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**An extension of relational methods in
mortality estimation**

Harald Hannerz

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An extension of relational methods in mortality estimation

Harald Hannerz¹

Abstract

Actuaries and demographers have a long tradition of using collateral data to improve mortality estimates. Three main approaches have been used to accomplish the improvement — mortality laws, model life tables, and relational methods. The present paper introduces a regression model that incorporates all of the beneficial principles from each of these approaches. The model is demonstrated on mortality data pertaining to various groups of Swedish people holding a life insurance policy.

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1. Introduction

A *mortality law* is a mathematical expression that describes mortality as a function of age. De Moivre made the first known contribution along this path in 1725. Many different expressions have been proposed since then (e.g. Gompertz 1825; Makeham 1867; Thiele 1872; Wittstein 1883; Pearson 1895; Perks 1932; Brillinger 1961; Heligman and Pollard 1980; Petrioli 1981; Mode and Busby 1982; Siler 1983; Anson 1988; Kostaki 1992; Hannerz 1999), and several reviews have been written on the subject (e.g. Benjamin and Haycocks 1970; Keyfitz 1982; Hartmann 1987; Kostaki 1988; Benjamin and Soliman 1993).

A *model life table* is simply a numerical table that gives death rates, probabilities for survival, expectations of life and other related information as a function of age. The model life tables are normally based on empirical observations in large populations believed to have reliable population and mortality data. The purpose of the model life tables is to substitute for unknown aspects of mortality in populations with limited or unreliable mortality data. The application of the models is based on the assumption that the age-pattern of mortality in the population under consideration resembles one of the life tables in the models (Adlakha 1972).

A *relational method* is a mathematical expression, which relates mortality in one population to that in others. In the description of relational methods as well as in the sequel of this paper, the following notations will be used: $F(x)$ is the cumulative distribution function (cdf), which gives the probability that a person will be dead x years after birth, $f(x) = DF(x)$ is the corresponding probability density function (pdf), and $m(x) = f(x)/(1-F(x))$ is the hazard rate. The odds for death within x years after birth, $F(x)/(1-F(x))$, will be called the cumulative distribution odds (CDO).

The classic relation, *proportional hazards*, which in mathematical terms may be expressed as

$$\frac{f_i(x)}{1 - F_i(x)} = \mathbf{a} \frac{f_j(x)}{1 - F_j(x)} \quad (1)$$

where the indices i and j refer to two different mortality experiences and \mathbf{a} is a constant, was first suggested as a method to relate mortality in one generation to that in others (Derrick 1927). The relation was used for the same purpose by demographers in the 1930's (Kermack, McKendrick and McKinlay 1934). It has also been used extensively by epidemiologists to relate the mortality experience of a studied population with that of the general population for the purpose of simplifying life expectancy calculations (Babigian and Odoroff 1969), and comparisons of mortality (e.g. Allebeck and Wistedt

1986). The relation was elaborated by Cox (1972) and has thereafter been known to many as ‘cox proportional hazards’. In the wake of the paper by Cox the interest for the relation among demographers was rekindled and it has subsequently been used in the study of human mortality (Vaupel, Manton and Stallard 1979; Vaupel and Yashin 1985) as well as in the construction of life tables for marriage dissolutions as a function of a variety of socio-demographic variables (Menken et al. 1981).

Equation (1) was criticised by Brass, who had observed that the ratios of the death rates in any two schedules were not constant with age but followed a more complicated course, in particular moving closer to unity for older people (Brass 1971). He also remarked that the discrepancy between (1) and reality could be quite staggering. With a set of life tables published by the United Nations (UN) (1955) he showed that the ratio of the death rate in a high mortality population to that in a low at ages 5-14 years may be twenty times as great as the corresponding value at ages 70-79 years (Brass 1969). The point of Brass’ criticism against the work of Derrick was not that the idea, that a simple relationship between different mortality patterns could be established, was wrong, but that the wrong transformation was used. Brass proposed that a more realistic relation would be obtained by

$$\frac{f_i(x)}{F_i(x)(1-F_i(x))} = \mathbf{b} \frac{f_j(x)}{F_j(x)(1-F_j(x))} \quad (2)$$

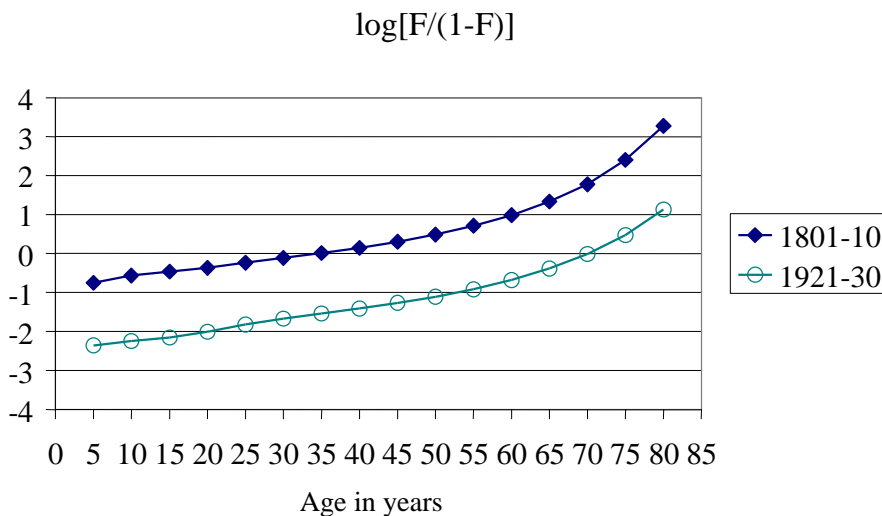
where i and j are defined as previously and \mathbf{b} is a constant. Integration of (2) would then yield the equation

$$\log \left[\frac{F_i(x)}{1-F_i(x)} \right] = \mathbf{a} + \mathbf{b} \log \left[\frac{F_j(x)}{1-F_j(x)} \right] \quad (3)$$

which states that the logarithmised CDO in one mortality schedule could be approximated by a linear function of the logarithmised CDO in another— a relation which has been applied extensively in studies of the demography in under-developed nations, particularly in Africa (Brass 1969). Brass also suggested that equation (3) would provide a useful tool for the projection of future mortality from past trends because the two parameters would be sufficiently few for efficient estimation from observations but flexible enough to describe a realistic range of possibilities (Brass 1971). In the verification of the system, Brass noted that \mathbf{b} seemed to be independent of the level of mortality. He also noted that \mathbf{b} often was close to unity and that, in

mortality changes in countries with a long series of life tables, **a** moved steadily with falling death rates while **b** fluctuated around 1 but had a strong tendency to return to this central value (Brass 1974). An illustration of this finding is given in Figure 1, where the logarithmised CDO with regard to the empiric mortality experiences of the male population of Sweden in the calendar periods 1801-10 and 1921-30 are given as a function of age. Although the median length of life had increased from 35 to 70 years between the two observation periods the curves of the logarithmised CDO of the two life tables are still approximately parallel. No smoothing has been attempted.

Figure 1: Logarithms of empirical cumulative distribution odds, with respect to the mortality experience in the male population of Sweden in the calendar periods 1801-10 and 1921-30. Data source:(Brass, 1971).



If we fix **b** in (3) at its central value, one, then we obtain the simpler relation, *proportional CDO's*, which mathematically may be expressed either as

$$\frac{F_i(x)}{1-F_i(x)} = \mathbf{a}' \frac{F_j(x)}{1-F_j(x)} \quad (4)$$

where \mathbf{a}' is equal to e^a in (3), or equivalently as

$$\frac{f_i(x)}{F_i(x)(1-F_i(x))} = \frac{f_j(x)}{F_j(x)(1-F_j(x))} \quad (5)$$

One can easily show that equation (4) also implies that

$$\mathbf{m}_i(x) = \frac{\mathbf{a}' \mathbf{m}_j(x)}{1-(1-\mathbf{a}')F_j(x)} \quad (6)$$

from which it is obvious that $\mathbf{m}_i(x)$ approaches $\mathbf{m}_j(x)$ monotonically as x approaches infinity, a feature that, according to the writings of Brass, might be more in agreement with experience than proportional hazards. Relation (4) has been used to project future life tables from past trends (Brass 1974). It has also been used to relate mortality among survivors of myocardial infarction to that of the general population for the purpose of increased precision in life expectancy estimates (Hannerz 1996a; 1996b).

It should be mentioned that Brass did not aim at perfect goodness-of-fit, but at a simple method to obtain decent approximations. A factor analysis of the data underlying the UN-tables of 1955 had indicated that at least three factors were needed to explain about 94% of the variation between the studied mortality schedules (Lederman and Breas 1959). These factors have been interpreted as (i) a factor governing the level of mortality; (ii) a factor governing the relationship between mortality in youth and adult life; and (iii) a factor governing mortality patterns at extreme ages (especially old age, 70+). It has also been postulated that two more factors are needed to explain the rest of the observed variation; (iv) a factor governing infant mortality; and (v) a factor governing the differences between male and female mortality schedules (Bourgois-Pichat 1962). The work of Brass together with that of Bourgois-Pichat later inspired the enlargement of equation (3) into two different four-parameter relations (Zaba 1979; Ewbank, Gomez de Leon and Stoto 1983).

The aim of the present work is (i) to investigate the appropriateness of the two single-parameter relations, proportional hazards and proportional CDO's, with respect to period mortality in a variety of subsets of the Swedish population, and (ii) to formulate and test a regression model intended to incorporate the beneficial principles

from each of the three ways previously used to improve mortality estimates — mortality laws, model life tables and relational methods.

2. Proportional hazards versus proportional CDO's: A comparison based on mortality in a variety of subsets of the male population of Sweden in 1983

Sometimes statisticians are confronted with the problem of estimating mortality in small populations or in populations where mortality data do not exist for all age strata. In these situations the mortality data of the sample alone may not be enough to obtain reasonable estimates of more than one parameter, and collateral data must be used to substitute for what we do not know. As a measure of relative mortality, the odds ratios (OR) formed by dividing CDO's in different groups will have an exact interpretation only if we regard closed populations in longitudinal studies. Most mortality studies concerns, however, only short periods, and most populations are open for non-death-related departures and/or entrance of new group members. From a mathematical viewpoint CDO's can still be estimated from observed hazard rates and the OR's could be used as a mortality index. The interpretation may however be questionable. As a method of graduating mortality for the purpose of life tables or estimation of life expectancies an assumption of proportionality in CDO's may still be useful and according to the work of Brass would often be more compatible with data than an assumption of proportional hazard rates. From this viewpoint, the present comparison could be regarded as a confirmatory analysis.

The material of the study was derived from the following three sources: a record-linkage between the Swedish national inpatient registry and the national cause of death registry 1978-83, mortality data from the Swedish Insurance Federation 1983, and mortality data from the publication "Statistical Abstracts of Sweden, 1983" (Statistics Sweden 1984). The observed outcome was deaths during the calendar year 1983.

The following subsets were regarded:

- "Men with a history of functional psychosis". This subset consists of men who had been psychiatric inpatients some time during the interval January 1978 - November 1982 with a diagnosis of functional psychosis (International Classification of Diseases, version eight (ICD-8)=295-299) as principal diagnosis.
- "Men with a history of drug abuse". This subset consists of men who had been psychiatric inpatients some time during the interval January

1978 - November 1982 with a diagnosis related to drug abuse (ICD-8=291, 303-304) as principal diagnosis.

- "Men with a history of acute myocardial infarction (AMI)". This subset consists of men who had been inpatients some time during the interval January 1978 - November 1982 with a diagnosis of AMI (ICD-8=410) as principal diagnosis.
- "Life insured men". This data set comes from the Swedish Insurance Federation.
- "Single men". The data on this subset was extracted from the publication "Statistical abstracts of Sweden 1983".
- "Married men". The data on this subset was extracted from the publication "Statistical abstracts of Sweden 1983".
- "Divorced men". The data on this subset was extracted from the publication "Statistical abstracts of Sweden 1983".

Deaths and person years at risk in the various subsets are given in Tables 1 and 2.

Table 1: *Risk masses and deaths in various subsets of the male Swedish population 1983*

Age	History of functional psychosis		History of drug abuse		History of AMI		Life insured	
	Person years	Deaths	Person years	Deaths	Person years	Deaths	Person years	Deaths
35-39	2130	13	6356	75	213	5	91237	78
40-44	1609	11	5574	112	460	9	89104	113
45-49	1338	28	5116	132	976	20	72900	136
50-54	1340	29	4963	167	1933	73	67289	256
55-59	1486	42	4387	168	3418	154	64661	396
60-64	1334	49	3286	169	4840	270	56179	581
65-69	1215	55	2239	137	5164	395	22818	405
70-74	1004	97	1241	103	5084	543	11338	401
75-79	656	82	500	56	3745	497	6889	320
80-84	253	44	98	12	2162	390	3668	288
85-89	67	13	15	3	998	168	1791	262
Sum:	12432	463	33775	1134	28993	2524	487874	3236

Table 2: Risk masses and deaths in various subsets of the male Swedish population 1983.

Age	All Swedish men		Single men		Married men		Divorced men	
	Person years	Deaths	Person years	Deaths	Person years	Deaths	Person years	Deaths
35-39	346213	518	93979	224	216870	191	34641	100
40-44	272269	660	45721	205	188437	286	36978	163
45-49	223777	822	28964	198	160896	387	32140	228
50-54	217892	1400	27106	301	159406	757	28184	313
55-59	228072	2287	28464	463	169454	1334	24567	411
60-64	239078	3969	29102	677	179568	2532	20483	525
65-69	204125	5530	23791	888	152074	3604	13901	557
70-74	177154	7832	21452	1176	125499	5000	9902	567
75-79	120785	8767	14895	1261	78222	5226	5168	502
80-84	66508	7837	8223	1031	36387	4079	2123	268
85-89	27158	5141	3108	629	11334	1947	648	130
Sum:	2123031	44763	324803	7053	1478146	25343	208734	3764

A series of graphs were made to study the appropriateness of the two relations. In creating the graphs, the hazard rate was assumed to be constant within each 5-year age category and was assessed by dividing the number of deaths by the risk mass in person years. In particular, for $i, j \in \{37.5, 42.5, \dots, 87.5\}$, the hazard rates were calculated by the formula $m_i = D_i / N_i$, where D_i is the number of deaths occurring in the age interval $(i-2.5, i+2.5)$ years and N_i is the number of person years at risk in that interval. The conditional distribution function F , for the i - and j -values in the above set, were thereafter approximated by the formula

$$F_j = F(j | X > 35) \cong 1 - \exp \left[- \left(\sum_{i < j} \frac{5D_i}{N_i} + \frac{5D_j}{2N_j} \right) \right].$$

Purely descriptive graphs based on the empirical hazard and distribution functions described above were created. Thereafter, the observed hazard ratios between various populations and the total population of Sweden were compared with the expected hazard ratios under the two hypotheses, proportional hazards and proportional CDO's. The following scheme was used to calculate the expected values. Let $m_{s,i}$ be the hazard rate and $F_{s,i}$ be the distribution function for the age i in the standard population (all Swedish men). Then, given proportional hazards, $m/m_{s,i} = g$ where g is a constant and,

given proportional CDO's, $m/m_i = b/[1-(1-b)F_{s,i}]$, where b is another constant. The parameter g was estimated by solving the equation

$$\sum D_i = g \sum \frac{D_{s,i} N_i}{N_{s,i}},$$

where D_i and N_i pertain to the study population while $D_{s,i}$ and $N_{s,i}$ pertain to the standard population, and the parameter b was estimated by solving the equation

$$\sum D_i = \sum \frac{b D_{s,i} N_i}{N_{s,i} [1 - (1 - b) F_{s,i}]}.$$

The results of the study are given in the Figures 2 through 9.

Figure 2: *Logarithms of empirical hazard rates in various subsets of the male Swedish population 1983. If proportional hazards were true then these below would be estimates of parallel lines. If proportional CDO's were true then these below would be estimates of lines that approach each other monotonically as age increases.*

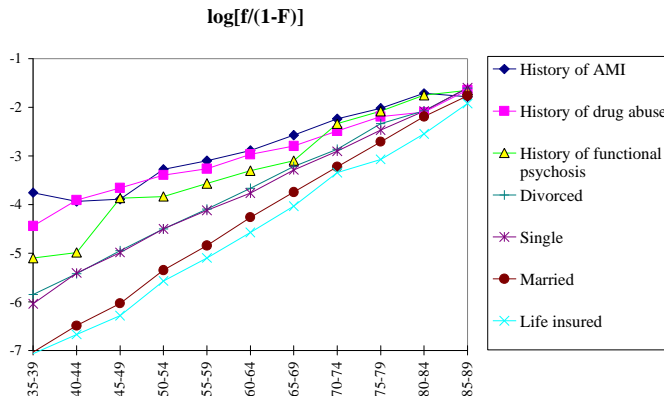


Figure 3: *Logarithms of empirical CDO's in various subsets of the male Swedish population 1983. If proportional CDO's were true then these below would be estimates of parallel lines. If proportional hazards were true then the distances between the lines below would increase monotonically with age.*

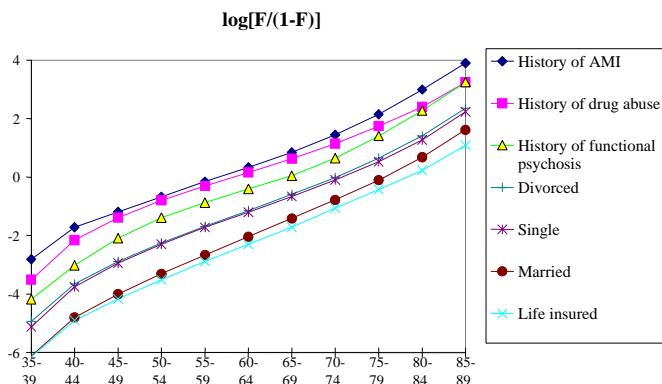


Figure 4: *Hazard ratios for various subsets of the male Swedish population in relation to the total male Swedish population 1983. If proportional hazards were true then the lines below would be estimates of parallel lines with constant values. If proportional CDO's were true then these below would be estimates of lines that approach unity, monotonically, as age increases.*

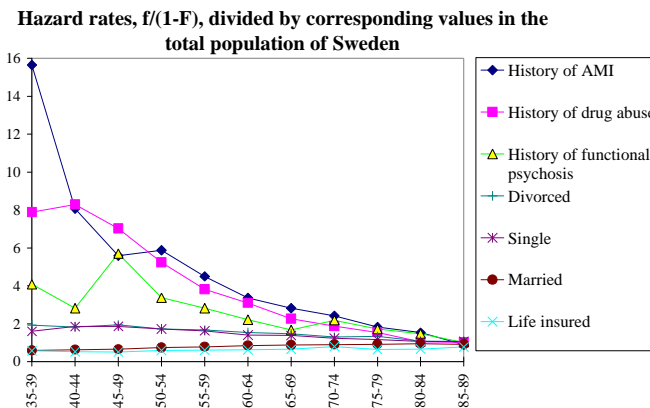


Figure 5: Hazard by distribution function, $f/(F(1-F))$, divided by corresponding values in the total male population of Sweden 1983. If proportional CDO's were true then $f/(F(1-F))$ would be independent of population, i.e. the equation for all of the lines below would be $y=1$.

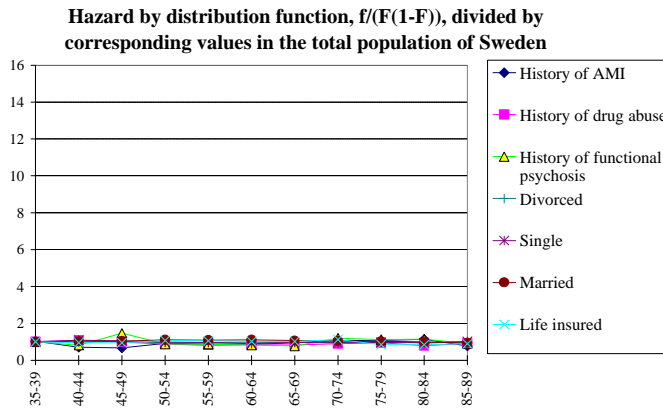


Figure 6: Observed and expected hazard ratios among men with a history of AMI (acute myocardial infarction), in relation to the total male population of Sweden 1983.

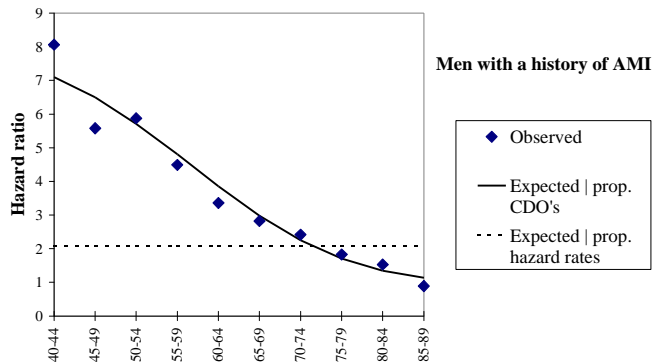


Figure 7: *Observed and expected hazard ratios among divorced men, in relation to the total male population of Sweden 1983.*

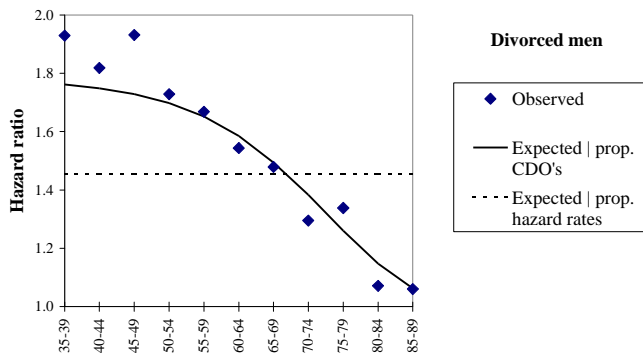


Figure 8: *Observed and expected hazard ratios among men with a history of functional psychosis, in relation to the total male population of Sweden 1983.*

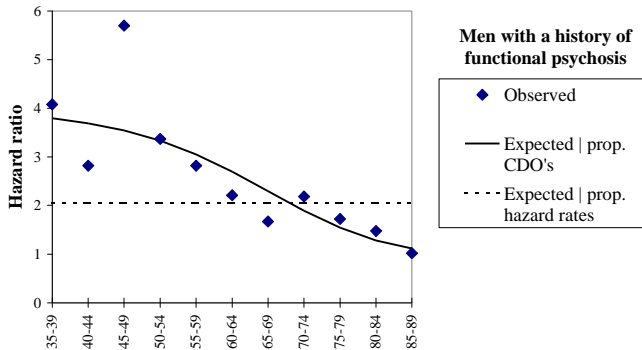
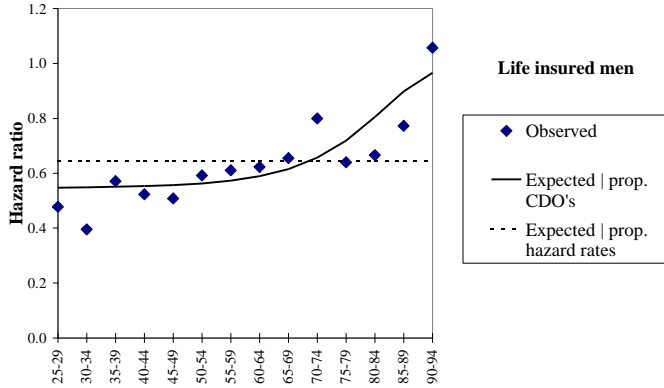


Figure 9: *Observed and expected hazard ratios among life insured men, in relation to the total male population of Sweden 1983.*



3. A reducible regression model: Formulation and testing

The Swedish Insurance Federation is an institution held in common by a number of Swedish insurance companies. One of the tasks of this institution is to provide estimates for mortality risks and life expectancies of life insured clients of the concerned companies. The tables below emanating from the insurance federation give risk masses, in person years, and deaths of a population of life insured individuals in 1982, divided into three groups on the basis of how long they had been insured.

Table 3: *Risk masses and deaths in cohorts of life insured Swedish males of 1982.*

Age	Insured since 11-∞ years		Insured since 6-10 years		Insured since 1-5 years	
	Person years	Deaths	Person years	Deaths	Person years	Deaths
15-19	80.33	0	565.92	0	1373.57	1
20-24	999.84	1	2189.36	1	4433.04	1
25-29	6070.61	9	5661.81	4	11714.54	2
30-34	16738.20	10	12166.34	7	23610.69	14
35-39	34922.37	40	20666.68	29	34152.53	22
40-44	35135.33	52	18588.22	29	26018.37	17
45-49	36493.85	102	14802.46	30	16895.60	27
50-54	42250.64	144	13015.72	32	11357.22	20
55-59	48379.96	291	10627.34	63	6661.93	36
60-64	44877.44	461	7365.84	54	3946.62	25
65-69	16273.61	321	3119.58	44	2827.88	37
70-74	8053.78	290	1644.68	55	1294.20	41
75-79	5572.67	286	710.33	30	315.47	16
80-84	3272.32	301	209.00	23	74.08	7
85-89	1717.01	255	37.00	1	9.50	1
90-94	217.48	42	6.50	0	1.00	0
Sum:	301055.44	2605	111376.78	402	144686.24	267

Table 4: *Risk masses and deaths in cohorts of life insured Swedish females of 1982.*

Age	Insured since 11-∞ years		Insured since 6-10 years		Insured since 1-5 years	
	Person years	Deaths	Person years	Deaths	Person years	Deaths
15-19	36.33	0	498.33	0	1151.66	0
20-24	868.68	0	1819.50	0	4531.08	1
25-29	4791.41	5	5370.41	2	8425.29	3
30-34	14056.37	9	9834.19	6	11336.49	1
35-39	26013.73	26	10971.96	7	13099.55	5
40-44	21636.16	19	7714.52	8	9099.53	6
45-49	18393.58	34	5760.26	4	5972.33	6
50-54	18577.74	68	5156.83	2	4159.51	5
55-59	16027.79	83	4221.09	10	2884.73	14
60-64	11716.04	64	3114.06	11	2112.13	7
65-69	4847.20	59	1524.40	13	1410.56	13
70-74	2953.41	52	954.83	14	756.17	7
75-79	1925.59	54	443.00	14	184.50	10
80-84	1152.66	94	147.58	5	72.75	6
85-89	635.99	55	39.00	4	13.50	1
90-94	71.57	13	3.00	1	1.00	0
Sum:	143704.25	635	57572.96	101	65210.78	85

In the present section, a regression model intended to incorporate the beneficial principles from each of the three ways previously used to improve mortality estimates — mortality laws, model life tables and relational methods — will be formulated, and tested on the above insurance data.

3.1 The ingredients of the regression model

The mortality for people in different ages within the cohorts of life insured people constitute potential collateral data to all other ages in the same cohort. To make use of this information we need a mortality law. The mortality in each of these cohorts constitutes collateral data for every other cohort. To make use of this information we need a relational method. Finally, we need to incorporate data from a standard life table to compensate for the scanty number of deaths in the extreme ages of the direct data sets.

3.1.1 The standard life table

It should be remembered that the incorporation of collateral data from a standard life table implies an assumption that at least some aspect of the age-pattern of mortality in the population under consideration resembles that of the standard. It is also evident that the higher the degree of similarity between the study population and the standard, the higher the advantage of its use. It has, for example, been concluded that some advantage would be gained by the use of standard life tables for each gender rather than one in common (Brass 1971), and that an African standard would be better than a European in the analysis of mortality in Africa (Zaba 1979). Another observation is that the standard itself can be improved by graduation (Hannerz 1999). In the present work, one life table for the total male population and one for the total female population of Sweden in 1982 will be used as standards for males and females respectively. The standard life tables will be graduated by the same mortality laws that will be used to graduate the mortality among the insurance policy holders.

3.1.2 The mortality law for the female population

The works of Brass, Zaba, and Ewbank et al. all imply that

$$G(x) = \log \left[\frac{F(x)}{1 - F(x)} \right] \quad (7)$$

is the choice transformation to work with in relational methods. The mortality law that will be used in the present work gives a five-parameter representation of the function $G(x)$. It was empirically derived by observations on the pattern of

$$g(x) = \frac{dG}{dx} = \frac{f(x)}{F(x)(1 - F(x))} = \frac{\mathbf{m}(x)}{F(x)} \quad (8)$$

in the female part of the Swedish population. The observed pattern suggested that $g(x)$ would be (i) inversely proportional to x^2 for small x (close to zero); (ii) approximately proportional to x throughout the age interval of the labour force (16-64 years) and; (iii) exponentially increasing in the very old ages (Hannerz 1999; 2001). A translation of the above scheme into the language of mathematics would produce a function consisting of three terms: a_1x^{-2} , a_2x and a_3e^{cx} . In this function a_1x^{-2} dominates in the left tail until a_2x takes over as the dominant term followed by a_3e^{cx} , which gradually drives

out the "straight line" and finally achieves a complete domination as x increases towards infinity. Hence,

$$g(x) = a_1x^{-2} + a_2x + a_3e^{cx} \tag{9}$$

where a_1 , a_3 and c are positive valued parameters and a_2 is a parameter that may be negative as long as $g(x)$ always is greater than or equal to zero, and

$$G(x) = \int g(x)dx = a_0 - a_1x^{-1} + \frac{a_2x^2}{2} + \frac{a_3}{c}e^{cx} . \tag{10}$$

By the analytical expressions given above for $g(x)$ and $G(x)$ we also obtain analytical expressions for the cumulative distribution function $F(x) = e^{G(x)}/[1+e^{G(x)}]$, the probability density function $f(x) = g(x)e^{G(x)}/[1+e^{G(x)}]^2$, the hazard function $m(x) = g(x) e^{G(x)}/[1+e^{G(x)}]$, and the survival function $l(x) = 1 - F(x) = [1+e^{G(x)}]^{-1}$.

Hannerz (2001) has shown that the mortality law described above nicely fits the population of Swedish females — a modern female population with low infant mortality and high life expectancy. The function has also been successfully used to graduate mortality among female survivors of acute myocardial infarction (Hannerz 1996a; 1996b), and stroke (Hannerz 1999).

3.1.3 The relational method for the female population

In harmony with Bourgois-Pichat's interpretation of the four factors in which female mortality schedules might differ from one another, we may use (10) to obtain the four-parameter relation

$$\log \left[\frac{F_i(x)}{1 - F_i(x)} \right] - \log \left[\frac{F_j(x)}{1 - F_j(x)} \right] = \mathbf{q}_0 + \mathbf{q}_1x^{-1} + \mathbf{q}_2 \frac{x^2}{2} + \mathbf{q}_3 \frac{e^{cx}}{c} \tag{11}$$

where the \mathbf{q}_i 's are parameters that denote mortality differences and c is a constant held in common by the two mortality schedules, i and j . If the parameter \mathbf{q}_1 in the above expression is different from zero then the two mortality schedules would differ with regard to infant mortality; if $\mathbf{q}_2 \neq 0$ then the schedules would differ in the relationship between mortality in youth and adult life; and if $\mathbf{q}_3 \neq 0$ then the schedules would differ in

the mortality pattern in the old ages. The parameter q_0 would finally determine the difference in the level of mortality. The simplest relation, proportional CDO's would be obtained if all parameters except q_0 were zero.

If we only consider mortality among people who are older than 15 years then the parameter q_1 in (11) becomes redundant. Hence, in the present work the reducible three-parameter relation

$$\log\left[\frac{F_i(x)}{1-F_i(x)}\right] - \log\left[\frac{F_j(x)}{1-F_j(x)}\right] = q_0 + q_2 \frac{x^2}{2} + q_3 \frac{e^{cx}}{c} \quad (12)$$

will be used to relate mortality between female mortality schedules.

3.1.4 The mortality law and relational method for the male population

Whereas the law given by (9) would be adequate for females, Hannerz (1999) has shown that it would be inadequate for male populations. He also showed that the only reason that the law does not fit male mortality schedules is a bulge in the hazard curve, associated with the passage into manhood. Such a bulge has been observed by many researchers and is often referred to as an accident hump (e.g. Heligman and Pollard 1980; Hartmann 1987; Kostaki 1992). If we assume that there is such a thing as a manhood trial—an added hazard intentionally or unintentionally imposed on males, either by themselves or by the environment, which is associated mainly with the passage into manhood—and that some males would die as a direct consequence of such a trial. Then, mortality among males could be divided into two types: mortality from manhood trials and mortality from other causes. A function that fits a male mortality experience would thus be obtained by the following scheme: Let $F_1(x)$ be the death distribution given that the subject will die a "natural" death and $F_2(x)$ be the distribution given that the death is caused by a "manhood trial". Let a be the probability that a death will be natural; then

$$F(x) = aF_1(x) + (1-a)F_2(x).$$

In the present work, the mixture distribution

$$\left\{ \begin{array}{l} F(x) = \mathbf{a}F_1(x) + (1-\mathbf{a})F_2(x) = \mathbf{a} \frac{e^{G_1(x)}}{1+e^{G_1(x)}} + (1-\mathbf{a}) \frac{e^{G_2(x)}}{1+e^{G_2(x)}} \\ G_1(x) = a_0 - \frac{a_1}{x} + \frac{a_2x^2}{2} + \frac{a_3e^{cx}}{c} \\ G_2(x) = a_4 - \frac{a_5}{x} + \frac{a_2x^2}{2} + \frac{a_3e^{cx}}{c} \end{array} \right. \quad (13)$$

will be used to model mortality among males. Hannerz (1999) has shown that (13) fits well when applied to the male population of Sweden. The function has also been successfully used to graduate mortality among people with mental disorders (Hannerz and Borgå 2000), and among male survivors of acute myocardial infarction (Hannerz 1996b), and stroke (Hannerz 1999).

According to Bourgois-Pichat, apart from the four factors needed to explain differences between female mortality schedules, there would be a fifth factor that governs the differences between male and female mortality. From (13) it is obvious that if we let \mathbf{a} be equal to 1 then the expression for the male mortality law would be identical to that of the females. Hence, if we were to associate one of the parameters in (13) with that fifth factor then \mathbf{a} seems to be the most natural choice. With respect to male mortality, Hannerz (1999) has shown that \mathbf{a} can vary between different schedules. Because of this fifth factor, the relational method for male mortality schedules in the present work will consider potential differences, not only in the parameters a_0 , a_2 and a_3 but also in the parameter \mathbf{a} .

3.2 Data procedure and model selection criteria

We used the likelihood ratio test to assess a smaller model against a larger one in which it is embedded. In our data, exact values did not exist for age nor for the number of x year old persons at risk of dying before reaching the age $x+1$. Therefore, all subjects in the age interval 15-19 years were considered to be exactly 17 years at the start of the observation period, the subjects in the age interval 20-24 years were considered to be 22 years etc., and " $m_x + D_x/2$ " (m_x is the person years at risk and D_x is the number of deaths in the age interval $[x, x+1)$) was used as a proxy for the number of persons at risk. The last mentioned approximation is standard (Ajne and Ohlin 1990; Keyfitz 1977; Perks 1932). We used the maximum likelihood method to estimate parameters.

3.3 Model selection for the female populations

A list of models will be given. For each model $-2\log L$ will be calculated. The first model will be the largest and all other will be submodels of the first one. The values of the parameters a_i , $i \in \{0,1,2,3\}$, and c were obtained in a mortality assessment of the total female population of Sweden 1982 and will be regarded as deterministic. The variables y_1 , y_2 and y_3 are indicators for the three different cohorts of insurance policy holders, i.e. $y_i = 1$ if the person belongs to cohort i .

Model W1: *Each cohort obeys its own distribution function.*

$$G(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \left(a_2 + \sum_{i=1}^3 \theta_{2,i} y_i \right) \frac{x^2}{2} + \left(a_3 + \sum_{i=1}^3 \theta_{3,i} y_i \right) \frac{e^{cx}}{c}$$

Number of estimated parameters = 9

$-2\log L = 9345.275$

Model W2: *The three cohorts are related to each other by proportional CDO's.*

$$G(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

Number of estimated parameters = 5

$-2\log L = 9348.589$ P-value = 0.507

Model W3: *The distribution function is independent of cohort.*

$$G(x, y_1, y_2, y_3) = a_0 + \theta_0 - \frac{a_1}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

Number of estimated parameters = 3

$$-2\log L = 9380.050 \quad P\text{-value} = 4.76 * 10^{-6}$$

Model W4: *The three cohorts are related to each other and to the population of Sweden, by proportional CDO's.*

$$G(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \frac{a_2}{2} x^2 + \frac{a_3 e^{cx}}{c}$$

Number of estimated parameters = 3

$$-2\log L = 9354.994 \quad P\text{-value} = 0.137$$

Model W5: *The three cohorts obey the same distribution function as the population of Sweden.*

$$G(x, y_1, y_2, y_3) = a_0 - \frac{a_1}{x} + \frac{a_2}{2} x^2 + \frac{a_3 e^{cx}}{c}$$

Number of estimated parameters = 0

$$-2\log L = 9450.581 \quad P\text{-value} = 1.32 * 10^{-18}$$

According to the above P-values, the smallest permissible model is W4, which contains three estimated parameters. To describe the mortality of the cohorts of female insurance policy holders we thus choose a model with the following interpretation: "The three cohorts are related to each other and to the population of Sweden, by proportional CDO's". Through the P-values we can also conclude that the mortality risks of the insurance policy holders differ significantly from the mean mortality risks of the Swedish population and that the mortality risks are not independent of cohort.

3.4 Model selection for the male populations

Model M1: *Each cohort obeys its own distribution function.*

$$G_1(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \left(a_2 + \sum_{i=1}^3 \theta_{2,i} y_i \right) \frac{x^2}{2} + \left(a_3 + \sum_{i=1}^3 \theta_{3,i} y_i \right) \frac{e^{cx}}{c}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + \left(a_2 + \sum_{i=1}^3 \theta_{2,i} y_i \right) \frac{x^2}{2} + \left(a_3 + \sum_{i=1}^3 \theta_{3,i} y_i \right) \frac{e^{cx}}{c}$$

$$H(x, y_1, y_2, y_3) = \left(\alpha + \sum_{i=1}^3 \beta_i y_i \right) \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + \left(1 - \alpha - \sum_{i=1}^3 \beta_i y_i \right) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 12

$$-2 \log L = 33557.150$$

Model M2: *The three cohorts are related by parallel $G_f(x)$'s.*

$$G_1(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

$$F(x, y_1, y_2, y_3) = (\alpha + \beta) \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + (1 - \alpha - \beta) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 6

$-2 \log L = 33567.311$ $P\text{-value} = 0.118$

Model M3: *The distribution function is independent of cohort.*

$$G_1(x, y_1, y_2, y_3) = a_0 + \theta_0 - \frac{a_1}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

$$F(x, y_1, y_2, y_3) = (\alpha + \beta) \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + (1 - \alpha - \beta) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 4

$-2 \log L = 33605.452$ $P\text{-value} = 8.64 * 10^{-8}$

Model M4: *The three cohorts are related to each other and to the population of Sweden, by parallel $G_i(x)$'s and do not differ from the general population with regard to the accident hump.*

$$G_1(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \frac{a_2}{2} x^2 + \frac{a_3 e^{cx}}{c}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + \frac{a_2}{2} x^2 + \frac{a_3 e^{cx}}{c}$$

$$F(x, y_1, y_2, y_3) = \alpha \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + (1 - \alpha) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 3

$$-2 \log L = 33580.906 \quad P\text{-value} = 4.70 * 10^{-3}$$

Model M5: *Same as model M2, but with $\mathbf{b}=0$.*

$$G_1(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + \left(\frac{a_2 + \theta_2}{2} \right) x^2 + \left(\frac{a_3 + \theta_3}{c} \right) e^{cx}$$

$$F(x, y_1, y_2, y_3) = \alpha \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + (1 - \alpha) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 5

$$-2 \log L = 33567.346 \quad P\text{-value} = 0.178$$

Model M6: Same as model M2, but with $\mathbf{b}=b_2=0$.

$$G_1(x, y_1, y_2, y_3) = a_0 + \sum_{i=1}^3 \theta_{0,i} y_i - \frac{a_1}{x} + a_2 \frac{x^2}{2} + (a_3 + \theta_3) \frac{e^{cx}}{c}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + a_2 \frac{x^2}{2} + (a_3 + \theta_3) \frac{e^{cx}}{c}$$

$$F(x, y_1, y_2, y_3) = \alpha \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + (1 - \alpha) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 4

$$-2 \log L = 33570.282 \quad P\text{-value} = 0.107$$

Model M7: The three cohorts obey the same distribution function as the population of Sweden.

$$G_1(x, y_1, y_2, y_3) = a_0 - \frac{a_1}{x} + \frac{a_2}{2} x^2 + \frac{a_3 e^{cx}}{c}$$

$$G_2(x, y_1, y_2, y_3) = a_4 - \frac{a_5}{x} + \frac{a_2}{2} x^2 + \frac{a_3 e^{cx}}{c}$$

$$F(x, y_1, y_2, y_3) = \alpha \frac{e^{G_1(x, y_1, y_2, y_3)}}{1 + e^{G_1(x, y_1, y_2, y_3)}} + (1 - \alpha) \frac{e^{G_2(x, y_1, y_2, y_3)}}{1 + e^{G_2(x, y_1, y_2, y_3)}}$$

Number of estimated parameters = 0

$$-2 \log L = 34441.945 \quad P\text{-value} < 10^{-100}$$

According to the above P-values, the smallest permissible model is M6, which contains four estimated parameters. To describe the mortality of the cohorts of male insurance policy holders we thus choose a model with the following interpretation: “The three cohorts are related to each other by proportional CDO’s but differ from the population of Sweden with respect to the mortality pattern in the old ages”. Through the P-values we can also conclude that the mortality risks of the insurance policy holders differ significantly from the mean mortality risks of the Swedish population and that the mortality risks are not independent of cohort.

4. Discussion

Bourgeois-Pichart (1962) asserted that five factors are needed to explain observed variation between mortality schedules. The present paper suggests a new relational method with one parameter to estimate for each of these factors. As part of a regression model, which also incorporates data from a standard life table, the method was demonstrated on mortality among life insured people in Sweden. The genders were treated separately and life-tables for the total Swedish male/female population were used as standards. The results confirm earlier observations that populations with considerable differences in the level of mortality can still be associated with a high degree of similarity with respect to the shape of the mortality curve. For each cohort of life insured females we needed only to estimate one parameter, and for the males four parameter estimates were enough to describe mortality in three cohorts.

The relational method that was used was intertwined with a law of mortality. Besides yielding an increased statistical precision, mortality laws often have the following advantages:

- The mortality data for each person can be treated as an individual record. Hence there is no need to aggregate the data into age groups, and individual-based ages and follow-up periods can be used in the analysis.
- A mathematical expression for the survival function is obtained with which probabilities for death can be calculated for any age and any risk period.

Not having to group the data would be especially useful in epidemiological studies where many prognostic variables might be studied simultaneously among people of different ages, who enter the study at different time points, but who are still followed up to a certain common end-of-the-study date. The advantages are of less importance in demographic research. Kostaki (1991) showed, however, that mortality laws provide a

convenient method to transform abridged life tables into unabridged ones. Another advantage is that smoothness of the curve is obtained, which makes the table look better. This feature is especially appreciated in the insurance industry in that insurance tables seem more reasonable when the premium rises steadily rather than irregularly with age (Keyfitz 1982). "... if the object of graduation be restricted to a practical instrument, there would be no doubt of the superiority of the curve-fitting method, yielding as it does a perfect smoothness impossible of attainment by any other method" (Perks 1932).

Mortality laws have, however, also been subject to some severe criticism: "Elaborate models requiring many constants to be determined are useless. This rules out the whole range of attempts to express mortality as a mathematical function of age from Gompertz onwards," (Brass 1974).

It must be admitted that the laws that were used in the present work would have been useless if they had been the only ingredients of the statistical model. It would, for example, have been clearly impossible to ascertain sensible estimates for all of the eight constants involved in the law that was used for the males if those estimates had to be based on nothing but one of the direct data sets given in Table 3. If, on the other hand, we regard mortality data from large populations, like the total population of Sweden that contains more than 4 million males, then the eight parameters are not too many, especially when compared to the un-graduated national life table, which in fact involves estimation of one parameter for each of one hundred age groups.

The key point of the present paper is that it is not necessary to choose between the application of a model life table, a relational method or a mortality law. You can use them all simultaneously.

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