Health-linked life annuites: combining protection and retirement income

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Agenda

- 1. Introduction & motivation
- 2. Life annuity design on the move
- 3. Health-linked life annuities: a general framework
- 4. Actuarial issues
- 5. Concluding remarks

1 INTRODUCTION & MOTIVATION

In current scenarios, many "traditional" products in the field of life & health insurance:

- do not fit the individual needs
- imply severe risks for insurance companies (and pension funds)

Significant individual needs:

- > post-retirement income (individual longevity risk)
- health-related protection, in particular at old and very old ages (expense risk and individual longevity risk)

Risks borne by the insurer:

- individual and aggregate longevity risk
- pricing and reserving risks due to
 - uncertainty in future mortality trends
 - ▷ poor statistical experience, in particular regarding high ages

Introduction & motivation (cont'd)

Looking at recent trends and proposals

- 1. Life annuities:
 - (a) from "investment" to longevity insurance \Rightarrow old-age life annuities
 - (b) extension of the rating principles \Rightarrow special-rate, or underwritten life annuities
 - (c) linking annuity benefits to aggregate longevity experience \Rightarrow *longevity-linked life annuities*
- 2. Long-term Care Insurance (LTCI): from stand-alone to *combo products*, e.g. including lifetime-related benefits

Note that:

- Approach 1(a) \Rightarrow restriction of the coverage (time frame)
- Approach 1(c) \Rightarrow possible reduction of the coverage (amount)
- Approach (2) \Rightarrow extension of the coverage (more needs)
- Approaches 1(b) and (2) ⇒ possible implementation via health-linked life annuities

Introduction & motivation (cont'd)

All the above trends and proposals should carefully be analyzed, also to capture new opportunities in product design

Main aim of this presentation: to provide a general framework, hopefully useful in exploring the broad (and evolving) range of recent or innovative products

2 LIFE ANNUITY DESIGN ON THE MOVE



How to design more attractive life annuities

Old-age life annuities

From life annuity as an investment to longevity insurance (with a "deductible")

- delayed annuitization
 - ▷ initial retirement period: drawdown from a fund
 - ▷ life annuity as insurance on the tail of the lifetime distribution
- ALDA (Advanced Life Delayed Annuity), proposed by Milevsky [2005]
- RCLA (Ruin Contingent Llfe Annuity), proposed by Huang et al. [2009]

Longevity-linked life annuities

Annual benefit linked to longevity experience (e.g., in a reference population)

Longevity risk sharing between annuity provider and annuitant

 \Rightarrow lower safety loading \Rightarrow better premium rates

A number of linking models proposed in the actuarial literature: see Olivieri and Pitacco [2019] and references therein

Remark

For a survey on technical aspects in life annuity products, see for example: Pitacco [2016a, 2017], and references therein

Among the earliest proposals of innovation in life annuity design, see for example: Wadsworth et al. [2001]

Health-linked life annuities

In the area of life & health insurance products

- immediate standard life annuities, and
- stand-alone long-term care insurance (LTCI) products

implement two "extreme" product designs, both with significant difficulties, from the client's as well as the insurer's perspective:

▷ life annuities: the "annuity puzzle" !

▷ stand alone LTCI: high premium for a "pure protection" product

Health-linked life annuities constitute "hybrid" (or "combo") products which can mitigate risks and disadvantages

3 HEALTH-LINKED LIFE ANNUITIES: A GENERAL FRAMEWORK

Our target:

- to recognize existing products and analyze their relevant features
- to provide hints for (possible) new products

Basic feature of a health-linked life annuity: for a given (single) premium Π , the benefit amount B(t) (either constant or varying throughout the policy duration) depends on the annuitant's health status (either at policy issue or throughout his/her lifetime)

In formal terms, the annuitant's health status can be represented by a stochastic process

$$\{H(t); t \ge 0\}$$

to be defined in terms of:

- ▷ values (e.g., in LTCI via ADL's or IADL's scoring)
- probabilistic structure (e.g., Markov or semi-Markov)

Examples of annuity products in the following

STANDARD LIFE ANNUITY (FLAT PROFILE)

$$B(t) = B = \frac{\Pi}{\ddot{a}_x} = f(\Pi); \ t = 1, 2, \dots$$

where Π = single premium

Health status not explicitly considered, but assumed very good (\Rightarrow annuitants' self-selection)

ANNUITIES WITH STATIC HEALTH-LINKING

$$B(t) = B = \phi(\Pi, H(0)); \ t = 1, 2, \dots$$

Underwritten ("special-rate") life annuity (flat profile)

The health status at policy issue is accounted for via underwriting Purpose:

$$\phi(\Pi, H(0)) > f(\Pi)$$

 \Rightarrow better annuity rate, in case of non-optimal health conditions Classification (according to increasing severity):

- Enhanced life annuities
- Impaired-life annuities
- Care annuities

ANNUITIES WITH DYNAMIC HEALTH-LINKING

 $B(t) = \psi(\Pi, H(t)); \ t = 1, 2, \dots$

Several examples, with related definitions of the health status, in particular:

- LTC annuities
 - ▷ stand-alone and combo products
- Disability annuities (e.g. Income Protection)
 possibly degree-related

Focus on LTC annuities

Stand-alone LTC degree-related annuity

$$B(t) = \begin{cases} 0 & \text{if } H(t) = \text{good} \\ b^{(1)} & \text{if } H(t) = \text{bad} \\ b^{(2)} & \text{if } H(t) = \text{very bac} \end{cases}$$

Health status expressed in terms of ADL's or IADL's

Problem: high sensitivity of actuarial values (premiums and reserves) w.r.t. biometric assumptions (disablement, possible recovery, mortality of disabled people)

Health-linked life annuities: a general framework (cont'd)

Enhanced pension (or life care pension)

$$B(t) = \begin{cases} b' & \text{if } H(t) = \text{good} \\ b'' & \text{if } H(t) = \text{bad} \end{cases}$$

The uplift b'' - b' can be financed by a reduction w.r.t. the basic pension

Advantage: lower sensitivity w.r.t. biometric assumptions

LTC annuity combined with old-age life annuity

$$B(t) = \begin{cases} b & \text{ if } H(t) = \operatorname{good} \land (t \ge t^*) \\ b' & \text{ if } H(t) = \operatorname{bad} \end{cases}$$

e.g., $t^* = 80$

An example of combo product, providing longevity insurance $\,\Rightarrow\,$ LTCI combined with ALDA

The disability state is assumed permanent \Rightarrow the two benefits are mutually exclusive

A death benefit can be added (\Rightarrow bequest motivation)

A NEW PROPOSAL

Combining static linking and dynamic linking can suggest interesting product designs

Doubly enhanced annuity

Formally, benefit given by:

$$B(t) = \Phi(\Pi, H(0), H(t)); \ t = 1, 2, \dots$$

Proposed by Ramsey and Oguledo [2019]

4 ACTUARIAL ISSUES

OUTLINE

Assume that an insurer is willing:

- to enlarge its life annuity portfolio by selling underwritten annuities (static health-linking)
 - b higher heterogeneity, partially observable via proxies provided by underwriting results
 - \triangleright larger portfolio size \Rightarrow better diversification via pooling (as regards idiosyncratic risk)
 - ▷ what about the "balance" ?
- to sell LTCI products (dynamic health-linking)
 - ouncertainty in biometric bases
 - what is the impact of "wrong" biometric bases on actuarial values (premiums and reserves), according to the product design ?
 - ▷ sensitivity analysis to assess the impact

IMPACT OF UNDERWRITTEN LIFE ANNUITIES ON THE PORTFOLIO RISK PROFILE

Risk classification based on a frailty model

A (potential) heterogeneous population split into classes (groups) of individuals with similar risk profile \Rightarrow each class with reduced heterogeneity (w.r.t. heterogeneity in the population)

Biometric assumption: constant multiplicative frailty model in terms of the force of mortality

$$\mu_x(z) = z \,\mu_x$$

For each individual *i* the frailty is a random variable $Z_x^{(i)}$

Assess approximately the individual frailty via medical examination (step of the underwriting process)

Define the group j, j = 1, 2, ..., J, as follows:

$$G_j = \{i : z_{j-1} < Z_x^{(i)} \le z_j\}$$

The probability distribution of the frailty in any given group can be assessed as a conditional distribution of the frailty for the whole population

Given the frailty distribution in group G_j and the survival function in the population, the survival function in group G_j can be derived

 \Rightarrow calculation of relevant actuarial values

Note: residual unobservable heterogeneity inside each group because of frailty

Numerical investigation

0	Frailty interval	Relative size at age 65 of group G_i	Expected value	Coefficient	Expected
Group		in the general population	of the frailty	of variation	lifetime
	$(z_{j-1}, z_j]$	$ ho_{j;65}$	$\mathbb{E}[Z_{65} G_j]$	$\mathbb{CV}[Z_{65} G_j]$	$\mathbb{E}[T_{65} G_j]$
G_1	(0, 1.038741]	60.121%	0.845593	15.243%	22.81
G_2	(1.038741, 1.307144]	30.111%	1.152338	6.479%	20.36
G_3	$(1.307144,\infty)$	9.769%	1.445866	8.736%	18.71
Population	$(0, \infty)$	100%	0.996594	23.308%	21.67

Groups (= Risk classes)

We consider six alternative portfolios (see Table):

- portfolios A E differ for the size of groups G_2 and G_3 , and possibly the total portfolio size
- portfolio F has the same size of A, but a different composition

Groups	Portfolio						
Creape	Α	В	С	D	Е	F	
G_1	1 000	1 000	1 000	1 000	1 000	500	
G_2	0	200	250	200	501	500	
G_3	0	0	0	50	162	0	
All	1 000	1 200	1 250	1 250	1 663	1 000	

Size and composition of alternative portfolios

Results presented in terms of present value of future benefits PV_t paid by the annuity provider, and in particular, to assess the risk profile in terms of:

- probability distribution (via stochastic simulation)
- > coefficient of variation (risk index)

$$\mathbb{CV}[PV_t] = \frac{\sqrt{\mathbb{Var}[PV_t]}}{\mathbb{E}[PV_t]}$$

Time t	Portfolio A	Portfolio B	Portfolio C	Portfolio D	Portfolio E	Portfolio F
0	1.30%	1.20%	1.17%	1.18%	1.04%	1.87%
5	1.48%	1.37%	1.34%	1.35%	1.19%	1.55%
10	1.75%	1.62%	1.60%	1.60%	1.39%	1.80%
15	2.10%	1.96%	1.91%	1.93%	1.70%	2.19%
20	2.64%	2.45%	2.41%	2.43%	2.17%	2.80%
25	3.55%	3.34%	3.31%	3.31%	3.04%	3.97%
30	5.62%	5.38%	5.32%	5.35%	4.96%	6.54%
35	11.10%	10.78%	10.78%	10.73%	10.28%	13.82%
40	32.19%	32.19%	32.19%	32.19%	31.40%	44.42%
45	136.25%	136.25%	136.25%	136.25%	136.25%	

Coefficient of variation of the present value of future benefits: $\mathbb{CV}[PV_t]$

Main findings and related interpretations

- Portfolio F: the highest riskiness
- Comparing F to A: same size, but in F more heterogeneity (groups G₁ and G₂) not counterbalanced by larger size ⇒ higher riskiness
- Portfolio E: high heterogeneity (groups G₁, G₂ and G₃) counterbalanced by the largest size ⇒ lowest riskiness, even lower than portfolio A, thanks to larger size
- \triangleright Higher degrees of heterogeneity \Rightarrow higher risk profile
- If matched by larger total portfolio size, risk profile can benefit from portfolio diversification (pooling effect)

For details, see: Olivieri and Pitacco [2016]

LTCI: A SENSITIVITY ANALYSIS

Uncertainty in technical bases, in particular biometric assumptions:

- probability of disablement, i.e. prob. of entering LTC state
- mortality of disabled people, i.e. mortality in LTC state

The following products addressed in the sensitivity analysis (see: Pitacco [2016b])

Stand-alone LTCI

(Product P1)

LTCI benefit: a lifelong annuity with predefined annual amount, from the LTC claim on

LTCI as an acceleration benefit in a whole-life assurance

(Product P2(s))

Annual LTC benefit = $\frac{\text{sum assured}}{s}$, paid for *s* years at most

Package including LTC benefits and lifetime-related benefits

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(Products P3a(x+n) and P3b(x+n))
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Benefits:

- (I) a lifelong LTC annuity, from the LTC claim on
- (II) a deferred life annuity from age x + n (e.g. x + n = 80), while the insured is not in LTC disability state
- (III) a lump sum benefit on death, alternatively given by
 - (IIIa) a fixed amount, stated in the policy
 - (IIIb) the difference (if positive) between a fixed amount and the total amount paid as benefit 1 and/or benefit 2

Benefits (I) and (II) are mutually exclusive

Enhanced pension (Life care pension)

(Product P4(b', b''))

LTC annuity benefit defined as an uplift with respect to the basic pension b

Uplift financed by a reduction (with respect to the basic pension b) of the benefit paid while the policyholder is healthy

- \triangleright reduced benefit b' paid as long as the retiree is healthy
- ▷ uplifted lifelong benefit b'' paid in the case of LTC claim Of course, b' < b < b''

Remark

For details on LTCI products, see for example: Pitacco [2014], and references therein

Biometric functions (needed)

Three-state model, one LTC state, no recovery

For an active (healthy) individual age x:

 $q_x^{aa} =$ prob. of dying before age x + 1

 $w_x = \text{ prob. of becoming invalid (disablement, i.e. LTC claim)}$ before age x + 1

For an invalid (in LTC state) age *x*:

 $q_x^i = \text{ prob. of dying before age } x+1$

Remark

No dependence on time elapsed since disability inception is allowed for

 \Rightarrow a Markov chain model is then adopted

Assumptions

 q_x^{aa} : life table (first Heligman-Pollard law) w_x : a specific parametric law $q_x^i = q_x^{aa} + extra-mortality$ (i.e. additive extra-mortality model)

Life table

First Heligman-Pollard law:

$$\frac{q_x^{aa}}{1 - q_x^{aa}} = a^{(x+b)^c} + d e^{-e \left(\ln x - \ln f\right)^2} + g h^x$$

In practice the following approximation can be used:

$$q_x^{aa} \approx \frac{g \, h^x}{1 + g \, h^x}$$

a	b	С	d	e	f	g	h
0.00054	0.01700	0.10100	0.00014	10.72	18.67	$2.00532\mathrm{E}\!-\!06$	1.13025

The first Heligman-Pollard law: parameters

$\overset{\circ}{e}_{0}$	$\overset{\mathrm{o}}{e}_{40}$	$\overset{\mathrm{o}}{e}_{65}$	Lexis	q_0^{aa}	q_{40}^{aa}	q_{80}^{aa}
85.128	46.133	22.350	90	0.00682	0.00029	0.03475

The first Heligman-Pollard law: some markers

Disablement (LTC claim)

Assumption by Rickayzen and Walsh [2002]:

$$w_x = \begin{cases} A + \frac{D - A}{1 + B^{C - x}} & \text{for females} \\ \left(A + \frac{D - A}{1 + B^{C - x}}\right) \left(1 - \frac{1}{3} \exp\left(-\left(\frac{x - E}{4}\right)^2\right)\right) & \text{for males} \end{cases}$$

Parameter	Females	Males	
A	0.0017	0.0017	
B	1.0934	1.1063	
C	103.6000	93.5111	
D	0.9567	0.6591	
E	n.a.	70.3002	

Parameters Rickayzen-Walsh



Probability of disablement (Males)

Extra-mortality

Assumption by Rickayzen and Walsh [2002]:

$$q_x^{i^{(k)}} = q_x^{[\text{standard}]} + \Delta(x, \alpha, k)$$

with:

$$\Delta(x, \alpha, k) = \frac{\alpha}{1 + 1.1^{50 - x}} \frac{\max\{k - 5, 0\}}{5}$$

where:

- parameter k expresses LTC severity category
 ▷ 0 ≤ k ≤ 5 ⇒ less severe ⇒ no impact on mortality
 ▷ 6 ≤ k ≤ 10 ⇒ more severe ⇒ extra-mortality
- parameter α (assumption by Rickayzen [2007])

lpha=0.10 if $q_x^{[ext{standard}]}=q_x^{aa}$ (mortality of insured healthy people)

Our (base) choice: $\alpha = 0.10$, k = 8; hence:

$$q_x^i = q_x^{aa} + \Delta(x, 0.10, 8) = q_x^{aa} + \frac{0.06}{1 + 1.1^{50 - x}}$$



Mortality assumptions (Males)

Sensitivity analysis concerning:

- probability of disablement, i.e. entering into LTC state
- extra-mortality of insureds in LTC state

Notation:

 $\Pi_x^{[PX]}(\delta, \lambda)$ = actuarial value (single premium) of product PX, according to the following assumptions:

• $\delta \Rightarrow$ disablement

$$\bar{w}_x(\delta) = \delta w_x$$

where w_x is given by the previous Eq. (assumption by Rickayzen and Walsh [2002])

• $\lambda \Rightarrow$ extra-mortality

$$\bar{\Delta}(x;\lambda) = \lambda \,\Delta(x,\alpha,k) = \Delta(x,\lambda \,0.10,8)$$

and hence:

$$q_x^i(\lambda) = q_x^{aa} + \bar{\Delta}(x;\lambda)$$

For products P1, P2, P3, normalize and define the ratio:

$$\rho_x^{[\mathrm{PX}]}(\delta,\lambda) = \frac{\Pi_x^{[\mathrm{PX}]}(\delta,\lambda)}{\Pi_x^{[\mathrm{PX}]}(1,1)}$$

For product P4, with given b and b'', normalize and define the ratio:

$$\rho_x^{[P4]}(\delta,\lambda) = \frac{b'(1,1)}{b'(\delta,\lambda)}$$

For all the products, we first perform *marginal* analysis, i.e. tabulating the functions:

 $\Pi_x^{[PX]}(\delta, 1)$ for P1, P2, P3, $b'(\delta, 1)$ for P4; $\rho_x^{[PX]}(\delta, 1)$ for P1, P2, P3, P4 $\Pi_x^{[PX]}(1, \lambda)$ for P1, P2, P3, $b'(1, \lambda)$ for P4; $\rho_x^{[PX]}(1, \lambda)$ for P1, P2, P3, P4

Sensitivity analysis: disablement assumption (parameter δ)



Ratios $\rho_x^{[\mathrm{PX}]}(\delta,1)$

Sensitivity analysis: extra-mortality assumption (parameter λ)



Ratios $ho_x^{[\mathrm{PX}]}(1,\lambda)$

Joint sensitivity analysis (parameters δ , λ)

Example 1

For the generic product PX, and a given age x, analyze the function:

$$z = \Pi_x^{[\mathrm{PX}]}(\delta, \lambda)$$

Example 2

For the generic product PX, and a given age x, find (δ, λ) such that:

$$\rho_x^{[PX]}(\delta, \lambda) = \rho_x^{[PX]}(1, 1) = 1$$
(*)

Eq. $(^{\ast})$ implies

• for products P1, P2, P3:

$$\Pi_x^{[\mathrm{PX}]}(\delta,\lambda) = \Pi_x^{[\mathrm{PX}]}(1,1)$$

• for product P4:

$$b'(\delta,\lambda)=b'(1,1)$$





Offset effect: isopremium lines

6 CONCLUDING REMARKS

When developing a new product:

- What benefit structure, e.g. what time profile of the health-linked benefits
- What rating model, in particular what information about the applicants should be taken into account (\Rightarrow rating classes)
- What probabilistic model
- What data

Starting from the bottom:

 Data are (almost) always a problem ⇒ sensitivity analysis can suggest adjustments in the product design

- Probabilistic model
 - does not constitute a problem by itself: Markov and semi-Markov multistate models capture whatever benefit structure
 - \triangleright its implementation can constitute a problem because of lack of data \Rightarrow approximations frequently needed
- Appropriate rating models can be suggested by recent proposals in the context of underwritten life annuities: a large variety of products, sharing the purpose of "tailoring" the premium rate
- Various benefit structures can be conceived, aiming at a higher flexibility of the benefit amount, in line with the annuitant's needs
 - suggestions e.g. from Income Protection policies, with reduction of benefit in case of partial recovery
 - complex claim settlement and monitoring then required

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Where links are provided, they were active as of the time this presentation was completed but may have been updated since then

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