1 if in subcohort; 0 if not case = 1 if a case; 0 if not age_enrollment = age at enrollment age_eof = age at end of follow-up (e.g. event time or censoring time) exp = exposure of interest covar2, covar3 = covariates of interest (coded as categories)

ID = identification variable

Subtype1=1 if case of disease subtype 1; 0 otherwise

Subtype2=1 if case of disease subtype 2; 0 otherwise
```

sampling_rate= number of particants in sub-cohort / number of participants in full eligible cohort sampling_rate_subtype1= number of case of subtype 1 selected / total number of subtype 1 cases sampling_rate_subtype2= number of case of subtype 2 selected / total number of subtype 2 cases

```
%LET epsilon=0.01; *or any number less than your smallest time unit;
%LET sampling rate=0.05; *for the example data set cc3;
%LET sampling rate subtype1=0.50; *50% of subtype1 selected;
%LET sampling rate subtype2=1; *100% of subtype2 selected;
*restructure data set so that cases in sub-cohort weighted differently
according to time (will appear as two entries);
DATA ccnew3;
     SET wcc.cc3;
     *selected cases within subcohort - contribute fully until just
     before diagnosis;
     IF subcohort=1 AND (subtype1=1 | subtype2=1) THEN DO;
           start = age enrollment;
           stop= age eof - ε
          event = 0; *considered a censored observation;
          wt= 1/&sampling rate;
           *inverse probability of sampling weight;
     OUTPUT;
     END:
     *cases contribute person-time right before event only if
     selected, contribute based on weights;
     IF (subtype1=1 | subtype2=1) THEN DO;
           start = age eof - ε
           stop = age eof;
          event = 1;
           IF subtype1=1 THEN wt=1/&sampling rate subtype1;
                ELSE IF subtype2=1 THEN wt=1/&sampling rate subtype2;
```

```
OUTPUT;
     END:
     *non-cases within subcohort - contribute full person time,
     censored;
     ELSE IF subcohort=1 AND subtype1=0 AND subtype2=0 THEN DO;
           start = age enrollment;
           stop = age eof;
          event = 0;
           wt= 1/&sampling rate;
     *inverse probability of sampling weight;
     OUTPUT;
     END:
RUN:
PROC PHREG DATA=ccnew3 covs(aggregate);
     CLASS covar1 covar2 covar3;
     MODEL (start, stop) *event(0) = exp covar1 covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN;
*subtype 1 only;
DATA ccnew3 1;
     SET wcc.cc3;
     *selected cases within subcohort - contribute fully until just
     before diagnosis;
     IF subcohort=1 AND subtype1=1 THEN DO;
           start = age enrollment;
           stop= age eof - ε
           event = 0; *considered a censored observation;
           wt= 1/&sampling rate; *inverse probability of sampling
           weight;
     OUTPUT;
     END;
     *cases contribute person-time right before event only if
     selected, contribute based on weights;
     IF subtype1=1 THEN DO;
           start = age eof - ε
           stop = age eof;
           event = 1;
           wt=1/&sampling rate subtype1;
     OUTPUT;
     END;
     *non-cases within subcohort - contribute full person time,
     censored;
     ELSE IF subcohort=1 AND subtype1=0 THEN DO;
           start = age enrollment;
           stop = age eof;
```

```
event = 0;
           wt= 1/&sampling rate;
           *inverse probability of sampling weight;
     OUTPUT;
     END;
RUN;
PROC PHREG DATA=ccnew3 1 covs(aggregate);
     CLASS covar1 covar2 covar3;
     MODEL (start, stop) *event(0) = exp covar1 covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN:
*subtype 2 only;
DATA ccnew3 2;
     SET wcc.cc3;
     *selected cases within subcohort - contribute fully until just
     before diagnosis;
     IF subcohort=1 AND subtype2=1 THEN DO;
           start = age enrollment;
           stop= age eof - ε
           event = 0; *considered a censored observation;
           wt= 1/&sampling rate;
           *inverse probability of sampling weight;
     OUTPUT;
     END:
     *cases contribute person-time right before event only if
     selected, contribute based on weights;
     IF subtype2=1 THEN DO;
           start = age eof - ε
           stop = age eof;
           event = 1;
           wt=1/&sampling rate subtype2;
     OUTPUT;
     END:
     *non-cases within subcohort - contribute full person time,
     censored;
     ELSE IF subcohort=1 AND subtype2=0 THEN DO;
           start = age enrollment;
           stop = age eof;
          event = 0;
           wt= 1/&sampling rate;
     *inverse probability of sampling weight;
     OUTPUT;
     END;
RUN;
PROC PHREG DATA=ccnew3 2 covs(aggregate);
```

```
CLASS covar1 covar2 covar3;
MODEL (start, stop) *event(0) = exp covar1 covar2 covar3;
WEIGHT wt;
ID ID;
HAZARDRATIO exp;
```

RUN;

	Full-cohort	Case-cohort
	n=2,983 cases	n=1,349 cases
	(2,197 Subtype 1,	(956 Subtype 1,
	393 Subtype 2)	393 Subtype 2)
Overall, HR (95% CI)	1.14 (1.04, 1.25)	
Subtype 1 or 2 only	1.11 (1.01, 1.23)	1.13 (0.96, 1.34)
Subtype 1, HR (95% CI)	1.07 (0.96, 1.20)	1.09 (0.90, 1.31)
Subtype 2, HR (95% CI)	1.33 (1.05, 1.68)	1.35 (1.04, 1.75)

Covariate and outcome-stratified case-cohort (Table, row 4) 100% of type I cases and y% of type 2 cases selected; Sub-cohort selection probabilities of x_A % (Group A) and x_B % (Group B) NOTE: This assumes that case status and subgroup status are selected independently; if not, weights can be re-calculated for each combination (= a product of the specified weights) Example: Oversampling for Black women and estrogen receptor-negative breast cancers

```
cc4 = a data set containing the case-cohort data, including the following variables subcohort = 1 if in subcohort; 0 if not case = 1 if a case; 0 if not age_enrollment = age at enrollment age_eof = age at end of follow-up (e.g. event time or censoring time) exp = exposure of interest covar1, covar2, covar3 = covariates of interest (coded as categories) ID = identification variable groupA=1 if in group A; 0 if in group B

Subtype1=1 if case of disease subtype 1; 0 otherwise

Subtype2=1 if case of disease subtype 2; 0 otherwise
```

sampling_rateA= number in sub-cohort from group A / number in full cohort from group A sampling_rateB= number in sub-cohort from group B / number in full cohort from group B sampling_rate_subtype1= number of case of subtype 1 selected / total number of subtype 1 cases sampling_rate_subtype2= number of case of subtype 2 selected / total number of subtype 2 cases

```
%LET epsilon=0.01; *or any number less than your smallest time unit;
%LET sampling rateA=0.08; *for the example data set cc4;
%LET sampling rateB=0.15; *for the example data set cc4;
%LET sampling rate subtype1=0.50; *50% of subtype1 selected;
%LET sampling rate subtype2=1; *100% of subtype2 selected;
*restructure data set so that cases in sub-cohort weighted differently
according to time (will appear as two entries);
DATA ccnew4;
     SET wcc.cc4;
     *selected cases within subcohort - contribute fully until just
     before diagnosis;
     IF subcohort=1 AND (subtype1=1 | subtype2=1) THEN DO;
           start = age enrollment;
           stop= age eof - ε
          event = 0; *considered a censored observation;
           IF groupA=1 THEN wt= 1/&sampling rateA;
                ELSE IF groupA=0 THEN wt=1/&sampling rateB;
           *inverse probability of sampling weight;
     OUTPUT:
     END;
     *cases contribute person-time right before event only if
     selected, contribute based on weights;
     IF (subtype1=1 | subtype2=1) THEN DO;
           start = age eof - ε
```

```
stop = age eof;
           event = 1;
           IF subtype1=1 THEN wt=1/&sampling rate subtype1;
                ELSE IF subtype2=1 THEN wt=1/&sampling rate subtype2;
     OUTPUT;
     END;
     *non-cases within subcohort - contribute full person time,
     censored;
     ELSE IF subcohort=1 AND subtype1=0 AND subtype2=0 THEN DO;
           start = age enrollment;
           stop = age eof;
           event = 0;
           IF groupA=1 THEN wt= 1/&sampling rateA;
                ELSE IF groupA=0 THEN wt=1/&sampling rateB;
           *inverse probability of sampling weight;
     OUTPUT;
     END;
RUN;
PROC PHREG DATA=ccnew4 covs(aggregate);
     CLASS covar2 covar3;
     MODEL (start, stop) *event(0) = exp groupA covar2 covar3;
     WEIGHT wt;
     ID ID;
     HAZARDRATIO exp;
RUN;
```

	Full-cohort	Case-cohort
All women		
Overall, HR (95% CI)	1.14 (1.04, 1.25)	
Group A or B; Subtype 1 or 2 only	1.09 (0.99, 1.21)	1.05 (0.89, 1.23)
Subtype 1, HR (95% CI)	1.07 (0.96, 1.20)	1.01 (0.84, 1.22)
Subtype 2, HR (95% CI)	1.33 (1.05, 1.68)	1.23 (0.94, 1.60)
Group A		
Overall, HR (95% CI)	1.12 (1.01, 1.23)	
Subtype 1 or 2 only	1.09 (0.98, 1.21)	1.06 (0.89, 1.25)
Subtype 1, HR (95% CI)	1.07 (0.96, 1.20)	1.04 (0.86, 1.25)
Subtype 2, HR (95% CI)	1.18 (0.90, 1.54)	1.15 (0.87, 1.53)
Group B		
Overall, HR (95% CI)	1.11 (0.81, 1.50)	
Subtype 1 or 2 only	1.10 (0.77, 1.58)	0.91 (0.51, 1.62)
Subtype 1, HR (95% CI)	0.92 (0.60, 1.40)	0.68 (0.32, 1.44)
Subtype 2, HR (95% CI)	2.02 (0.99, 4.14)	1.98 (0.88, 4.47)

*subtype-specific estimates computed as for example 3;

Case-independent designs (Table, row 5) v% cases and z% of non-cases included in case-cohort sample; want to measure the association between previously measured exposure ("exp") and a second exposure ("exp2"), independent of case status Example: Lawrence et al. (2020) Association of neighborhood deprivation with epigenetic aging using four clock methodologies. *JAMA Open*

Sampling_rate_cases= number of selected cases / total number of cases sampling_rate_subcohort= number selected into subcohort / total number in cohort

```
%LET sampling rate cases=1; *for the example data set cc5 (all cases);
%LET sampling rate noncases=0.049; *5% of non-cases selected into
subcohort;
DATA wcc.cc5;
     SET wcc.cc5;
     IF case=1 THEN wt= 1/&sampling rate cases;
           ELSE IF case=0 THEN wt= 1/&sampling rate noncases;
     *create indicator versions of covariates;
     IF covar1=1 THEN covar1 1=1;
           ELSE covar1_1=0;
     IF covar1=2 THEN covar1 2=1;
           ELSE covar1 2=0;
     IF covar1=3 THEN covar1 3=1;
           ELSE covar1 3=0;
     IF covar2=1 THEN covar2 1=1;
           ELSE covar2 1=0;
     IF covar2=2 THEN covar2 2=1;
           ELSE covar2 2=0;
     IF covar2=3 THEN covar2 3=1;
           ELSE covar2 3=0;
     IF covar3=2 THEN covar3 2=1;
           ELSE covar3 2=0;
     IF covar3=3 THEN covar3 3=1;
           ELSE covar3 3=0;
RUN;
PROC REG DATA=wcc.cc5;
     MODEL exp2 = exp age enrollment covar1 1 covar1 2 covar1 3
     covar2 1 covar2 2 covar2 3 covar3 2 covar3 3/ CLB;
     WEIGHT wt;
RUN;
QUIT;
```

	Full-cohort	Case-cohort
β (95% CI)	1.34 (1.21, 1.47)	1.62 (1.23, 2.02)