

13th DHC 2019

January 23-24-25
Papendal, Arnhem

HOVON • NVvH

Dutch Hematology Congress



Primary therapy and relative survival in primary myelofibrosis: a population-based study among 1,599 patients diagnosed in The Netherlands, 2001-2016

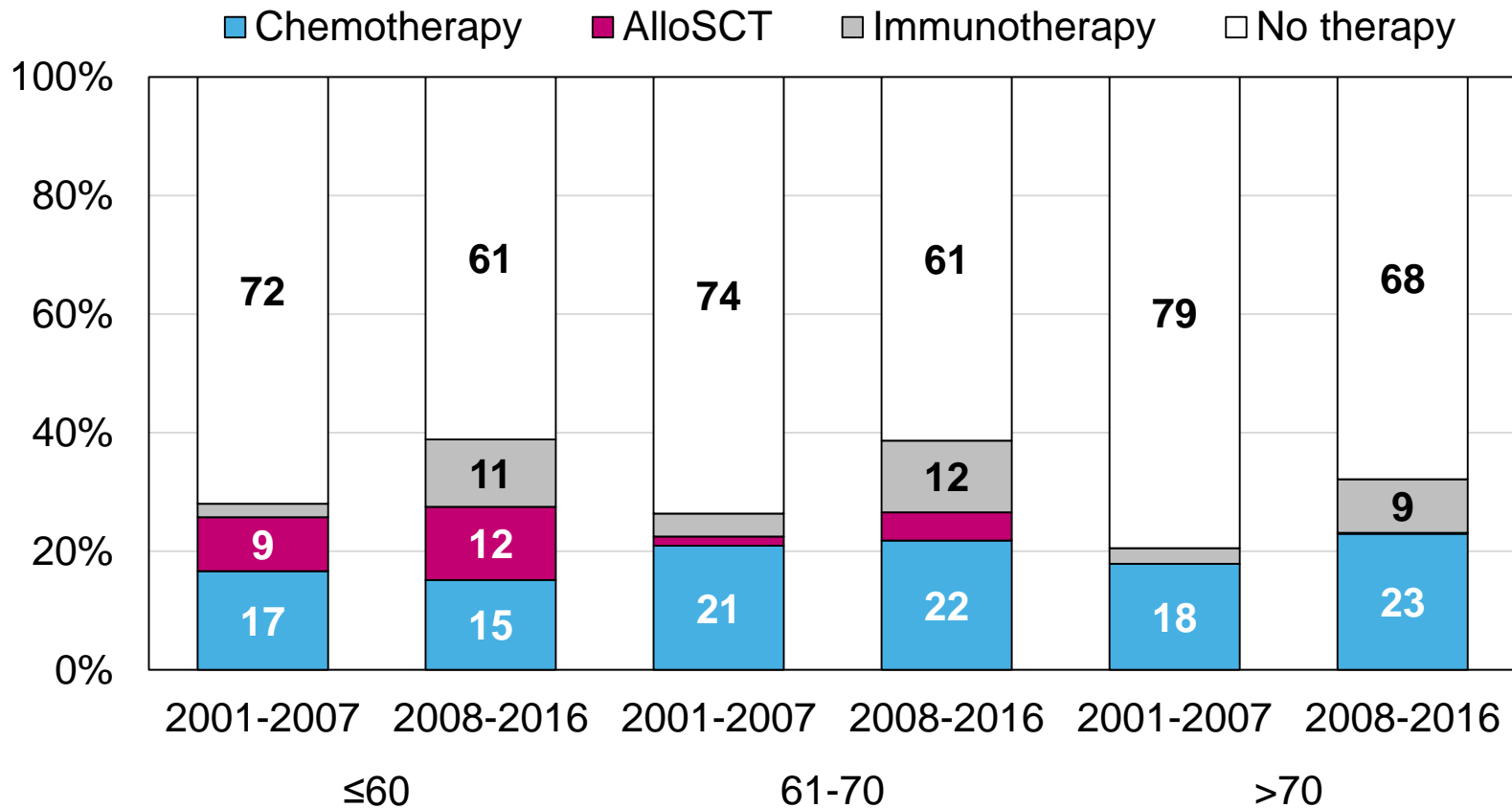
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13th Dutch Hematology Congress | Papendal, Arnhem | January 23, 2019

Primary therapy of patients with pMF

According to age at diagnosis and period of diagnosis



Age at diagnosis and period of diagnosis

Proportions are depicted in the bars

Primary therapy of patients with pMF

Data from 401 patients diagnosed during 2014-2016

No anti-neoplastic therapy (N = 223; 56%)

Chemotherapy (N = 89; 22%)

- 98%: hydroxyurea (n = 88)
- 2%: melfalan (n = 1)

Immunotherapy (N = 71; 18%)

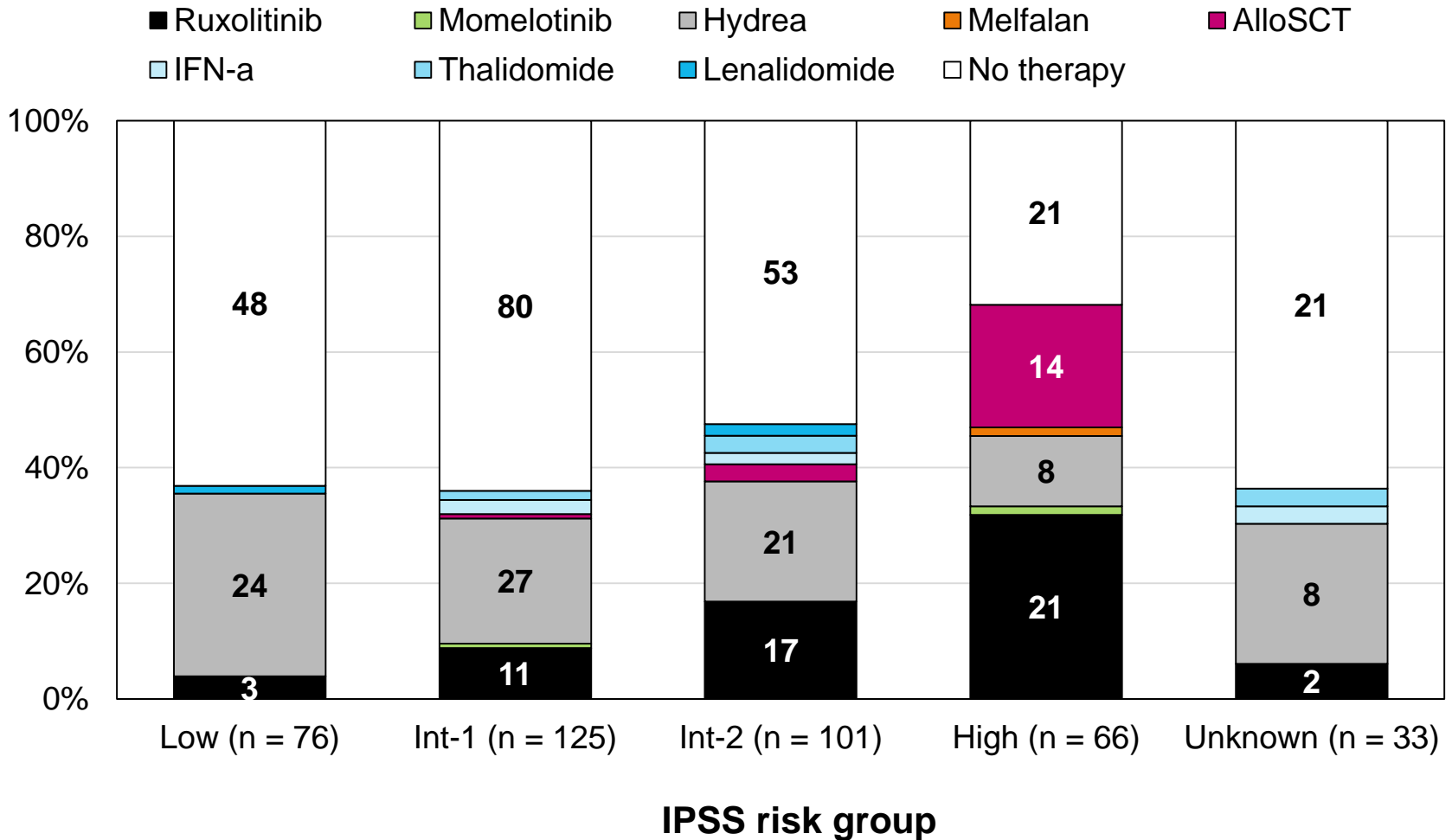
- 75%: ruxolitinib (n = 54)
- 9%: pegylated-IFN- α (n = 6)
- 9%: thalidomide (n = 6)
- 4%: lenalidomide (n = 3)
- 4%: momelotinib (n = 2)

AlloSCT (N= 18; 4%)

- 72%: RIC-alloSCT (n = 13)
- 28%: myeloablative alloSCT (n = 5)

Primary therapy of patients with pMF

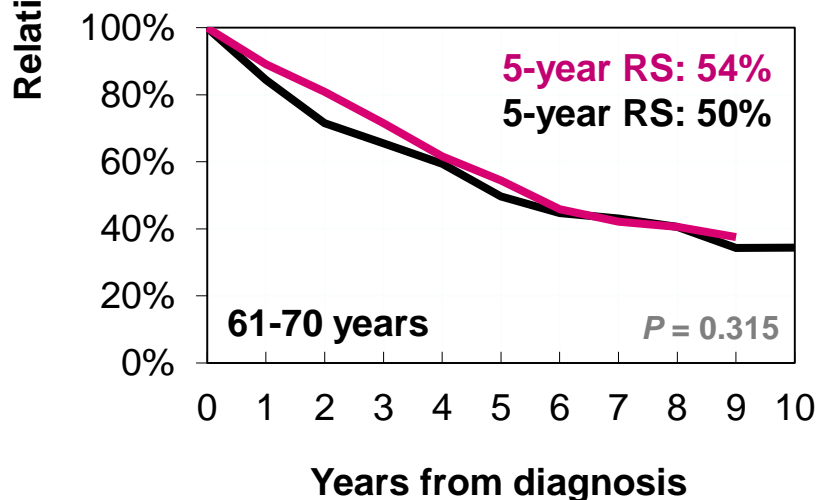
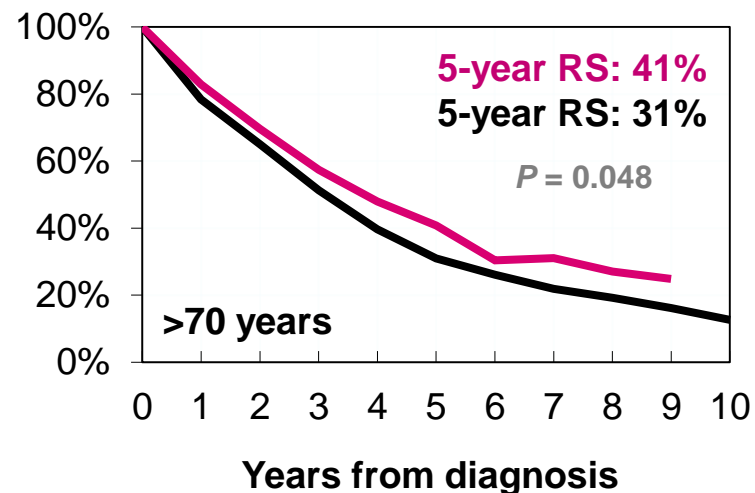
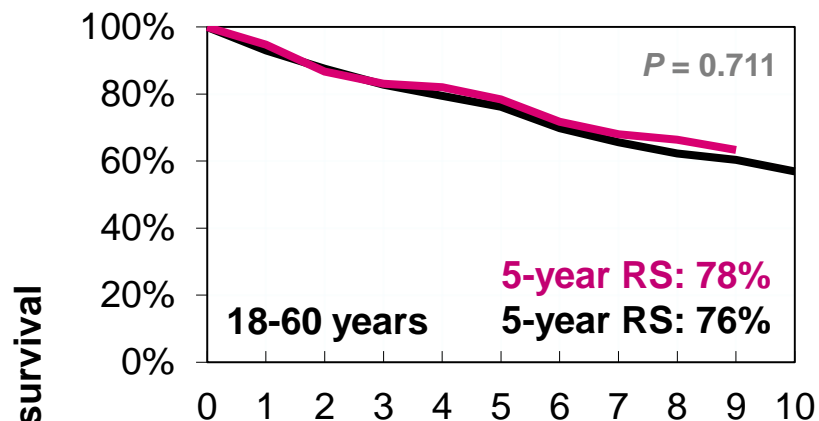
Data from 401 patients diagnosed during 2014-2016



Absolute numbers are depicted in the bars

Relative survival of pMF

According to age at diagnosis and period of diagnosis



— 2001-2007 — 2008-2015

- The majority of patients received no anti-neoplastic therapy
 - Application of immunotherapy increased modestly
- Patients with pMF experience considerable excess mortality
 - Current treatment practices might be inadequate to improve outcome
 - Survival improved in >70 years: better management or earlier diagnosis?
- Future population-based research is needed to assess the impact of improved management and novel agents on pMF outcome
 - For example, broader application of RIC-alloSCT and ruxolitinib
 - The latter registered for routine use in The Netherlands as from 2014

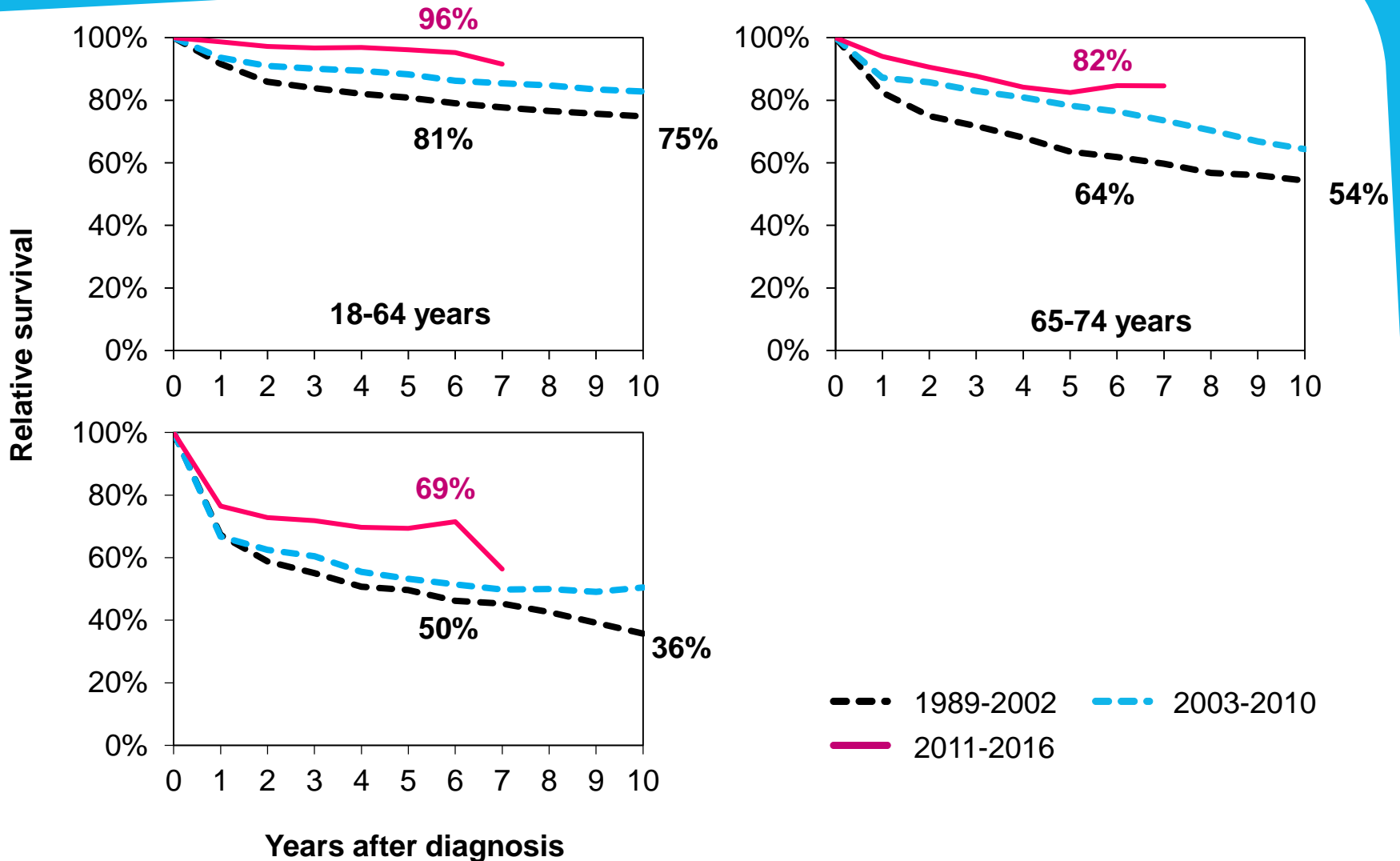
Survival continues to increase in diffuse large B-cell lymphoma: a population-based analysis among patients diagnosed in the Netherlands between 1989 and 2016

Müjde Durmaz,¹ Djamila E. Issa,^{2,3} Otto Visser,⁴ Sonja Zweegman,² Marie José Kersten,⁵ Pieterella J. Lugtenburg,⁶ Martine E.D. Chamuleau,² Avinash G. Dinmohamed^{1,6,7}

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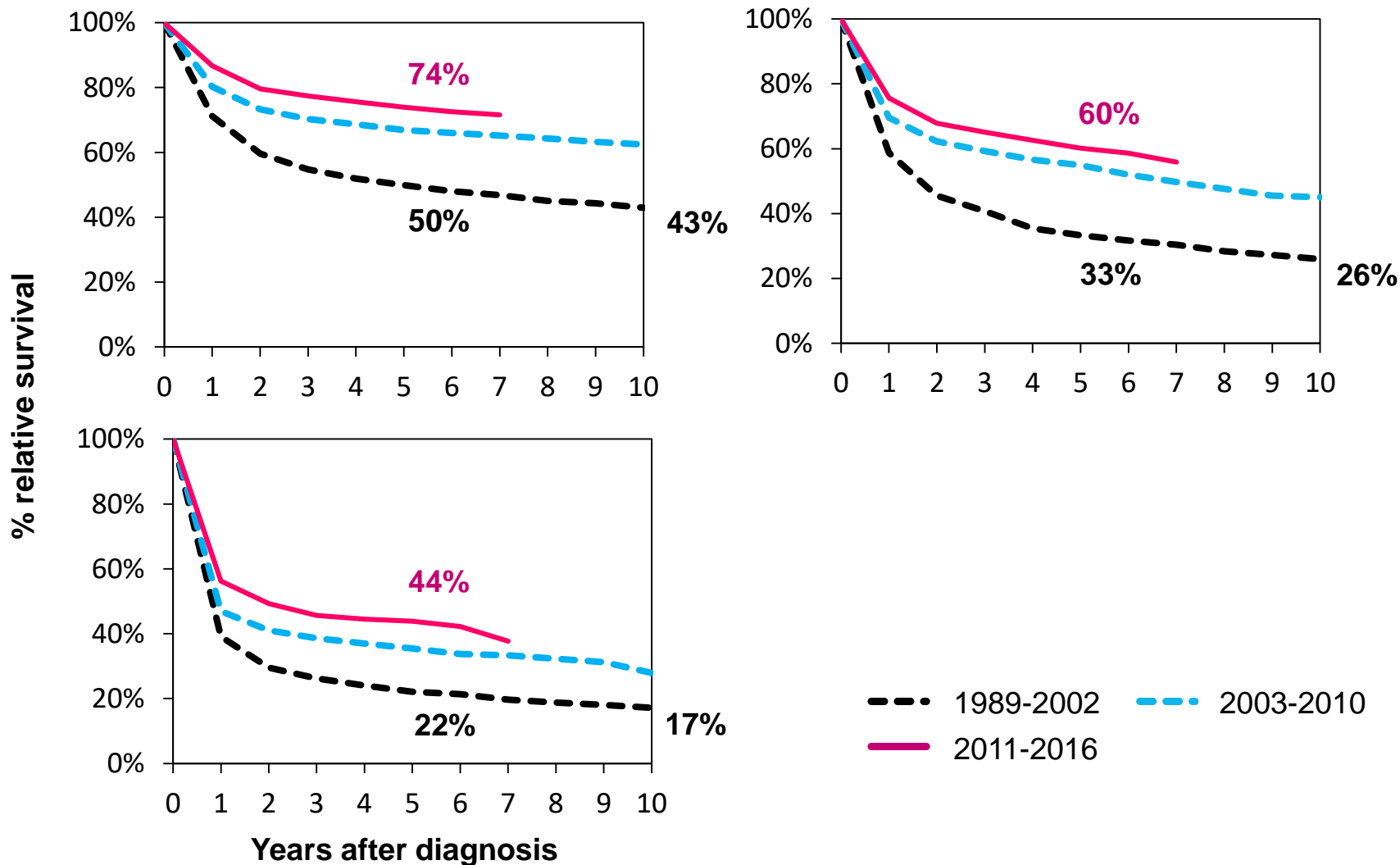
Relative survival of stage I DLBCL

According to age at and calendar period of diagnosis



Relative survival of DLBCL stage II-IV

According to age at and calendar period of diagnosis



- In contemporary clinical practice, survival continues to increase among patients with DLBCL in The Netherlands
 - Attributed to changes in therapy over time
 - Especially among patients with stage II-IV DLBCL
 - Also, advances in supportive care, staging techniques, and risk-adapted therapy might have accounted for the improvement
- The current population-based study provides a benchmark to assess survival with novel treatment strategies for DLBCL

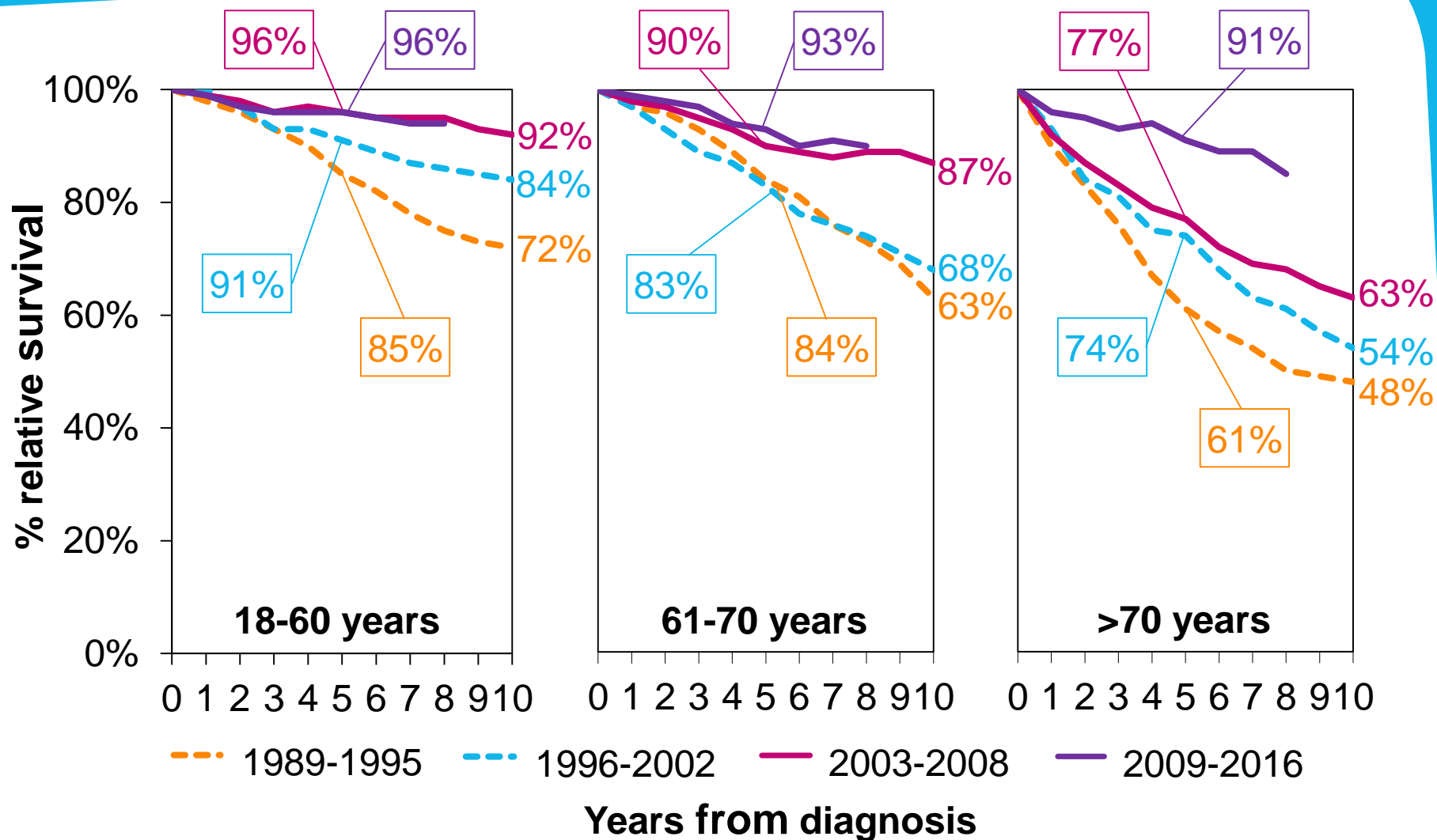
Incidence, primary therapy, and survival of follicular lymphoma (FL): a population-based analysis in the Netherlands, 1989-2016

Manette A.W. Dinnessen,¹ Marjolein W.M. van der Poel,² Sanne H. Tonino,³ Otto Visser,⁴ Marie José Kersten,³ Pieterella J. Lugtenburg,⁵ Avinash G. Dinmohamed^{1,5,6}

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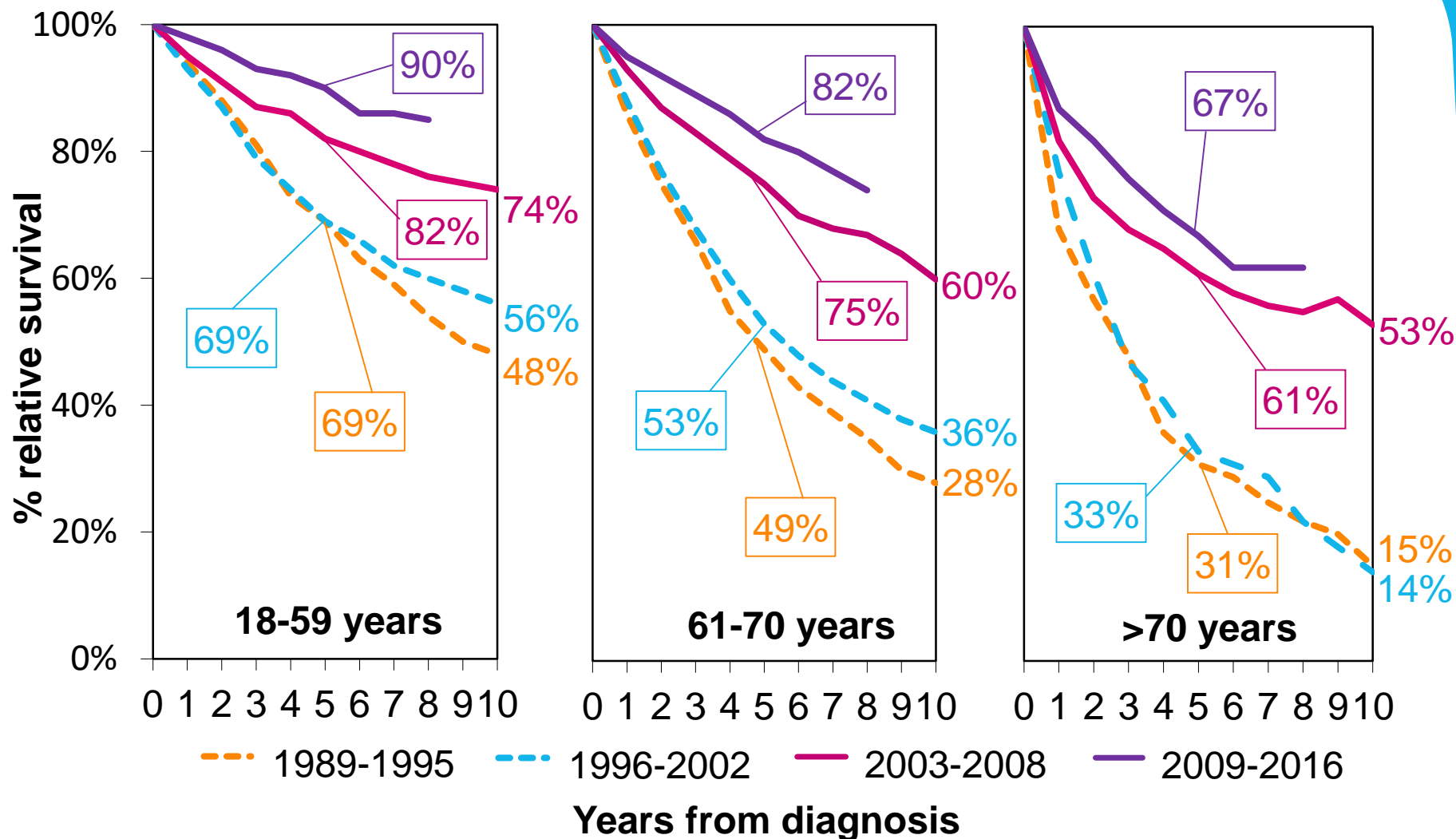
Relative survival of limited-stage FL

According to age at and period of diagnosis



Relative survival of advanced-stage FL

According to age at and period of diagnosis



- Survival increased across both stage and all age groups
 - Most pronounced in older age groups and advanced-stage FL
- For limited-stage FL, the effect of age on survival seems to have diminished over time
 - Excess mortality was as low as 10% in in the most current period
- For advanced-stage FL, there was a massive jump in improvement of survival after the introduction of rituximab

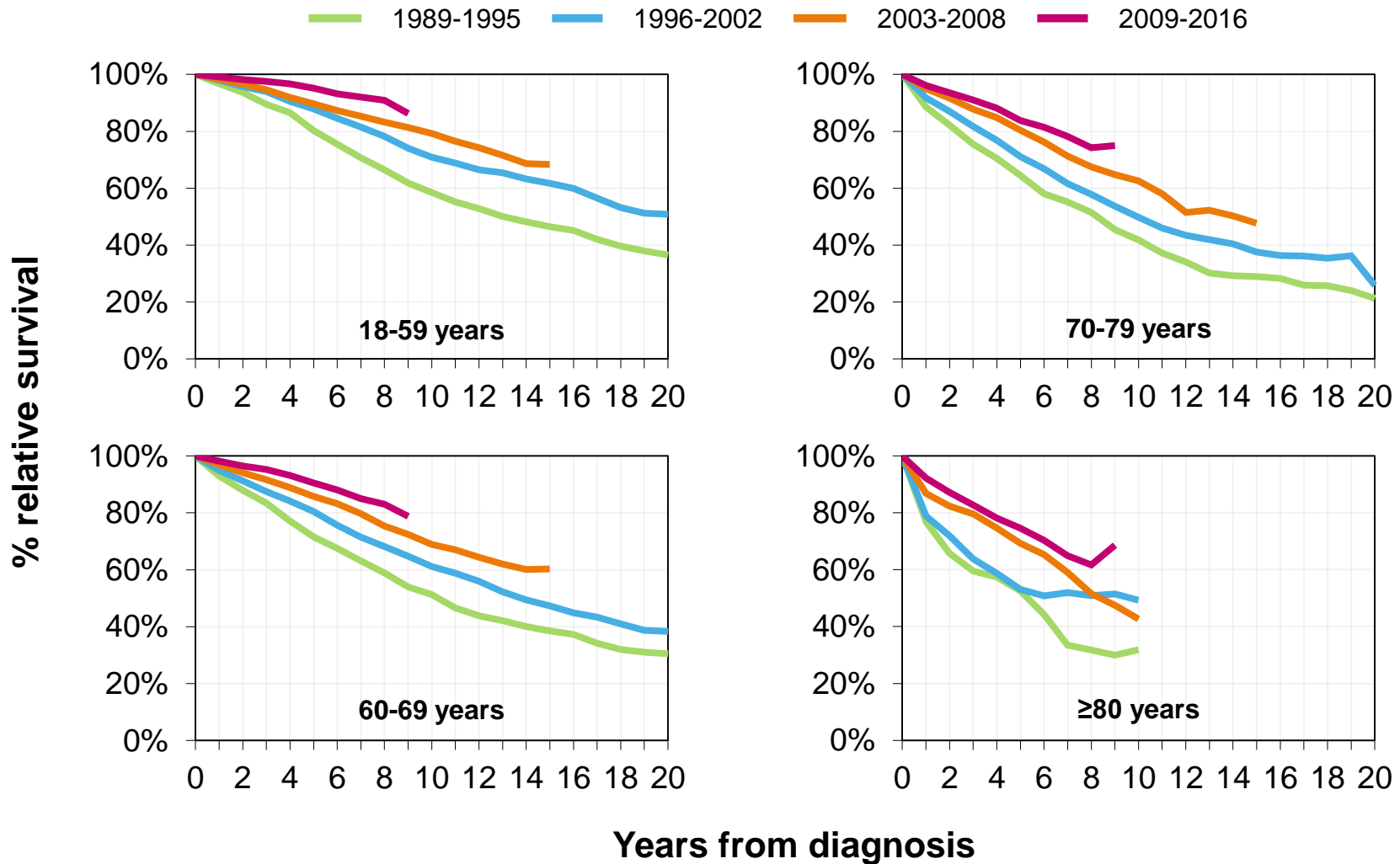
Survival continues to increase in CLL: a population-based analysis among 20,324 patients diagnosed in the Netherlands between 1989 and 2016

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Relative survival of CLL

According to age at and calendar period of diagnosis



- In this large, nationwide, population-based study, 5- and 10-year RS improved over time among patients with CLL across all age groups
- Advances in supportive care and ameliorated management might have accounted for the improvement
 - Earlier CLL detection might also artificially increase survival estimates
 - However, this would have only marginally biased our results, as the overall ASR remained comparatively steady since the period 2003-2008
- Kinase inhibitors and anti-apoptotic agents may further improve RS
 - The current study thus provides a benchmark for future research

Treatment and survival of patients with primary plasma cell leukemia: a population-based study among 192 patients in The Netherlands, 1989 to 2016

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Niels W.C.J. van de Donk,³ **Avinash G. Dinmohamed**,¹⁻³

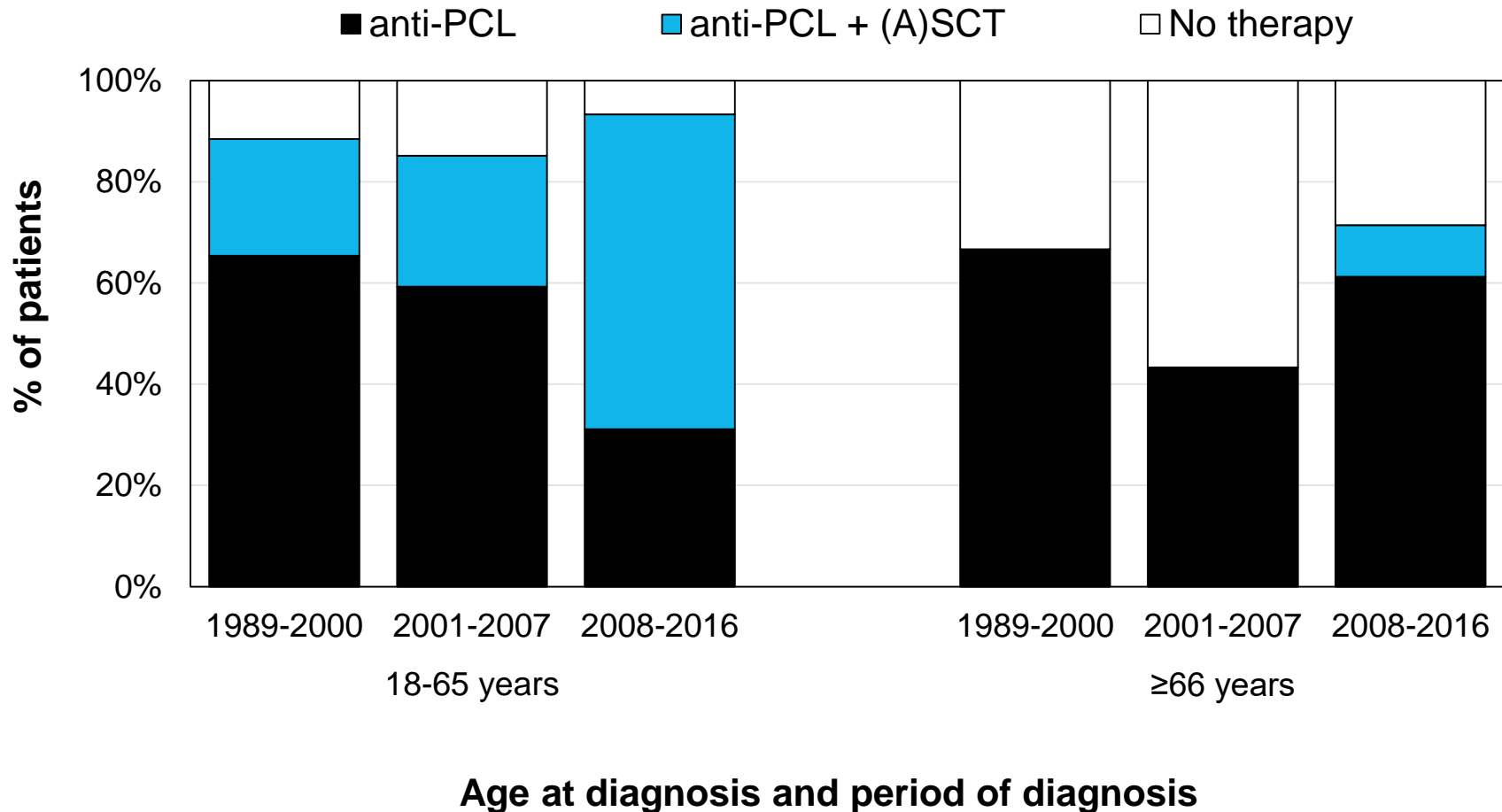
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Patient characteristics at diagnosis

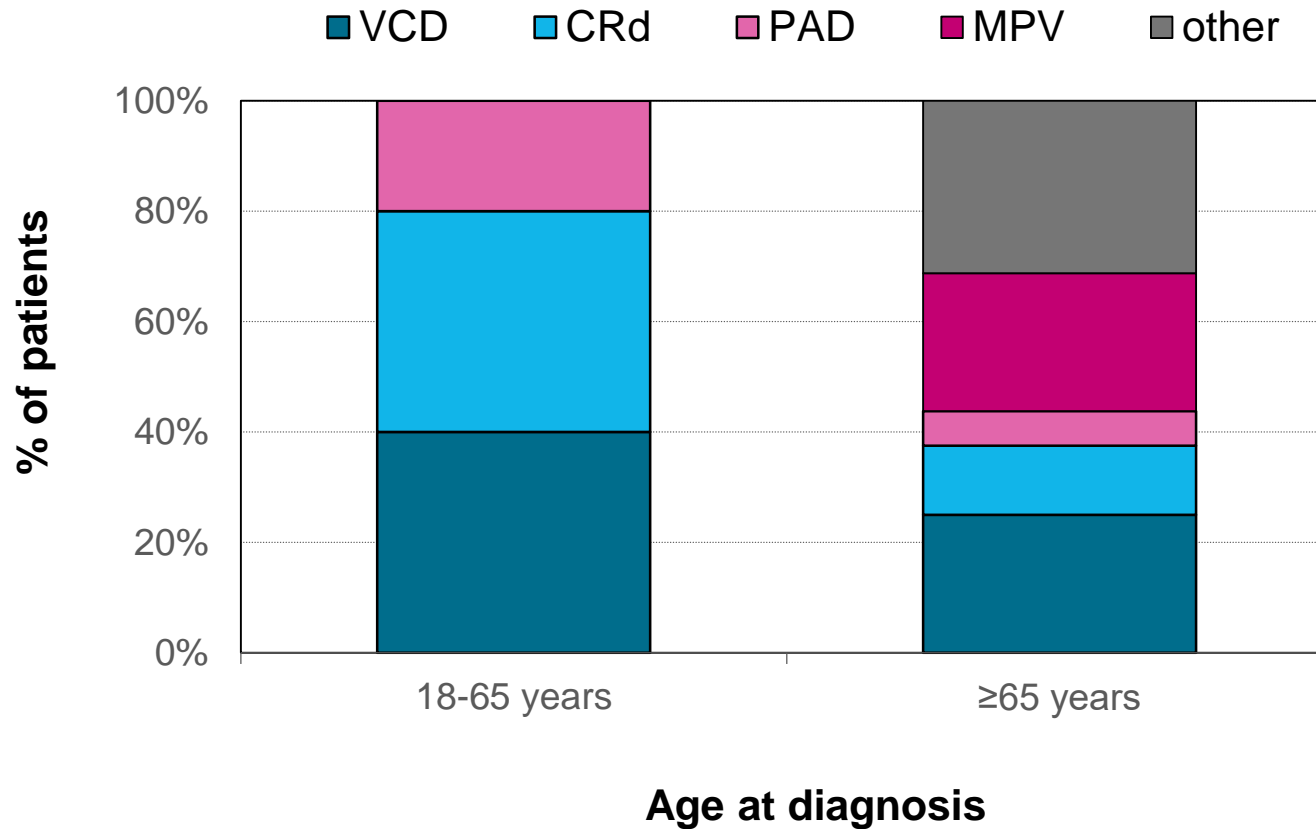
Characteristics	Calendar period			Total	
	1989-2000	2001-2007	2008-2016	No.	
Total No. of patients	41	57	94	192	
Sex					
Male	61%	47%	51%	100	52%
Female	39%	53%	49%	92	48%
Age, years					
Median (IQR)	63 (51-71)	66 (57-76)	66 (60-75)	65 (58-74)	
18-65	63%	47%	48%	98	51%
≥66	37%	53%	52%	94	49%
Hospital type of diagnosis					
non-academic	78%	75%	76%	146	76%
academic	22%	25%	24%	46	24%
Previous malignancy					
No	93%	95%	85%	172	90%
Yes	7%	5%	15%	20	10%

Abbreviation: IQR, inter quartile range

Primary therapy of pPCL in The Netherlands



Primary therapy of pPCL in The Netherlands, 2014-2016

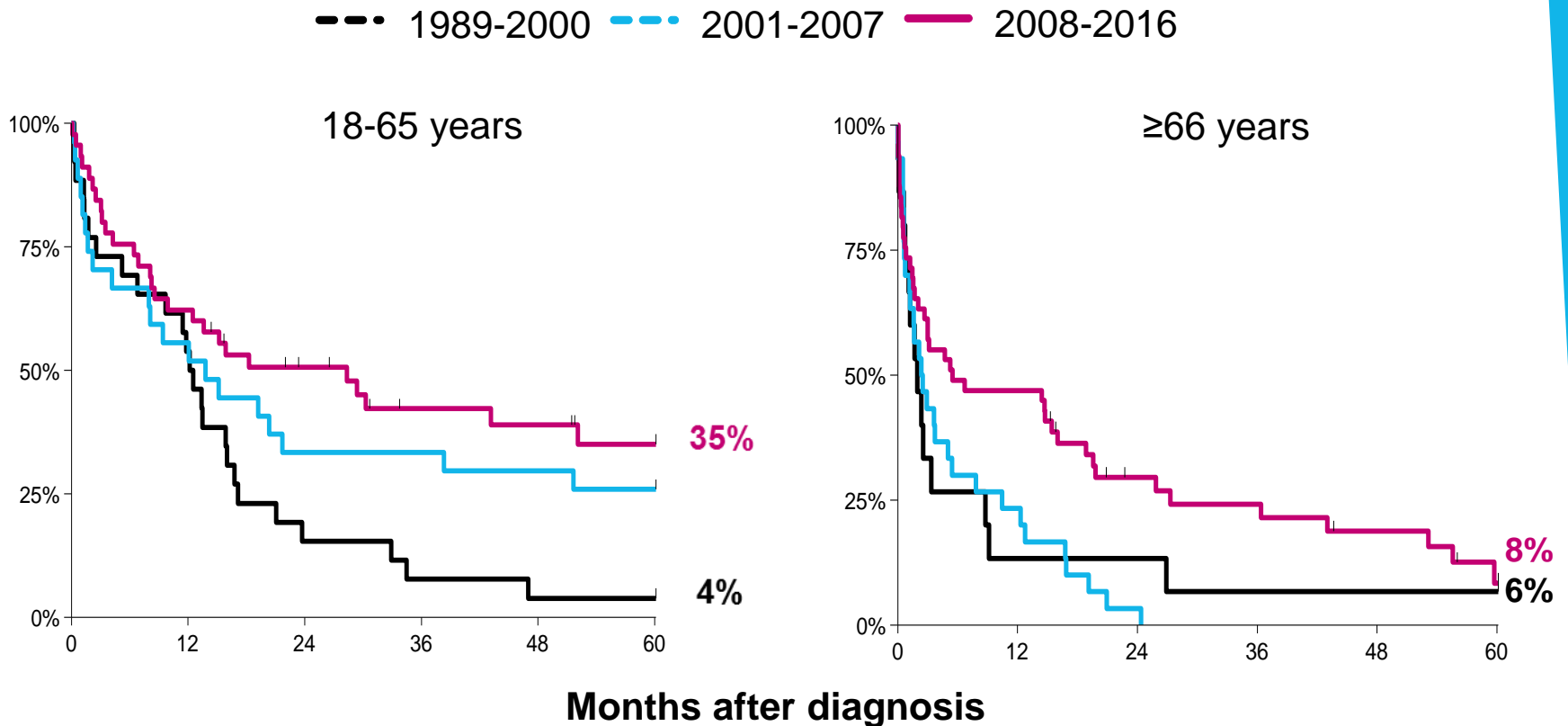


Overall survival of pPCL in The Netherlands

	18-65 years			≥66 years	
Median OS (months)	12.2	28.4	Median OS (months)	2.0	5.5

--- 1989-2000 - - - 2001-2007 — 2008-2016

Overall survival of pPCL in The Netherlands



- pPCL is a very rare malignancy
- Application of (A)SCT increased substantially over time
 - for patients aged ≤ 65 , but also for the elderly
- Overall survival improved greatly over time for patients ≤ 65 years
 - largely explained by changes in the application of therapy over time
- Therapeutic advances gradually translate into benefits for patients with pPCL, particularly among patients aged ≤ 65

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