OCTOBER

SHINE A LIGHT ON 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

VISIT
WWW.RETT.CA

ORSA since 1991
Ontario Rett Syndrome Association
Building Healthy Tomorrows

We are lit Purple to Support RETT SYNDROME