

Supplementary material

Table 1. Functional imaging techniques.

Method	Pros	Cons	Source
99mTc-MDP bone scintigraphy	Easily available. Can help define a region for a biopsy.	Non-specific and low sensibility: incomplete visualization of deposits; not possible to distinguish periarticular amyloid deposition and arthropathies of different origins.	Sethi 1990 Floege 2001 Ketteler 2001
Gallium-67 SPET	Detection of inflammatory lesions. Helpful for differentiating active from inactive deposits and for distinguishing articular-periarticular lesions from bone lesions.	These techniques label inflammatory changes and are not specific for amyloid-induced arthropathy.	
Thallium-201 SPET			
SAP scintigraphy (123I labelled)	The tracer accumulates in B2M amyloid deposits.	Uncertainties about specificity and sensibility: low accumulation in hips and shoulders. Spleen frequently labelled, while it is usually spared by B2M amyloidosis.	Tan 1999
β_2 m scintigraphy: β_2 m labelled 131I β_2 m labelled 111In rh β_2 m labelled 111In	High specificity demonstrated by tracer enrichment in amyloid fibrils. Good correlation between clinical/radiological and scan findings. In comparison with 131I : improved image quality and sensitivity. reduction in radiation exposure. -Safe and stable protein source -Scan specificity maintained -Enhanced sensibility -Lower radiation exposure	Elevated radiation exposure and low resolution of deposits in small joints. Not reliable in the presence of residual renal function.	Schäffer 2000 Ketteler 2001
FDG-PET	Can demonstrate areas of amyloid deposition through peri-articular inflammatory changes. Widely available.	Non-specific, does not allow discrimination between DRA and other causes of periarticular and articular inflammation.	Kecler-Pietrzik 2014 Piccoli 2017

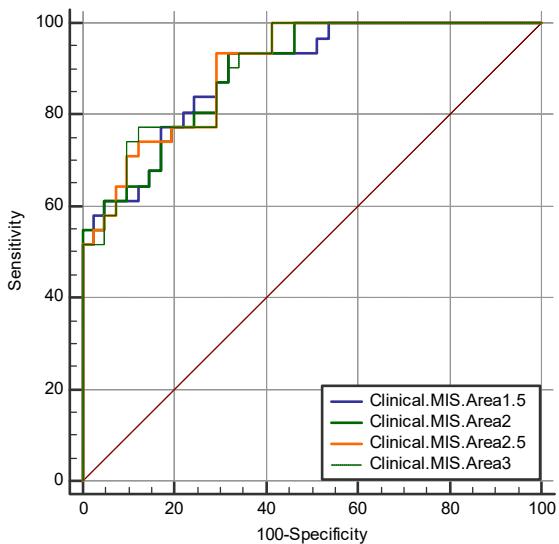
Supplementary Table 1: scintigraphic and nuclear medicine imaging methods employed for the detection of dialysis-related amyloidosis.

References

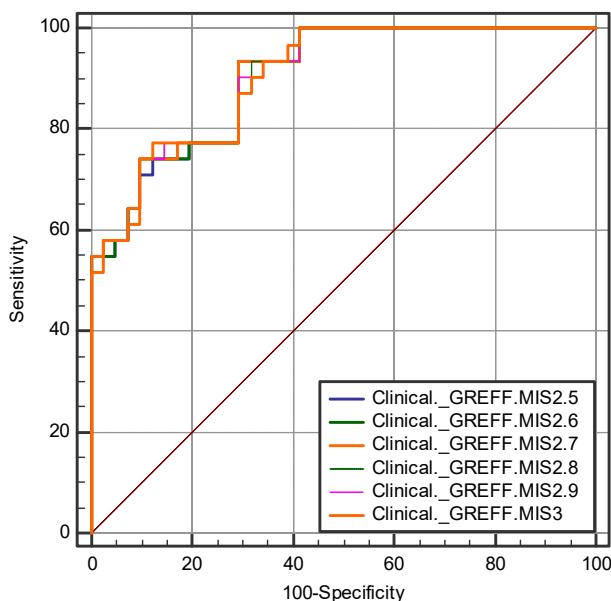
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Supplementary Figures



Variables	AUC (CI)
Clinical.MIS.Area1.5	0.895 (0.800 - 0.955)
Clinical.MIS.Area2	0.896 (0.802 - 0.956)
Clinical.MIS.Area2.5	0.903 (0.810 - 0.960)
Clinical.MIS.Area3	0.904 (0.811 - 0.961)



Variables	AUC (CI)
Clinical.GREFF.MIS2.5	0.903 (0.810 - 0.960)
Clinical.GREFF.MIS2.6	0.904 (0.811 - 0.961)
Clinical.GREFF.MIS2.7	0.906 (0.813 - 0.962)
Clinical.GREFF.MIS2.8	0.904 (0.811 - 0.961)
Clinical.GREFF.MIS2.9	0.903 (0.810 - 0.960)
Clinical.GREFF.MIS3	0.904 (0.811 - 0.961)

Figure S1. a) Empirical identification of the combined score. The MIS, multiplied by different factors, is added to the clinical amyloid score; b) Empirical identification of the combined score.

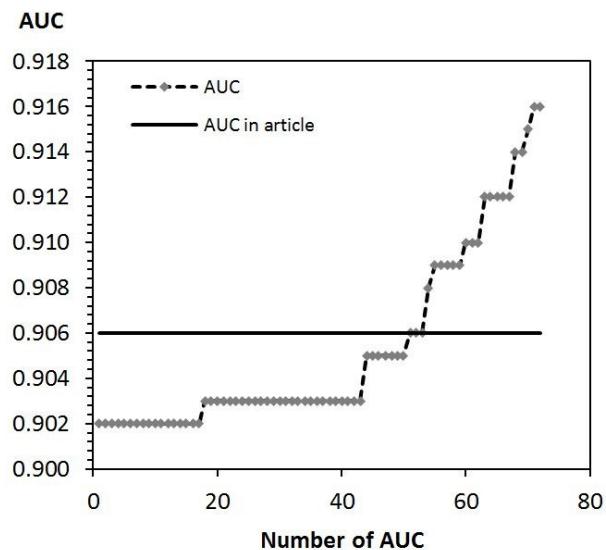


Figure S2. Leave-one-out validation.