

DISEASE STABILISATION IS IMPORTANT TO PEOPLE LIVING WITH SMA



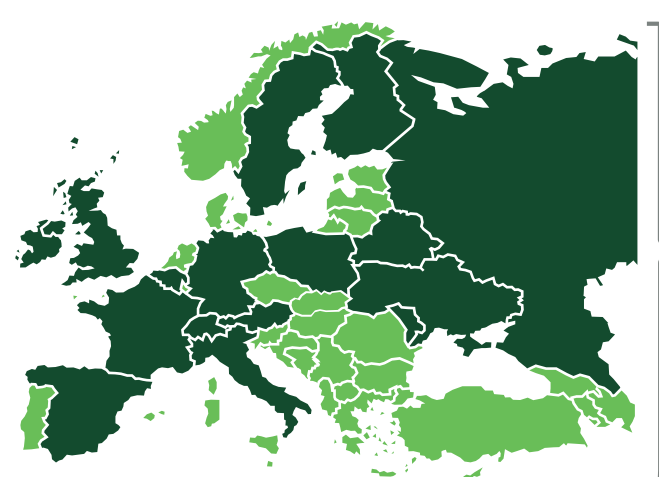
WITHOUT INTERVENTION, PEOPLE LIVING WITH SPINAL MUSCULAR ATROPHY (SMA) WILL SEE PROGRESSION OF THEIR DISEASE OVER TIME...^{1,2}



...SO IT'S IMPORTANT TO **STABILISE AND MAINTAIN MOTOR FUNCTION** FOR AS LONG AS POSSIBLE.

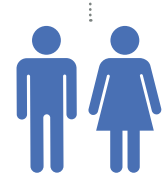
HERE IS WHAT PEOPLE LIVING WITH SMA AND THEIR CAREGIVERS THINK...

16 COUNTRIES



822 PARTICIPANTS

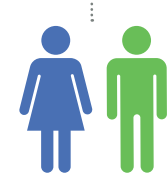
52% (425) OF THE REPRESENTED PEOPLE LIVING WITH SMA WERE ADULTS AGED 20 AND ABOVE



436 PEOPLE LIVING WITH SMA



383 PARENTS/ CAREGIVERS



3 UNDISCLOSED

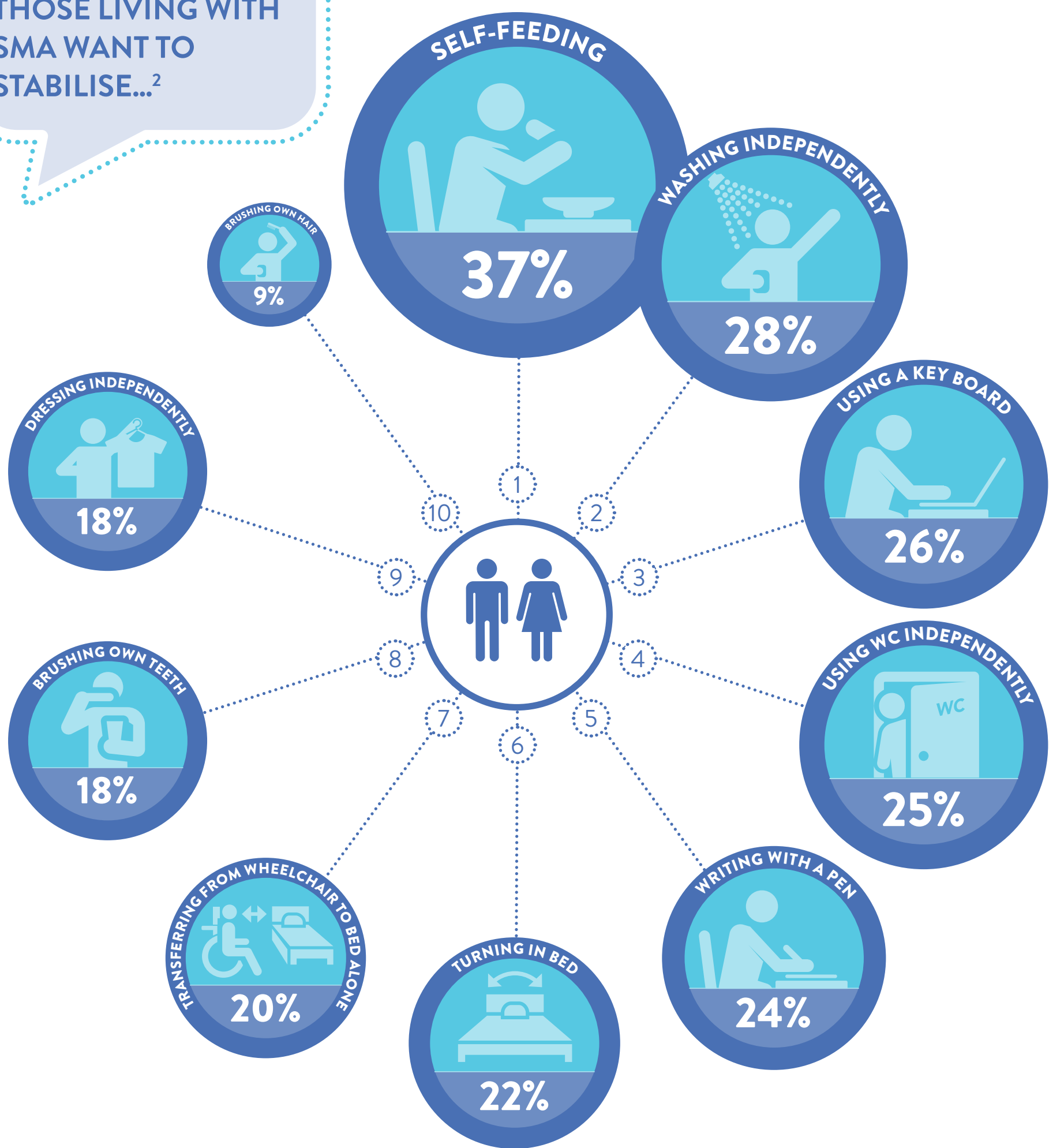


A recent survey of people affected by type II/III SMA described the impact on their quality of life and their expectations relating to care options²

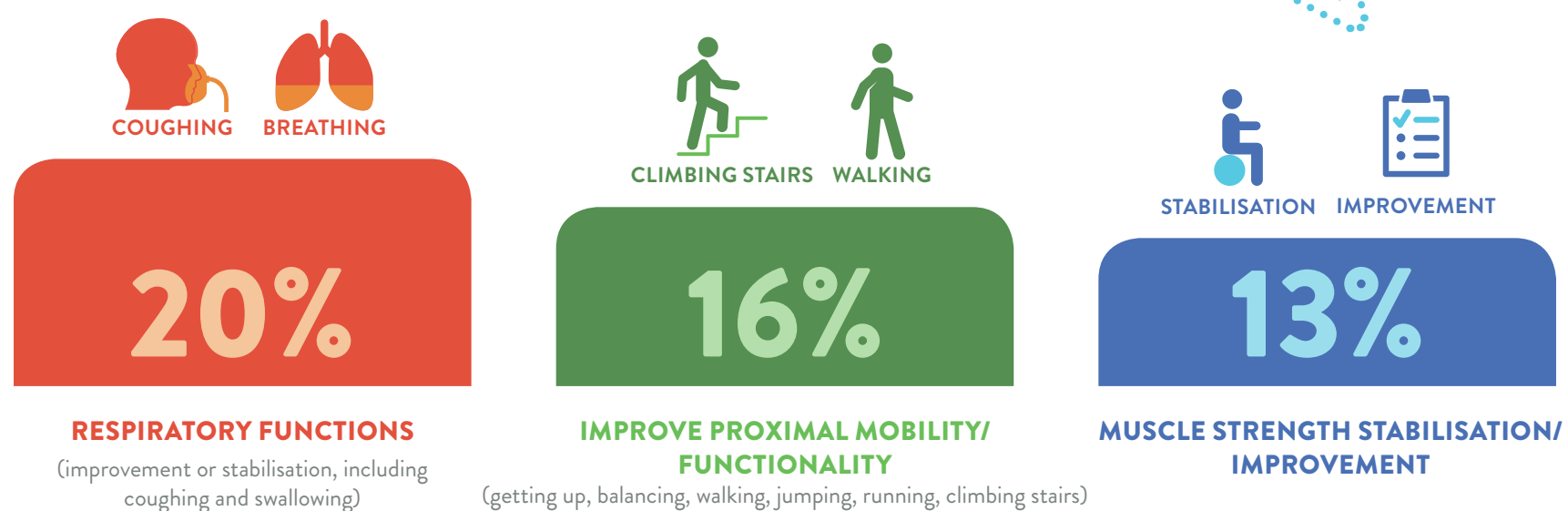
DISEASE STABILISATION IS IMPORTANT TO PEOPLE LIVING WITH SMA. SEE WHAT IT MEANS TO THEM...²

81% OF PARTICIPANTS FELT THAT WOULD REPRESENT MAJOR PROGRESS

SEE WHICH ABILITIES THOSE LIVING WITH SMA WANT TO STABILISE...²



THEIR PERSONAL EXPERIENCE WITH SMA DICTATES WHAT THEY WANT OUT OF CARE. SEE WHAT THEY FOUND IMPORTANT...²



TALK TO YOUR DOCTOR TODAY!
TO DISCUSS OPTIONS FOR DISEASE STABILISATION.

