

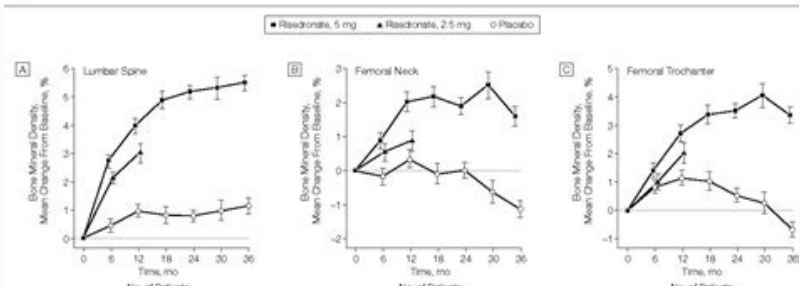
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Risedronate davis pdf



	N50 (n=10)					N50 (n=10)					N50 (n=10)				
Residual F _{1,9}	730	266	233	405	405	766	272	140	354	457	738	272	140	354	457
Residual F _{2,9}	340	207	443	305	305	340	207	443	305	305	340	207	443	305	305
Error	762	273	237	446	446	762	273	237	446	446	762	273	237	446	446

After each experiment SS, MM, the data of 6, 12, 18, and 30 months were performed at a subset of study centers. In A, P < .05 vs baseline for all time points and groups except the placebo group at 6 months; P < .05 vs placebo for all time points in the 2.5-mg and 5-mg-metastudies groups. In B, P < .05 vs baseline for all time points and groups except the placebo group between 24 and 30 months; P < .05 vs placebo for all time points for the 5-mg-metastudy group and at all time points for the 2.5-mg group. In C, P < .05 vs baseline for all time points and groups except the placebo group at 30 months; P < .05 vs placebo for all time points in the 5-mg-metastudy group except 6 months.

Upper GI tract adverse event	NA (%) of patients			
	Fleicos (n=508)	Rosuvastatin, 5 mg (n=9023)	RR (95% CI)	P value ^a
Any upper GI tract adverse event	1492 (29.6)	1497 (28.0)	1.01 (0.94-1.07)	1
Dyspepsia	170 (3.3)	148 (2.8)	0.98 (0.82-1.21)	0.94
Abdominal pain	462 (9.2)	664 (9.2)	1.01 (0.88-1.14)	>99
Altered bowel habit	453 (9.0)	453 (9.0)	1.01 (0.88-1.14)	>99
Diarrhoea	170 (3.3)	166 (3.3)	0.98 (0.82-1.21)	0.94
GI tract disorder	1914 (37.9)	1382 (27.7)	0.71 (0.64-0.79)	<23
Heartburn	282 (5.6)	115 (2.3)	0.41 (0.35-0.47)	<27
Ischaemia	92 (1.8)	86 (1.7)	0.94 (0.74-1.26)	71
Ischaemic colitis	51 (1.0)	110 (2.1)	0.47 (0.34-0.64)	<23
Stomach ulcer	41 (0.8)	48 (1.0)	1.18 (0.78-1.78)	46
GI tract haemorrhage	27 (0.5)	45 (0.9)	1.22 (0.78-1.90)	46
Stomach ulcer haemorrhage	18 (0.4)	18 (0.4)	1.00 (0.55-1.82)	>99
Subacute chest pain	16 (0.3)	18 (0.4)	1.01 (0.49-2.07)	>99
Stomach ulcer perforation	2 (0.0)	17 (0.3)	0.32 (0.03-5.54)	>99
Stomach ulcer	9 (0.2)	13 (0.3)	0.82 (0.39-1.73)	>99
Stomach ulcer haemorrhage	11 (0.2)	6 (0.1)	0.52 (0.17-1.53)	24
Stomach ulcer	7 (0.1)	9 (0.2)	1.02 (0.42-2.11)	>99
Stomach ulcer haemorrhage	7 (0.1)	8 (0.2)	1.15 (0.42-3.17)	>99
Stomach ulcer	7 (0.1)	8 (0.2)	1.02 (0.42-2.45)	>99
GI tract haemorrhage	2 (0.0)	2 (0.0)	0.45 (0.03-5.77)	22
Stomach ulcer haemorrhage	3 (0.1)	3 (0.1)	1.00 (0.24-4.29)	>99
GI tract haemorrhage	3 (0.1)	2 (0.0)	0.40 (0.02-6.45)	>99
Stomach ulcer haemorrhage	3 (0.1)	2 (0.0)	0.40 (0.02-6.45)	>99
Stomach ulcer	3 (0.1)	0 (0.0)	NA	>99
Stomach ulcer perforation	3 (0.1)	0 (0.0)	NA	>99
Stomach ulcer	3 (0.1)	0 (0.0)	NA	>99

†The Cox regression model was used to estimate the RR of patients with any upper GI tract adverse event between treatment groups. The Fisher exact test was used to compare the treatment groups with respect to the percentages of patients who experienced individual upper GI tract adverse events.

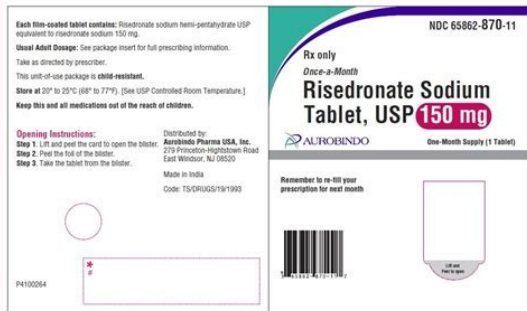
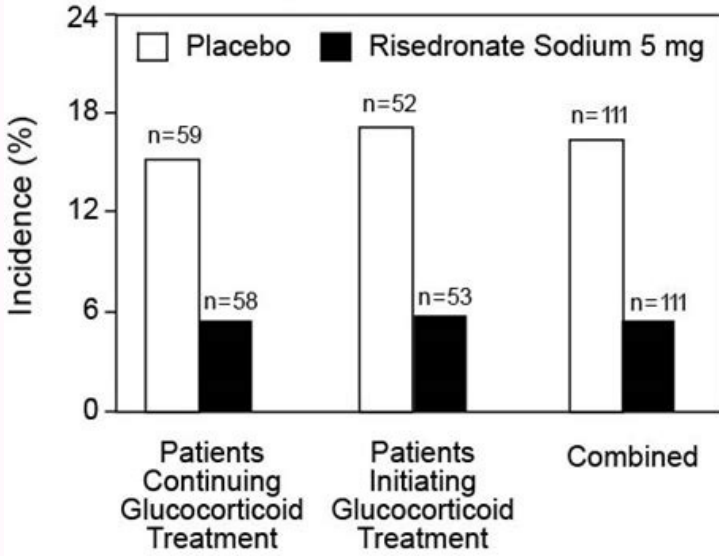


Figure 5
Incidence of Vertebral Fractures in Patients
Initiating or Continuing Glucocorticoid Therapy



Intraday precision	Concentration found (µg/mL)	Recovery (%)	Concentration added (µg/mL)
	3.974	99.35	4.0
	5.929	98.81	6.0
	8.030	100.40	8.0
		99.52 ± 0.81	
		0.81	
		0.47	
	0.995	99.50	1.0
	4.054	101.35	4.0
	5.968	99.47	6.0
		100.11 ± 1.08	
		1.08	
		0.62	

of three separate assays.

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06/01/2021 : But highly educated individuals also can be tripped up by poorly written directions, says study coauthor Terry Davis, professor of medicine at Louisiana State University Health Sciences Center at Shreveport. Take dosing recommendations as one example, she says. "If it's three times a day, that doesn't tell you when to take it," Davis says. Learn more about VITAMIN D uses, effectiveness, possible side effects, interactions, dosage, user ratings and products that contain VITAMIN D. Enter the email address you signed up with and we'll email you a reset link. Evert AB, Boucher JC, Cypress M, Dunbar SA, Franz MJ, Mayer-Davis EJ, Neumiller JJ, Nwankwo R, Verdi CL, Urbanski P, Yancy WS Jr. Nutrition therapy recommendations for the management of adults with diabetes. Diabetes Care 2013;36:3821-42. [PubMed abstract] Lima MDL, Cruz T, Pousada JC, Rodrigues LE, Barbosa K, Canguçu V.

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