



## Strength In Numbers: Facts and figures about ANCA-associated vasculitis (AAV)

AAV is a rare, severe condition affecting small blood vessels...<sup>1-4</sup>

**3** types of AAV:<sup>3</sup>

**GPA**  
**MPA**  
**EGPA**



**1 in 10,000** people  
affected by AAV  
in Europe<sup>5,6</sup>



**60%**  
**MALE**<sup>5,7</sup>



**40%**  
**FEMALE**<sup>5,7</sup>

Average age of  
diagnosis: **57 years**<sup>8</sup>

...impacting a range of areas in your body, making diagnosis difficult.<sup>8,9</sup>

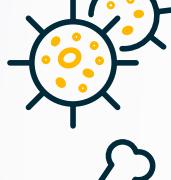


**9** different organs  
or body parts that  
can be affected<sup>10</sup>



**6** month delay in  
diagnosis is experienced  
by 1/3<sup>rd</sup> of patients<sup>9</sup>

Serious health issues are caused by disease and treatment...<sup>11-14</sup>



**7x** higher risk  
of infection<sup>11</sup>



**65%** increased  
cardiovascular risk<sup>13</sup>



**8x** higher risk  
of osteoporosis<sup>12</sup>



**26%** of patients have  
severe kidney issues  
after 3 years<sup>14</sup>

...and AAV has a big impact on quality of life...<sup>15,16</sup>



**20%** of working age AAV  
patients were unemployed  
due to their condition<sup>16</sup>



**50%** of AAV patients thought  
their careers were hindered  
due to their condition<sup>16</sup>

...but support is available for anyone affected by AAV

**7** patient  
association  
groups across  
Europe



# #StrengthInNumbers

ANCA, anti-neutrophil cytoplasmic antibody; EGPA, eosinophilic granulomatosis with polyangiitis; GPA, granulomatosis with polyangiitis; MPA, microscopic polyangiitis.

References

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