

Thank you for completing this important information about the way you are conducting outreach for the National ALS Registry. Please submit a form monthly for your chapter or district.

Org Code

District or Chapter Name

Clinic, Support Group, Seminar, Other group activities

	Activity	Date	# Registry information kits distributed	# people talked to about the Registry	# people helped sign-up for the Registry	City and State
Total		X				X

Social Media and local mailings (e.g., tweet, email blast, newsletter)

	Activity	Date	#	City and State
Total		X		X

Fundraising event (e.g., walk, golf tournament)

	Type of event	Date	# attendees	Materials distributed or displayed	City and State
Total		X			X

Outreach calls made where the Registry was discussed

# calls	City and State

Total

	X

CDC estimates the average public reporting burden for this collection of information as 5 minutes per response, including the time for reviewing instructions, searching existing data/information sources, gathering and maintaining the data/information needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0923-0041).