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AIM AND OBJECTIVES



To establish patients' adherence to evolocumab therapy, a Protein convertase subtilisin/kexin type 9 (PCSK9) inhibitor, and to analyse the reduction of patients' LDL cholesterol (LDL-C) levels.

MATERIALS AND METHODS





N = 139		Group of adherence	Reduction percentage after 12 weeks (IQR)	Patients with at least 50% reduction (%)
79 males (57.25%) Age: 62.97 v.o. (TOR 15.53)		1	-69.18 (26.69)	71 (78.79%)
Posology: 140 mg/2 weeks (100%)		2	-68.64 (28.89)	23 (76.67%)
		3	-54.56 (44.69)	11 (61.11%)
ADHERENCE DISTRIBUTION Existing literature data				
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13%		Phase III	clinical trial (N)	Reduction percentage (CI 95%)
13%	► >90% [1]	Phase III 20110114	clinical trial (N) MENDEL-2 (614)	Reduction percentage (CI 95%) -58 (-60, -55)
13% 22%	 >90% [1] 75% - 90% [2] 	Phase III 20110114 20110115 L	clinical trial (N) MENDEL-2 (614) APLACE-2 (1896)	Reduction percentage (CI 95%) -58 (-60, -55) -64 (-66, -62)
13% 22% 65%	>90% [1] 75% - 90% [2] <75% [3]	Phase III 20110114 20110115 L 20110117 RL	clinical trial (N)MENDEL-2 (614)APLACE-2 (1896)JTHERFORD2 (329)	Reduction percentage (CI 95%) -58 (-60, -55) -64 (-66, -62) -63 (-66, -59)

CONCLUSIONS



- Low adherence seems to decrease LDL-C reduction capacity
 - ✓ These results would support the possibility of decreasing the frequency of administration, favouring the adherence to treatment and reducing costs

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