Monoclonal Gammopathy of Undetermined Significance (MGUS) and Smoldering Multiple Myeloma (SMM): The Key To Neoplasia

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Disclosures for Robert A. Kyle

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Data Monitoring Committee



Merck

Monoclonal Gammopathy of Undetermined Significance: Prevalence ≥50 Years

Site of Study	Subjects	With M-proteins
	(number)	(number, %)
Sweden	3,674	59 (1.6%)
France	17,968	303 (1.7%)
USA	1,200	15 (1.25%)







MGUS: Olmsted County, Minnesota Inclusion Criteria

> Serum M-spike <3.0 g/dL

 Bone marrow plasma cells <10% (if done)

 No evidence of other B-cell disorders

No end-organ damage



MGUS: Olmsted County, Minnesota Prevalence Study

 Olmsted County residents ≥50 years: 28,038

 Serum samples obtained from population: 77%

Kyle et al: New Engl J med 354:1362, 2006



MGUS: Olmsted County, Minnesota

			<u>Posi</u>	<u>tive</u>
	Ν	%	Ν	%
Male	9,469	44	350	3.7
Female	11,994	5 6	344	2.9
Total	21,463		694	3.2
			$\mathbf{p} = 0$.0006

Kyle et al., New Engl J Med, 2006, 354:1362



MGUS: Olmsted County, Minnesota

M-protein

P1118008-35

Age	No.	No.	%
50 – 59	8,373	141	1.7
60 – 69	6,019	178	3.0
70 – 79	4,508	205	4.6
≥80	2,563	170	6.6
Total	21,463	694	3.2
≥ 70	7,071	375	5.3
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Prevalence of MGUS According to Age





Kyle et al., New Engl J Med, 2006, 354:1362

MGUS OLMSTED COUNTY MN

Conclusions:

- Prevalence rate remained almost constant throughout collection, suggesting that patients who frequently seek medical care are at little or no greater risk for MGUS than those who do not.
- The prevalence was 4-fold higher in persons ≥80 years of age than those age 50-59 years.



MGUS OLMSTED COUNTY MN

Conclusions (continued)

- The prevalence was 2-fold higher than from the literature in persons ≥50 years of age and almost twice that previously reported in persons ≥70 years of age.
- MGUS is one of the most common pre-malignant disorders in the general population ≥50 years of age.



MGUS Precedes MM

N = 77,469

N

71

Developed MM



MGUS Present Before DX MM



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Virtually all patients with multiple myeloma have a previously recognized Mprotein (MGUS).





How many patients with MGUS are recognized during clinical practice at **Mayo Clinic?**



MGUS: Prevalence in Olmsted County, MN vs Clinical Detection



CP1291127-6

Prevalence vs Detection of MGUS

	Prev	Prevalence		detected
Age	Actual (%)	Clinically detected (%)	Yes (%)	No (%)
50	1.3	0.1	8	92
60	2.3	0.3	13	87
70	3.8	8.0	21	79
80	5.8	1.9	33	67
90	7.4	2.3	31	69
≥50	3.2	0.7	22	78

MAYO CLINIC Kyle et al., Blood, 2007, 110:79a



How long has MGUS been present when it is recognized?



Duration of MGUS for A Patient At Age 70



CP1291127-11

Monoclonal Gammopathy of Undetermined Significance (MGUS): Conclusions

- Only 22% with known MGUS have been recognized clinically
- 28% recognized as MGUS at age 70 have had it >20 years
- Median duration of MGUS prior to its recognition = 11 years



Monoclonal Gammopathy of Undetermined Significance Natural History in 241 Cases

241 patients with a protein in the serum but initially no evidence of multiple myeloma, macroglobulinemia, amyloidosis, or lymphoma 1956-1970 were followed up.

Am J Med 64:814, 1978



MGUS

Status at Follow-Up 1-39 Years (241 Cases)

		Follow Person-yea Media	-up rs 3,579 an
oroup	Description	No.	%
1	No substantial increase of M-protein (benign)	14	6
2	Increase M-protein (≥3 g/dL)	25	10
3	Died of unrelated causes	138	57
4	Development of myeloma, macroglobulinemia, amyloidosis, etc	64	27
Fotal		241	100

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MGUS

Development of Myeloma or Related Disease in 64 Patients with MGUS

			Inter disea	val to se (yr)	
	No.	%	Median	Range	
Multiple myeloma	44 *	69	10.6	1-32	
Macroglobulinemia	7	11	10.3	4-16	
Amyloidosis	8	12	9.0	6-19	
Lymphoproliferative disease	5	8	0.8	4-19	
Total *Dx of myeloma made after	<mark>64</mark> 20-yr F-L	<mark>100</mark> J in 10 pt	10.4	1-32	

Mayo Clinic Proceed 79:859, 2004 Kyle et al.





Mayo Clinic Proceedings 2004; 79:859, Kyle et al.



Conclusion

All patients with an apparently benign monoclonal gammopathy must be followed indefinitely



	MGUS SE Minnesota Jan 1, 1960-Dec 31,	1994	
	n=1,384		
	Male (%)	54	
	Age (med years)	72	
	<40 years (%)	1.7	
	M-spike (g/dL-med)	1.2	
MAYO CLINIC MAYO CLINIC	Kyle, et al., New Engl J Med, 346:564, 2002		CP1118008-24

MGUS SE MINNESOTA

Duration of Follow-up Person Years 11,009 0-35 **Range years Median years** 15.4 **Deaths (70%)** 963



MGUS SE MINNESOTA

Relative Risk of Progression

	Obs	Exp*	RR
Multiple Myeloma	75	3	25
Lymphoma	19	7.8	2.4
Amyloidosis	10	1.2	8.4
Macroglobulinemia	7	0.2	46
CLL	3	3.5	0.9
Plasmacytoma	1	0.1	8.5
Total	115	15.8	7.3

* Iowa SEER Registry

MAYO CLINIC MAYO CLINIC Kyle, et al., New Engl J Med, 346:564, 2002



Full Progression or Death



CP971723-6

Relative Risk of Full Progression by Serum M-Spike Size



CP999081-2

MGUS and Free Light Chain (FLC)

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		RR	Risk	of Prog
	Ν	95% CI	2	0 yr
				%
			Absolute	Competing risk
M-protein < 1.5 g/dl, IgG, Normal FLC	449	1	5	2
1 risk factor, abn	420	5.4	21	10
2 risk factors, abn	226	10.1	37	18
3 risk factors, abn	53	20.8	58	27

Rajkumar, et al., Blood; 106:1148, 2005



MGUS SE Minnesota Summary

- MGUS patients more likely to die of unrelated disease than to progress
- Myeloma accounts for 65% of progression
- Risk of progression is 1%/year
- Risk of progression associated with size and type of M-protein and FLC

MGUS SE Minnesota Conclusion

All MGUS patients must be monitored forever



Smoldering Multiple Myeloma

- Serum M-spike ≥ 3 g/dl and/or
- Bone marrow plasma cells ≥ 10%
- No end organ damage

Kyle RA and Greipp PR, NEJM, 302:1347, 1980.



Smoldering Multiple My	elom	<u>3</u>
Mayo Clinic 1970 – 1	994	
	Ν	%
Serum M-protein ≥ 3 g/dl and Bone marrow plasma cells ≥ 10%	106	38
Serum M-protein < 3 g/dI and Bone marrow plasma cells ≥ 10%	143	52
Serum M-protein ≥ 3 g/dl and	27	10
Bone marrow plasma cells < 10%		
MAYO CLINIC Kyle et al., NEJM 356:2582, 2007	276	100

Smoldering Multiple Myeloma

Progression

	N	%	Expected No. Pts	R.R.
Multiple myeloma	157	57	0.3	522
Primary amyloid (AL)	5	2	0.1	50
Total	162	59		

Kyle et al., NEJM 356:2582, 2007



Smoldering Multiple Myeloma

Time to progression	Median years	% progression at 15 years
Serum M-spike ≥ 3 Bone marrow plasma cells ≥ 10	2	87
Serum M-spike < 3 Bone marrow plasma	8	70
cells ≥ 10		
Serum M-spike ≥ 3 Bone marrow plasma cells < 10	19	39
Total (N = 276) p= <0.0	001 5	73
	356:2582, 2007	

Progression to Multiple Myeloma or Amyloid



Progression to MM or AL



Smoldering Multiple Myeloma Role of Free Light Chain (FLC)

N Relative Risk

- Serum M protein ≥ 3 g/dL & BMPC ≥ 10%
 - FLC ratio < 0.125 or > 8 78 2.06

Serum M protein < 3 g/dL & BMPC ≥ 10%

FLC ratio < 0.125 or > 8 82 1.72

Dispenzieri et al: Blood 111:785, 2008





