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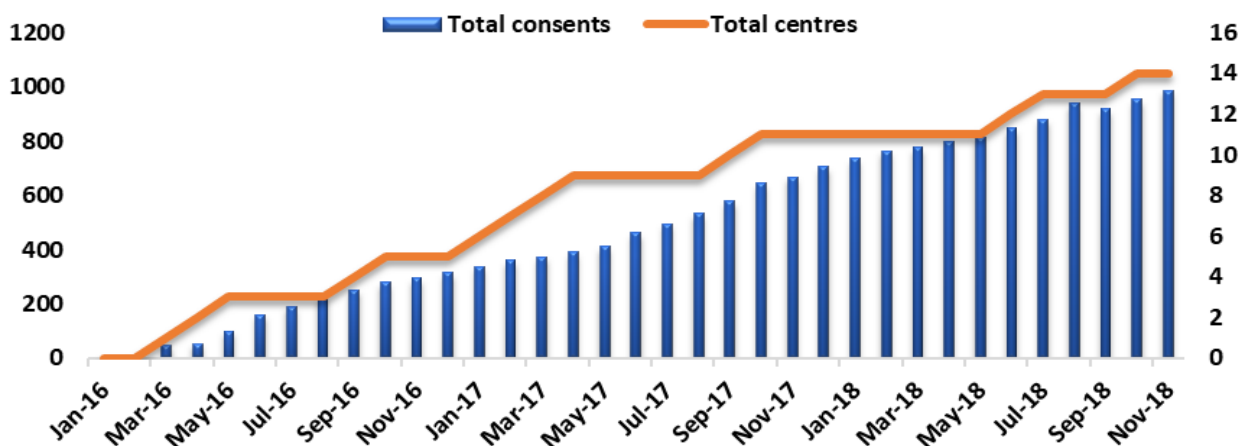
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The MND Register is a population register which aims to collect information about every person with MND in England, Wales and Northern Ireland. Recording every person with MND allows us to work out the number of people diagnosed with MND per year, how many people currently have the disease and how this is changing with time.

MND Register participants and recruiting centres



Results so far...

Preliminary findings were recently presented at the 29th MNDA International Symposium in Glasgow. The symposium is the largest medical and scientific conference for MND where the latest advances in research are presented. Results from the SEALS (South East ALS) register were compared with some of our data. Results have found a consistency between the two registers. This suggests our current data collection method is viable. We will continue to expand and collect data for people living with MND in England, Wales and Northern Ireland.

Data collection period		2015-2018
Total number of cases analysed		655
Mean age of onset (SD)		62.60 yrs (11.90)
Sex ratio F:M		1:1.5 (261:394)
Site of onset- area of the body that symptoms where first noticed.	Bulbar (speech problems)	24% (154 people)
	Spinal (weakness in a limb)	66% (434 people)
	Respiratory (breathing problems)	1% (5 people)
	Other	2% (16 people)
	Missing	7% (46 people)
MND subtype -type of MND.	ALS	90% (587 people)
	PLS	6% (43 people)
	PMA	1% (8 people)
	Other	2% (17 people)

