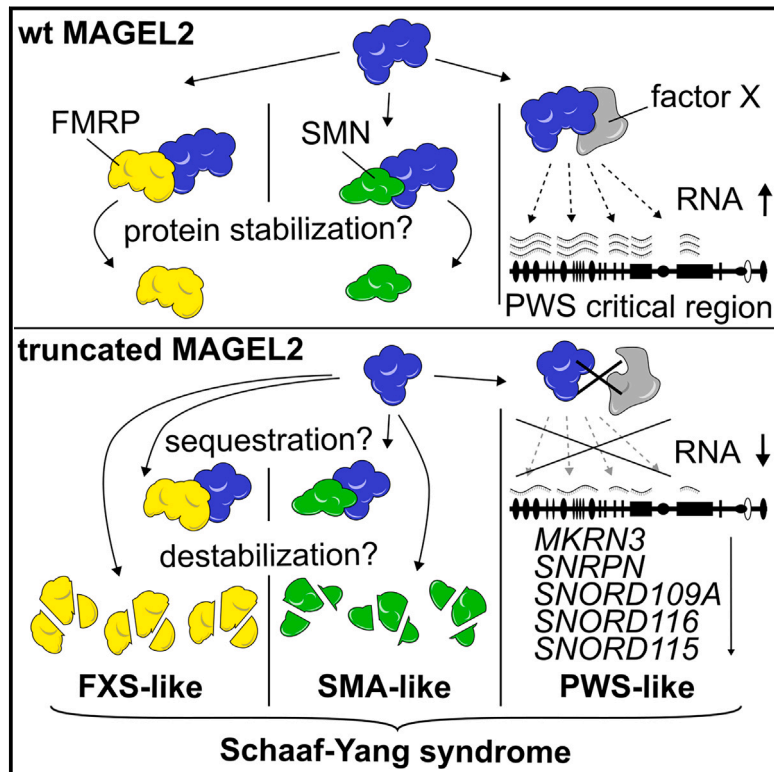


# Truncated variants of MAGEL2 are involved in the etiologies of the Schaaf-Yang and Prader-Willi syndromes

## Graphical abstract



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**Mutations within *MAGEL2* from chromosomal region 15q11–q13 cause Schaaf-Yang syndrome, which is phenotypically related to Prader-Willi syndrome, caused by deletion of the *SNORD116* cluster within the same locus. We correlate mutations within *MAGEL2* to spinal muscular atrophy and autism and also demonstrate its influence on the abundance of *SNORD116*.**

# Truncated variants of MAGEL2 are involved in the etiologies of the Schaaf-Yang and Prader-Willi syndromes

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## Summary

The neurodevelopmental disorders Prader-Willi syndrome (PWS) and Schaaf-Yang syndrome (SYS) both arise from genomic alterations within human chromosome 15q11–q13. A deletion of the *SNORD116* cluster, encoding small nucleolar RNAs, or frameshift mutations within *MAGEL2* result in closely related phenotypes in individuals with PWS or SYS, respectively. By investigation of their subcellular localization, we observed that in contrast to a predominant cytoplasmic localization of wild-type (WT) *MAGEL2*, a truncated *MAGEL2* mutant was evenly distributed between the cytoplasm and the nucleus. To elucidate regulatory pathways that may underlie both diseases, we identified protein interaction partners for WT or mutant *MAGEL2*, in particular the survival motor neuron protein (SMN), involved in spinal muscular atrophy, and the fragile-X-messenger ribonucleoprotein (FMRP), involved in autism spectrum disorders. The interactome of the non-coding RNA *SNORD116* was also investigated by RNA-CoIP. We show that WT and truncated *MAGEL2* were both involved in RNA metabolism, while regulation of transcription was mainly observed for WT *MAGEL2*. Hence, we investigated the influence of *MAGEL2* mutations on the expression of genes from the PWS locus, including the *SNORD116* cluster. Thereby, we provide evidence for *MAGEL2* mutants decreasing the expression of *SNORD116*, *SNORD115*, and *SNORD109A*, as well as protein-coding genes *MKRN3* and *SNRPN*, thus bridging the gap between PWS and SYS.

## Introduction

Prader-Willi syndrome (PWS [MIM: 176270]) and Schaaf-Yang syndrome (SYS [MIM: 615547]) are two related rare genetic neuro-developmental disorders emerging from mutations in the maternally imprinted locus on paternal chromosome 15q11–q13.<sup>1</sup> This region harbors paternally expressed genes exclusively, i.e., *MKRN3* (MIM: 603856), *MAGEL2* (MIM: 605283), *NDN* (MIM: 602117), *PWRN1* (MIM: 611215), *NPAP1* (MIM: 610922), *SNRPN* (MIM: 182279), *SNHG14* ([MIM: 616259]; its introns contain *SNORD107*, *SNORD64*, *SNORD108*, *SNORD109A*, *SNORD109B*, *SNORD115*, and *SNORD116*), and *IPW* (MIM: 601491).<sup>2</sup>

PWS appears in 1 in 16,000 to 1 in 21,000 live births<sup>3</sup> and is caused by either (1) a 5–6 Mb *de novo* deletion within the paternal Prader-Willi critical region (65%–75% of the cases), (2) maternal uniparental disomy 15 (20%–30% of the cases), or (3) imprinting center defects (1%–3% of the cases).<sup>1,4</sup> Clinical features of PWS are developmental delay, hypotonia, endocrine dysfunction, intellectual disability, and in particular hyperphagia resulting in severe obesity.<sup>5</sup>

Recent studies showed that micro-deletions of 71–200 kb in size, encompassing the entire *SNORD116* cluster

only, are sufficient to cause PWS.<sup>6–10</sup> *SNORD116* belongs to the class of box C/D small nucleolar RNAs (snRNAs). Together with all other snRNAs from the PWS region, *SNORD116* was initially discovered in our lab by an RNA-seq approach for non-coding RNAs for which we coined the term RNomics.<sup>11,12</sup> Canonical box C/D snRNAs guide 2'-O-methylation of precursor ribosomal and small nuclear RNAs within the nucleolus by base-pairing.<sup>13,14</sup> The majority of snRNAs are embedded in intronic regions of RNA Polymerase II transcribed protein-coding<sup>15,16</sup> or long non-coding RNA (lncRNA) genes.<sup>17,18</sup> Subsequent to intron splicing and debranching, processing and maturation of pre-snRNAs involves 5'- and 3'-end exonucleolytic trimming<sup>19,20</sup> and binding to snRNP proteins.<sup>17,21,22</sup>

Both *SNORD116* and *SNORD115* are uniquely organized within the PWS critical region in tandem repetitive multi-copy gene arrays of 30<sup>23</sup> and 48<sup>6</sup> copies, respectively, share high sequence homology within their cluster<sup>11</sup> and are post-transcriptionally processed from their long non-coding host *IC-SNURF-SNRPN* transcript.<sup>24</sup> In contrast to canonical snRNAs, we were able to demonstrate that both *SNORD116* and *SNORD115* snRNAs lack obvious sequence complementarity within their antisense element toward canonical target pre-rRNAs or snRNAs.<sup>11</sup>

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<https://doi.org/10.1016/j.ajhg.2024.05.023>

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Molecular functions of orphan mammalian *SNORD116* and its host gene still remain to be fully elucidated. Human PWS phenotypes were partially recapitulated in mice carrying a deletion of the paternal *Snord116* cluster.<sup>25,26</sup> Although these mice did not develop obesity, they showed retarded growth, failure to thrive, and reduced feeding.<sup>25,26</sup> In addition, hyperphagia, increased anxiety, and reduced motor learning were also observed in mice with PWS.<sup>25</sup> The flanking non-coding exons of the *SNORD116* copies were shown to be processed into a functional lncRNA, designated as lncRNA *HG116* which was suggested to regulate transcription of several target genes in a diurnal manner.<sup>27</sup> Furthermore, rhythmic methylation of CpG islands was disrupted in a *Snord116* deletion mouse model.<sup>28</sup>

SYS is a very rare disease with a prevalence of <1 in 1,000,000<sup>29</sup> and was originally described by Schaaf et al.,<sup>30</sup> when they reported four individuals with nonsense or frameshift indel mutations within *MAGEL2* who displayed PWS or PWS-like phenotypes. Individuals with SYS show developmental delay, mild to profound intellectual disability, neonatal hypotonia with respiratory distress and central apnea, joint contractures, and social behavioral characteristics.<sup>1,31–33</sup> Moreover, autism spectrum disorder or autistic features are diagnosed in 75%–80% of affected individuals as well as hyperphagia resulting in subsequent obesity which can develop as SYS-affected individuals get older.<sup>1,31–33</sup>

Up until now, several mutations within the single exon gene *MAGEL2* have been described with a correlation between the severity of phenotypes and the genotype of affected individuals.<sup>31,32,34</sup> Persons carrying a *MAGEL2* c.1996delC (p.Gln666Serfs\*36) mutation passed away during gestation or hours after birth.<sup>31,32,35</sup> The c.1996dupC is not only the most prevalent but also the *MAGEL2* mutation resulting in the most severe phenotype.<sup>32</sup>

*MAGEL2* (melanoma antigen L2) belongs to the MAGE protein family of RING E3 ubiquitin ligase regulators.<sup>36–38</sup> It was initially shown to specifically bind to the RING-type E3 ubiquitin transferase TRIM27 and the ubiquitin-specific protease USP7 to form a MAGE-RING E3 ligase complex, termed MUST, which is recruited by the retromer to early and recycling endosomes to facilitate F-actin polymerization.<sup>37–39</sup> However, additional protein interaction partners have also been identified, implicating *MAGEL2* in other biological functions.<sup>40</sup>

*Magel2*-null mice were shown to recapitulate some PWS phenotypes, such as growth retardation, excessive weight gain,<sup>41</sup> and differences in circadian regulation.<sup>42</sup> In addition, rats with paternal *Magel2* truncation exhibit changes in body composition, cardiac structure and function, breathing, as well as sociability and anxiety-like behavior, also observed in SYS-affected individuals.<sup>43</sup>

Because of their similar phenotypes as well as their common genomic location, in this study we investigated a potential functional cross-talk between both PWS and SYS. To this end, we first cloned WT *MAGEL2* as well as

mutated *MAGEL2* c.1996dupC (p.Gln666Profs\*47) (GenBank: NM\_019066.5) as mCherry- or GFP-fusion proteins, resulting in the WT and truncated p.Gln666Profs\*47 variant and addressed their subcellular localization. In addition, we attempted to identify novel protein interaction partners by either BioID (miniTurbo-fusion proteins) or co-immunoprecipitation (CoIP; GFP-fusion proteins). To identify potential common interaction partners between *MAGEL2* and *SNORD116*, we further analyzed the interactome of *SNORD116* by RNA-CoIP.

Our data showed an involvement of *MAGEL2* in biological processes such as RNA stabilization, chromosome condensation, and transcription. Based on these findings, we investigated the involvement of WT and truncated *MAGEL2* on the expression of selected genes from the PWS locus in small molecule neural precursor cells (smNPCs) derived from persons suffering from SYS and compared them to healthy control subjects. Interestingly, the expression of most genes within the PWS locus—including *SNORD116*—was found to be down-regulated in SYS cell lines carrying the *MAGEL2* c.1996dupC (see above) or the *MAGEL2* c.1802delC (p.Pro601Glnfs\*101) mutation (GenBank: NM\_019066). Taken together, our results thus suggest a functional link between SYS and PWS.

## Material and methods

### Cloning of reporter plasmids

Plasmids were designed with the Benchling software (<https://benchling.com>). Plasmid constructs were cloned by Gibson Assembly or mutagenesis PCR (NEB) and Oligonucleotides for Gibson Assembly and mutagenesis PCR were designed with the online NEBuilder Assembly Tool 2.0 and NEBaseChanger Tool (NEB), respectively. Gene fragments and oligonucleotides were purchased from IDT.

Human *SNORD116-3* (GenBank: LN847959) and human *SNORD100* (GenBank: LN847822) were designed with a T7 promoter sequence in the 5' end and flanked by BamHI and EcoRI restriction sites. snoRNA genes were amplified by PCR from gene fragments (IDT), digested by the respective enzymes, and ligated into a PUC19 vector. Single-exon *MAGEL2* (ENSG00000254585) was amplified by PCR from human genomic DNA. Due to the high GC-content within the coding sequence of *MAGEL2* (>70% in the proline-rich region), the PCR was prone to premature termination. To circumvent this issue, the sequence was divided into three equal parts for amplification. Amplicons were analyzed by gel electrophoresis, and PCR products with the correct length were gel excised and purified using the Monarch DNA Gel Extraction Kit (NEB).

The *MAGEL2* sequence was cloned into the Low0 vector backbone employing Gibson Assembly (NEB). A plasmid harboring the *MAGEL2* c.1996dupC mutation, resulting in the truncated *MAGEL2* p.Gln666Profs\*47 variant, was generated via Gibson Assembly. Furthermore, various plasmids for the expression of WT *MAGEL2* and p.Gln666Profs\*47 fusion proteins were generated. To this purpose, either mCherry, GFPspark, or miniTurbo<sup>44</sup> were cloned N-terminally to the respective *MAGEL2* variant by Gibson Assembly, thereby preserving the endogenous 3' UTRs of *MAGEL2* and *MAGEL2* c.1996dupC transcripts. Plasmids were transformed

into homemade chemically competent NEB 5- $\alpha$  *E. coli*, and plasmid DNA of selected clones was purified by the Monarch Plasmid Miniprep Kit (NEB) and sequenced by Eurofins Mix2Seq (Eurofins). Positive clones were re-cultured and DNA was extracted employing the QIAGEN Plasmid Midi Kit (QIAGEN).

### Cell culture

HEK293T (ATCC CRL-3216) or Neuro-2a (ATCC CCL-131) (N2a) cells were grown in pyruvate-depleted, 4.5 g/L glucose- and L-glutamine-containing DMEM medium (Gibco) supplemented with 100 U/mL penicillin/streptomycin and 10% heat-inactivated FCS (Gibco).

HeLa cells (HeLa ACC57, DSMZ) were cultured in 4.5 g/L glucose and L-glutamine containing DMEM medium (Sigma-Aldrich) supplemented with 100 U/mL penicillin/streptomycin and 10% FCS (Thermo Fisher Scientific).

Induced pluripotent stem cell (iPSC) lines were cultured on hESC-qualified Matrigel-coated plates (Corning) with StemMACS iPS-Brew (Miltenyi), passaged with Accutase (Sigma-Aldrich), and supplemented with 10  $\mu$ M Y-27632 (Miltenyi) for 24 h. The cells were cryopreserved in knock-out serum replacement (Thermo Fisher Scientific) supplemented with 10% DMSO (Roth) and stored at  $-80^{\circ}\text{C}$  until further usage.

smNPCs were cultured with N2B27 medium (1:1 mixture of DMEM/F12 and Neurobasal [Gibco] supplemented with 0.5% N-2 [Gibco], 1% B-27 minus Vitamin A [Gibco], 1 $\times$  GlutaMAX [Gibco], 3  $\mu$ M CHIR 99021 [Axon Medchem], 0.5  $\mu$ M Purmorphamine [Miltenyi], and 150  $\mu$ M ascorbic acid [Sigma]). The cells were passaged with Accutase (Sigma-Aldrich) and cryopreserved in knock-out serum replacement (Thermo Fisher Scientific) supplemented with 10% DMSO (Roth) and stored at  $-80^{\circ}\text{C}$  until further usage. All cell lines were maintained at  $37^{\circ}\text{C}$ , 5%  $\text{CO}_2$ , and saturated humidity culturing conditions.

### Healthy and Schaaf-Yang syndrome proband cell lines and iPSC reprogramming

Human control fibroblast lines AG, FCE02, FCE10, and FCE11 were obtained from the Coriell Institute for Medical Research. All the procedures were in accordance with the Office for Human Research Protections, Department of Health and Human Services ("DHHS") regulations for the protection of human subjects (45 CFR Part 46). Fibroblasts were cultured with fibroblast medium (DMEM [Thermo Fisher Scientific], 15% FBS [Thermo Fisher Scientific], 1 $\times$  non-essential amino acids [Sigma]). The medium was changed every second day and the cells were passaged using Trypsin-EDTA solution (Sigma). The cells were cryopreserved in 90% FBS supplemented with 10% DMSO (Roth) and stored at  $-80^{\circ}\text{C}$  until further usage. Fibroblasts were reprogrammed into iPSCs employing the CytoTune iPS 2.0 Sendai Reprogramming Kit (Thermo Fischer Scientific) according to the manufacturer's guidelines with some modifications. In brief, 40,000 fibroblasts were plated per 24 well a day prior to the viral transduction. The cells were transduced with the recommended viral multiplicity of infection (MOI). After 24 h the viral medium was removed and the medium was renewed every second day. On day 7 the cells were replated on hESC-qualified Matrigel-coated 6-well plates (Corning). From day 8 onwards, the fibroblast medium was replaced with StemMACS iPS-Brew (Miltenyi) and was renewed daily. Around day 12, iPSC putative colonies started to appear and once they reached the appropriate size they were manually isolated and transferred on hESC-qualified Matrigel-coated 48

wells (Corning). The colonies were further expanded, characterized, and eventually cryopreserved in knock-out serum replacement (Thermo Fisher Scientific) supplemented with 10% DMSO (Roth) and stored at  $-80^{\circ}\text{C}$  until further usage. The pluripotency of the cells was confirmed by immunofluorescence staining for pluripotency-associated markers and differentiation of the cells into the three germinal layers (STEMdiff Trilineage Differentiation Kit, Stemcell Technologies). Moreover, the absence of Sendai-associated transgene expression was confirmed via RT-qPCR.

Human SYS fibroblast lines were obtained from skin biopsies of SYS-affected individuals carrying the *MAGEL2* c.1802delC or *MAGEL2* c.1996dupC mutations and healthy control individuals (BA cell line). From respective individuals under human research protocol H-34578 of the Institutional Review Board of Baylor College of Medicine, Houston, USA, proper informed consent was obtained. Fibroblasts were reprogrammed into iPSCs by the Baylor College of Medicine Human Stem Cell Core (HSCC) using the CytoTune-iPS 2.0 Sendai Reprogramming Kit (Life Technologies), following the manufacturer's protocol. Following transduction, cells were grown in high-glucose DMEM supplemented with 10% fetal bovine serum and 1 $\times$  MEM non-essential amino acids (Life Technologies) for 5 days, in TeSR-E7 (StemCell) for 8 days, and in TeSR-E8 (StemCell) until day 21 when pluripotent colonies were manually picked. Immunofluorescence and RT-qPCR for pluripotency markers were performed on all iPSC lines as described before.<sup>45</sup> Each iPSC line was shown to have a normal karyotype.

### smNPC derivation

smNPC lines were derived from iPSCs following the protocol of Reinhardt et al.<sup>46</sup> Briefly, upon 50%–70% confluency, iPSCs were mechanically dissociated with 0.5 mM EDTA (Gibco) and transferred to low-adherence plates in order to form embryoid bodies. After two days, the media was changed to N2B27 medium (1:1 mixture of DMEM/F12 and Neurobasal [Gibco] supplemented with 0.5% N-2 [Gibco], 1% B-27 minus Vitamin A [Gibco], 1 $\times$  GlutaMAX with 10  $\mu$ M SB-431542 [MedChem Express], 1  $\mu$ M Dorsomorphin [Miltenyi], 3  $\mu$ M CHIR 99021 [Axon Medchem], and 0.5  $\mu$ M Purmorphamine [Miltenyi]). On day 4, media was changed to N2B27 medium supplemented with 3  $\mu$ M CHIR 99021, 0.5  $\mu$ M Purmorphamine, and 150  $\mu$ M Ascorbic acid (Sigma-Aldrich). On day 6, embryoid bodies were triturated with a pipette and seeded as a monolayer on Growth Factor Reduced Matrigel-coated wells (Corning). Subsequently, the cells were further expanded and homogenized by high passaging ratios. After 5–6 passages, the cell morphology appeared homogeneous and the cells were further maintained with N2B27 medium on Growth Factor Reduced Matrigel-coated wells. Successful induction of the neural precursor fate was validated by immunofluorescence stainings for NPC markers.

### Leptomycin B treatment

50,000 HeLa cells were seeded in a 12-well plate containing glass cover slips. 24 h after seeding, cells were transiently transfected with a plasmid encoding either Low0\_GFPspark\_MAGEL2 or Low0\_GFPspark\_MAGEL2 c.1996dupC using PEI (PolySciences). 48 h after transfection cells were left untreated or treated with 20 nM Leptomycin B (Cell Signaling) for 20 h. Cells were fixed with 4% paraformaldehyde, permeabilized with 0.1% Triton X-100, and stained for DAPI. Coverslips were mounted on microscopy slides using polyvinylalcohol mounting medium (Sigma-Aldrich). Images were acquired with a 40 $\times$  objective (Plan Apo  $\lambda$  40 $\times$ ) using a Spinning Disc Eclipse Ti-2 microscope.



Fiji was used for image analysis to determine the nuclear cytosolic ratio of Low0\_GFPspark\_MAGEL2 or Low0\_GFPspark\_MAGEL2 p.Gln666Profs\*47. DAPI staining was used to distinguish the nucleus from the cytosol. A region of interest was selected in the nucleus and a region of similar size was selected in the cytosol. The mean fluorescent intensity of these regions was measured and the ratio of nuclear to cytosolic signal was calculated. Data were statistically analyzed employing the Wilcoxon test.

### Immunofluorescence staining

15 mm VWR No.1 coverslips (VWR) were coated in 12-well plates (Greiner Bio-One) with Poly-D-Lysine (Sigma-Aldrich) for 5 min, washed twice with DPBS (Gibco), and dried for at least 2 h. Plates were optionally stored at 4°C up to two weeks. Wells were washed again with DPBS and 10<sup>5</sup> HEK293T or N2A cells were seeded 24 h before transfection. Per expression vector, 750 ng of DNA were transfected with 1.5 µL Metafectene Pro (Biontex).

24 h after transfection, the cultivation medium of the transfected HEK293T or N2A cells was aspirated and the coverslips were washed twice with DPBS. All subsequent steps were conducted in the dark. Cells were incubated in 4% paraformaldehyde in 1× cytoskeletal buffer (CB; 10 mM Pipes [pH 6.8], 150 mM NaCl, 5 mM EGTA [pH 9], 5 mM glucose, 5 mM MgCl<sub>2</sub>) for 8 min at RT and washed twice in 1× CB. At this point, cells were either stored in 1× CB at 4°C or immediately subjected to the immunofluorescence steps described below.

For EEA1 and RAB11A staining, the CB was discarded and cells were incubated in IF blocking buffer (2% gelatin [w/v], 50 mM NH<sub>4</sub>Cl, 1× CB, 0.025% Saponin [w/v]; supplemented with BSA) for at least 45 min at RT, followed by incubation with the primary antibody in IF blocking buffer for 1 to 2 h at RT. For stainings in HEK293T cells, rabbit α-EEA1 (Abcam, ab109110) or rabbit α-RAB11A (Cell Signaling, #2413S) were used in a 1:750 or 1:50 dilution, respectively. Endogenous proteins in N2a cells were stained with α-EEA1 (Abcam, ab109110) and rabbit α-RAB11A (Thermo Fischer, #715300) were used in a 1:500, 1:50 dilution, respectively. Coverslips were washed 6 times with IF washing buffer (50 mM NH<sub>4</sub>Cl, 1× CB) for 5 min at RT and incubated in CoraLite488 goat α rabbit IgG (Proteintech, SA00013-2) diluted in 1:250 in IF blocking buffer for 40 to 60 min at RT. After 6 washing steps in IF washing buffer for 5 min at RT, cells were counterstained with DAPI (f.c. 1 µg/mL) in IF washing buffer for 1 min and subsequently washed 3 times in IF washing buffer for 5 min. Finally, coverslips were mounted in 20 µL Mowiol on microscope slides and stored light protected at 4°C until microscopy.

For staining of endogenous FMRP and SMN, the fixed N2a cells (see above) were incubated in 0.2% Triton X-100 in DPBS and subsequently washed three times in DPBS for 5 min. After blocking of unspecific binding in 5% BSA (Roth) in DPBS-T (0.1% Tween 20 [Roth] in DPBS) (BB) for at least 1 h at RT, the primary antibody was incubated in BB for 1–1.5 h at RT. α-SMN (Proteintech, #11708) or α-FMRP (Cell Signaling, #4317) antibodies were applied in a 1:400 or 1:50 dilution, respectively. After washing the cells thrice in DPBS-T for 5 min, the secondary CoraLite488 goat α rabbit IgG antibody (Proteintech, SA00013-2) was incubated in a 1:250 dilution in BB for 40–60 min at RT. After three consecutive washing steps in DPBS-T for 5 min at RT, DAPI (f.c. 1 µg/mL) was applied in DPBS-T for 1 min at RT. The coverslips were washed three times in DPBS-T and mounted as described above. Antibodies used are listed in [Table S1](#).

### Confocal microscopy and image processing

Confocal fluorescence microscopy was performed on the Zeiss LSM980 Airyscan 2 microscope using the ZEN 3.5 software (Carl Zeiss Microscopy GmbH) in the Airyscan super resolution mode. Images were acquired as Z-stacks with an LD LCI Plan-Apochromat 63×/1.2 objective with Immersol G glycerine immersion oil (Zeiss) and using an LSM T-PMT. Excitation wavelength was preset by the software and is stated for DAPI = 405 nm, mCherry = 561 nm, and CoraLite488 = 488 nm. Deconvolution of images was performed employing the Classic Maximum Likelihood Estimation (CMLE) algorithm in the Huygens Software (Scientific Volume Imaging) and for further image processing and figure compilation the software Fiji<sup>47</sup> was used.

### In vivo proximity biotinylation

2.5 × 10<sup>6</sup> HEK293T cells were seeded in 100 mm dishes. After 24 h, cells were transiently transfected with 5 µg of plasmids harboring *MAGEL2* or *MAGEL2* c.1996dupC with an N-terminally fused HA-miniTurbo and 20 µL of Metafectene Pro. 48 h post transfection, the culture medium was discarded and cells were synchronized in starvation medium (DMEM, supplemented with 1% Pen/Strep) supplemented with 200 µM (f.c.) biotin for 4 h. HEK293T cells were carefully washed once with 1× DPBS and scraped in fresh DPBS. After centrifugation at 500 × g for 5 min, cells were resuspended in 500 µL RIPA lysis buffer (50 mM Tris-HCl [pH 7.4]; 1% Igepal CA-630; 0.25% SDS, 150 mM NaCl; 1 mM EDTA [pH 8.0]; 1× cOmplete [Roche]) according to Hutter et al.<sup>48</sup> and incubated on ice for 10 min. Subsequently, the cell debris was pelleted for 10 min at 14,000 × g and the protein concentration was determined by Bradford assay employing RotiQuant (Roth). Affinity capture of biotinylated proteins was performed as described before.<sup>48</sup> Briefly, 600 µg of total protein lysate was diluted in 500 µL RIPA lysis buffer supplemented with 60 µL of Streptavidin Magnetic Beads (NEB) and incubated overnight at 4°C under constant rotation. For mass spectrometric analysis, beads with biotinylated proteins were washed twice with RIPA lysis buffer.

### Protein co-immunoprecipitation

2.5 × 10<sup>6</sup> HEK293T cells were seeded in 100 mm dishes and grown for 24 h. Subsequently, cells were transfected with 5 µg of Low0\_GFPspark\_MAGEL2 or Low0\_GFPspark\_MAGEL2 c.1996dupC plasmids and 20 µL of Metafectene Pro (Biontex) and incubated for 48 h. HEK293T cells were washed with ice-cold DPBS (Gibco), scraped in 500 µL ice-cold homemade Pierce IP Lysis Buffer (25 mM Tris-HCl [pH 7.4]; 150 mM NaCl; 1 mM EDTA; 1% Igepal CA-630; and 5% glycerol; supplemented with 1× [f.c.] cOmplete protease inhibitor [Roche]) and incubated on ice for at least 15 min. Lysates were sonicated twice with 5 pulses at 0.5 cycles and 30 mA (Hielscher) on ice and 5% were retained for later SDS-PAGE analysis. 20 µL Affi-Prep Protein A slurry (BioRad) were washed three times in 200 µL Pierce IP Lysis Buffer at 1,000 × g for 1 min at 4°C (centrifugation setting for all steps) and then incubated with 500 µL cell lysate on a rotation wheel for 90 min at 4°C. 20 µL of 25% covalently coupled anti-GFP (Roche) Affi-Prep Protein A slurry (BioRad) were washed thrice in 200 µL Pierce IP lysis buffer (supplemented with cOmplete protease inhibitor [Roche]). 25 µL of precleared cell lysate were saved for later analysis and the remains was added to the washed anti-GFP-Protein A beads and incubated overnight at 4°C under rotation. Samples were centrifuged and supernatant was stored. Beads were washed thrice in 1 mL IP Lysis Buffer (with cOmplete

protease inhibitor [Roche]), twice in 1 mL 0.5× DPBS, twice in 1 mL 0.1× DPBS, and 3 times in 1 mL MilliQ-water. Co-immunoprecipitated proteins were eluted from beads twice with 2% formic acid for 15 min. At the second elution step, samples were vortexed after 10 min. After adjusting the volume of protein mixtures to 200 µL with 2% formic acid, 20 µL were saved for later SDS-PAGE and samples were analyzed by mass spectrometry.

When samples were prepared for Western blot analysis, cell seeding, transfection, cell lysis, and pre-clearing was conducted as described above or  $6.25 \times 10^6$  HEK293T cells were seeded in 150 mm dishes. For that purpose, all following reagents and buffers were upscaled 2.5-fold accordingly—the following protocol refers to  $2.5 \times 10^6$  starting cell number for 100 mm dishes.

Pre-cleared IP cell lysates were incubated with 2.5 ng/µL (f.c.) mouse anti-GFP antibody (Roche, #11814460001) at 4°C overnight under rotation. 20 µL Affi-Prep Protein A slurry (BioRad) were washed three times with 200 µL IP lysis buffer, supplemented with cOmplete protease inhibitor (Roche). Subsequently, they were incubated with the cell lysate/GFP-antibody mixture at 4°C for 90 min. After three wash steps in 1 mL IP lysis buffer, supplemented with cOmplete protease inhibitor (Roche), bound proteins were eluted from the beads in at least 1.5 bead volumes 1.5× Lämmli buffer at 95°C for 5 min with periodical vortexing.

### RNA co-immunoprecipitation

snoRNA genes and a scrambled RNA (amplified from the AmpR gene) sequence harboring a 5'-T7 promoter sequence were amplified by PCR on respective plasmids using Phusion High-Fidelity DNA Polymerase (Thermo Fisher Scientific) (for primers and RNA sequences refer to [Table S2](#)). Amplicons were checked on a 1% Agarose gel and purified by Monarch PCR & DNA Cleanup Kit (5 µg) (NEB). Transcripts were generated employing the HiScribe T7 High Yield RNA Synthesis Kit (NEB). After template DNA was digested with 4 U RQ1 DNase (Promega), T7 transcripts were PAGE purified from a 8% denaturing polyacrylamide gel by UV shadowing and passively eluted in gel elution buffer (300 mM NaCl; 0.2% sodium dodecyl sulfate; 60 mM NaOAc [pH 5.2]) under agitation overnight at 4°C as described by Hoernes et al.<sup>49</sup> Samples were incubated at 60°C for 1 h and stored on ice until RNA was purified. This particular step was performed using the Monarch RNA Cleanup Kit (50 µg) (NEB). Transcripts were biotinylated by use of Pierce RNA 3' End Biotinylation Kit (Thermo Fisher Scientific) according to the manufacturer's protocol. Thereby, a single biotinylated cytidine (bis)phosphate is ligated to the 3'-end of the respective transcripts in presence of DMSO at 16°C overnight. Biotinylated RNAs were purified utilizing Monarch RNA Cleanup Kit (10 µg) (NEB). Ligation efficiency was determined by northern blot analysis and dot blotting, followed by detection employing the Pierce Chemiluminescent Nucleic Acid Detection Module Kit (Thermo Fisher Scientific).

For LC-MS/MS analysis, HEK293T cells were washed with ice-cold DPBS (Gibco), scraped in ice-cold homemade Pierce IP Lysis Buffer (25 mM Tris-HCl [pH 7.4]; 150 mM NaCl; 1 mM EDTA; 1% Igepal CA-630; and 5% glycerol; supplemented with 1× [f.c.] cOmplete protease inhibitor [Roche]) and incubated for 5 min on ice. Cell debris was pelleted by centrifugation at  $13,000 \times g$  for 10 min at 4°C. Protein concentration was determined by Bradford assay with ROTIQuant (Roth).

40 pmol of biotinylated RNA were renatured at 65°C for 10 min, at RT for 20 min, and stored on ice until they were immobilized on 20 µL Dynabeads M-280 Streptavidin (Thermo Fisher Scientific) in

30 min incubation time, otherwise following the manufacturer's protocol. Immobilized RNA-bead mixture was washed two times in 1× Protein-RNA binding buffer (20 mM Tris [pH 7.5]; 150 mM NaCl; 2 mM MgCl<sub>2</sub>; 0.1% Tween-20); pull-down was performed by a modified protocol from Pierce Magnetic RNA-Protein Pull-Down Kit (Thermo Fisher Scientific). Binding was performed in a 200 µL RNA-protein binding reaction containing 1 µg/µL protein lysate, 1× Protein-RNA binding buffer, 15% glycerol, 0.4 U/µL of RNase inhibitor (Promega), and 1× cOmplete protease inhibitor (Roche) at 4°C for 1 h on a rotation wheel. Beads were washed thrice with 200 µL IP wash buffer (20 mM Tris [pH 7.5], 150 mM NaCl, 0.1% Tween-20) and thrice with 500 µL 50 mM Tris (pH 7.5).

Sample preparation for western blot analysis had modifications, i.e., only 20 pmol of biotinylated RNA were used and all other reagents were downscaled accordingly. Transfection of HEK293T with Low0\_GFPspark, Low0\_GFPspark\_MAGEL2, or Low0\_GFPspark\_MAGEL2 c.1996dupC plasmids as well as cell lysate preparation were conducted as described above. After Bradford analysis, all three lysates were mixed in equal parts according to their protein concentration and 100 µg of this lysate mix were used for immunoprecipitation. After the third wash step with IP wash buffer, proteins were eluted in 40 µL 1× Lämmli buffer at 95°C for 5 min.

### LC-MS/MS analysis

Proteins from BioID and RNA-CoIP were reduced with dithiothreitol, alkylated with iodoacetamide, and digested with trypsin (Promega) according to Branon et al.<sup>44</sup> Then samples were purified using Pierce C18 Tips, 100 µL (Thermo Fisher Scientific) according to the manufacturer's instructions prior to nano LC-MS analysis.

Samples from Protein-CoIP were dried in a speedvac and dissolved in 33 µL 100 mM NH<sub>4</sub>HCO<sub>3</sub> buffer (pH 8.0). Subsequently 33 µL of 10 mM DTT in 100 mM NH<sub>4</sub>HCO<sub>3</sub> were added and incubated at 56°C for 30 min. Thereafter 1.0 µg trypsin (Promega) was added to the sample and incubated at 37°C for 6 h. Finally, 33 µL of 55 mM IAA dissolved in 100 mM NH<sub>4</sub>HCO<sub>3</sub> were added and incubated at room temperature for 20 min in the dark.

Digested samples were analyzed using an UltiMate 3000 nano-HPLC system coupled to a Q Exactive HF mass spectrometer (Thermo Fisher Scientific) equipped with a Nanospray Flex ionization source. The peptides were separated on a homemade fritless fused-silica microcapillary column (75 µm i.d. × 280 µm o.d. × 16 cm length) packed with 2.4 µm reversed-phase C18 material (Reprosil). Solvents for HPLC were 0.1% formic acid (solvent A) and 0.1% formic acid in 85% acetonitrile (solvent B). The gradient profile was as follows: 0–4 min, 4% B; 4–57 min, 4%–35% B; 57–62 min, 35%–100% B; and 62–67 min, 100% B. The flow rate was 250 nL/min.

BioID samples were analyzed with a Q Exactive HF mass spectrometer. The instrument was operating in the data-dependent mode selecting the top 20 most abundant isotope patterns with charge >1 from the survey scan with an isolation window of 1.6 mass-to-charge ratio (m/z). Survey full-scan MS spectra were acquired from 300 to 1,750 m/z at a resolution of 60,000 with a maximum injection time (IT) of 120 ms and automatic gain control (AGC) target 1e6. The selected isotope patterns were fragmented by higher-energy collisional dissociation (HCD) with normalized collision energy of 28 at a resolution of 30,000 with a maximum IT of 120 ms and AGC target 5e5.

Samples from Protein-CoIP and RNA-CoIP were analyzed with an Orbitrap Eclipse mass spectrometer. The instrument

equipped with a field asymmetric ion mobility spectrometer (FAIMS) interface was operating in the data-dependent mode with compensation voltages (CV) of  $-45$  and  $-65$  and a cycle time of 1 s. Survey full-scan MS spectra were acquired from 375 to 1,500  $m/z$  at a resolution of 240,000 with an isolation window of 1.2 mass-to-charge ratio ( $m/z$ ), a maximum injection time (IT) of 50 ms, and automatic gain control (AGC) target 400,000. The MS2 spectra were measured in the Orbitrap analyzer at a resolution of 15,000 with a maximum IT of 22 ms, and AGC target of 50,000. The selected isotope patterns were fragmented by higher-energy collisional dissociation with normalized collision energy of 28.

Data analysis was performed using Proteome Discoverer 2.5 (Thermo Fisher Scientific) with search engine Sequest. Depending on the input sample type, the raw files were searched against *Homo sapiens* or *Mus musculus* uniprot databases. Precursor and fragment mass tolerance was set to 10 ppm and 0.02 Da, respectively, and up to two missed cleavages were allowed. Carbamidomethylation of cysteine was set as static modification, and oxidation of methionine was set as variable modifications. Peptide identifications were filtered at 1% false discovery rate.

### Western blot analysis

For analyses of co-immunoprecipitation experiments, proteins were denatured for 5 min at  $95^{\circ}\text{C}$  and separated on gradient SDS-PAGE gels in tris-glycine running buffer (24.8 mM Tris, 191.8 mM Glycine, 0.5% SDS) in a Novex Invitrogen XCell SureLock Electrophoresis System (Thermo Fisher Scientific).

Separated proteins were transferred onto a 0.45  $\mu\text{m}$  PVDF membrane (Thermo Fisher Scientific) in  $1\times$  transfer buffer (24.8 mM Tris, 191.8 mM glycine, 10% ethanol, 0.08% SDS) in a Novex Invitrogen XCell II Blot Module 1 (Thermo Fisher Scientific) wet blotting apparatus at 20 V, 100 mA for 1 to 4 h.

The membrane was blocked in blocking buffer ( $1\times$  TBS-T: 137 mM NaCl, 27 mM KCl, 25 mM Tris [pH 7.4], 0.1% Tween 20; supplemented with 5% BSA [Roth]) for at least 1 h at RT and incubated with the primary antibody diluted in blocking buffer for 1–2 h at RT or overnight at  $4^{\circ}\text{C}$  under constant rotation. Membranes were washed thrice in TBS-T for 5 min at RT and incubated with the secondary antibody conjugated to horseradish peroxidase in locking solution for 1 h at RT. Membranes were washed again 3 times in TBS-T for 5 min and signals were detected employing Pierce ECL chemiluminescent substrate (Thermo Fisher Scientific) and Kodak BioMax MS film. Antibodies used for western blot analysis are listed in [Table S3](#).

### Northern blot analysis

Total smNPC RNA was isolated employing TRI Reagent (Sigma-Aldrich) according to the manufacturer's protocol. For the detection of MAGEL2 and MAGEL2 c.1996dupC mRNAs, 10  $\mu\text{g}$  of total RNA were loaded onto a 1.2% denaturing agarose gel in  $1\times$  RP buffer (40 mM MOPS, 10 mM sodium acetate, 2 mM EDTA [pH 7.0]) containing 2.17% formaldehyde and the gel was run in RP buffer containing 2.17% formaldehyde at 100 V for 50 min. The gel was washed 2 times in  $10\times$  SSC (1.5 M NaCl, 150 mM sodium citrate [pH 7.2]) for 15 min. RNA was transferred onto an Amersham Hybond- $\text{N}^+$  nylon membrane (GE Healthcare) employing passive capillary blotting in  $10\times$  SSC overnight. Subsequently, the membrane was rinsed with  $2\times$  SSC, shortly air dried, and the RNA was cross-linked to the membrane with 254 nm at 0.12 Joule in a UV cross-linker (Stratagene).

Digoxigenin (DIG)-labeled probes for detection were produced by PCR employing a dNTP mix containing a DIG-UTP:dTTP ratio of 1:2 (for primers refer to [Table S4](#)).  $M2_{-5'}$  and  $M2_{-3'}$  probes ([Table S4](#)) anneal within the 5' and 3' portion of the MAGEL2 transcript, respectively. After pre-hybridization in high-SDS solution (7% SDS, 50% deionized formamide, 20% blocking solution, 10% blocking reagent [Roche],  $1\times$  maleic acid buffer [100 mM maleic acid, 150 mM NaCl, 100 mM NaOH (pH 7.5)]), 1% sodium lauryl sarcosinate) at  $42^{\circ}\text{C}$  for 4 h, cross-linked membranes were hybridized with 15 mL high-SDS solution containing DIG-labeled probes at  $42^{\circ}$  overnight. Subsequently, the membranes were washed three times in  $1\times$  SSC supplemented with 0.1% SDS at RT for 10 min and two times at  $65^{\circ}\text{C}$  for 15 min. 5 min incubation in DIG wash buffer (99.7%  $1\times$  maleic acid buffer, 0.3% Tween 20) were followed by DIG-P2 (90%  $1\times$  maleic acid buffer, 10% blocking reagent [Roche]) incubation for 30 min, before alkaline phosphatase conjugated anti-DIG antibody solution (99.99% DIG-P2, 0.01% anti-DIG antibody [Roche]) was applied for 30 min at RT.

Membranes were washed once in DIG wash buffer for 5 min, twice for 10 min at RT, and incubated in DIG-P3 buffer (0.1 M Tris, 0.1 M NaCl, 50 mM  $\text{MgCl}_2$  [pH 9.5]) for 5 min, before the DIG CSPD substrate solution (99% DIG-P3, 1% CSPD [Roche]) was applied for additional 5 min at RT. Signals were detected applying Kodak BioMax MS films (Kodak) to the membranes at  $37^{\circ}\text{C}$  for approximately 2 h and developed.

For the detection of ncRNAs, 15  $\mu\text{g}$  of total RNA were separated on 8% denaturing polyacrylamide gels (acrylamide:bisacrylamide = 29:1, 7 M urea in  $1\times$  TBE) at 120 to 370 V for 2 to 3 h. After ethidium bromide staining for 10 min, the RNA was transferred onto Amersham Hybond- $\text{N}^+$  membranes employing the Trans-Blot SD Semi-Dry Transfer Cell (Bio-Rad) at 15 V and 400 mA for 45 min. The RNA was UV-crosslinked at 0.12 kJ in a UV crosslinker (Stratagene) and ncRNAs of interest were detected by 5'-radioactively labeled oligonucleotides. Therefore, oligos (in total 1.2  $\mu\text{M}$  f.c.) were labeled with 1  $\mu\text{Ci}/\mu\text{L}$  [ $\gamma$ - $^{32}\text{P}$ ]-ATP (Hartmann Analytic) by 0.8 U/ $\mu\text{L}$  T4 PNK (NEB) for at least 30 min at  $37^{\circ}\text{C}$  and denatured at  $96^{\circ}\text{C}$  for 1 min. After equilibration of cross-linked membranes in hybridization buffer (178 mM  $\text{Na}_2\text{HPO}_4$ , 822 mM  $\text{NaH}_2\text{PO}_4$ , 7% (v/v) SDS [pH 6.2]), the labeling mix was added for overnight incubation at  $42^{\circ}\text{C}$  under constant rotation. Blots were washed in wash buffer I (300 mM NaCl, 34 mM sodium citrate, 0.1% (w/v) SDS [pH 7]) for 10 min and wash buffer II (15 mM NaCl, 1.7 mM sodium citrate, 0.1% (w/v) SDS [pH 7]) for 5 min and rinsed in Aqua dest. Signals were detected employing a phosphorimager and Typhoon FLA 9500 (GE Healthcare).

If necessary, blots were stripped in 0.5% (w/v) SDS at  $65^{\circ}\text{C}$  for 1 h, rinsed with Aqua dest., incubated in Aqua dest. at  $65^{\circ}\text{C}$  for 30 min, and rinsed again with Aqua dest. At least 19 *SNORD116* copies were detected with three probes (5'-GGACCTCAGTTCC GATGA-3' for copies 1–3, 6, 8, and 9; 5'-GGACCTCAGTTTCGAC GAG-3' for copies 12 and 16–22; copy 10 with 2 mismatches; and 5'-GGACCTCAGCTCACAGAA-3' for copies 25–27, 29, and 30). 5'-GGGCCTCAGCGTAATCCT-3' was used to detect several copies of *SNORD115* (1, 4–16, 21, 24–26, 29, 30, 33, 34, 36, and 38–44). 5.8S rRNA (5'-GCAATTCACATTAATTCTCGCAGCTAGC-3') and U6 snRNA (5'-TATGGAACGCTTCACGAATTTG-3') were used as controls.

### RT-qPCR

167 ng/ $\mu\text{L}$  total RNA was reverse transcribed using the SuperScript IV VIL0 Master Mix with ezDNase Enzyme (Thermo Fisher



Scientific) according to the manufacturer's protocol. 2  $\mu$ L of 1/100 diluted cDNA was amplified with 3  $\mu$ L Luna Universal qPCR Master Mix (NEB) and 200 nM forward and reverse primers (Table S5) each in 6  $\mu$ L reaction volume in 384-well plates using QuantStudio 5 Real-Time PCR System (Applied Biosystems). After quality control (QC, i.e., melt-curve analysis and outlier removal—samples were analyzed only when at least 2 of 3 technical replicates passed QC), primer efficiencies were calculated from amplification curves using web-based LinRegPCR (<https://www.gear-genomics.com/rdml-tools/>).<sup>50</sup> Inter-plate variation was corrected employing RDML-Analyze from rdml-tools.<sup>51</sup> RefFinder<sup>52,53</sup> was used to find the most stable housekeeping genes *GAPDH*, *TBP*, and *HPRT*. Percentage changes were calculated by subtracting the median fold changes (as calculated by Ruijter et al.<sup>54</sup> of SYS and control cell lines. For comparison Wilcoxon test was used.

## Computational analyses

R and R Studio was used for most computational analyses. For general data processing the packages *rio* and *tidyverse*<sup>55</sup> were used. Mass spectrometric-identified protein abundances were normalized to mean total protein abundances per condition, followed by removal of proteins discovered with only one respective peptide or no available abundance. Only proteins with a reviewed status in the uniprot database and present in at least two replicates in at least one condition were included in the analysis. Proteomics analysis was conducted by the DEP package.<sup>56</sup> In brief, abundances of single replicates were normalized using variance stabilizing transformation (VSN). Missing value imputation was performed per sample by mixed imputation. Proteins present in two of three or one of three replicates were defined as missing at random (MAR) or missing not at random (MNAR), respectively. MARs were imputed by maximum likelihood estimation and MNARs were imputed with zero. For differential enrichment analysis, DEP's "test\_diff" function was modified, so that FDR was adjusted for the *p* value instead of t-statistic. Proteins with an FDR < 0.05 were assumed statistically significant.

Before GO enrichment analysis, the datasets from differential enrichment analysis were cleansed of general contaminant proteins by use of the CRAPome database.<sup>57</sup> For proximity-based *in vivo* biotinylation, the CRAPome database was filtered for cell/tissue type "HEK293," subcellular fractionation "total cell lysate" and "total cell lysate and chromatin," epitope tag "BirA\*-FLAG," and affinity approach "Streptavidin." For GFP-fusion protein CoIP, we used the filters cell/tissue type "HEK293," subcellular fractionation "total cell lysate and chromatin," and affinity approach "anti-GFP mouse." For each gene, spectral counts (designated as "PSMs"—number of peptide spectrum matches—in Proteome Discoverer 2.5) were averaged across the three replicates and across the retrieved datasets from the CRAPome database. Subsequently, per BioID and GFP-CoIP only proteins with at least twice as many spectral counts than the control dataset were used for further analysis. As for RNA-CoIPs, no public contaminant lists are available, so we included a scrambled RNA control in the experiments. Because it was not possible to quantify bait amount in our experimental design, we conducted GO enrichment analysis with proteins which were found in at least two of three replicates, whereupon interactors of scrambled control were removed from obtained interactor lists of *SNORD100* or *SNORD116* pull-downs employing interactivenn.<sup>58</sup> Gene ontologies were analyzed using the EnrichR R package which connects to the enrichr web-

server.<sup>59–61</sup> Upon GO enrichment analysis functional categories were sorted by combined score<sup>60</sup> and top 5 hits per condition were depicted.

## Statistics

Plots were generated with ggplot2<sup>62</sup> or ggpubr. R-package rstatix was used for statistical analyses.

## NLS and NES prediction tool

The computational prediction of potential nuclear localization signal (NLS) motifs for human MAGEL2 WT and the truncated p.Gln666Profs\*47 variant was performed using the cNLS Mapper online tool<sup>63</sup> with a cut-off score of 2.0 as well as the NLStradamus online tool<sup>64</sup> with a prediction cut-off of 0.5 in the 4 state HMM static model. Potential nuclear export signal (NES) motif prediction was performed using the LocNES server.<sup>65</sup>

## Results

### The truncated p.Gln666Profs\*47 variant accumulates predominantly within the nucleus while WT MAGEL2 shuttles between the cytoplasm and the nucleus

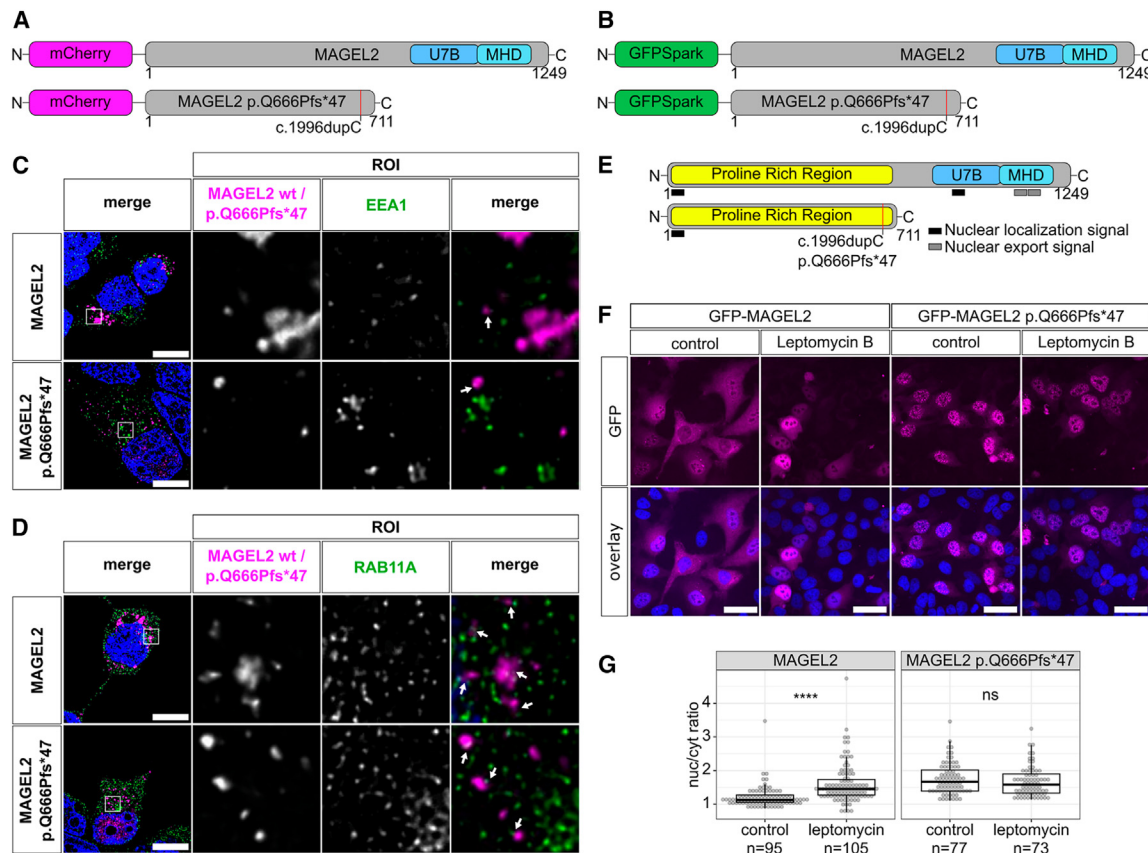
Human MAGEL2 is a 1,249 amino acid protein with a MAGE homology domain (MHD) and a USP7 binding site (U7BS) in the C-terminal domain, located at residues 1,027 to 1,195 and 820 to 1,034, respectively.<sup>36,37,40</sup> The N-terminal region (aa 1–819) is an intrinsically disordered domain containing 28% proline, 15% alanine, and 11% glutamine residues.<sup>40</sup>

The *MAGEL2* c.1996dupC frameshift mutation represents the most commonly found pathogenic variant among individuals with SYS and shows the second most severe phenotype, with the only exception of the *MAGEL2* c.1996dupC mutation which has been reported to be associated with pre- or perinatal lethality.<sup>32</sup>

In the present study, we initially investigated the subcellular localization of the predominant, truncated p.Gln666Profs\*47 variant and its WT counterpart. Subsequently, we aimed to decipher functional interaction partners of WT and mutant MAGEL2 proteins and to investigate molecular mechanisms which might be connected to the expression and/or function of *SNORD116* snoRNAs as well as protein-coding genes from the PWS locus. To this end, we first generated expression vectors in which mCherry or GFP tags were fused in frame to the N terminus of the WT or the c.1996dupC frameshift mutant *MAGEL2* sequence (Figures 1A and 1B) to maintain transcription of the full-length mRNA. Thereby, the inserted c.1996dupC mutation resulted in the truncated MAGEL2 p.Gln666Profs\*47 variant due to a premature stop codon (Figures 1A and 1B).

MAGEL2 has previously been shown to bind USP7 and TRIM27 proteins, thus forming the MUST complex.<sup>36,37,66</sup> The trimer was shown to facilitate F-actin polymerization and retrograde recycling on endosomes by ubiquitination of the WASH complex.<sup>36,37,66</sup> Similar to the actin nucleation promoting factor WASH containing complex SHRC





**Figure 1. Subcellular localization of MAGEL2 WT and p.Gln666Profs\*47**

(A and B) MAGEL2/p.Gln666Profs\*47 fusion proteins employed for investigation of their subcellular localization, tagged with either an N-terminal mCherry (A) or GFPspark (B).

(C and D) Confocal immunofluorescence microscopy of recombinant fusion proteins WT mCherry-MAGEL2 or mCherry-p.Gln666Profs\*47 (magenta) in transiently transfected HEK293T cells, 24 h after transfection. Nuclei were counterstained with DAPI (blue). The scale bar is 10  $\mu$ m. White arrows indicate sites of spatial proximity between WT MAGEL2 and p.Gln666Profs\*47 with endogenous early endosomes (C) and recycling endosomes (D), which were visualized with anti-EEA1 and anti-RAB11A antibodies (green), respectively. At least 5 cells were imaged in three independent replicates and representative cells are depicted.

(E) MAGEL2 WT and truncated p.Gln666Profs\*47 with computationally predicted putative NLS (black boxes) and NES (gray boxes) motifs.

(F and G) HeLa cells were transfected with the GFP-tagged wild-type MAGEL2 or MAGEL2 c.1996dupC mutant. 48 h after transfection cells were treated with leptomycin B (20 nM) or left untreated for 20 h. Nuclei were counterstained with DAPI (blue).

(F) HeLa cells with and without leptomycin B treatment. Cells from three independent transfection experiments were used for quantification. The scale bar is 40  $\mu$ m.

(G) Quantification of the nuclear/cytoplasmic ratio of individual cells from three independent experiments. Wilcoxon tests were calculated employing rstatix.

(WASH regulatory complex),<sup>67–69</sup> the MUST complex is actively recruited to endosomal subdomains by the retromer subunit VPS35.<sup>36,37,66</sup>

Hence, we first addressed a potential co-localization between an mCherry-tagged WT or truncated MAGEL2 protein with the early- and recycling-endosomal markers EEA1 and RAB11A, respectively. Therefore, HEK293T cells were transiently transfected with fluorescently tagged WT and truncated MAGEL2 expression vectors. WT and truncated MAGEL2 were randomly distributed within the cytoplasm, where the p.Gln666Profs\*47 variant predominantly appeared in vesicular-like structures (Figures 1C and 1D). However, both proteins were only rarely in spatial proximity to EEA1 (Figure 1C) but were located adjacent to RAB11A (Figure 1D).

Interestingly, during these analyses we observed a considerably higher nuclear accumulation of the p.Gln666Profs\*47 mutant compared to WT MAGEL2 (Figures 1C and 1D). Based on these findings, we hypothesized that truncated MAGEL2 p.Gln666Profs\*47 might be subject to either an increased nuclear import or a reduced nuclear export kinetic in comparison to its wild-type counterpart. Indeed, by employing computational prediction tools, a putative bipartite nuclear localization signal (NLS) motif was indicated, i.e., 4-LSKNLGDSSPPAEAPKPPVYSRPTVLMRAP-33, at the N-terminal end of WT MAGEL2 and p.Gln666Profs\*47 (Figure 1E; Table S6). Additionally, a putative monopartite NLS, i.e., 883-GKATRKKKHLE-893, and two nuclear export signal (NES) motifs, i.e., 1,107-RPKFGLLMVVLSLIF-1,121 and 1,127-VREDLIFNLFKLG-1,141, were

predicted within the C-terminal half of WT MAGEL2, which were absent in the truncated p.Gln666Profs\*47 variant (Figure 1E, Tables S6 and S7).

To further investigate the nuclear export of MAGEL2 proteins, we transiently transfected HeLa cells with WT GFP-MAGEL2 or p.Gln666Profs\*47 expression vectors (Figure 1B) and subsequently blocked the nuclear export of MAGEL2 proteins by inhibiting the export factor karyopherin CRM1 (exportin 1) with leptomycin B.<sup>70,71</sup> As observed in HEK293 cells, also in HeLa cells the p.Gln666Profs\*47 variant showed an increased nuclear localization compared to WT MAGEL2 (Figures 1F and 1G). However, some nuclear localization was also observed for WT MAGEL2 (Figures 1C, 1D, and 1F). Consistent with our hypothesis, leptomycin B treatment resulted in a significant accumulation of WT MAGEL2 within the nucleus compared to the untreated control ( $W = 2017$ ,  $p = 3.7 \times 10^{-13}$ ) (Figures 1F and 1G), while truncated p.Gln666Profs\*47 localization was not affected by CRM1 inhibition ( $W = 3123$ ,  $p = 0.24$ ) (Figures 1F and 1G).

#### ***In vivo* proximity labeling and co-immunoprecipitation reveal protein interaction partners for MAGEL2 WT and the p.Gln666Profs\*47 variant**

To decipher potential pathogenic roles of MAGEL2 frame-shift mutants in SYS-affected individuals, we considered three different hypotheses to link the truncated MAGEL2 proteins to the SYS phenotype: firstly, the different distribution of WT and the truncated MAGEL2 protein might lead to differences in the composition of their interactomes explaining a potential gain or loss of function of MAGEL2 p.Gln666Profs\*47 within the nucleus or cytoplasm, respectively. Secondly, the truncated form of MAGEL2, consisting mainly of the intrinsically disordered proline-rich N-terminal region,<sup>40</sup> might *per se* promote interactions with novel binding partners. Lastly, MAGEL2 p.Gln666Profs\*47 might still be able to bind canonical interaction partners via its N-terminal region but would lack a functional response due to loss of U7BS and MHD in the C-terminal region.

In a recent study, interacting proteins of over-expressed WT MAGEL2 and the C-terminal portion of MAGEL2 have been identified in HEK293T cells by Sanderson et al.<sup>40</sup> By employing a “subtractive” approach, i.e., comparing proteins binding to full-length MAGEL2 versus proteins binding to the C-terminal portion, also N-terminal protein binding partners were predicted.<sup>40</sup> In contrast, in our study we directly employed the original truncated N-terminal p.Gln666Profs\*47 variant and compared its interactome to that of full-length WT MAGEL2 to identify MAGEL2 WT and p.Gln666Profs\*47 interaction partners. To achieve this aim, we generated vectors for the expression of biotin ligase miniTurbo<sup>44</sup> (Figure 2A) or GFPspark (Figure 1B) WT MAGEL2 and p.Gln666Profs\*47 fusion proteins.

By applying different proteomic approaches, we wanted to cover the entire spectrum of MAGEL2 interaction part-

ners as comprehensively as possible, but also aimed to pinpoint candidates that specifically bind either to the WT or the truncated MAGEL2 proteins. By employing BioID, all proteins in proximity of about 10 nm to the biotin ligase are known to be biotinylated<sup>73</sup> prior to the pull-down. In contrast, by GFP-CoIP, only direct interaction partners and potentially their respective complexes are co-precipitated. Therefore, the interactome identified by BioID was expected to be larger than that obtained by GFP-CoIP (Figure 2B).

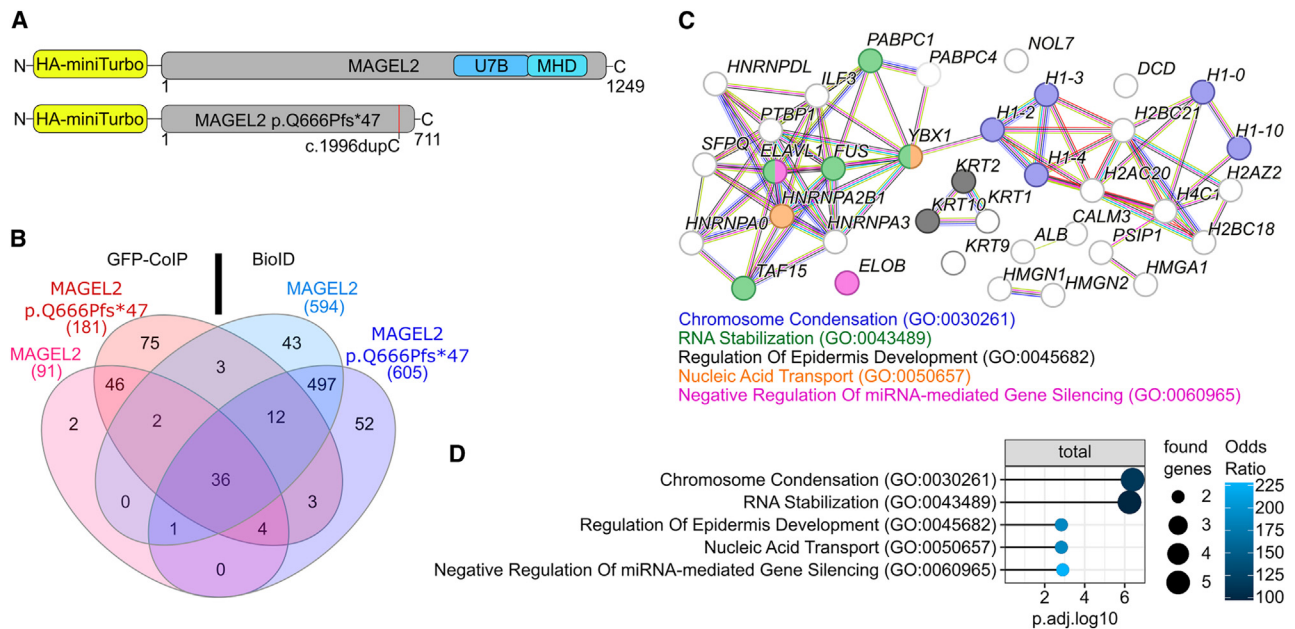
#### **MAGEL2 WT and the p.Gln666Profs\*47 variant share interactomes involved in RNA stabilization and chromosome condensation**

First, we investigated common interaction partners for MAGEL2 WT and p.Gln666Profs\*47, which were present in at least two out of three replicates (Figures 2B–2D). Previous analysis of the full-length MAGEL2 interactome suggested that the protein might be involved in various aspects of RNA metabolism.<sup>40</sup> Using BioID and GFP-CoIP, our proteomics analyses revealed 36 common protein interaction partners for WT and truncated MAGEL2 (Figures 2B–2D), of which only YBX1 and PAPBC1 have been previously described to be in proximity to the C-terminal or full-length MAGEL2, respectively.<sup>40</sup> These proteins clustered into two major groups in protein-protein association network analysis<sup>72</sup> (Figure 2C). The first cluster contained PABPC1, YBX1, FUS, ELAVL1, TAF15, and HNRNPA0, which are all implicated in RNA stabilization, while histones H1.0, H1.2, H1.3, H1.4, and H1.10 in the second cluster are involved in chromosome condensation (Figures 2C and 2D). Additionally, HMGA1, SFPQ, TAF15, ILF3, HMGN1, YBX1, and FUS represent a subset of common protein interaction partners involved in the regulation of transcription (Figure 2C; Table S8).

#### **Mutations in MAGEL2 might result in loss of function or gain of function in RNA metabolism processes**

Next, we aimed to identify differences in the interactomes of MAGEL2 WT and p.Gln666Profs\*47 and analyzed proteins overlapping in GFP-CoIP and BioID for each protein variant (Figure 3A). As expected, fewer proteins were identified employing GFP-CoIP compared to BioID (Figure 3A) due to the general experimental differences between the two approaches, whereby p.Gln666Profs\*47 interacted with more proteins than WT MAGEL2 (Figure 3A). The interactome of WT MAGEL2 was mainly found to be involved in chromosome condensation and organization (Figure 3B), whereas mRNA stabilization was one of the top hits of the proteins bound to p.Gln666Profs\*47 (Figure 3B).

Subsequently, we focused on 2-fold enriched proteins per pull-down set to identify proteins that were either proximal to or would directly interact with either WT MAGEL2 or p.Gln666Profs\*47. Hence, these analyses might also indicate potential novel gain or loss of functions of the truncated MAGEL2 variant (Figure 3C). The enriched proteins identified by the MAGEL2 BioID



**Figure 2. MAGEL2 WT and p.Gln666Profs\*47 show high affinity for proteins involved in RNA metabolism and chromosome condensation**

(A) Fusion-protein constructs for BioID.

(B–D) Analyses were carried out for proteins identified in at least two of three replicates per experiment.

(B) Combined Venn diagram interaction partners of MAGEL2 WT and p.Gln666Profs\*47 identified by BioID and GFP-CoIP.

(C) Protein-protein association networks (gene names are depicted)<sup>72</sup> of the 36 common interaction partners identified in (B), colored by the top 5 biological processes in (D).

(D) Top 5 hits by combined score<sup>60</sup> of gene enrichment analysis employing enrichR for biological processes (2023) in either BioID or GFP-CoIP. Only GO terms with an adjusted *p* value <0.05 were included.

were mostly involved in transcription regulation (Figures 3C and S1C). In contrast, proteins in proximity to MAGEL2 p.Gln666Profs\*47 were implicated in mRNA stabilization and snRNA metabolism, while direct interaction partners of MAGEL2 p.Gln666Profs\*47 identified by LC-MS/MS were involved in cell metabolism, i.e., pyruvate metabolism and gluconeogenesis (Figure 3C; Table S9). Included in that group of proteins was the DNA- and RNA-binding protein KHSRP,<sup>74</sup> which was also enriched in MAGEL2 p.Gln666Profs\*47 employing BioID (Figure 3A).

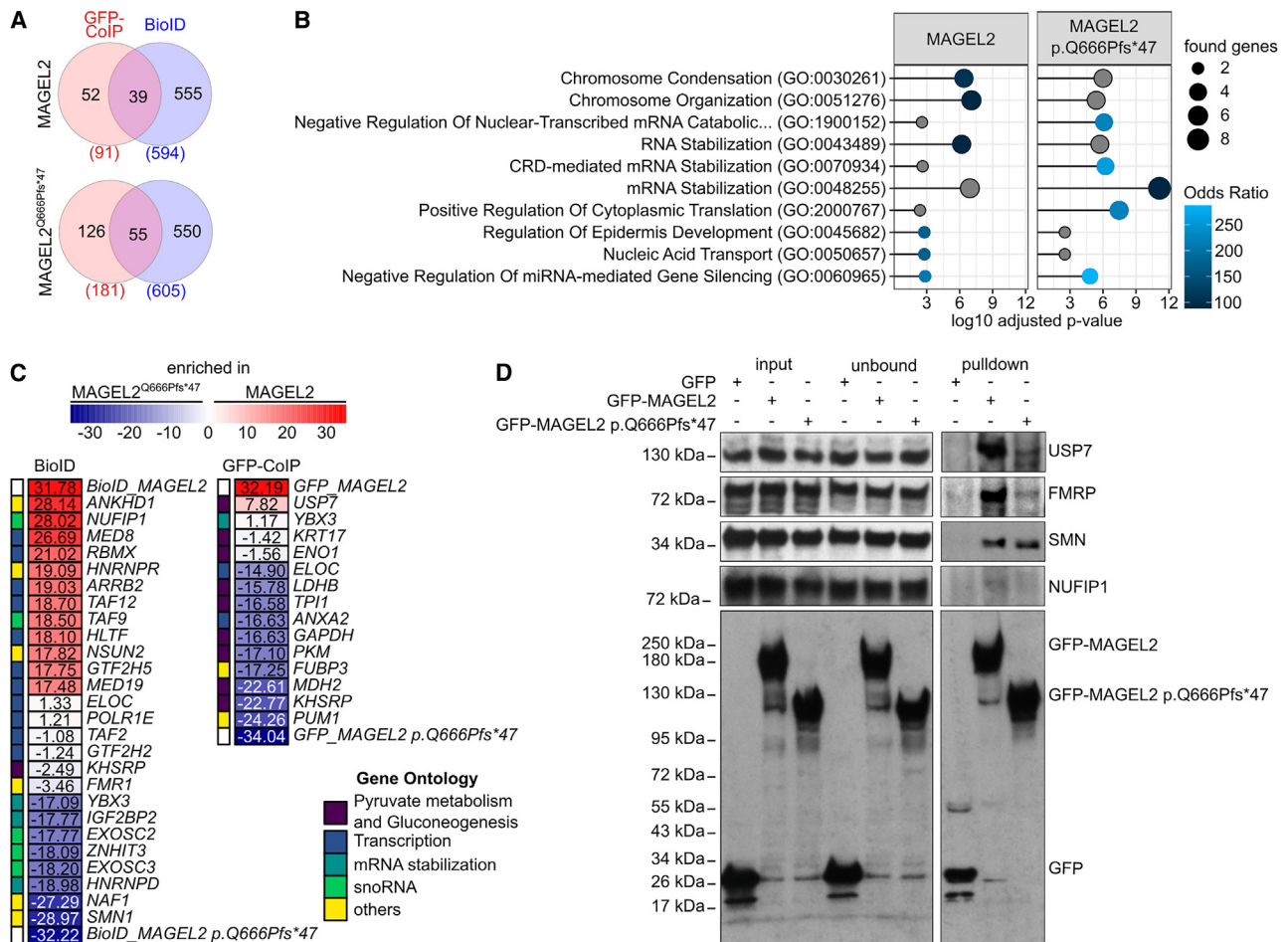
In addition, NUFIP1 was enriched in the WT MAGEL2 BioID (Figure 3C). However, NUFIP1 was on the threshold of detectability in only one of three replicates of western blot analyses upon GFP-CoIP with WT GFP-MAGEL2 (Figure 3D). SMN and FMRP enrichment was observed in the MAGEL2 p.Gln666Profs\*47 BioID approach (Figure 3C). Importantly, subsequent western blot analysis confirmed FMRP and SMN to be specifically co-immunoprecipitated with WT GFP-MAGEL2 and GFP-p.Gln666Profs\*47 (Figure 3D). In this context, FMRP showed a higher affinity to WT MAGEL2 in comparison to the truncated protein (Figure 3D). Hence, our data strongly indicate that FMRP and SMN are indeed proximal interaction partners of WT and truncated MAGEL2 proteins, whereas NUFIP1 is most likely an indirect binding partner of WT MAGEL2 (Figures 3C and 3D).

USP7, which has been previously described as a functional binding partner of MAGEL2,<sup>66</sup> was found to be enriched in the WT MAGEL2 GFP-CoIP in both LC-MS/MS and western blot analyses (Figures 3A and 3B). Intriguingly, USP7 was also slightly detectable in the MAGEL2 p.Gln666Profs\*47 variant (Figure 3B), although the annotated U7BS is absent in the truncated mutant (Figure 2A).<sup>36</sup>

#### MAGEL2 WT and p.Gln666Profs\*47 co-localize with SMN and FMRP in mouse N2a cells

Considering that WT and truncated MAGEL2 variants were specifically binding to SMN and FMRP in our GFP-CoIPs (Figure 3D), we next asked the question of whether both MAGEL2 variants colocalized with SMN and FMRP in neuronal cells. To that aim, we transiently transfected the mCherry-MAGEL2 and p.Gln666Profs\*47 constructs into mouse N2A cells (Figures 4A and 4B). In the majority of the transfected cells, we observed colocalization of both MAGEL2 variants with SMN and FMRP in distinct speckles present in the cytoplasm. Analogous to the experiments presented before in HEK293T cells (Figures 1B and 1C), we also investigated whether MAGEL2 variants were found on endosomes in N2A cells. Indeed, transiently transfected mCherry-MAGEL2 and p.Gln666Profs\*47 partially colocalized with EEA11 and RAB11A (Figures 4C and 4D). Interestingly, the association of MAGEL2 with early and recycling endosomes appears preferentially in neuronal cells





**Figure 3. MAGEL2 WT and p.Gln666Profs\*47 interact with proteins involved in RNA metabolism and gene expression as well as disease related SMN (spinal muscular atrophy) and FMRP (fragile-X syndrome)**

(A) Venn diagrams of protein interaction partners for MAGEL2 WT and p.Gln666Profs\*47 identified by BioID or GFP-CoIP.  
 (B) Gene enrichment analysis for Biological Processes (2023) in either BioID or GFP-CoIP employing enrichR. Shades of blue depict top 5 hits by combined score.<sup>60</sup> Gray circles indicate that the respective GO term is among the top hits in another condition. GO terms with an adjusted *p* value > 0.05 were excluded.  
 (C) Heatmap for enriched proteins (gene names are depicted) in MAGEL2 WT and p.Gln666Profs\*47 by BioID and GFP-CoIP. GO term enrichment analyses for Biological Process and KEGG Pathways from Figure S1 were grouped to broader functional terms as shown in Table S9 are written in the respective colors (Gene Ontology). Gene ontology "others" include novel interaction partners of interest which could not be assigned to the remaining annotated terms.  
 (D) Western blots of proteins which co-immunoprecipitated with WT GFP-MAGEL2 or GFP-p.Gln666Profs\*47 (*n* = 3). 5% of the lysate were used in input fractions.

and was less pronounced in HEK293T cells (Figures 1C and 1D). Of note, p.Gln666Profs\*47 showed less overlap with RAB11A compared to WT MAGEL2 (Figure 4B).

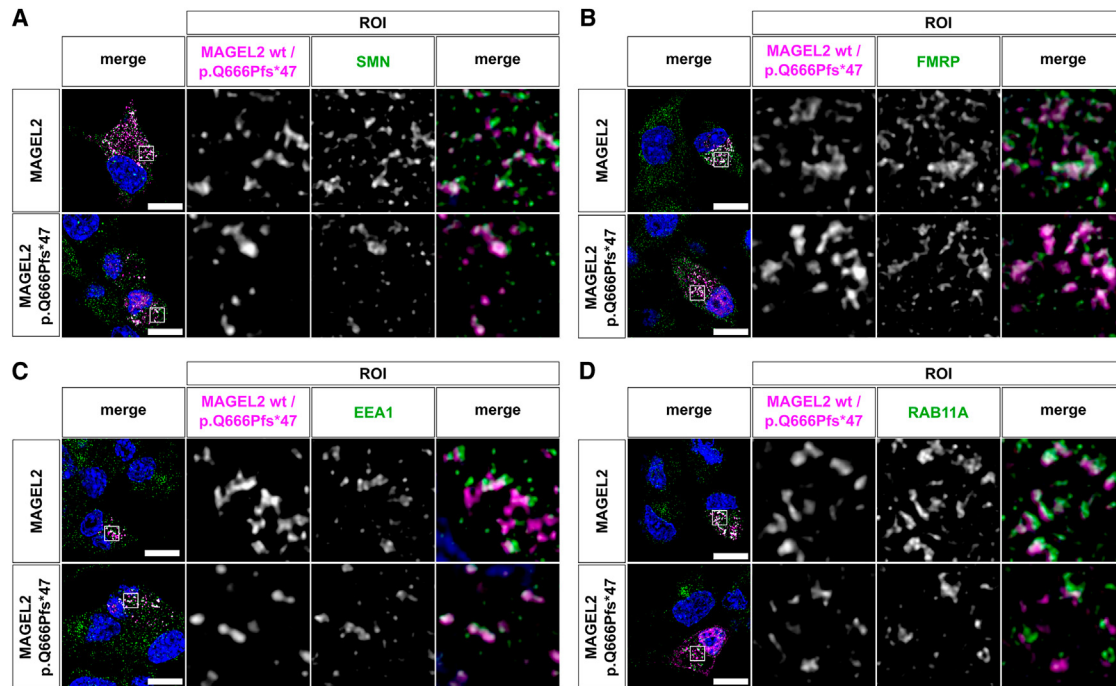
### Protein interactomes of MAGEL2 and the SNORD116 snoRNA

Due to the phenotypic similarities between SYS and PWS and the genomic proximity of the respective affected genes, i.e., MAGEL2 and SNORD116,<sup>1</sup> we also sought to find a functional link between MAGEL2 and SNORD116 snoRNAs. Therefore, we employed an RNA-centric based pull-down approach, employing 3'-biotinylated SNORD116 (copy 3), the canonical SNORD100 snoRNA, and a scrambled RNA control (SCR) followed by LC-MS/MS or western blot analyses for identification of protein interaction partners (Figure 5).

From these analyses, a direct interaction between SNORD116 and WT MAGEL2 or p.Gln666Profs\*47 was not corroborated since WT and mutant MAGEL2 were present in all co-IP samples including SNORD100 and SCR controls, which might indicate a more general affinity of mutant and WT MAGEL2 to various RNA species (Figure S2). Because SNORD100 and the SCR control sample were found to be more efficiently 3'-biotinylated than SNORD116 (Figure S3), we decided to analyze proteins present in at least two of three experiments by LC-MS/MS and subtracted proteins identified in the SCR sample. Thereby, we aimed to exclude non-specific RNA binding proteins (Figure 5A).

GO term analysis revealed that box C/D snoRNP assembly and ribosome biogenesis are within the top 5 GO biological processes of the 97 proteins bound to SNORD116 and canonical SNORD100 and validated our pull-down





**Figure 4. MAGEL2 WT and p.Gln666Profs\*47 co-localize with EEA1, RAB11, SMN and FMRP in mouse Neuro-2a cells**

Mouse Neuro-2a cells were transiently transfected with WT mCherry-MAGEL2 or mCherry-p.Gln666Profs\*47 (magenta) fusion proteins and subcellular localization was analyzed employing confocal immunofluorescence microscopy 24 h after transfection. Nuclei were counterstained with DAPI (blue). The scale bar is 10  $\mu$ m. SMN (A), FMRP (B), endogenous early endosomes (C), and recycling endosomes (D) were visualized with anti-SMN, anti-FMRP, anti-EEA1, and anti-RAB11A antibodies (green), respectively. A white signal in the overlap images indicated signal overlap or co-localization. At least 3 transfected cells were imaged in three independent replicates and representative cells are depicted.

approach (Figure 5A and 5B). Moreover, GO term analysis of co-immunoprecipitated proteins from either *SNORD100* or *SNORD116* RNA-CoIPs resulted in the same top GO terms (Figure 5B and 5C), which indicates that orphan *SNORD116* snoRNAs assemble into canonical snoRNPs.

To more precisely pinpoint specific *SNORD116* protein interaction partners, the RNA co-IP was also performed employing mouse N2a cell lysates (Figure S4). Among the ten common interaction partners co-immunoprecipitated with both snoRNAs (*SNORD116* and *SNORD100*) in both cell types, we identified box C/D snoRNP assembly proteins NUFIP1, RUVBL1, and RUVBL2 (Figures S4A and S4C).<sup>75</sup> In contrast, specific *SNORD116* interactors, overlapping in human and mouse, were identified as MRPL1, WDR82, IGHMBP2, KHDRBS1, STRAP, and NXF1 (Figures S4A and S4B). In summary, our data did not indicate an overlapping *SNORD116* and MAGEL2 interactome, as no common proteins could be identified by the MAGEL2 BioID and GFP-CoIP and by the *SNORD116* RNA-CoIP (Figures 5C and 5D).

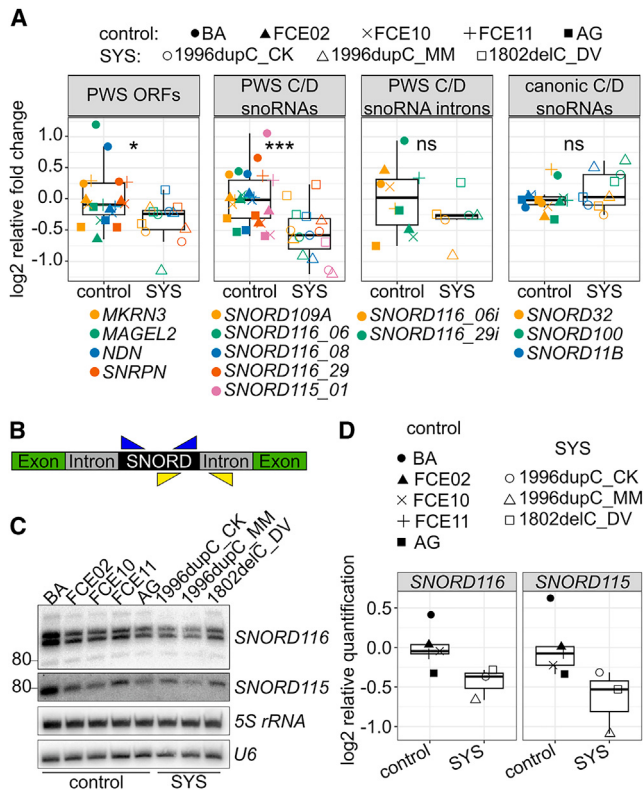
#### Mutations in MAGEL2 reduce expression of *SNORD116* snoRNAs in SYS proband cell lines

Even in the absence of a common interactome between MAGEL2 and *SNORD116* snoRNAs, our results do not necessarily exclude a common pathophysiological mechanism between SYS and PWS. Since we hypothesized a po-

tential involvement of MAGEL2 in transcriptional regulation and RNA stabilization, we assessed the influence of truncated MAGEL2 mutants on the expression of *SNORD116* snoRNAs involved in the etiology of PWS.<sup>6–10</sup> Hence, we employed RT-qPCR and northern blot analysis to investigate the abundance of *SNORD116* in small molecule neural progenitor cells (smNPCs) derived from healthy and SYS iPSCs, paternally harboring the WT or mutant (c.1996dupC or c.1802delC) *MAGEL2* alleles, respectively.

Indeed, RT-qPCR analysis revealed that expression of *SNORD116* ( $W = 110$ ,  $p = 0.0101$ ) (Figure S5A), as well as *SNORD115*, and of *SNORD109A* snoRNAs were downregulated by 31%, 42%, and 28% in smNPCs, respectively (Figure S5B). Collectively, the abundance of these box C/D snoRNAs within the PWS gene locus was about 32% lower in SYS smNPCs ( $W = 313$ ,  $p = 0.000254$ ) (Figure 6A). Because of the high sequence similarity of the different *SNORD116* and *SNORD115* snoRNA copies within their respective gene location,<sup>6</sup> we were able to apply RT-qPCR analysis only to three or one copy of the *SNORD116* and *SNORD115* clusters, respectively (Figures 6A, S5A, and S5B). Hence, we also investigated abundances of both snoRNAs by northern blot analysis employing specific probes which detected multiple copies of *SNORD116* (19 of 30) and *SNORD115* (30 of 49) (Figure 6C). In agreement with qPCR analysis, quantification of northern blot signals





**Figure 6. Influence of *MAGEL2* frameshift mutations on gene expression, assessed in SYS smNPCs compared to healthy controls**

(A) RT-qPCR analysis of genes of interest (GOI) within and outside chromosome 15q11–q13. Relative expression was calculated with primer efficiency normalized NO values (linRegPCR<sup>50</sup>). NO values were normalized to the geometric mean of housekeeping genes *TBP*, *GAPDH*, and *HPRT* and to the geometric mean of healthy control samples. Genes were combined into groups depending on type and genomic localization. Wilcoxon tests were applied employing rstatix.

(B) Scheme of primer position to detect snoRNAs before (yellow) and after (blue) splicing and processing.

(C) Northern blot analysis of 19 and 30 copies of *SNORD116* and *SNORD115*, respectively.

(D) Northern blot bands were quantified employing Fiji and normalized to geometric mean areas of *5S rRNA* and *U6* snRNA signals.

whereas the abundance of *NDN* (−2%) or *MAGEL2* (−8%) was unchanged or only slightly lower, respectively (Figure S5B).

We finally addressed the question of whether in smNPCs *MAGEL2* c.1996dupC and *MAGEL2* c.1802delC mRNAs might exhibit different sizes since both frameshift mutants harbor elongated 3′ UTRs of more than 2 kb (in contrast to 433 bases in the WT transcript). We hypothesized that the abundance of the intronless mutant *MAGEL2* transcripts would be decreased due to EJC-independent nonsense-mediated decay (NMD). This control mechanism assures the degradation of mRNAs exhibiting a premature stop codon resulting in unusually long (~1 kb) and unstructured 3′ UTRs.<sup>77</sup> However, by northern blot analysis, employing probes hybridizing either to the 3′ or 5′ end of

the *MAGEL2/MAGEL2* c.1996dupC transcripts, we showed that *MAGEL2* c.1996dupC and *MAGEL2* c.1802delC mRNAs are not subject to NMD. These findings are consistent with mRNA sizes of *MAGEL2* mutants being indeed identical to that observed for WT *MAGEL2* mRNA (Figure S6).

## Discussion

Since the discovery of truncated *MAGEL2* protein variants and their involvement in the etiology of the neurodevelopmental disorder SYS,<sup>30</sup> most of the past research has focused on their molecular functions and their involvement in disease.<sup>36,40,66,78,79</sup> Due to the phenotypic similarities between SYS and PWS,<sup>1</sup> we sought to elucidate possible links between both diseases at a molecular level, i.e., to link *MAGEL2* mutations to genes involved in PWS.

Our data show an accumulation of the predominant p.Gln666Profs\*47 variant within the nucleus in HEK 293T and HeLa cells, whereas WT *MAGEL2* accumulated within the nucleus only upon CRM1 inhibition. This suggests that WT *MAGEL2* is dynamically shuttled between the cytoplasm and the nucleus. These observations imply functions of *MAGEL2* within the nucleus and are corroborated by our proteomics data, which link *MAGEL2* to regulators of transcription. This also suggests yet undescribed functions of both protein variants within the nucleus. Previously, the group by Urreizti and coworkers described a predominant nuclear localization of a truncated *MAGEL2* variant.<sup>80</sup> In another study, Centeno-Pla et al.<sup>81</sup> have recently reported a predominant nuclear localization of six truncated *MAGEL2* variants. Among these, the variants with the strongest nuclear accumulation associated with arthrogyrosis multiplex congenita (AMC [MIM: 618947]), indicating a genotype-phenotype correlation dependent on the level of nuclear localization of truncated *MAGEL2*.<sup>81</sup> In summary, the above findings are thus consistent with increased nuclear localization of all truncated *MAGEL2* mutants investigated so far due to the loss of the predicted C-terminal NES motifs upon truncation of the protein.

In the cytoplasm of HEK293T cells, mCherry-tagged *MAGEL2* WT and p.Gln666Profs\*47 were found to be adjacent to endogenous RAB11A-positive recycling endosomes, whereas only a small fraction of both proteins was found proximate to endogenous EEA1-positive endosomes. In addition, we observed the truncated *MAGEL2* protein to be organized in vesicular-like structures in the cytosol. In contrast, in N2A cells, both *MAGEL2* variants showed a strong colocalization with EEA1 and RAB11A indicating that the mechanisms controlling the subcellular localization of *MAGEL2* might be partially cell type specific. A recent study showed that F-actin and ArpC5 were reduced in retromer subunit VPS35-positive endosomes in PWS and SYS stem cell-derived neurons.<sup>79</sup> Together with our data, this indicates that the N-terminal



region of MAGEL2 might be required for recognition and binding by VPS35, which also recruits the SHRC complex through an interaction with FAM21.<sup>36</sup> Hence, our findings indicate that MAGEL2 association with the retromer might persist even in the presence of the truncated MAGEL2 mutants, thereby displaying compromised MUST assembly (i.e., loss of the functional U7BS and MHD regions).

Since SYS phenotypes are not resembled by a complete loss of the MAGEL2,<sup>32,82,83</sup> a few possible scenarios have been raised in the past to explain the SYS disease. On the one hand, it was proposed that transcription of the imprinted maternal allele might become leaky upon loss of the paternal allele.<sup>1,32,84</sup> On the other hand, truncated proteins have been linked to gain-of-function phenotypes<sup>85,86</sup> and also neomorphic effects have been proposed for truncated MAGEL2 variants.<sup>1</sup> Consistent with these observations, our results suggest gain or loss of cellular functions of truncated MAGEL2 mutants within the nucleus or the cytoplasm, respectively. In addition, WT MAGEL2 dynamically shuttles between the cytoplasm and the nucleus, and therefore might indicate novel functions within the nucleus.

MAGEL2 is an intrinsically disordered protein (IDP) exhibiting over 85% of intrinsically disordered protein regions (IDPR).<sup>40</sup> Previous studies have shown that highly structured proteins are related to integral membrane proteins or enzymes, while signaling and regulation—including differentiation, transcription regulation, DNA condensation, and mRNA processing—are mainly orchestrated by intrinsically disordered proteins.<sup>87,88</sup>

Sanderson and coworkers have conducted a biotin proximity-labeling approach employing HEK293 cells stably expressing the full-length or a C-terminal MAGEL2 mutant.<sup>40</sup> In these analyses they have demonstrated that the N-terminal portion of MAGEL2 might be implicated in RNA metabolism processes. In a “subtractive” approach, they assigned hits that were found in full-length but not the C-terminal MAGEL2 conditions as N-terminal interaction partners.

By studying interaction partners of MAGEL2 WT and p.Gln666Profs\*47 employing BioID and CoIP, our GO term analysis revealed their potential function in RNA stabilization as well as chromosome condensation. Taken together, these data suggest that the N-terminal half of MAGEL2 might be required for the interaction with regulators of RNA metabolism.<sup>40</sup> Association of KHSRP and FUBP3 with the N-terminal protein portion of MAGEL2 were shown in the present study and by Sanderson et al.<sup>40</sup> The absence of U7BS and MHD might lead to loss of function of the pathogenic truncated MAGEL2 variants, for example through the deregulation of targeted proteins by ubiquitination or de-ubiquitination.

The deubiquitinase USP7 was found as an interactor of MAGEL2 WT and the p.Gln666Profs\*47 variant, confirming the direct binding of those proteins as part of the MUST complex.<sup>66</sup> The interaction is decreased in p.Gln666Profs\*47 compared to WT MAGEL2, which sug-

gests that part of the SYS phenotype could potentially be related to decreased association of these two proteins. This is particularly interesting, as heterozygous loss-of-function variants of *USP7* are the cause of Hao-Fountain syndrome (HAFIOUS [MIM: 616863]), a neurodevelopmental disorder that clinically manifests developmental delay/intellectual disability, autism spectrum disorder, muscular hypotonia.<sup>89</sup> This suggests that the two genes and conditions represent not only a molecular but also a pathophysiological and clinical continuum.<sup>89</sup>

Some proteins, such as ENO1, GAPDH, PKM, and LDHA, have been previously identified as binding partners for the C-terminal MAGEL2 variant,<sup>40</sup> while in our study these proteins were enriched in GFP-MAGEL2 p.Gln666Profs\*47 by co-immunoprecipitation. Discrepancies between the results from the two studies might arise from the different experimental setups. Sanderson and colleagues<sup>40</sup> generated a C-terminal MAGEL2 variant which exhibited a 62 amino acid overlap with the C terminus of the truncated MAGEL2 p.Gln666Profs\*47 protein. Hence, this portion of the protein might harbor binding sites for the above-mentioned proteins.

Based on our study, it is unlikely that MAGEL2 and *SNORD116* share a common interactome or directly form an RNP complex. Still, we were able to identify interaction partners for both MAGEL2 variants as well as for *SNORD116*. WT and truncated MAGEL2 were found both as potential interaction partners of SMN, FMRP, KHSRP, and FUS. Intriguingly, SMN was reported to associate with FMRP,<sup>90,91</sup> KHSRP,<sup>92</sup> and FUS.<sup>90,93,94</sup>

SMN has been shown to shuttle between the cytoplasm and the nucleus,<sup>95</sup> which we also observed for the WT MAGEL2 protein, and has been implicated in snRNP assembly, mRNA trafficking, and local translation, and was also observed to influence mitochondria and bioenergetic pathways.<sup>93</sup> Mutations or the homozygous loss of *SMN1* have been reported to be the main cause of spinal muscular atrophy (SMA [MIM: 253300]).<sup>96–98</sup>

Silencing of *FMR1* by DNA methylation of expanded CGG triplet repeats within the 5' UTR leads to the loss of FMRP and causes the fragile X syndrome (FXS [MIM: 300624]).<sup>99,100</sup> FMRP (fragile-X-messenger ribonucleoprotein) is an RNA-binding protein, which was initially shown to inhibit translation *in vitro* and in *Xenopus laevis* oocytes.<sup>101</sup> It was recently described to regulate activity-dependent local translation in CA1 neurons by repressing translation<sup>102</sup> and thereby is a crucial regulator in neuronal mRNA metabolism, neuronal development, axonal growth, and synaptic plasticity.<sup>103,104</sup>

KHSRP (also designated as FUBP2) functions as a DNA- and RNA-binding protein which was previously described to be involved in degradation of AU-rich element (ARE)-containing mRNAs and splicing.<sup>74,105,106</sup> Recent findings by Olguin and colleagues indicated that KHSRP is a regulator of brain development and function.<sup>74</sup> Therefore, MAGEL2 might be an integral component of these respective complexes.



Due to phenotypic similarities, SMA should be considered as a differential diagnosis for SYS<sup>33</sup> and PWS.<sup>107</sup> Moreover, intellectual disability, behavioral symptoms, and autism spectrum disorder are shared symptoms of FXS<sup>99,100</sup> and SYS.<sup>1,31–33</sup> It is noteworthy that there are also case reports of individuals with FXS exhibiting a Prader-Willi phenotype (i.e., hyperphagia and obesity), which was likely caused by reduced cytoplasmic FMRP interacting protein (CYFIP1) expression.<sup>108</sup> Vice versa, haploinsufficiency of *Cyfp1* and *Fmr1* knockout mice have been shown to share a common phenotype.<sup>109</sup> Interestingly, heterozygous loss of *CYFIP1* appears in individuals with type I deletions of the PWS critical region, underlining the complexity between the phenotypic similarities of PWS and FXS.<sup>109</sup> Consequently, downstream molecular pathways of the potential MAGEL2 targets FMRP and SMN might be negatively regulated in SYS-affected individuals.

Besides modulating the (de-)ubiquitination of WASH,<sup>36</sup> MAGEL2 has also been shown to control the circadian rhythm via cryptochrome 1 (CRY1) stabilization mediated by USP7-dependent de-ubiquitination.<sup>78</sup> In addition, previous studies revealed that the ubiquitin proteasome system (UPS) targets and degrades SMN<sup>110</sup> and FMRP through a phosphorylation-induced ubiquitination cascade.<sup>111</sup> Together with our data on the specific co-immunoprecipitation and co-localization of SMN and FMRP with WT and truncated MAGEL2 variants, we suggest a model where MAGEL2 regulates the stability of both proteins. Future studies should examine whether the truncation of MAGEL2 leads to decreased levels of SMN and FMRP. This might indeed explain some of the common phenotypes observed between individuals with SYS and FXS<sup>1,31–33,99,100</sup> or between individuals with SYS and SMA.<sup>33</sup>

Utilizing RNA-CoIP, we were able to identify protein interaction partners for *SNORD116* RNAs which differed from canonical *SNORD* snoRNA proteins.<sup>14</sup> In these analyses, IGHMBP2 and STRAP specifically co-immunoprecipitated with *SNORD116*. Mutations in IGHMBP2 are linked to spinal muscular atrophy, type 1 (SMARD1 [MIM: 604320]),<sup>112–117</sup> which shows symptoms such as hypotonia, feeding problems, and poor suck.<sup>118</sup> In addition, more than 30% and 50% of individuals with PWS die of respiratory failure<sup>119</sup> or respiratory diseases,<sup>120</sup> respectively. This links PWS to SMARD1, where life-threatening respiratory distress/respiratory failure was shown to appear within 13 months of age.<sup>112</sup>

MAGEL2 WT and p.Gln666Profs\*47 both interacted with RNA stabilization associated factors YBX1, ELAVL1, TAF15, PABPC1, and FUS. RNA stabilizers function mainly in post-transcriptional control of RNAs, ranging from splicing, polyadenylation, mRNA stabilization, and mRNA localization to translation.<sup>121</sup> For example, the m<sup>5</sup>C reader YBX1 recruits ELAVL1 and thereby preserves its target mRNA from degradation.<sup>122</sup> Strikingly, m<sup>5</sup>C was reported as an important dynamically regulated RNA modification during brain development and is enriched

in genes for synaptic plasticity regulation.<sup>123</sup> Our data thus indicate that MAGEL2 might also act as an essential component in RNA stabilization, possibly by fine-tuning the interaction of the m<sup>5</sup>C reader YBX1 with ELAVL1 and thereby affecting neuronal plasticity. Interestingly, ELAVL1 was also found to bind to *SNORD115* from the PWS critical region.<sup>124</sup>

By linking of MAGEL2 to gene expression and RNA stabilization, we hypothesized that mutations in *MAGEL2* might lead to altered gene expression or transcript abundance in SYS smNPCs (exhibiting c.1996dupC and c.1802delC mutations), in particular within the PWS critical region. Indeed, we observed reduced transcript abundances of most genes from the PWS region, in particular *SNORD116* (i.e., a reduction by about 31%). In addition, RNA levels of *SNORD115*, *SNORD109A*, *MKRN3*, and *SNRPN* were also decreased. A fairly moderate reduction of transcript abundances for RNA transcripts from the PWS region might not be considered sufficient for the observed phenotypes in SYS. However, as demonstrated in the case of miRNAs, downregulation of expression of their mRNA targets has been reported to be within a similar range.<sup>125</sup> This indicates that even moderate changes in mRNA expression can result in severe disease phenotypes, in particular if several genes are affected. In addition, the moderate reduction of *SNORD116* abundance in probands with SYS, compared to individuals with PWS, might explain at least some of the missing clinical features not observed in SYS, such as severe obesity.

In contrast to the above genes, abundances of *NDN* from the PWS critical region and canonical box C/D snoRNAs located outside the PWS region were found not to be affected in SYS smNPCs. The observed reduced abundance of transcript levels of genes from the PWS critical region might thus arise, for example, from an aberrant interaction of mutant MAGEL2 proteins with regulators of transcription or RNA stability.

MAGEL2 mRNA levels differed only slightly between SYS and healthy smNPCs. Together with the identification of novel protein interaction partners, the stability of the truncated MAGEL2 mRNA in SYS proband cell lines as well as its expression in transfection experiments underline that this variant likely leads to gain-of-function effects in probands with SYS, as also suggested previously.<sup>1</sup> We thus propose that the preferential sequestration of the truncated variant to the nucleus plays a fundamental role in the pathophysiology of the disease.

Recent case reports have described individuals with PWS exhibiting atypical microdeletions within 15q11–q13 whose *SNORD116* cluster was not deleted.<sup>126,127</sup> In contrast, several previous studies have reported that an entire deletion of the *SNORD116* cluster was sufficient to cause PWS.<sup>6–10</sup> These contradictory case reports underline the complexity of genes involved in the etiology of PWS. Our study demonstrates functions of MAGEL2 in the regulation of RNA metabolism, especially of genes within the PWS critical region. It thus might be conceivable that in

PWS-affected individuals with atypical microdeletions which do not result in the deletion of the *SNORD116* cluster,<sup>126,127</sup> the expression of *SNORD116* might still be downregulated. Therefore, we propose that examination of *SNORD116* transcript abundance by RT-qPCR might be included as a tool for the diagnosis of PWS in the future, e.g., from NPCs differentiated from fibroblasts derived from individuals with PWS.

## Conclusion

In summary, we have provided evidence that MAGEL2 shuttles between the cytoplasm and the nucleus and that it interacts with proteins involved in transcription regulation and RNA metabolism. Consistent with these findings, we observed the abundances of genes expressed from the PWS critical region, including *SNORD116*, to be reduced in SYS smNPCs harboring the c.1996dupC and c.1802delC *MAGEL2* mutations. Hence, MAGEL2 might indeed be involved in the transcription of genes from the PWS locus thus resembling some but not all features of PWS. In addition, among other proteins, we identified SMN and FMRP proteins as binding partners for MAGEL2 variants, both causing diseases linked to known phenotypic features of individuals with SYS or PWS. Hence, although the precise roles of *SNORD116* and MAGEL2 protein complexes and their pathways in the etiology of SYS and PWS yet remain to be fully determined, our data provide a first step in the elucidation of their role in these pathologies, in particular in their relation to other motor neuron diseases. A perturbed interaction of mutant MAGEL2 with interaction partners such as SMN and FMRP and/or its influence on *SNORD116* expression might thus explain some of the observed phenotypes in SYS and PWS diseases. Hence, our study sets the stage for further investigation into the molecular mechanisms of both diseases and might aid in the generation of therapeutic drugs for future therapeutic options.

## Data and code availability

The datasets generated during this study are available via figshare (doi: <https://doi.org/10.6084/m9.figshare.c.7217859>). If there is interest in the raw Airyscan immunofluorescence images, please contact the corresponding author.

## Supplemental information

Supplemental information can be found online at <https://doi.org/10.1016/j.ajhg.2024.05.023>.

## Acknowledgments

This research was funded by the Austrian Science Funds (FWF ZFP326120 and FWF TAI 411-B) to A.H. We would like to thank Dr. Martin Offterdinger (Institute of Biooptics, Medical University of Innsbruck) for his technical support. We also thank Beate Abt and Alexander Kühbacher (Institute of Molecular Biology, Medical

University of Innsbruck) for technical support. We thank Matthias Erlacher, Utz Fischer, and Andreas Janecke for helpful discussions and critically reading the manuscript. We thank Anna and her parents for inspiration to initiate this study. We dedicate this paper to Jürgen Brosius, on the occasion of his 75th birthday, and Harry Noller, both leading pioneers in RNA research.

## Author contributions

A.H., conceptualization, funding acquisition, supervision, writing – review & editing, project administration, validation; D.H., investigation, writing – original draft, formal analysis, data curation, visualization, validation, writing – review & editing; A.V., investigation, validation, writing – review & editing; A.E., resources, writing – review & editing; A.S., resources, writing – review & editing; M.S.-C., resources, writing – review & editing; H.F., validation, writing – review & editing; V.R., investigation, verification, writing – review & editing; M.S., resources, writing – review & editing; C.P.S., resources, writing – review & editing; L.A.H., validation, writing – review & editing; L.K., resources, writing – review & editing; B.S., resources, writing – review & editing; F.E., resources, writing – review & editing; S.G., validation, writing – review & editing; M.E.G.d.A., validation, writing – review & editing. All authors contributed to editing and revising and approved the final version manuscript.

## Declaration of interests

The authors declare no competing interests.

Received: April 16, 2024

Accepted: May 29, 2024

Published: June 21, 2024

## Web resources

enrichR, <https://cran.r-project.org/web/packages/enrichR/index.html>

GenBank, <https://www.ncbi.nlm.nih.gov/genbank/>

Ggpubr, <https://CRAN.R-project.org/package=ggpubr>

OMIM, <https://www.omim.org/>

R, <https://www.R-project.org/>

R Studio, <http://www.rstudio.com/>

Rio, <https://cran.r-project.org/web/packages/rio/index.html>

rstatix, <https://CRAN.R-project.org/package=rstatix>

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