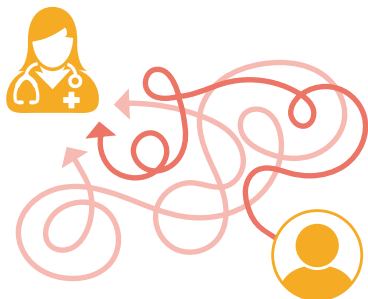
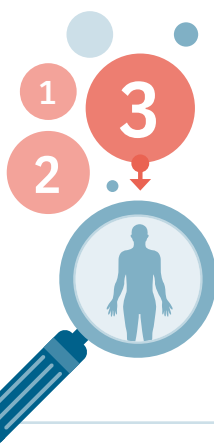


THE AL AMYLOIDOSIS DIAGNOSIS CHALLENGE

DID YOU KNOW?...



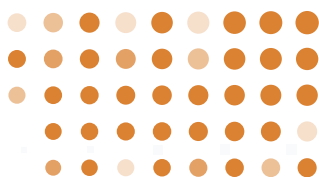
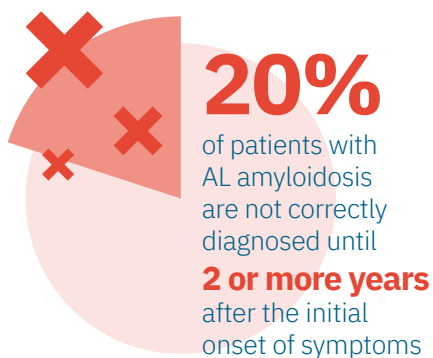
Patients with AL amyloidosis may experience **lengthy** and **confusing** pathways to diagnosis



On average, patients report seeing at least **3 different types of doctor** before being diagnosed



It can take over **1 year** and many visits to healthcare professionals before a patient is diagnosed

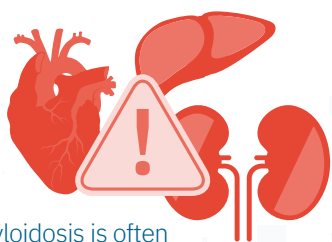
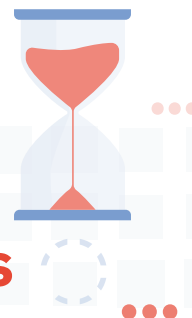


AL amyloidosis is a rare disease, meaning awareness is low

The median time from first consultation to diagnosis is

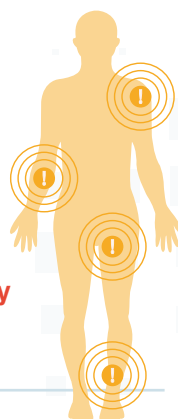
441 days

(according to a study in the Netherlands)



AL amyloidosis is often not diagnosed or recognised until **irreversible organ damage** has occurred

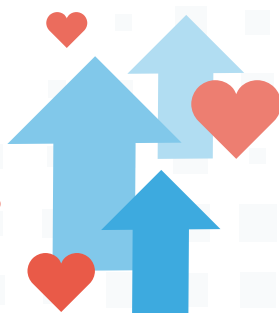
Symptoms of AL amyloidosis often develop slowly, are non-specific, vary widely and can present anywhere in the body (except the brain), **making it easy to suspect a more common condition**



Early diagnosis of AL amyloidosis enables **earlier treatment and prevents further organ damage**



Earlier diagnosis can **improve survival, quality of life and emotional well-being of AL amyloidosis patients**



MP^e
Myeloma
Patients
Europe

YEAR OF ACTION ON
DIAGNOSIS