SHINE PURPLE FOR A CURE

FOR RETT SYNDROME

RETT SYNDROME IS A RARE GENETIC NEUROLOGICAL DISORDER

1 IN 10,000 ARE AFFECTED BY RETT

CAUSED BY A MUTATION OF THE MECP2 GENE ON THE X CHROMOSOME

THE HALLMARK OF RETT IS NEAR CONSTANT REPETITIVE HAND MOVEMENTS

THERE IS CURRENTLY NO CURE

WWW.RETT.CA

ORSA since 1991 Ontario Rett Syndrome Association Building Healthy Tomorrows

OCTOBER 29TH #LightCanadaPurple RETT SYNDROME AWARENESS