Altered Expression of Ion Channels in White Matter Lesions of Progressive Multiple Sclerosis: What Do We Know About Their Function?

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Despite significant advances in our understanding of the pathophysiology of multiple sclerosis (MS), knowledge about contribution of individual ion channels to axonal impairment and remyelination failure in progressive MS remains incomplete. Ion channel families play a fundamental role in maintaining white matter (WM) integrity and in regulating WM activities in axons, interstitial neurons, glia, and vascular cells. Recently, transcriptomic studies have considerably increased insight into the gene expression changes that occur in diverse WM lesions and the gene expression fingerprint of specific WM cells associated with secondary progressive MS. Here, we review the ion channel genes encoding K+, Ca2+, Na+, and Cl− channels; ryanodine receptors; TRP channels; and others that are significantly and uniquely dysregulated in active, chronic active, inactive, remyelinating WM lesions, and normal-appearing WM of secondary progressive MS brain, based on recently published bulk and single-nuclei RNA-sequencing datasets. We discuss the current state of knowledge about the corresponding ion channels and their implication in the MS brain or in experimental models of MS. This comprehensive review suggests that the intense upregulation of voltage-gated Na+ channel genes in WM lesions with ongoing tissue damage may reflect the imbalance of Na+ homeostasis that is observed in progressive MS brain, while the upregulation of a large number of voltage-gated K+ channel genes may be linked to a protective response to limit neuronal excitability. In addition, the altered chloride homeostasis, revealed by the significant downregulation of voltage-gated Cl− channels in MS lesions, may contribute to an altered inhibitory neurotransmission and increased excitability.

Keywords: multiple sclerosis, progressive, white matter, lesions, ion channels, transcriptome
INTRODUCTION

Multiple sclerosis (MS) is an inflammatory demyelinating disease of the central nervous system (CNS) affecting more than 2 million people worldwide. MS lesions in CNS white matter (WM) are multiple focal areas of myelin loss accompanied by inflammation, gliosis, phagocytic activity, and axonal damage (Compston and Coles, 2008; Kuhlmann et al., 2017; Filippi et al., 2018; Rommer et al., 2019). Available MS therapies have little benefit for secondary-progressive MS (SPMS) patients, who develop progressive disability after a disease course characterized by inflammatory attacks. Therefore, promoting neuroprotection and remyelination are important therapeutic goals to prevent irreversible neurological deficits and permanent disability.

Ion channels play a fundamental role in maintaining WM integrity and regulating function of axons, interstitial neurons (Sedmak and Judas, 2021), glia, and vascular cells. Dysregulation of ionic homeostasis in the WM during demyelination is decisive for axonal damage and cell death and may interfere with tissue repair processes (Boscia et al., 2020). Furthermore, MS may involve an acquired channelopathy (Waxman, 2001; Schatting et al., 2014). Hence, selectively targeting ion channels in WM represents an attractive strategy to overcome axonal and glial impairment and prevent disease progression.

Recently, transcriptomic studies have considerably increased our insight into gene expression changes occurring in the MS brain (Elkjaer et al., 2019; Jakel et al., 2019; Schirmer et al., 2019). Aiming at identifying the ion channel genes governing WM dysfunction in SPMS brain, we analyzed the recent bulk RNA-sequencing (RNA-seq) datasets by using the MS-Atlas (Elkjaer et al., 2019; Frisch et al., 2020). We put a special emphasis on the distribution of shared and unique genes encoding ion channels in chronic active (CA), active (AL), inactive (IL), and remyelinating (RL) lesions, and normal-appearing white matter (NAWM) compared to control WM (Figures 1A,B, Table 1). We identified uniquely expressed ion channel genes: 34 genes in CA, 9 in IL, 1 in AL, as well as 2 genes in all lesions and NAWM (Figures 1, 2, Table 1). The CA lesions displayed the highest number of upregulated ion channels genes while downregulated ion channels genes were more consistently found in ILs (Figure 1C). Next, we explored recent single-nuclei RNA-seq (snRNA-seq) datasets to identify the expression of dysregulated ion channel genes in cell clusters in the WM of control and SPMS brain (Jakel et al., 2019; Tables 1, 2, Figure 3).

The goal of the present review is to discuss the current knowledge on the expression and function of ion channels that turned out to be significantly and uniquely dysregulated in WM lesions of SPMS brain. We summarize the information in the context of human MS and the related experimental models (Tables 1–3, Figure 4).

K⁺ CHANNELS

Voltage-Gated K⁺ Channels (Kᵥ)

Kᵥ channels are composed of four α-subunits that assemble as homo- or hetero-tetramers to form a membrane pore. Forty human genes encode for Kᵥ α-subunits representing 12 families. Kᵥ1–Kᵥ4 (Shaker, Shaw, Shab, and Shal), Kᵥ7 (KCNQ), and Kᵥ10–Kᵥ12 (eag, erg, and elk) α-subunits produce functional channels, while Kᵥ5, Kᵥ6, Kᵥ8, and Kᵥ9 fail to produce currents when expressed alone in heterologous expression system and are considered modulatory subunits for Kᵥ2-subfamily. The diversity of Kᵥ channels is further increased by the ability of α-subunits to combine with auxiliary subunits, which regulate gating properties.

**Kᵥ1.1, Kᵥ1.2, and Kᵥ1.4 (KCNA1, KCNA2, and KCNA4)**

Kᵥ1.1, Kᵥ1.2, and Kᵥ1.4 (KCNA1, KCNA2, and KCNA4) KCN genes encode for low-threshold voltage-activated Kᵥ (Kᵥ1.1–1.8) channels, of which Kᵥ1.1–Kᵥ1.6 are expressed in the brain (Chittajallu et al., 2002; Vautier et al., 2004; Vacher et al., 2008; Rasmussen and Trimmer, 2019). Kᵥ1 channels display little/no inactivation, resulting in sustained delayed rectifier K⁺ currents, with the exception of Kᵥ1.4, which underlies transient A-type K⁺ current.

**Neurons**

Kᵥ1.1 expression is highest in the brainstem, while Kᵥ1.4 > Kᵥ1.2 represent the main Kᵥ1 subunits in the hippocampus (Trimmer, 2015). Kᵥ1.1 channels, in association with Kᵥ1.2, cluster in the juxtaparanodal regions of axons under the myelin sheath and regulate action potential (AP) propagation and neural excitation (Wang et al., 1993; Trimmer and Rhodes, 2004; Ovsepian et al., 2016). Mutations of Kᵥ1 channels result in hyper-excitability, episodic ataxia, myokymia, and epilepsy (Allen et al., 2020).

**Glia**

Mouse astrocytes express low levels of Kᵥ1.1, Kᵥ1.2, and Kᵥ1.4 transcripts (Smart et al., 1997), but Kᵥ1.2 and Kᵥ1.4 expression is high in reactive rat astrocytes (Akhtar et al., 1999). Kᵥ1.1 transcripts and proteins are highly expressed in C6 glioma cells. Rodent oligodendrocyte precursor cells (OPCs) express Kᵥ1.1, Kᵥ1.2, and Kᵥ1.4 transcripts (Attali et al., 1997; Chittajallu et al., 2002; Falcao et al., 2018; Batiuk et al., 2020) but only Kᵥ1.4 and low level of Kᵥ1.2 proteins (Attali et al., 1997; Schmidt et al., 1999). In OPCs and astrocytes, the Kᵥ1 subunits regulate cell growth and cell cycle progression, e.g., Kᵥ1.4 overexpression in vitro increases OPCs proliferation (Schmidt et al., 1999) while deletion decreases it (Gonzalez-Alvarado et al., 2020). Recent RNA-seq did not detect Kᵥ1.1, Kᵥ1.2, and Kᵥ1.4 in mouse microglia (Hammond et al., 2019), but earlier studies found Kᵥ1.1 and Kᵥ1.2 mRNAs and/or proteins in BV2 microglia,
Expression and Function in MS

Bulk RNA-seq found upregulation of Kv1.1, Kv1.2, and Kv1.4 transcripts in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq detected significant Kv1.2 expression in neuronal clusters, slight increase of Kv1.4 transcripts in neuronal but not glial clusters, and no Kv1.1 transcript (Tables 1, 2; Jakel et al., 2019).

The CA lesion is characterized by ongoing tissue damage and, functionally, Kv1.2 upregulation in CA lesions may be a hallmark of axonal damage. While recent data found that KCNA1 gene is downregulated during demyelination in the cuprizone model (Martin et al., 2018), in animal models of MS, Kv1.2 (and also Kv1.1) ectopically redistributes to nodes and internodes of WM axons (McDonald and Sears, 1969; Wang et al., 1995; Sinha et al., 2006; Jukkola et al., 2012; Zoupi et al., 2013; Kasrati et al., 2015), while in human MS, the dislocation of Kv1.2 channels is associated with paranodal pathology, particularly in NAWM regions, and contributes to axonal dysfunction (Howell et al., 2010; Gallego-Delgado et al., 2020). The upregulated and redistributed Kv1.2 and Kv1.1 channels may hyperpolarize the axonal resting membrane potential (Vrest), elevate the amount of depolarization necessary for AP initiation, and impair AP conduction (Wang et al., 1995; Sinha et al., 2006; Jukkola et al., 2012). Pharmacological inhibition of Kv1.1 and Kv1.2 channels, e.g., with 4-aminopyridine, enhances axonal conduction and improves MS symptoms (Lugaresi, 2015).

It is difficult to speculate regarding Kv1.4 function in MS because data are not consistent. In animal models of MS and spinal cord injury (SCI), this developmentally restricted subunit re-appears/increases in OPCs, OLs, and astrocytic processes around lesion sites (Herrero-Herranz et al., 2007; Jukkola et al., 2012), but not in WM axons or microglia (Edwards et al., 2002; Jukkola et al., 2012). Mice lacking Kv1.4 exhibit reduced myelin loss in the spinal cord WM during EAE but no change of demyelination/remyelination in the corpus callosum in the cuprizone model (Gonzalez-Alvarado et al., 2020). However, it is unclear whether function of Kv1.4 subunits is relevant for glial cells in human MS because snRNA-seq barely detected Kv1.4 transcripts in glia clusters (Table 2).

Kv2.1 and Kv2.2 (KCNB1 and KCNB2)

Kv2 channels (encoded by KCNB1 and KCNB2 genes) mediate high-voltage-activated slowly inactivating delayed rectifier K+ currents (Guan et al., 2007). Kv2.1 channels can assemble with electrically silent Kv5 subunits, resulting in greater variability of Kv2 currents (Guan et al., 2007; Trimmer, 2015; Johnson et al., 2019).

Neurons

High-density clusters of Kv2.1 and Kv2.2 localize to soma, proximal dendrites, and axonal initial segment (AIS). Kv2
TABLE 1 | Expression and distribution of unique and overlapping genes coding for ion channels within SPMS lesions.

| Protein | Gene | Bulk lesion* | Fold change Up (+)/down (−) regulated (compared to control WM)* | Current type/conductance | Highly expressed in WM clusters of human brain* |
|---------|------|--------------|---------------------------------------------------------------|--------------------------|-----------------------------------------------|
| **K⁺ channels** | | | | | |
| Kv 1.1 | KCNA1 | CA | +1.42 | Delayed rectifier | |
| Kv 1.2 | KCNA2 | CA | +1.06 | Delayed rectifier | neuron2 |
| Kv 1.3 | KCNA3 | AL, CA, IL | +1.67 (AL); +1.35 (CA); +1.34 (IL) | Delayed rectifier | |
| Kv 1.4 | KCNA4 | CA | +1.34 | A-type | |
| Kv 1.5 | KCNA5 | AL, RL | +0.86 (AL); +1.36 (RL) | Delayed rectifier | |
| Kv 2.2 | KCNB2 | CA | +1.56 | Delayed rectifier | Neuron1, 2, 3, 4, 5 |
| Kv 2.1 | KCNB1 | CA | +1.26 | Delayed rectifier | Neuron1, 2, 3 |
| Kv 3.3 | KCNC3 | CA | +0.87 | A-type | |
| Kv 3.4 | KCNC4 | AL, IL | +0.81 (AL); +0.72 (IL) | A-type | |
| Kv 4.2 | KCND2 | CA | +0.95 | A-type | OPC, COP, neuron1,3 |
| Kv 4.3 | KCND3 | AL, CA, IL | +0.63 (AL); +0.86 (CA); +0.93 (IL) | A-type | neuron1, 2, 3 |
| Kv 6.1 | KCNQ1 | AL, RL | +2.72 (AL); +3.7 (RL) | Modifier of Kv2 | |
| Kv 7.1 | KCNQ1 | AL, CA | +0.91 (AL); +0.75 (CA) | M-type | |
| Kv 7.2 | KCNQ2 | CA | +0.75 | M-type | neuron1, 2 |
| Kv 7.3 | KCNQ3 | CA | +0.85 | M-type | ImOGLs, neuron1, 2, 3, 5, microglia/macrophages |
| Kv 7.4 | KCNQ4 | AL, CA, IL, RL | +1.19 (AL); +0.92 (CA); +1.36 (IL); +2.22 (RL) | M-type | |
| Kv 7.5 | KCNQ5 | CA | +1.69 | M-type | Neuron1, 2, 3, 5 |
| Kv 8.1 | KCNQ6 | CA | +1.48 | | |
| Kv 9.2 | KCNQ7 | CA | +0.90 | | |
| Kv 9.3 | KCNQ8 | AL, IL, RL, NAWM | −2.72 (AL); −1.5 (IL); −1.98 (RL); −0.71 (NAWM) | Modifier of Kv2 | |
| Kᵥ 10.1/EAG1 | KCNH1 | CA, IL | +0.81 (CA); +0.93 (IL) | Delayed rectifier | Neuron1, 2, 3 |
| Kᵥ 10.2/EAG2 | KCNH1 | CA, IL | +1.38 | Delayed rectifier | Neuron2 |
| Kᵥ 11.3/ERG3 | KCNH7 | CA | +1.38 | Delayed rectifier | Neuron1, 2, 3, 5 |
| Kᵥ 12.1/ELK1 | KCNH8 | AL, CA, IL, RL, NAWM | −1.25 (AL); −1.4 (CA); −2.05 (IL); −2.38 (RL); −0.62 (NAWM) | Delayed rectifier | Oligo3, Oligo4, Oligo6 |
| TREK1 | KCNK2 | CA | +1.03 | Leak, two pore | |
| TWIK2 | KCNK6 | AL, IL | +1.57 (AL); +0.82 (IL) | Leak, two pore | |
| TREK2 | KCNK10 | AL | −0.66 | Leak, two pore | |
| Kᵥ 2.1 | KCNJ1 | AL, CA, IL, RL, NAWM | +0.69 (AL); +0.87 (CA); +0.7 (IL) | Calcium-Activated | OPC, neuron1, 2, 3, 5, microglia/macrophages |
| Kᵥ 2.3 | KCNJ3 | IL | −0.7 | Calcium-Activated | Astrocytes1 |
| Kᵥ 6.1 | KCNT1 | CA | +1.24 | Sodium-Activated | | |
| Kᵥ 11.2 | KCNT2 | CA, IL | +0.92 (CA); +1.15 (IL) | Sodium-Activated | Neuron1, 2, 3, pericytes, vascular smooth cells |
| Kᵥ 2.1 | KCNJ2 | AL, CA, IL, RL | −0.54 (AL); −0.48 (CA); −0.54 (IL); −0.92 (RL) | Inward rectifier | |
| Kᵥ 3.4 | KCNJ5 | AL, CA, RL, NAWM | +2.58 (AL); +1.56 (CA); +1.9 (RL); +1.53 (NAWM) | Inward rectifier | |
| Kᵥ 3.2 | KCNJ6 | CA | +1.34 | Inward rectifier | Neuron1, 2, 3 |
| Kᵥ 6.1 | KCNJ8 | AL, IL | +0.74 (AL); +0.71 (IL) | Inward rectifier | |
| Kᵥ 3.3 | KCNJ9 | CA, RL | −0.52 (CA); −0.9 (FL) | Inward rectifier | |
| Kᵥ 4.1 | KCNJ10 | IL, RL | −1.06 (IL); −1.09 (FL) | Inward rectifier | Oligo5 |
| Kᵥ 5.1 | KCNJ16 | CA | +1.27 | Inward rectifier | |
| **Na⁺ channels** | | | | | |
| Naᵥ 1.1 | SCN1A | CA | +1.12 | TTX-sensitive | OPC, COP, neuron1, 2, 3, 4, 5 |

(Continued)
| Protein | Gene | Bulk lesion | Fold change Up (+)/down (−) regulated (compared to control WM) | Current type/ conductance | Highly expressed in WM clusters of human brain |
|---------|------|-------------|---------------------------------------------------------------|---------------------------|-----------------------------------------------|
| Na,1,2  | SCN2A| CA          | +1.1                                                          | TTX-sensitive             | Neuron1, 2, 3, 4, 5                          |
| Na,1,3  | SCN3A| CA          | +0.87                                                         | TTX-sensitive             | OPC, neuron1, 2, 3, 5                        |
| Na,1,6  | SCN8A| CA          | +1.15                                                         | TTX-sensitive             | Neuron1, 2, 3, 5                             |
| Na,1,9  | SCN11A| IL         | −1.16                                                         | TTX-resistant            |                                               |
| Ca²⁺ channels |  |             |                                                                |                           |                                               |
| Ca,1,2  | CACNA1C| CA        | +0.56                                                         | L-type                    | Neuron1, 2, 3, 5, pericytes                   |
| Ca,1,3  | CACNA1D| CA        | +0.57                                                         | L-type                    | Neuron1, 3                                  |
| Ca,2,1  | CACNA1A| CA        | +0.64                                                         | P/Q-type                  | OPC, neuron1, 2                              |
| Ca,2,3  | CACNA1E| CA        | +0.97                                                         | P/Q-type                  | Neuron1, 2, 5                               |
| Ca,3,1  | CACNA1G| IL         | +1.8                                                          | T-type                    |                                               |
| Ca,3,2  | CACNA1H| CA        | +1.12                                                         | T-type                    |                                               |
| Ca,3,3  | CACNA1I| CA        | +1.03                                                         | T-type                    |                                               |
| Ryanodine |      |             |                                                                |                           |                                               |
| Ryr2    | RYR2 | CA         | +0.85                                                          | Ca²⁺ Release channel      | Neuron1, 2, 3                               |
| Ryr3    | RYR3 | IL         | −0.76                                                          | Ca²⁺ Release channel      | Astrocytes1                                  |
| TRP channels |  |             |                                                                |                           |                                               |
| TRPC1   | TRPC1| AL, IL, RL  | −0.5 (AL); −0.48 (IL); −0.85 (RL)                             | Ca²⁺-permeable cation channel |                                               |
| TRPM2   | TRPM2| IL         | +0.92                                                          | Ca²⁺-permeable cation channel |                                               |
| TRPM3   | TRPM3| IL, RL     | −1.09 (IL); −0.98 (RL)                                        | Ca²⁺-permeable cation channel | Astrocytes1, neuron1                         |
| TRPM6   | TRPM6| CA, IL, RL  | −0.99 (CA); −1.06 (IL); −1.08 (RL)                            | Ca²⁺-permeable cation channel |                                               |
| TRPP1   | PKD2 | IL         | −0.48                                                          | Ca²⁺-permeable cation channel |                                               |
| TRPP3   | PKD2L2| CA        | −0.58                                                          | Ca²⁺-permeable cation channel |                                               |
| TRPV1   | TRPV1| CA         | −1.04                                                          | Ca²⁺-permeable cation channel |                                               |
| TRPV3   | TRPV3| AL, CA, IL, RL | −0.51 (AL); −0.72 (CA); −0.5 (IL); −0.74 (RL) | Ca²⁺-permeable cation channel |                                               |
| TRPV5   | TRPV5| AL, CA, IL, RL | −1.4 (AL); −1.67 (CA); −1.72 (IL); −2.02 (RL) | Ca²⁺-permeable cation channel |                                               |
| TRPV6   | TRPV6| AL, CA, IL, RL, NAWM | −1.77 (AL); −1.97 (IL); −1.32 (CA); −2.23 (RL); 0.86 (NAWM) | Ca²⁺-permeable cation channel |                                               |
| Cl⁻ channels |  |             |                                                                |                           |                                               |
| CLC-2   | CLCN2| CA         | −0.57                                                          | Inward rectification      |                                               |
| CLC-4   | CLCN4| AL, IL, RL  | −0.79 (AL); −0.73 (IL); −1.03 (RL)                            | Cl⁻/H⁺ antiporter         |                                               |
| CLC-7   | CLCN7| CA         | −0.72                                                          | Cl⁻/H⁺ antiporter         |                                               |
| Connexins and pannexins |  |             |                                                                |                           |                                               |
| Cx43    | GJA1 | AL, CA, RL  | +1.53 (AL); +1.12 (CA); +1.19 (RL)                            | Monovalent and divalent ions | Astrocytes1, astrocytes2                     |
| Cx32    | GJB1 | AL, CA, IL, RL | −1.6 (AL); −1.5 (CA); −1.85 (IL); −2.44 (RL) | Monovalent and divalent ions | Oligo5                                       |
| CX37    | GJA4 | IL         | +1.19                                                          | Monovalent and divalent ions | Pericytes                                    |
| Cx47    | GJC2 | AL, CA     | −1.62 (AL); −1.74 (CA)                                        | Monovalent and divalent ions |                                               |

(Continued)
channels influence AP duration during high-frequency firing and regulate neuronal excitability (Guan et al., 2007). Kv2.1 mutations are associated with neonatal encephalopathy epilepsies and neurodevelopmental delays (Torkamani et al., 2014; Thiffault et al., 2015; de Kovel et al., 2017).

**Glia**

RNA-seq detected KCNB1 gene in mouse OPC and microglia (Falcao et al., 2018; Hammond et al., 2019).

**Expression and Function in MS**

Bulk RNA-seq revealed upregulation of Kv2.1 and Kv2.2 transcripts in CA lesions of SPMS brain (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq found Kv2.1 and Kv2.2 in neuronal clusters (Tables 1, 2; Jakel et al., 2019). During EAE, Kv2.1 protein expression was downregulated in spinal cord motor neurons (Jukkola and Gu, 2015). Remarkably, Kv2.1 channels exist as freely dispersed conducting channels, or form electrically silent somatodendritic clusters (Schulien et al., 2020). Upregulated clustered Kv2.1 channels promote functional coupling of L-type Ca2+ channels in plasma membrane to ryanodine receptors (RyRs) of the endoplasmic reticulum (ER) (Deutsch et al., 2012; Kirmiz et al., 2018; Vierra et al., 2019) and may modulate intracellular Ca2+ level contributing to cell damage, while dispersal of Kv2.1-clusters blocks apoptogenic K+ currents and provides neuroprotection (Sesti et al., 2014; Justice et al., 2017). Hence, to elucidate the functional role of Kv2 upregulation in MS (Table 1), it will be important to determine whether it reflects an increase in clustered or dispersed Kv2 channels.

Kv3.3 (KCNC3)

The KCNC3 gene encodes for the Kv3.3 subunit, which, together with Kv3.1, Kv3.2, and Kv3.4, belongs to the Kv3 channel subfamily (Shaw). The Kv3.3 and Kv3.4 mediate transient A-type K+ currents, while Kv3.1 and Kv3.2 mediate sustained K+ currents.

**Neurons**

Kv3 channels localize to axonal and somatodendritic domains, and play a critical role in regulating AP firing at high frequency (Rasmussen and Trimmer, 2019). KCNC3 mutations result in spinocerebellar ataxia type-13 and cerebellar neurodegeneration (Rasmussen and Trimmer, 2019).

**Glia**

Cortical and hippocampal astrocyte cultures express Kv3.3 and Kv3.4 mRNAs and proteins (Bekar et al., 2005; Boscia et al., 2017). KCNC3 mRNA was detected in mouse OPCs and microglia (Larson et al., 2016; Falcao et al., 2018).

**Expression and Function in MS**

Bulk RNA-seq showed significant upregulation of Kv3.3 in CA lesions (Figure 2, Table 1), while snRNA-seq revealed its predominant distribution in neuronal clusters (Table 2; Jakel et al., 2019). Kv3.3 may play a detrimental role in MS because it increases in injured WM axons during EAE progression in mice and in human MS lesions (Jukkola et al., 2017), and the deletion of Kv3.1, which forms hetero-tetramers with Kv3.3, reduced EAE severity in mice (Jukkola et al., 2017).
FIGURE 2 | The expression profile of the ion channel genes uniquely expressed in different lesion types. (A) Left panel: The Venn diagram represents the number of overlapping and lesion-specific differentially expressed genes coding for ion channels in chronic active (CA), active (AL), inactive (IL), and remyelinating (RL) lesions and in normal-appearing white matter (NAWM) compared to control white matter. Right panel: The heatmap shows two genes, coding for ion channels KCNH8 and TRPV6 that are significantly altered in all lesion types compared to control white matter. Scale bar indicates fold changes. (B) The Venn diagram, the heatmap, and the scale bar show the single ion channel gene, KCNK10, which is uniquely downregulated in active lesion (AL). (C) The Venn diagram, the heatmap, and the scale bar show the eight genes coding for ion channels that are uniquely significantly differentially dysregulated in inactive lesion (IL). (D) The Venn diagram, the heatmap, and the scale bar show the 33 genes coding for ion channels that are significantly and differentially dysregulated compared to control white matter in chronic active lesion (CA). The red box in Venn diagrams marks the genes that are specifically dysregulated in the corresponding type of lesion.
| Protein | Gene      | Neuron | Astrocyte | OPC | COP | ImOLG | Oligo | Microglia | Pericyte |
|---------|-----------|--------|-----------|-----|-----|-------|-------|-----------|----------|
| K⁺ channels |           |        |           |     |     |       |       |           |          |
| Kv1.1   | KCNA1     | n.d    | n.d       | n.d | n.d | n.d   | n.d   | n.d       | n.d      |
| Kv1.2   | KCNA2     | +      | +/−       | +/− | +/− | −     | +/−   | −          | −        |
| Kv1.4   | KCNA4     | +/-    | −         | −   | −   | −     | −     | −          | −        |
| Kv2.1   | KCNB1     | +      | +/−       | +/− | +   | +/−   | +/−   | −          | +        |
| Kv2.2   | KCNB2     | ++     | −         | +/− | +/− | +/−   | −     | −          | −        |
| Kv3.3   | KCNC3     | +      | +/−       | −   | +/− | −     | +/−   | −          | −        |
| Kv4.2   | KCND2     | ++     | ++        | +++ | ++ | −     | +     | +/−       | −/−      |
| Kv7.2   | KCNQ2     | +      | +/−       | +   | +/− | −/−   | −/−   | −/−       | +/−      |
| Kv7.3   | KCNQ3     | +      | +         | +   | +   | ++    | +/−   | ++        | +/−      |
| Kv7.5   | KCNQ5     | ++     | +/−       | +   | +   | +/−   | +/−   | +/−       | +/−      |
| Kv8.1   | KCNV1     | +      | −         | −   | +/− | +/−   | −     | −          | −        |
| Kv9.2   | KCNS2     | +      | −         | −   | +/− | −     | −     | −          | −        |
| Kv10.2/EG2 | KCNH5 | +     | +/−       | +/− | +   | +     | +/−   | −          | −        |
| Kv11.2/ERG3 | KCNH7 | +++   | +         | −   | +   | +/−   | −     | +/−       | −        |
| Kv12.1/ELK1 | KCNH8 | +     | +++      | ++  | +++ | +/−   | ++   | +/−       | +/−      |
| TREK1   | KCNK2     | +      | +/−       | +   | +/− | −/−   | −     | −/−       | −        |
| TREK2   | KCNK10    | +      | +/−       | +   | +/− | +/−   | +/−   | −/−       | −        |
| KᵥCa₂.³ | KCNN3     | +      | +         | +   | +   | +     | +/−   | +/−       | +/−      |
| KᵥNa₂.¹ | SCN1A     | ++     | +         | +++ | ++ | +/−   | −     | −          | −        |
| KᵥNa₂.¹ | SCN2A     | ++     | +         | +   | +   | +     | +/−   | −/−       | −/−      |
| KᵥNa₂.¹ | SCN3A     | ++     | +/−       | +   | +   | +     | +/−   | +/−       | +/−      |
| KᵥNa₂.¹ | SCN8A     | n.d    | n.d       | n.d | n.d | n.d   | n.d   | n.d       | n.d      |
| KᵥNa₂.¹ | SCN11A    | n.d    | n.d       | n.d | n.d | n.d   | n.d   | n.d       | n.d      |
| Ca²⁺ channels |          |        |           |     |     |       |       |           |          |
| Cav1.2  | CACNA1C   | +++    | +         | +   | +   | +/−   | −     | +/−       | +/−      |
| Cav1.3  | CACNA1D   | ++     | +/−       | +   | +   | +/−   | +     | −          | −        |
| Cav2.1  | CACNA1A   | +++    | +         | +++ | ++ | +/−   | +/−   | +/−       | +/−      |
| Cav2.3  | CACNA1E   | ++     | +/−       | +   | +   | +/−   | +     | +/−       | +/−      |
| Cav3.1  | CACNA1G   | +      | −         | +/− | −   | −     | −     | −          | −        |
| Cav3.2  | CACNA1H   | n.d    | n.d       | n.d | n.d | n.d   | n.d   | n.d       | n.d      |
| Cav3.3  | CACNA1I   | −      | +/−       | +   | +   | −     | −     | −          | −        |
| Ryanodine |          |        |           |     |     |       |       |           |          |
| Ryr2    | RYR2      | +++    | +/−       | +   | +   | +     | +/−   | +/−       | +/−      |
| Ryr3    | RYR3      | +      | +++       | +   | +   | +/−   | +/−   | +/−       | +/−      |
| TRP channels |        |        |           |     |     |       |       |           |          |
| TRPM2   | TRPM2     | +      | +/−       | +   | +   | +/−   | −     | +/−       | +/−      |
| TRPP1   | PKD2      | +      | +         | +   | +   | +     | +     | +/−       | +/−      |
| TRPP3   | PKD2L2    | n.d    | n.d       | n.d | n.d | n.d   | n.d   | n.d       | n.d      |
| TRPV1   | TRPV1     | +/-    | −         | +/− | −   | −     | +/−   | −/−       | −/−      |
| TRPV6   | TRPV6     | n.d    | n.d       | n.d | n.d | n.d   | n.d   | n.d       | n.d      |
| CI⁻ channels |          |        |           |     |     |       |       |           |          |
| Cl⁻2    | CLCN2     | +/-    | +/−       | +/− | +/− | +/−   | +/−   | +/−       | +/−      |
| Cl⁻7    | CLCN7     | +      | +         | +   | +   | +     | +     | +/−       | +/−      |
| Connexins |          |        |           |     |     |       |       |           |          |
| Cx37    | GJA4      | −      | −         | −   | −   | −     | −     | −         | −        |

(Continued)
Kv4.2 (KCND2)
The Kv4.2 gene encodes for the Kv4.2 subunit that (together with Kv4.1 and Kv4.3) is a member of the Kv4 channel subfamily (Shal) and is highly expressed in the brain (Alfaro-Ruiz et al., 2019). Kv4 channels activate at subthreshold potentials and then inactivate and recover rapidly. They mediate transient A-type K+ current (Bahring et al., 2001; Birnbaum et al., 2004).

Neurons
Kv4.2 subunits are highly expressed in soma and dendrites of hippocampal neurons and interneurons. They regulate the threshold for AP initiation and repolarization, frequency-dependent AP broadening, and AP back-propagation (Nerbonne et al., 2008). Kv4.2 mutations are associated with infant-onset epilepsy and autism.

Glia
Kv4.2-transcripts were found in mouse astrocytes (Bekar et al., 2005) and OPCs, but only at very low levels in microglia (Falcao et al., 2018; Hammond et al., 2019; Batiuk et al., 2020).

Expression and Function in MS
Bulk RNA-seq found significant Kv4.2 upregulation in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq reported significant expression of Kv4.2 transcripts in neuronal, OPCs, and committed OPCs (COP) clusters (Table 2; Jakel et al., 2019). Kv4.2 subunit may contribute to oligodendrocyte dysfunction in SPMS brain because dysregulated KCND2 transcripts are associated with oligodendrocyte dysfunction in mental illnesses (Vasistha et al., 2019).

Kv7.2, Kv7.3, and Kv7.5 (KCNQ2, KCNQ3, and KCNQ5)
The KCNQ genes encode for Kv7.1–Kv7.5 (KCNQ1–KCNQ5) family members that underlie a voltage-gated non-inactivating outward K+ current, known as M current (I_M).

Neurons
The Kv7.2/3 or Kv7.3/5 hetero-tetramers represent the dominant subunit composition in neurons (Wang et al., 1998; Cooper et al., 2000; Kharkovets et al., 2000), while Kv7.4/Kv7.5 is dominant in

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TABLE 2 | Continued

| Protein | Gene | Neuron | Astrocyte | OPC | COP | ImOLG | Oligo | Microglia | Pericyte |
|---------|------|--------|-----------|-----|-----|-------|-------|-----------|---------|
| Pannexin |      |        |           |     |     |       |       |           |         |
| Px1     | PANX1| n.d    | n.d       |     |     | n.d   | n.d   | n.d       | n.d     |
| Catsper |      |        |           |     |     |       |       |           |         |
| CATSPERG| CATSPERG| +/– | +/– | – | +/– | +/– | +/– | – | – |
| CATSPERE| CATSPERE| n.d | n.d | n.d | n.d | n.d | n.d | n.d | n.d |

*snRNA-seq from the white matter of individuals with SPMS and non-neurological controls. The information is based on the snRNA-seq from the WM of individuals with SPMS and non-neurological control subjects (Jakel et al., 2019), and data are collected from the database available at https://ki.se/mbb/oligodendrocytes/. Expression levels are based on the mean normalized expression counts (log-scale) per cluster.

Kv7.2 and Kv7.3 subunits co-cluster with NaC channels at AIS and nodes of Ranvier in rodent somatosensory cortex and spinal cord WM and gray matter (GM) (Pan et al., 2006; Cooper, 2011; Battefeld et al., 2014). Kv7.5 localizes to soma and dendrites of cortical and hippocampal neurons and contributes to afterhyperpolarization currents (Tzingounis et al., 2010). The Kv7 channels stabilize V_rest, influence neuronal subthreshold excitability, and regulate spike generation (Jentsch, 2000; Miceli et al., 2008). By reducing the steady-state inactivation of nodal NaC channels, the Kv7 channels increase the availability of transient NaC currents at nodes of Ranvier, thereby accelerating the AP upstroke and elevating short-term axonal excitability (Hamada and Kole, 2015). In the perisomatic region, Kv7 channels counteract the persistent NaC current and restrain repetitive firing (Pan et al., 2006; Cooper, 2011). Variants of KCNQ2/KCNQ3 or KCNQ4 genes cause developmental/epileptic disorders and hearing loss (Soldovieri et al., 2011; Miceli et al., 2013).

Glia
KCNQ3 gene is expressed in spinal cord WM astrocytes (Devaux et al., 2004), while KCNQ5 is expressed in rat retinal astrocytes (Caminos et al., 2015). The KCNQ2-5 mRNAs and proteins were detected in rat cortical OPCs and microglia cultures, while differentiated oligodendrocytes showed weak KCNQ4 expression (Wang et al., 2011; Vay et al., 2020).

Expression and Function in MS
Bulk RNA-seq found upregulation of KCNQ2-3-5 transcripts in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq reported significant expression of KCNQ2-3-5 transcripts in neuronal clusters and KCNQ3 expression in immune oligodendroglia (ImOLG) and microglia/macrophages clusters (Tables 1, 2; Jakel et al., 2019). Kv7.3 upregulation may reflect increased necessity of the channels along the axons because Kv7.3 subunit extensively redistributes to internodes of acute and chronically demyelinated GM axons in the cuprizone model (Hamada and Kole, 2015). It is tempting to speculate that Kv7 upregulation may be beneficial during MS. First, Kv7 channels may increase the availability of transient NaC current via membrane hyperpolarization supporting AP conduction in demyelinated axons (Battefeld et al., 2014).
Second, Kv7 channels may mitigate inflammation-induced neuronal excitability because, following LPS exposure, the I\textsubscript{M} inhibition underlies hyperexcitability of hippocampal neurons that is reversed by a nonselective Kv7-opener retigabine (Tzour et al., 2017). Although retigabine also exerts neuroprotective effects in several neurodegenerative conditions (Boscia et al., 2006; Nodera et al., 2011; Wainger et al., 2014; Bierbower et al., 2015; Li et al., 2019; Vigil et al., 2020; Wu et al., 2020), a clinical trial with retigabine analog flupirtine failed to demonstrate neuroprotective effects during MS (Dorr et al., 2018). Furthermore, blockade of Kv7 channels with XE-991 inhibited migration of LPS-treated pro-inflammatory microglia \textit{in vitro} (Vay et al., 2020), suggesting that these channels may promote the pro-inflammatory role of microglia also during MS. Hence, neuronal and glial Kv7 channels may have diverse functions during MS.

### FIGURE 3 | Distribution of uniquely dysregulated genes encoding ion channels in SPMS lesions. Schematic representation of active (AL), chronic active (CA), inactive (IL), and remyelinating (RL) lesions, and normal-appearing white matter (NAWM). Upregulated (blue) and downregulated (red) ion channels encoded by uniquely dysregulated genes are listed according to their expression in the lesions and in neuronal, oligodendrocyte precursor cells (OPCs), committed OPCs (COPs), oligodendrocytes (Oligo), microglia, immune oligo (ImOLG), astrocyte, pericyte, and unknown clusters. GM, gray matter; WM, white matter; v, brain ventricle. Gray areas indicate active inflammatory lesion, white areas indicate demyelinated inactive lesions, red spot indicates tissue damage, red arrow indicates axonal dysfunction. Source icon is from Biorender.com.

| Lesion features | Active | Chronic Active | Inactive | Remyelinating | NAWM |
|----------------|--------|----------------|----------|---------------|-------|
| • inflammation | • demyelination | • hypocellularity in the core | • demyelination | • OPC proliferation | • axonal abnormalities |
| • demyelination | • axonal damage | • microglia/macrophages, OPCs, and astrocytes accumulation in the rim | • hypocellularity | • OPC differentiation | • myelin abnormalities |
| • axonal damage | | • remyelination failure | • axonal damage | | • microglia activation |
| • glia activation | | | • glia scar | | |
| • phagocytic activity | | | • angiogenesis | | |

| Clusters | Genes |
|----------|-------|
| Neuronal | Kv1.1, Kv1.2, Kv1.4, Kv2.1, Kv2.2, Kv3.3, Kv4.2, KCNQ2, KCNQ3, KCNQ5, KCNV1, KCNS2, EAQ2, ERG3, TREK1, KNa1.1, Kir3.2, Nav1.1, Nav1.2, Nav1.3, Nav1.8, Cav1.2, Cav2.1, Cav2.3, Cav1.3, Cav3.2, Cav3.3, RYR2 |
| OPC | Kv4.2, Kir5.1, TREK1, Nav1.1, Nav1.3, Cav2.1 |
| COP | Kv4.2, Nav1.1 |
| Microglia | KCNQ3 |
| ImOLG | KCNQ3 |
| Astrocyte | KCa2.3, RYR3 |
| Pericyte | Cav1.2, Cx37 |
| Unknown | TREK2, TRPV6, TRPV1, TRPV6, PKD2L2, CLC-2, CLC-7, CATSPERG, Px1, Nav1.9, TRPV6, PKD2, CATSPER |

Second, Kv7 channels may mitigate inflammation-induced neuronal excitability because, following LPS exposure, the \( I_{M} \) inhibition underlies hyperexcitability of hippocampal neurons that is reversed by a nonselective Kv7-opener retigabine (Tzour et al., 2017). Although retigabine also exerts neuroprotective effects in several neurodegenerative conditions (Boscia et al., 2006; Nodera et al., 2011; Wainger et al., 2014; Bierbower et al., 2015; Li et al., 2019; Vigil et al., 2020; Wu et al., 2020), a clinical trial with retigabine analog flupirtine failed to demonstrate neuroprotective effects during MS (Dorr et al., 2018). Furthermore, blockade of Kv7 channels with XE-991 inhibited migration of LPS-treated pro-inflammatory microglia \textit{in vitro} (Vay et al., 2020), suggesting that these channels may promote the pro-inflammatory role of microglia also during MS. Hence, neuronal and glial Kv7 channels may have diverse functions during MS.
### TABLE 3 | Expression and role of unique dysregulated ion channels in experimental models of MS.

| Gene/protein | Distribution, localization | Cellular functions during physiological conditions | WM in MS models |
|--------------|---------------------------|---------------------------------------------------|-----------------|
| **KCN1/Kv1.1** | JPN of myelinated axons | Regulate AP propagation and neural excitability | Redistribution to internodes and nodal segments, upregulation |
| Microglia, astrocyte (t), OPCs (t) | | | Hyperpolarise axonal V<sub>rest</sub>, affect AP threshold, impair AP conduction |
| **KCN2/Kv1.2** | JPN of myelinated axons | Regulate AP propagation and neural excitability | Redistribution to internodes and nodal segments, upregulation |
| Reactive astrocyte, microglia, OPC | | | Hyperpolarise axonal V<sub>rest</sub>, affect AP threshold, impair AP conduction |
| **KCN4/Kv1.4** | Axons (HP) | Regulate AP propagation and neural excitability | Upregulation in astrocytes and OPCs around EAE lesions |
| | | | Deficiency ameliorated EAE course in KO mice, but have no effect on demyelination/remyelination in the cuprizone model |
| **KCNB1/Kv2.1** | Reactive astrocyte, OPCs | Proliferation | Unknown in WM |
| Soma, proximal dendrites, AIS | | Influence AP duration during high frequency firing, regulate neuronal excitability | Downregulation in motor neurons of GM spinal cord during EAE |
| Microglia, OPCs (t) | | | Unknown |
| **KCNB2/Kv2.2** | Soma, proximal dendrites, AIS | Influence AP duration during high frequency firing, regulate neuronal excitability | Unknown |
| Not detected in glia | | | Unknown |
| **KCN3/Kv3.3** | Axons, somatodendritic compartment | Regulate AP firing at high frequency | Upregulation in some injured WM axons |
| Astrocyte, microglia (t), OPCs (t) | | | Unknown |
| **KCN2/Kv7.2** | AIS, nodes of Ranvier OPCs, microglia | Stabilize V<sub>rest</sub>, regulate activity of Na<sub>V</sub>-channels, accelerate AP upstroke, influence neuronal subthreshold excitability, regulate spike generation, and repetitive firing | Unknown |
| **KCNQ2/Kv7.3** | AIS, nodes of Ranvier Microglia (pro-inflammatory), OPCs, astrocyte (t) | Stabilize V<sub>rest</sub>, regulate activity of Na<sub>V</sub>-channels, accelerate AP upstroke, influence neuronal subthreshold excitability, regulate spike generation and repetitive firing | Unknown |
| **KCNQ5/Kv7.5** | Soma, dendrites | Co-assemble with K<sub>v</sub>2.1, reduce K<sub>v</sub>2.1 current density which may lead to AP broadening and hyper-synchronized high-frequency firing | Unknown |
| Astrocyte, OPCs, microglia | | | Unknown |
| **KCNV1/Kv8.1** | Unknown | Co-assemble with K<sub>v</sub>2.1 | Unknown |
| Oligo lineage (t) | | | Unknown |
| **KCN2/Kv9.2** | Unknown | Co-assemble with K<sub>v</sub>2.1 | Unknown |
| Oligo lineage (t) | | | Unknown |
| **KCNH5/EAG2** | Unknown | Dampen excitability, stabilize V<sub>rest</sub> | Unknown |
| Astrocyte (t), OPCs (t) | | | Unknown |
| **KCNH7/ERG3** | Unknown | Dampen excitability, stabilize V<sub>rest</sub> | Unknown |
| Astrocyte (t), OPCs (t), microglia (t) | | | Unknown |
| **KCNH8/ELK1** | Unknown | Dampen excitability, stabilize V<sub>rest</sub> | Unknown |
| OPCs (t) | | | Unknown |

(Continued)
| Gene/protein | Distribution, localization | Cellular functions during physiological conditions | WM in MS models |
|-------------|----------------------------|--------------------------------------------------|-----------------|
| **KCNK2/TREK1** | Axons, and node of Ranviers in afferent myelinated nerve | Contribute to "leak" K⁺-current, help establishing and maintaining V<sub>rest</sub>, regulate neuronal excitability, ensure AP repolarization at nodes of Ranvier in afferent myelinated fibers | Unknown | Deficiency aggravates EAE course in KO mice |
|              | Astrocyte, microglia (t) OPcs (t) | Contribute to passive membrane K⁺ conductance, glutamate release | Unknown | Channel activation reduces CNS immune cell trafficking across BBB and attenuate EAE course |
| **KCNK10/TREK2** | Unknown | Contribute to "leak" K⁺-current, help establishing and maintaining V<sub>rest</sub>, regulate neuronal excitability, ensure AP repolarization at nodes of Ranvier in afferent myelinated fibers | Unknown | Unknown |
| **KCNT1/K<sub>i</sub>4.1** | Soma, axons Astrocytes (t) | Regulate the generation of slow afterhyperpolarization, firing patterns, and setting and stabilizing the V<sub>rest</sub> | Unknown | Unknown |
| **KCNN3/K<sub>Ca</sub>2.3** | Dendrites, AIS Astrocyte, microglia, oligo lineage (t) | Regulate AP propagation and neuronal excitability, contribute to maintaining Ca<sup>2+</sup>-homeostasis K⁺ buffering in astrocytes | Unknown | Unknown |
| **KCNN3/K<sub>Ca</sub>2.3** | Somatodendritic compartment | K⁺-homeostasis, maintenance of V<sub>rest</sub>, hyperpolarization, control of AP firing and neuronal excitability, inhibition of excitatory neurotransmitter release | Unknown | Unknown |
| **KCNJ6/K<sub>i</sub>3.2** | Astrocye, oligo lineage (t) | Silent channel when combined with K<sub>i</sub>2.1. When combined with K<sub>i</sub>4.1, build channels with larger conductance and greater pH-sensitivity. Plays a role in synaptic transmission | Unknown | Unknown |
| **KCNJ16/K<sub>i</sub>5.1** | Somatodendritic compartment, dendritic spines Astrocyte, oligo lineage (t) | Silent channel when combined with K<sub>i</sub>2.1. When combined with K<sub>i</sub>4.1, build channels with larger conductance and greater pH-sensitivity. Plays a role in synaptic transmission | Unknown | Unknown |
| **SCN1A/Na<sub>v</sub>1.1** | Somatodendritic compartment, AIS, nodes of Ranvier Microglia, astrocyte, OPcs (t) | Salutatory conduction, maintenance of sustained firing, control of excitability Microglia phagocytosis, cytokine release | Increase or no change; localize along the demyelinated regions | Unknown |
| **SCN2A/Na<sub>v</sub>1.2** | AIS, immature nodes of Ranvier, along the non-myelinated axons Astrocyte, pre-oligodendrocytes | Back-propagation of AP into the somatodendritic compartment, may support slow spike propagation Oligodendroglia maturation | Increase of diffuse distribution along demyelinated axons in various mouse models; no change in myelin-deficient rat | Unclear. Suggested: preservation of AP propagation, or axonal damage |
| **SCN3A/Na<sub>v</sub>1.3** | Somatodendritic compartment, along the axons including myelinated fibers Astrocyte oligo lineage (t) | AP initiation and propagation, proliferation and migration of cortical progenitors | No change in the optic nerve | Unknown |
| **SCN8A/Na<sub>v</sub>1.6** | AIS, nodes of Ranvier; low density on cell soma, dendritic shafts, synapses Astrocyte, microglia oligo (t) | AP initiation and propagation, neuronal excitability Decrease at the nodes of Ranvier, increase of diffuse distribution along the damaged axons, no change at AIS Upregulated in microglia/macrophages during EAE | Unknown | May trigger Na⁺ increase in axoplasm, reversal of NCX, and intra-axonal Ca<sup>2+</sup> overload. Deletion improves axonal health during EAE |
| **SCN11A/Na<sub>v</sub>1.9** | Soma, proximal processes Negligible in all glial cells (t) | Regulate excitation, control activity-dependent axonal elongation, mediate sustained depolarizing current upon activation of muscarinic receptors | Unknown | Unknown |
| Gene/protein | Distribution, localization | Cellular functions during physiological conditions | WM in MS models |
|-------------|---------------------------|-------------------------------------------------|----------------|
| CACNA1C/ CaV1.2 | Somatodendritic compartment (synaptically, extrasynaptically), axons, axonal terminals (extrasynaptically), pioneer axons during development Astrocite, oligo lineage, reactive microglia | Synaptic modulation, propagation of dendritic Ca\(^{2+}\) spikes, regulation of glutamate receptor trafficking, CREB phosphorylation, coupling of excitation to nuclear gene transcription, modulation of long-term potentiation, neurites growth and axonal pathfinding during development Astrogliosis OPCs development and myelination | Unknown | Unknown. Suggested: Neurodegeneration because L-type VGCCs blockers attenuate mitochondrial pathology in nerve fibers and axonal loss Deletion in astrocyte reduces cell activation and pro-inflammatory mediators release in the cuprizone model Deletion in OPCs reduced remyelination in the cuprizone model |
| CACNA1D/ CaV1.3 | Somatodendritic compartment, axonal cylinders Astrocyte, microglia oligo lineage | Pacemaking activity, spontaneous firing, Ca\(^{2+}\)-dependent post-burst after-hyperpolarization, Ca\(^{2+}\)-dependent intracellular signaling pathways, regulation of morphology of dendritic spines and axonal arbores Oligodendrocyte-axon signaling, release of pro-inflammatory mediators by microglia | Unknown | Unknown. Suggested: neuroprotection because L-type VGCCs blockers attenuate mitochondrial pathology in nerve fibers and axonal loss |
| CACNA1A/ CaV2.1 | Axonal synaptic terminals, axonal shafts in WM, somatodendritic compartment Reactive astrocyte OPCs, premyelinating oligo, microglia (t) | Neurotransmitter release at neuronal and neuron-glia synapses, regulation of BK and SK channels, control of neuronal firing, regulation of gene expression, local Ca\(^{2+}\) signaling, and cell survival Calcium influx in oligo upon neuronal activity | Unknown | Unknown |
| CACNA1E/ CaV2.3 | Dendritic spines, axonal terminals Astrocyte, oligodendrocyte | Neurotransmitter release, synaptic plasticity, regulation of BK, SK, and K\(_{V}4.2\) channels | Unknown | Unknown |
| CACNA1G/ CaV3.1 | Somatodendritic compartment, AIS Astrocyte (t) oligo lineage | Generation and timing of APs, regulation of neuronal excitability, rhythmic AP bursts in thalamus, neuronal oscillations, neurotransmitter release | Unknown | T-cells from KO mice show decreased cytokine release Deficiency in KO mice inhibits the autoimmune response in the EAE model |
| CACNA1H/ CaV3.2 | Somatodendritic compartment, AIS Astrocyte oligo lineage | Generation and timing of APs, regulation of neuronal excitability, rhythmic AP bursts in thalamus, neuronal oscillations, neurotransmitter release | Unknown | Unknown |
| CACNA1I/ CaV3.3 | Somatodendritic compartment | Generation and timing of APs, regulation of neuronal excitability, rhythmic AP bursts in thalamus, neuronal oscillations, neurotransmitter release | Unknown | Unknown |
| RyR2 | Along ER (also in axons) Astrocyte, oligo lineage | Ca\(^{2+}\) release from the ER into the cytoplasm, vesicle fusion, neurotransmitter release, synaptic plasticity, growth cone dynamics | Unknown | Unknown |
| RyR3 | Along ER (also in axons) Astrocyte, OPCs, oligodendrocytes | Ca\(^{2+}\) release from the ER into the cytoplasm, vesicle fusion, neurotransmitter release, synaptic plasticity, growth cone dynamics Astrocyte motility OPCs development | Unknown | Unknown |
| TRPV1 | Soma, post-synaptic dendritic spines, synaptic vesicles Astrocyte, microglia, oligodendrocytes | Regulation of Ca\(^{2+}\)-signaling, synaptic plasticity | Suggested a main role in regulating microglia inflammatory response | Both detrimental and beneficial effects have been described in EAE disease |

(Continued)
| Gene/protein | Distribution, localization | Cellular functions during physiological conditions | WM in MS models |
|-------------|---------------------------|--------------------------------------------------|-----------------|
| TRPV6       | Unknown Astrocyte (t)     | Unknown                                         | Unknown         |
| TRPM2       | Soma and neurites in neural cultures Microglia, astrocyte (t), oligodendrocyte (t) | Contribute to synaptic plasticity and play an inhibitory role in neurite outgrowth Microglia activation and generation of proinflammatory mediators | Upregulated in monocyte-lineage cells TRPM2 deficiency reduce monocyte infiltration in EAE |
| PKD2/TRPP1  | ER, primary cilia, and plasma membrane Astrocyte (t), microglia (t), oligo lineage (t) | Maintenance of Ca\(^{2+}\)-homeostasis, cell proliferation | Unknown         |
| PKD2L2/ TRPP3 | Unknown Astrocyte (t), microglia (t) | Unknown                                         | Unknown         |
| CLCN2/CLC-2 | Plasma membranes, intracellular membranes Astrocyte, OPCs, microglia | Maintenance of low intracellular Cl\(^{-}\) level, control of cell volume homeostasis, regulation of GABA\(_A\)R-mediated synaptic inputs, regulation of neuronal excitability Interacts with AQP4 in astrocytes, regulates OPCs differentiation, contribute to volume regulation and phagocytosis in microglia | Unknown         |
| CLCN7/CLC-7 | Lysosomes Microglia, astrocyte (t), oligo lineage (t) | Suggested function in the neuronal endo-lysosomal pathway Regulate lysosomal acidification in activated microglia | Unknown         |
| GJA4/CX37   | Largely expressed in vascular cells | Regulate vasomotor activity, endothelial permeability, and maintenance of body fluid balance | Unknown         |
| PANX1/Px1   | Soma, dendrites, axons Astrocyte, OPCs microglia | Paracrine and autocrine signaling, ATP-sensitive ATP release in complex with P2X7-Rs, intercellular propagation of Ca\(^{2+}\)-waves, cell differentiation, migration, synaptic plasticity, memory | Unknown Panx-1 induced ATP release and inflammasome activation contribute to WM damage during EAE Inhibition of Panx1 using pharmacology or gene disruption delays and attenuates disease course in EAE and cuprizone model |
| CATSPERG    | Unknown Oligo lineage (t) Microglia (t) | Unknown                                         | Unknown         |
| CATSPERE    | Unknown Oligo lineage (t) Microglia (t) | Unknown                                         | Unknown         |

**AHP**, afterhyperpolarization; **AIS**, axon initial segment; **AP**, action potential; **BK**, big-conductance Ca\(^{2+}\)-activated K\(^{+}\)-channels; **ER**, endoplasmatic reticulum; **GABA\(_A\)R**, ionotropic gamma-aminobutyric acid A receptor; **HP**, hippocampus; **JPN**, juxtaparanodal regions; **NCX**, Na\(^{+}\)/Ca\(^{2+}\) exchanger; **SCI**, spinal cord injury; **SK**, small-conductance Ca\(^{2+}\)-activated K\(^{+}\)-channels; **SSCx**, somatosensory cortex; **t**, transcripts; **V\(_{rest}\)**, resting membrane potential.

**K\(_v\)8.1 and K\(_v\)9.2 (KCNV1 and KCNS2)**

**Neurons**

KCNV1 and KCNS2 genes encode for electrically silent (K\(_v\)S) K\(_v\)8.1- and K\(_v\)9.2 subunits that assemble into hetero-tetrameric channels with K\(_v\)2 subunits (Bocksteins, 2016). A number of channelopathies is ascribed to K\(_v\)S subunits (Salinas et al., 1997a; Liu et al., 2016; Allen et al., 2020), pointing to their important physiological role.

**Glia**

KCNV1 and KCNS2 transcripts were found in oligodendrocyte lineage cell (Marques et al., 2016).

**Expression and Function in MS**

Bulk RNA-seq showed upregulation of KCNV1 and KCNS2 genes in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq detected KCNV1 and KCNS2 in neuronal clusters (Table 2; Jakel et al., 2019). Co-assembly between K\(_v\)8.1 and K\(_v\)2.1 reduces K\(_v\)2.1 current density (Hugnot et al., 1996; Castellano et al., 1997): the high stoichiometry of the K\(_v\)8.1 subunit suppresses surface expression and favors retention of heteromeric channels in the ER (Salinas et al., 1997b). Neurons with reduced K\(_v\)2.1-mediated currents demonstrate broadened APs (Du et al., 2000) underlying hyper-synchronized high-frequency firing observed during epilepsy. Hence, upregulated K\(_v\)S subunits in CA lesions may influence the localization of...
clustered K_v2 subunits in SPMS brain and affect AP firing and/or propagation.

**Eag2, erg3, and elk1 (KCNH5, KCNH7, and KCNH8)**

KCNH genes encode for K_v10–K_v12 subfamilies, all orthologs of the Drosophila ether-ágo-go (EAG) channels. They include two eag (K_v10), three eag-related (erg/K_v11), and three eag-like (elk/K_v12) K^+ channels that can form heteromeric channels within each subfamily (Rasmussen and Trimmer, 2019).

**Neurons**

All EAG channels are expressed in the CNS neurons (Ludwig et al., 2000; Papa et al., 2003; Zou et al., 2003), but only erg-mediated currents have been verified using suitable blockers (Bauer and Schwarz, 2018).

**Glia**

RNA-seq detected KCNH5, KCNH7, and KCNH8 expression in mouse OPCs (Falcao et al., 2018). KCNH5 and KCNH7 genes were found in astrocytes (Batiuk et al., 2020), while only the KCNH7 gene was detected in mouse microglia (Hammond et al., 2019). Erg-type currents were reported in neopallial microglia cultures (Zhou et al., 1998) and hippocampal astrocytes (Emmi et al., 2000; Papa et al., 2003).

**Expression and Function in MS**

Bulk RNA-seq detected increased KCNH5(eag2) and KCNH7(erg3) transcripts in CA lesions and downregulation of KCNH8(elk1) transcript in all lesions and NAWM (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq found significant expression of KCNH5 and KCNH7 transcripts in neuronal clusters and KCNH8 in mature oligodendrocyte clusters (Tables 1, 2; Jakel et al., 2019). The functional role of eag2, erg3, and elk1 during MS may be related to altered neuronal excitability. Indeed, human eag1 and eag2 gain-of-function mutations underlie severe neurological disorders associated with epileptic seizures (Allen et al., 2020). The erg channels that are active at subthreshold potentials stabilize the V_rest and dampen excitability (Fano et al., 2012). Erg3 knockdown in mice increases intrinsic neuronal excitability and enhances seizure susceptibility, while treatment with erg activator reduces epileptogenesis (Xiao et al., 2018). Erg3 expression is decreased...
in the brain of epilepsy patients. Remarkably, association of KCNHi7(erg) intronic polymorphisms with MS pathogenesis was speculated although never substantiated (Martinez et al., 2008; Couturier et al., 2009).

**Two-Pore Domain K⁺ Channels (K2P)**

K2P K⁺ channels are encoded by 15 KCNK genes, stratified into six subfamilies: TWIK, TASK (TWIK-related acid-sensitive), TREK (TWIK-related arachidonic acid activated), THIK (tandem pore domain halothane-inhibited), TALK (TWIK-related alkaline pH-activated), and TRESK (TWIK-related spinal cord) K⁺ channels (Enyedi and Czirjak, 2010). K2P K⁺ channels contribute to “leak” K⁺ current, helping to establish and maintain V<sub>rest</sub> (Enyedi and Czirjak, 2010).

**TREK1 and TREK2 (KCNK2 and KCNK10)**

**Neurons and Glia**

KCNK2 and KCNK10 genes encode for TREK-1 and TREK-2 channels, which are expressed in neurons, astrocytes, and OPC (Hervieu et al., 2001; Talley et al., 2001; Falcao et al., 2018). Only TREK-1 transcripts were detected in microglia (Hammond et al., 2019). In astrocytes, TREK channels contribute to passive conductance and glutamate release (Zhou et al., 2009; Woo et al., 2012). TREK-1 and TREK-2 may be activated by a wide range of physiological and pathological stimuli reminiscent of inflammatory environment including membrane stretch, heat, intracellular acidity, and cellular lipids (Ehling et al., 2015).

**Expression and Function in MS**

Bulk RNA-seq found upregulated TREK-1 transcripts in CA lesions, but a divergent modulation was observed for TREK-2 mRNAs in ALs (Figure 2, Tables 1, 2; Elkjaer et al., 2019; Frisch et al., 2020). KCNK2 and KCNK10 transcripts were detected in neuronal and oligodendrocyte clusters, but scarcely observed in astrocytes (Table 2; Jakel et al., 2019). TREK-1 upregulation in CA lesions most likely reflects a protective response because TREK-1 plays a neuroprotective role during neurological diseases, including MS (Djillani et al., 2019). TREK-1 reduces neuronal excitability by hyperpolarizing the membrane potential (Honore, 2007) and is required for rapid AP repolarization at the node of Ranvier in mammalian afferent myelinated nerves, while TREK-1 loss-of-function retards nerve conduction and impairs sensory responses in animals (Kanda et al., 2019). Treatment of mice with TREK-1 activators, riluzole (Gilgun-Sherki et al., 2003), or alpha-linolenic acid attenuates EAE course (Blondeau et al., 2007), while these effects are reduced in TREK-1<sup>−/−</sup> mice (Bittner et al., 2014). TREK-1 function is also important for non-neuronal cells because aggravated EAE course in TREK-1<sup>−/−</sup> mice is associated with increased numbers of infiltrating T cells and higher endothelial expression of ICAM1 and VCAM1 (Bittner et al., 2013), and TREK-1 is reduced in the microvascular endothelium in inflammatory MS brain lesions (Bittner et al., 2013).

TREK-2 downregulation in AL, a lesion type characterized by myelin breakdown and infiltration by inflammatory cells (Elkjaer et al., 2019; Frisch et al., 2020), may contribute to reduced glutamate and K⁺ buffering and neuronal over-excitation because TREK-2 helps maintain the membrane potential and low extracellular glutamate and K⁺ level during ischemia (Gnatenco et al., 2002; Rivera-Pagan et al., 2015).

**Na⁺- and Ca²⁺-Activated K⁺ Channels**

**K<sub>Na</sub>1.1 (KCNT1)**

**Neurons**

The KCNT1 and KCNT2 genes encode for Slack and Slick K⁺ channels that are activated by Na⁺ influx (Bhattacharjee and Kaczmarek, 2005). They localize to soma and axons of neurons (Bhattacharjee et al., 2002; Brown et al., 2008; Rizzi et al., 2016) and are involved in the generation of slow after-hyperpolarization, regulation of firing patterns, and setting and stabilizing the V<sub>rest</sub> (Franceschetti et al., 2003). Alterations in KCNT1 and KCNT2 genes are linked to early-onset epileptic encephalopathies and Fragile-X-syndrome (Kim and Kaczmarek, 2014).

**Glia**

RNA-seq detected KCNT1 gene in mouse astrocytes (Batiuk et al., 2020).

**Expression and Function in MS**

Bulk RNA-seq showed K<sub>Na</sub>1.1 upregulation in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). SnRNA-seq detected KCNT1 in neuronal clusters (Table 2). KCNT1 function in MS may be related to myelination/demyelination because severely delayed myelination occurs in patients with KCNT1 mutations (Vanderver et al., 2014). Furthermore, KCNT1 is a causative gene in infants with hypomyelinating leukodystrophy showing WM alterations (Arai-Ichinoi et al., 2016), and KCNT1 mutations occur in infant epilepsy associated with delayed myelination, thin corpus callosum, and WM hyper-intensity in MRI (McTague et al., 2013; Shang et al., 2016; Borlot et al., 2020).

**K<sub>Ca</sub>2.3, SK3 (KCNN3)**

The KCNN3 gene encodes for the SK3 subunit of small-conductance Ca²⁺-activated K⁺ channels (SK channels). They mediate Ca²⁺-gated K⁺ current and thus couple the increase in intracellular Ca²⁺ concentration to hyperpolarization of the membrane potential.

**Neurons**

SK3 channels are found on dendrites and AIS (Abiraman et al., 2018). They play a role in AP propagation and regulation of neuronal excitability (Stocker, 2004). They protect against excitotoxicity by maintaining Ca²⁺ homeostasis after NMDA receptor activation (Dolga et al., 2011).

**Glia**

RNA-seq detected intense KCNN3 expression in mouse astrocytes (Batiuk et al., 2020), confirming earlier studies, which showed labeling of GFAP⁺ processes in the supraoptic nucleus for SK3 channels and suggested the role of SK3 in astrocytic K⁺ buffering (Armstrong et al., 2005). Oligodendrocyte lineage cells express low levels of KCNN3 mRNA (Falcao et al., 2018), while mouse microglia does not express KCNN3 (Hammond et al., 2015).
et al., 2019). However, rat microglia in culture expresses the SK3 subunit, which is increased upon microglia activation with LPS (Schlichter et al., 2010). SK3 activation inhibited microglia proliferation, inflammatory IL-6 production, and morphological transformation to macrophages, while blocking SK3 in microglia-reduced neurotoxicity (Dolga et al., 2012).

**Expression and Function in MS**

Bulk RNA-seq showed significant and unique downregulation of KCNN3 in ILS (Elkjaer et al., 2019; Frisch et al., 2020). SnRNA-seq revealed high KCNN3 expression in astrocyte clusters (Figure 2, Table 1; Jakel et al., 2019). ILS consist of large demyelinated areas devoid of macrophages but filled with scar-forming astrocytes showing reduced ability to buffer glutamate and K+ (Compston and Coles, 2008; Kuhlmann et al., 2017; Filippi et al., 2018; Schirmer et al., 2018). Hence, KCNN3 downregulation in MS may reflect altered function of astrocytes, e.g., K+ buffering (Armstrong et al., 2005), contributing to axonal hyper-excitability and death.

**Inward Rectifier K+ Channels (Kir)**

KCNJ gene family encodes Kir channels and comprises 16 subunits of Kir1–9 subfamilies categorized into four groups: (1) classical (Kir2.x); (2) G-protein-gated (Kir3.x); (3) ATP-sensitive (Kir6.x); and (4) K+-transport channels (Kir1.x, Kir4.x, Kir5.x, Kir7.x) (Hibino et al., 2010). At a comparable driving force, Kir channels allow greater influx than efflux of K+ ions. Their high open probability at negative transmembrane voltages makes them well-suited to set the Vrest and to control cell excitability.

**Kir3.2 (KCNJ6)**

KCNJ6 gene encodes for Kir3.2 subunits, also known as G-protein-gated Kir (GIRK2) channels that are effectors for Gl/o-dependent signaling and mediate outward K+ current.

**Neurons**

Kir3.1/Kir3.2 hetero-tetramers are found in the somatodendritic compartment of neurons. Activation of GIRK channels is mediated by G-protein-coupled receptors including muscarnic, metabotropic glutamate, somatostatin, dopamine, endorphins, endocannabinoids, etc. GIRK channels are important for K+ homeostasis and maintenance of Vrest near the K+ equilibrium potential. GIRK current hyperpolarizes neuronal membrane reducing spontaneous AP firing and inhibiting neurotransmitter release (Luscher and Sleisinger, 2010). GIRK signaling contributes to learning/memory, reward, pain, anxiety, schizophrenia, addiction, and other processes (Mayfield et al., 2015). Kir3.2 mutations in mice lead to a loss of K+ selectivity and increased Na+ permeability of the channel, resulting in the weaver phenotype (Liao et al., 1996; Surmeier et al., 1996).

**Glia**

Astrocytes and Müller cells express Kir3 channels (Raap et al., 2002). Kir3.2 transcripts were detected in the mouse optic nerve (Papanikolaou et al., 2020) and oligodendrocyte lineage (Falcao et al., 2018), but not in microglia (Hammond et al., 2019).

**Expression and Function in MS**

RNA-seq revealed KCNJ6 upregulation in the CA lesions (Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq predominantly found KCNJ6 transcripts in neuronal clusters (Jakel et al., 2019; Table 1). The functional role of Kir3.2 channels in MS may be related to membrane hyperpolarization and compensation of excessive neuronal excitability driving neurodegeneration.

**Kir5.1 (KCNJ16)**

KCNJ16 gene encodes for Kir5.1 subunit, which forms an electrically silent channel when combined with Kir2.1 (Derst et al., 2001; Pessia et al., 2001), but is functional when combined with Kir4.1 (Konstas et al., 2003). Clustering of heteromeric Kir4.1/Kir5.1 and homomeric Kir5.1 channels on plasmalemma involves the anchoring protein PSD-95 (Tanemoto et al., 2002; Brasko et al., 2017). Heteromeric Kir4.1/Kir5.1 channels exhibit larger channel conductance, greater pH sensitivity, and different expression patterns if compared to Kir4.1 homomers (Tanemoto et al., 2000; Tucker et al., 2000; Pessia et al., 2001; Hibino et al., 2010).

**Neurons**

In cultures, Kir5.1 immunoreactivity was detected in somatodendritic compartments where PSD-95 immunoreactivity was also localized. The Kir5.1/PSD-95 complex may exist at dendritic spines in vivo and play a role in synaptic transmission (Tanemoto et al., 2002).

**Glia**

Kir5.1 mRNA is two-fold higher in OPCs (NG2+ glia) vs. astrocytes (Zhang et al., 2014), and mouse brain microglia expresses Kir5.1 transcript too (Hammond et al., 2019). Kir5.1 expression in oligodendrocytes and astrocytes depends on its association with Kir4.1: loss of Kir4.1 reduces Kir5.1, suggesting that altered expression/distribution of Kir5.1 may contribute to the phenotype of Kir4.1 knockout mice (Brasko et al., 2017; Schirmer et al., 2018). The oligodendroglial Kir5.1/Kir4.1 channels are important for K+ clearance (Poopalasundaram et al., 2000; Neusch et al., 2001), long-term maintenance of axonal function, and WM integrity (Kelley et al., 2018; Schirmer et al., 2018). In astrocytes, Kir5.1/Kir4.1 channels contribute to chemoreception, spatial K+ buffering, and breathing control (Mulkey and Wenker, 2011).

**Expression and Function in MS**

Bulk RNA-seq revealed Kir5.1 upregulation in CA lesions (Table 1; Elkjaer et al., 2019; Frisch et al., 2020). SnRNA-seq detected Kir5.1 in OPCs clusters and scarcely in astrocytes. The KCNJ16 gene is upregulated during demyelination and axon remyelination in mouse cuprizone model (Martin et al., 2018). Upregulation of Kir5.1 may reflect the role of the oligodendroglial Kir4.1/Kir5.1 channels in K+ clearance during MS and may represent a mechanism to compensate Kir4.1 reduction in MS brain (Schirmer et al., 2014). Alternatively, Kir5.1 upregulation may underlie reduced Kir4.1 function in MS because presence of Kir5.1 subunit confers loss of functional activity to Kir4.1/Kir5.1 channels under oxidative stress (Jin et al., 2012).
Voltage-Gated Na\(^+\) Channels (Na\(_{\text{V}}\))

In the mammalian brain, Na\(_{\text{V}}\) are composed of α-subunit (260 kDa) and one or several β-subunits (β1–β4, of 33–36 kDa) (Goldin et al., 2000). The α-subunit forms the channel pore and acts as a voltage sensor; β-subunits play a modulatory role and influence voltage dependence, gating kinetics, and surface expression of the channel (Goldin et al., 2000; Yu and Catterall, 2003; Namadurai et al., 2015). The nine Na\(_{1.1–1.9}\) α-subunits are encoded by the corresponding genes SCN1A–SCN5A and SCN8A–SCN11A. In addition, Na\(_{\text{V}}\) isoform was described, which is encoded by the SCN6/7A gene.

Na\(_{1.1}\) (SCN1A)

Neurons

Na\(_{1.1}\) channels localize to the somatodendritic compartment of principal neurons and AIS of GABAergic interneurons, spinal cord motor neurons, and retinal neurons (Ogiwara et al., 2007; Duflocq et al., 2008; Dumenieu et al., 2017). Na\(_{1.1}\) channels are also present at the nodes of Ranvier of the cerebellar WM, fimbria, corpus callosum, and spinal cord WM (Ogiwara et al., 2007; Duflocq et al., 2008; O’Malley et al., 2009). They play a role during salutatory conduction along myelinated axons and are essential for maintaining the sustained firing of GABAergic interneurons and Purkinje cells, thus controlling the excitability of neuronal networks (Duflocq et al., 2008; Dumenieu et al., 2017). Mutations in Na\(_{1.1}\) channels result in various types of epilepsy and reduced volume of brain GM and WM (Lee et al., 2017; Scheffer and Nabbott, 2019).

Glia

Human astrocytes show negligible immunolabelling for Na\(_{1.1}\) and no upregulation in the WM of MS patients (Black et al., 2010). Transcriptome analysis revealed low level of SCN1A in mouse cortical and hippocampal astrocytes (Batiuk et al., 2020). RNA-seq detected SCN1A in oligodendrocytes and OPCs throughout the CNS (Larson et al., 2016; Marques et al., 2016; Falcao et al., 2018). The functional role of Na\(_{1.1}\) channels in astrocytes and oligodendroglia remains unknown. Transcriptome studies have not detected SCN1A in microglia prepared from brain homogenates (Hammond et al., 2019), but Na\(_{1.1}\) protein was found in microglia derived from neonatal rat mixed glial cultures (Black et al., 2009). Na\(_{1.1}\) channels may be involved in regulation of phagocytosis and/or release of IL-1α, IL-β, and TNF-α from microglia (Black et al., 2009). The Na\(_{1.1}\) mRNA was detected in astrocytoma, oligodendroglioma, and glioblastoma samples from patients where these channels may contribute to the pathophysiology of brain tumors (Schrey et al., 2002).

Expression and Function in MS

Bulk RNA-seq detected upregulation of SCN2A gene in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq showed abundant SCN2A expression in neuronal clusters (Tables 1, 2; Jakel et al., 2019). The upregulation may reflect re-expression of Na\(_{1.2}\) protein, in line with previous reports showing diffuse distribution of Na\(_{1.2}\) channels along the demyelinated axons in human MS lesions within optic nerve and spinal cord (Craner et al., 2009). Axonal Na\(_{1.2}\) channels may contribute to preservation of AP propagation and re-establishment of myelin sheathes (Como et al., 2006), as it occurs during development. On the other hand, Na\(_{1.2}\) channels may promote axonal damage by increasing the intracellular Na\(^+\) concentration that triggers reversal of Na\(^+\)/Ca\(^{2+}\) exchanger (NCX) and Ca\(^{2+}\) overload in the axoplasm (Friese et al., 2014; Schattling et al., 2016). In line with this, human gain-of-function mutation in the mouse SCN2A gene triggers axonal damage, neurodegeneration, disability, and lethality in the mouse model of MS (Schattling et al., 2016). Expression of “developmental” Na\(_{1.2}\) channels in axons was also found in animal models of EAE (Craner et al., 2003; O’Malley et al., 2009), while in the spinal cord, these channels clustered at the nodes of Ranvier and localized along the demyelinated regions (O’Malley et al., 2009). SCN1A upregulation in human MS may reflect the necessity of the channel for redistribution along the demyelinated axons and support of AP propagation.

Na\(_{1.2}\) (SCN2A)

Neurons

The Na\(_{1.2}\) channels localize to the AIS, immature nodes of Ranvier, and in non-myelinated axons during early development. As nervous system matures, Na\(_{1.2}\) channels are replaced by Na\(_{1.6}\) channels (Boiko et al., 2001; Osorio et al., 2005; Dumenie et al., 2017), although in some neurons, they remain into adulthood. Na\(_{1.2}\) channels of the AIS control back-propagation of APs into the somatodendritic compartment, while Na\(_{1.6}\) channels are being placed at distal parts of the AIS control initiation and propagation of AP into the axon (Boiko et al., 2003; Hu et al., 2009). Na\(_{1.2}\) channels are also diffusely distributed along non-myelinated axons in the adult CNS where they may support slow spike propagation (Arroyo et al., 2002; Dumenieu et al., 2017).

Glia

Na\(_{1.2}\) protein was found in rat astrocytes isolated from the spinal cord and optic nerve (Black et al., 1995), but only limited Na\(_{1.2}\) expression was observed in human astrocytes in control and MS tissue (Black et al., 2010). The RNA-seq detected SCN2A expression in oligodendrocytes and OPCs (Larson et al., 2016; Marques et al., 2016). Knockdown of Na\(_{1.2}\) in pre-oligodendrocytes of the auditory brainstem resulted in reduced number and length of cellular processes and decreased MBP level, indicating that Na\(_{1.2}\) channels are important for structural maturation of myelinating cells and myelination (Berret et al., 2017). Microglia expresses no/little functional Na\(_{1.2}\) channels (Black et al., 2009; Pappalardo et al., 2016; Hammond et al., 2019).

Expression and Function in MS

Bulk RNA-seq detected upregulation of SCN2A gene in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq showed abundant SCN2A expression in neuronal clusters (Tables 1, 2; Jakel et al., 2019). The upregulation may reflect re-expression of Na\(_{1.2}\) protein, in line with previous reports showing diffuse distribution of Na\(_{1.2}\) channels along the demyelinated axons in human MS lesions within optic nerve and spinal cord (Craner et al., 2009b). Axonal Na\(_{1.2}\) channels may contribute to preservation of AP propagation and re-establishment of myelin sheathes (Como et al., 2006), as it occurs during development. On the other hand, Na\(_{1.2}\) channels may promote axonal damage by increasing the intracellular Na\(^+\) concentration that triggers reversal of Na\(^+\)/Ca\(^{2+}\) exchanger (NCX) and Ca\(^{2+}\) overload in the axoplasm (Friese et al., 2014; Schattling et al., 2016). In line with this, human gain-of-function mutation in the mouse SCN2A gene triggers axonal damage, neurodegeneration, disability, and lethality in the mouse model of MS (Schattling et al., 2016). Expression of “developmental” Na\(_{1.2}\) channels in axons was also found in animal models of
MS, i.e., in adult Shiverer mice that lack myelin (Westenbroek et al., 1992; Boiko et al., 2001), in transgenic mice overexpressing proteolipid protein that initially have normal myelination but then lose myelin (Rasband et al., 2003), and in the demyelinated optic nerve and spinal cord during EAE (Craner et al., 2003, 2004a; Herrero-Herranz et al., 2008). However, other data showed that in chronic spinal cord MS lesions, Na\textsubscript{v}1.2 channels localize on astrocytic processes surrounding the axons rather than on axons themselves (Black et al., 2007), and Na\textsubscript{v}1.2 expression/distribution was unchanged in the spinal cord of myelin-deficient rats (Arroyo et al., 2002).

**Na\textsubscript{v}1.3 (SCN3A)**

**Neurons**

Na\textsubscript{v}1.3 channels are highly expressed in rodent and human CNS throughout the embryonic development (Black and Waxman, 2013). Some studies reported that their expression decreases during the first weeks after birth, while others found Na\textsubscript{v}1.3 immunoreactivity in GM and/or WM of adult rat and human brain (Whitaker et al., 2001; Lindia and Abbadi, 2003; Thimmmapaya et al., 2005; Cheah et al., 2013). Na\textsubscript{v}1.3 channels mainly localize to the somatodendritic compartment of neurons but were also detected along the axons including myelinated fibers where they may contribute to initiation and propagation of APs (Whitaker et al., 2001; Lindia and Abbadi, 2003; Cheah et al., 2013; Wang et al., 2017). In the developing brain, Na\textsubscript{v}1.3 channels regulate proliferation and migration of cortical progenitors that do not fire APs (Smith et al., 2018).

**Glia**

The mRNA and Na\textsubscript{v}1.3 protein were detected in astrocytes (Black et al., 1995). RNA-seq demonstrated SCN3A expression in oligodendroglial cells and suggested higher expression in OPCs vs. mature oligodendrocytes (Larson et al., 2016; Marques et al., 2016). Na\textsubscript{v}1.3 expression in microglia was negligible or absent (Black et al., 2009; Hammond et al., 2019). Heterogeneous expression (from weak to strong) of Na\textsubscript{v}1.3 mRNA occurred in human astrocytoma, oligodendrogliomas, and glioblastomas (Schrey et al., 2002). Functions of Na\textsubscript{v}1.3 channels in glia remain unknown.

**Expression and Function in MS**

Bulk mRNA-seq reported upregulation of SCN3A gene in the CA lesions (Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq found significant SCN3A expression in neuronal and OPCs clusters (Jakel et al., 2019; Tables 1, 2). SCN3A upregulation during MS may reflect augmented expression of Na\textsubscript{v}1.3 protein in axons that is necessary for supporting/re-establishment of AP propagation in injured WM, because increased Na\textsubscript{v}1.3 levels are known to be associated with higher neuronal firing. For instance, mRNA and Na\textsubscript{v}1.3 protein were upregulated in spontaneously epileptic rats (Guo et al., 2008), and expression in hippocampal neurons of a novel coding variant SCN3A-K354Q resulted in enhanced Na\textsubscript{v}1.3 currents, spontaneous firing, and paroxysmal depolarizing shift-like depolarizations of the membrane potential (Estacion et al., 2010).

**Na\textsubscript{v}1.6 (SCN8A)**

**Neurons**

Na\textsubscript{v}1.6 channels cluster at high-density at the AIS and nodes of Ranvier of GM and WM axons, but can be also located on the soma, dendrites, and synapses although at a lower density (Caldwell et al., 2000; Dumenieu et al., 2017; Johnson et al., 2017; Eshed-Eisenbach and Peles, 2020). The expression level of Na\textsubscript{v}1.6 channels is low during development, but significantly increases as the nervous system matures (Boiko et al., 2001; Osorio et al., 2005; Dumenieu et al., 2017). In the adult CNS, Na\textsubscript{v}1.6 channels are the major Na\textsuperscript{+} channels responsible for initiation and propagation of APs (Boiko et al., 2003; Hu et al., 2009). Loss of Na\textsubscript{v}1.6 activity results in decreased neuronal excitability, while gain-of-function mutations potentiate excitability (O’Brien and Meisler, 2013). SCN8A mutations in mice result in ataxia, tremor, and dystonia; in humans, SCN8A haploinsufficiency is associated with intellectual disability, while hyperactivity can contribute to pathogenesis of epileptic encephalopathy (O’Brien and Meisler, 2013; Meisler, 2019).

**Glia**

RNA-seq detected SCN8A transcripts in mouse oligodendrocyte lineage (Marques et al., 2016), but they were negligible in microglia (Hammond et al., 2019). Immunoreactivity for Na\textsubscript{v}1.6 was observed in cultured spinal cord astrocytes and in brain microglia in vitro and in situ (Reese and Caldwell, 1999; Black et al., 2009; Black and Waxman, 2012; Hossain et al., 2013), but their functional role is unknown.

**Expression and Function in MS**

Bulk mRNA-seq found upregulation of SCN8A gene in CA lesions (Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq did not detect SCN8A transcripts (Jakel et al., 2019) (Tables 1, 2). Upregulation of SCN8A may reflect increased diffuse distribution of the channels along the demyelinated axons; it may be important for remyelination but may also contribute to axonal damage. Re-distribution of Na\textsubscript{v}1.6 channels, in parallel to their loss from the nodes of Ranvier, was reported previously in chronic, active, and inactive MS plaques within cerebral hemisphere, cerebellum, and spinal cord WM tissue from MS patients (Craner et al., 2004b; Black et al., 2007; Howell et al., 2010; Bouafia et al., 2014), as well as in several CNS regions affected by demyelination in animal models, including optic nerve and spinal cord WM (Craner et al., 2003, 2004a,b; Hassen et al., 2008; Howell et al., 2010). Expression of Na\textsubscript{v}1.6 channels is disrupted at the nodes of Ranvier of WM axons in Shiverer mice that lack compact myelin (Boiko et al., 2001, 2003), and in transgenic mice overexpressing proteolipid protein that initially have normal myelination but then lose myelin (Rasband et al., 2003). During EAE in animals, Na\textsubscript{v}1.6 co-localizes with NCX and may contribute to persistent Na\textsuperscript{+} influx, increased Na\textsuperscript{+} level in the axoplasm, reversal of NCX, and intra-axonal Ca\textsuperscript{2+} overload leading to axonal damage (Craner et al., 2004a). Interestingly, robust increase in Na\textsubscript{v}1.6 expression was detected also in microglia/macrophages and was associated with microglia activation and phagocytosis in human MS brain and in the EAE model (Craner et al., 2005). SCN8A deletion resulted in reduced...
inflammation and improved axonal health during EAE (Alrashdi et al., 2019). Hence, microglial Na\textsubscript{V}1.6 may contribute to the pathophysiology of MS as well, yet, snRNA-seq did not detect SCN8A in WM glia clusters (Tables 1, 2).

**Na\textsubscript{V}1.9 (SCN11A)**

**Neurons**

Although Na\textsubscript{V}1.9 channels are mainly expressed in sensory ganglia neurons (Wang et al., 2017), Na\textsubscript{V}1.9 mRNA and/or protein were detected in somat and/or proximal processes of neurons in the olfactory bulb, hippocampus, cerebellar cortex, supraoptic nucleus, and spinal cord of rodents and humans (Jeong et al., 2000; Blum et al., 2002; Subramanian et al., 2012; Wetzel et al., 2013; Black et al., 2014; Kurowski et al., 2015). Information regarding axonal labeling for Na\textsubscript{V}1.9 is lacking. Na\textsubscript{V}1.9 channels regulate excitation in hippocampal neurons in concert with BDNF and TrkB, control activity-dependent axonal elongation in spinal cord motoneurons, and mediate sustained depolarization upon current activation of M1 muscarinic receptors in cortical neurons (Blum et al., 2002; Subramanian et al., 2012; Kurowski et al., 2015). It is uncertain whether, similar to their role in the PNS (Cummins et al., 1999; Wang et al., 2017), Na\textsubscript{V}1.9 channels contribute to the regulation of V\textsubscript{rest} and AP threshold in the CNS neurons.

**Glia**

Very little expression of Na\textsubscript{V}1.9 channels occurs in astrocytes, myelinating glia, and microglia (Marques et al., 2016; Pappalardo et al., 2016).

**Expression and Function in MS**

Bulk mRNA-seq showed SCN11A downregulation in ILs (Elkjaer et al., 2019; Frisch et al., 2020). By contrast, snRNA-seq did not detect SCN11A mRNA (Jakel et al., 2019; Tables 1, 2). Functional consequence of SCN11A downregulation in MS is unknown.

**Voltage-Gated Ca\textsuperscript{2+} Channels (VGCCs)**

The VGCCs are composed of \(\alpha\)-, \(\beta\)-, \(\delta\)-, and \(\gamma\)-subunits (Catterall, 2011; Zamponi et al., 2015). The pore-forming \(\alpha\)-subunit determines channel activity, whereas other subunits are auxiliary and regulate function of \(\alpha\)-subunit. In mammalian cells, 10 different \(\alpha\)-subunits, encoded by different genes, classify into three subfamilies: Ca\textsubscript{V}1, Ca\textsubscript{V}2, and Ca\textsubscript{V}3 (Catterall, 2011; Zamponi et al., 2015; Alves et al., 2019). Depending on the pharmacological properties and activation voltage of Ca\textsuperscript{2+} currents, five different types of VGCCs are distinguished: L-type, N-type, P/Q-type, R-type, and T-type.

**L-Type VGCCs**

The \(\alpha\)-subunit of L-type VGCCs is encoded by CACNA1S (Ca\textsubscript{V}1.1), CACNA1C (Ca\textsubscript{V}1.2), CACNA1D (Ca\textsubscript{V}1.3), or CACNA1F (Ca\textsubscript{V}1.4) genes. High sensitivity to dihydropyridine modulators distinguishes L-type Ca\textsuperscript{2+} channels from other types of VGCCs. In the CNS, mainly Ca\textsubscript{V}1.2 and Ca\textsubscript{V}1.3 subunits are expressed (Lipscombe et al., 2004; Zamponi et al., 2015), but Ca\textsubscript{V}1.1 subunit was detected in human and rat basal ganglia where it is co-expressed with RyRs in GABAergic neurons (Takahashi et al., 2003).

**Ca\textsubscript{V}1.2 (CACNA1C)**

**Neurons**

Ca\textsubscript{V}1.2 channels account for 89% of all Ca\textsuperscript{2+} currents mediated by L-type VGCCs in the brain (Alves et al., 2019; Enders et al., 2020). In hippocampal neurons, Ca\textsubscript{V}1.2 channels localize to somatodendritic compartment being placed at synapses or extra-synaptically (Joux et al., 2001; Hoogland and Saggau, 2004; Obermaier et al., 2004; Tippens et al., 2008; Ortmann and Striessnig, 2016), as well as to axons and/or extrasympathetic regions of axonal terminals (Tippens et al., 2008). Within the WM, Ca\textsubscript{V}1.2 channels were identified in the developing rat pioneer axons and the follower axons projecting through the optic nerve, corpus callosum, anterior commissure, lateral olfactory tract, corticofugal fibers, thalamocortical axons, and the spinal cord (Ouardouz et al., 2003; Huang et al., 2012).

Ca\textsubscript{V}1.2 channels open upon membrane depolarization beyond \(-30\) mV, and mediate direct Ca\textsuperscript{2+} entry from the extracellular space into the cytoplasm. In addition, they may act as voltage sensors, transducing membrane depolarization to the RyRs activation and subsequent Ca\textsuperscript{2+} release from the ER via the mechanism of Ca\textsuperscript{2+}-induced Ca\textsuperscript{2+} release (CICR) (Ouardouz et al., 2003; Micu et al., 2016; Vierra et al., 2019). Clustering and functional coupling of plasmalemmal Ca\textsubscript{V}1.2 channels to RyRs of the ER is mediated by the Kv\textsubscript{2.1} channels (Vierra et al., 2019).

Neuronal Ca\textsubscript{V}1.2 channels are involved in synaptic modulation, propagation of dendritic Ca\textsuperscript{2+} spikes, regulation of glutamate receptor trafficking, CREB phosphorylation, coupling of excitation to nuclear gene transcription, modulation of long-term potentiation, spatial learning, and fear response (Hofmann et al., 2014; Hopp, 2021). During brain development, spontaneous Ca\textsuperscript{2+} transients mediated by Ca\textsubscript{V}1.2 channels regulate neurite growth and axonal pathfinding (Huang et al., 2012; Kamijo et al., 2018). Genetic variations in CACNA1C gene are associated with Timothy syndrome, Brugada syndrome, epilepsy, depression, schizophrenia, and autism spectrum disorders (Bhat et al., 2012; Bozarth et al., 2018).

**Glia**

Ca\textsubscript{V}1.2 channels are expressed in cultured astrocytes and mediate Ca\textsuperscript{2+} transients upon direct Ca\textsuperscript{2+} entry and/or subsequent activation of RyRs (D’Ascenzo et al., 2004; Du et al., 2014; Cheli et al., 2016b). Ultrastructural studies found Ca\textsubscript{V}1.2 proteins also in hippocampal astrocytes (Tippens et al., 2008). In vitro, Ca\textsubscript{V}1.2 channels contribute to the mechanism of astrogliosis (Du et al., 2014; Cheli et al., 2016b), and in mouse models of Alzheimer’s disease, they were detected in reactive astrocyte associated with A\(\beta\)-positive plaques (Willis et al., 2010; Daschil et al., 2013).

Ca\textsubscript{V}1.2 mRNA and/or protein are expressed in oligodendrocytes and their progenitors (Agrawal et al., 2000; Paez et al., 2009, 2012; Fulton et al., 2010; Haberlandt et al., 2011; Cheli et al., 2016a; Larson et al., 2016; Marques et al., 2016; Santiago Gonzalez et al., 2017; Paez and Lyons, 2020; Pitman et al., 2020). Ca\textsubscript{V}1.2 channels may regulate proliferation, migration, survival, or differentiation of OPCs, and myelination (Cheli et al., 2015, 2016a; Paez and Lyons, 2020; Pitman et al., 2020).
2020). In human cultured OPCs, static magnetic stimulation augmented CaV1.2 mRNA expression, intracellular Ca\(^{2+}\) levels, and OPC differentiation (Prasad et al., 2017), suggesting a causal relationship between these processes.

Functional expression of CaV1.2 channels in microglia is still debated (Hopp, 2021). Sequencing data showed no/low CACNA1C expression in microglia (Hammond et al., 2019), and no CaV1.2 was found in cultured microglia even upon stimulation with TNF-\(\alpha/\)IFN-\(\gamma\) (Schampel et al., 2017). However, increased immunolabelling for \(\alpha\)1C-subunit of L-type VGCCs was observed in reactive microglia during excitotoxicity in rat hippocampus (Espinosa-Parrilla et al., 2015).

**Expression and Function in MS**

Bulk RNA-seq detected increased CACNA1C expression in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq showed significant CACNA1C expression in neuronal and pericyte clusters (Jakel et al., 2019; Tables 1, 2), while low expression in OPCs and astrocyte clusters. In mouse models of MS, application of L-type VGCCs blockers reduces brain and spinal cord WM damage, decreases mitochondrial pathology in nerve fibers, attenuates axonal loss, increases oligodendrocyte survival, and promotes remyelination (Brand-Schieber and Werner, 2004; Schampel et al., 2017; Ingwersen et al., 2018; Zamora et al., 2020). These findings suggest that CaV1.2 channels contribute to damage during MS. However, expression and activity of CaV1.2 channel increased in OPCs within the demyelinated lesions in the mouse corpus callosum after cuprizone treatment (Paez et al., 2012), and deletion of CaV1.2 specifically in OPCs resulted in reduced myelination and lower MBP and MOG expression (Santiago Gonzalez et al., 2017). Hence, activity of L-type channels in oligodendroglial lineage is crucial for remyelination in this MS model, but it is unclear whether oligodendroglial CaV1.2 channels also play a role during MS in humans. Upregulation of CaV1.2 channels in pericytes may reflect altered microcirculation in MS, in analogy to the role of L-type VGCCs in pericytes outside the brain (Hashitani and Mitsui, 2019).

**CaV1.3 (CACNA1D)**

**Neurons**

CaV1.3 channels localize primarily in neuronal cell bodies and dendrites in GM (Hell et al., 1993; Zhang et al., 2005) but were also found in the developing rat optic nerve, corpus callosum (Huang et al., 2012), and axons in spinal dorsal columns of adult rats where they form clusters with RyR2s (Ouardouz et al., 2003). CaV1.3 channels activate at the membrane potential of \(-55\) mV (Lipscombe et al., 2004) and are important players in generating the pacemaking activity and spontaneous firing (Zuccotti et al., 2011). CaV1.3 channels control Ca\(^{2+}\)-dependent post-burst after-hyperpolarization in CA1 pyramidal neurons, and their activity may trigger Ca\(^{2+}\)-dependent intracellular signaling pathways (Gamelli et al., 2011; Striessnig et al., 2014). CaV1.3 channels may contribute to the mechanisms of memory because their increased expression correlates with memory loss during aging while their inhibition improves age-related memory deficits (Veng et al., 2003). Deletion of CaV1.3 channels results in increased firing rates of amygdala neurons (probably caused by a reduced slow component of post-burst after-hyperpolarization) and underlies altered fear consolidation in CaV1.3 knockout mice (McKinney et al., 2009). CaV1.3 channels are important for formation of cellular architecture: their various splice variants regulate morphology of dendritic spines while their deletion results in reduced morphology of axonal arbors (Hirtz et al., 2012; Stanika et al., 2016).

**Glia**

CaV1.3 mRNA and/or protein were detected in cultured or freshly isolated rat brain astrocytes; CaV1.3 channels may mediate intracellular Ca\(^{2+}\) increase directly and via Ca\(^{2+}\)-mediated activation of RyRs (Latour et al., 2003; Yan et al., 2013; Du et al., 2014; Enders et al., 2020). CaV1.3 expression increases in reactive astrocytes after status epilepticus in mice, suggesting that role in initiation, maintenance, or spread of seizures (Xu J. H. et al., 2007). Yet, other studies have not found CaV1.3 channels in astrocytes (D’Ascenzo et al., 2004).

CaV1.3 channels are expressed in cortical and hippocampal OPCs where they, in concert with other Ca\(^{2+}\) channels, may mediate Ca\(^{2+}\) entry from the extracellular space and/or trigger CICR from the ER (Haberlandt et al., 2011; Cheli et al., 2015). Knockdown of CaV1.3 reduces Ca\(^{2+}\) influx but does not affect expression level of myelin proteins, proliferation, or morphological differentiation of OPCs (Cheli et al., 2015). In the adult rat spinal cord WM, CaV1.3 channels are expressed by APC-positive oligodendrocytes, may mediate oligodendrocyte-axon signaling, and/or contribute to Ca\(^{2+}\)-dependent injury following trauma (Sukiasyan et al., 2009). Static magnetic stimulation may alter CaV1.3 gene expression level in human cultured OPCs (Prasad et al., 2017), suggesting that external manipulations may be a useful approach to modulate L-type VGCCs in oligodendroglial cells during diseases.

RNA-seq detected CACNA1D gene (and its various splice variants) in microglia (Hammond et al., 2019), and its expression increased upon microglia activation (Espinosa-Parrilla et al., 2015). CaV1.3 channels regulate synthesis and release of pro-inflammatory substances from microglia, e.g., NO and TNF-\(\alpha\) (Espinosa-Parrilla et al., 2015).

**Expression and Function in MS**

Bulk RNA-seq showed CACNA1D upregulation in CA lesions (Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq detected significant expression of CACNA1D in neuronal clusters (Jakel et al., 2019; Tables 1, 2). Administration of L-type VGCCs blockers resulted in multiple beneficial effects in animal MS models (see above), suggesting that CaV1.3 channels, perhaps in concert with CaV1.2 channels, contribute to tissue damage during MS.

**P/Q-Type VGCCs**

**CaV2.1 (CACNA1A)**

The CACNA1A gene encodes the pore-forming \(\alpha\)1-subunit of P/Q-type (CaV2.1) VGCCs. Sensitivity to \(\omega\)-Agatoxin distinguishes Ca\(^{2+}\) currents mediated by these channels.
Neurons

Cav_{2.1} channels localize on axonal synaptic terminals and play a fundamental role in neurotransmitter release: their direct interaction with the SNARE proteins and synaptotagmin is required for positioning the docked synaptic vesicles near the Ca^{2+} channels for fast vesicular exocytosis (Rettig et al., 1996; Zamponi et al., 2015; Mochida, 2019). Cav_{2.1} channels are also present at somatodendritic compartments of neurons (Catterall, 2000; Zamponi et al., 2015; Mochida, 2019) where they co-localize with BK and SK channels and provide Ca^{2+} for activation of these channels (Berkefeld et al., 2006; Indriati et al., 2013; Irie and Trussell, 2017). Ca^{2+} enters through the Cav_{2.1} channels and triggers further Ca^{2+} release from the intracellular stores upon activation of RyRs on the ER (Berkefeld et al., 2006; Indriati et al., 2013; Irie and Trussell, 2017). These mechanisms control neuronal firing even in the millisecond time scale (Irie and Trussell, 2017). Somatodendritic Cav_{2.1} channels regulate gene expression, local Ca^{2+} signaling, and cell survival (Pietrobon, 2010).

Cav_{2.1} channels are also present in the WM, i.e., corpus callosum and developing optic nerve (Alix et al., 2008; Nagy et al., 2017). In the optic nerve, Cav_{2.1} channels are transiently clustered in the axolemma at the sites where the underlying vesicular and tubular elements are fusing with the axonal membrane (Alix et al., 2008). Some of these sites later become nodes of Ranvier, and mutations of the α_{1A}-subunit result in malformation of the nodes of Ranvier (Alix et al., 2008). In the corpus callosum, Cav_{2.1} channels mediate fast release of glutamatergic vesicles at axon-OPC synapses, and blockade of these channels in slices reduces release at axon-glia synapses by 88% (Nagy et al., 2017).

Cav_{2.1} channels may play a role in nociception because inflammatory and neuropathic pain is altered in mice with deletion of Cav_{2.1} channels (Pietrobon, 2010). Mutations in the CACNA1A gene underlie familial hemiplegic migraine type 1, spinocerebellar ataxia type 6, and episodic ataxia type 2, and may suggest that Cav_{2.1} channels may play a role in nociception because inflammatory and neuropathic pain is altered in mice with deletion of Cav_{2.1} channels (Pietrobon, 2010; Rajakulendran et al., 2012; Izquierdo-Serra et al., 2020).

Glia

RT-PCR detected α_{1A}-subunit in mouse cortical astrocytes in culture, but Cav_{2.1} channels did not mediate Ca^{2+} entry into astrocytes (Cheli et al., 2016b). Exposure of mouse primary astrocytes to β-Amyloid did not affect Cav_{2.1} transcript levels (Daschil et al., 2014). However, increased expression of Cav_{2.1} channels was observed in reactive astrocytes after status epilepticus in mice, suggesting their role in initiation, maintenance, or spread of seizures (Xu J. H. et al., 2007). Cav_{2.1} channels are expressed in hippocampal OPCs, and in pre-myelinating oligodendrocytes of the brainstem (Haberlandt et al., 2011; Barron and Kim, 2019). In brainstem oligodendrocytes, opening of Cav_{2.1} channels is triggered upon depolarization mediated by glutamate (via AMPA receptors) or high K^{+}, as well as upon electrical stimulation of axons (Barron and Kim, 2019), suggesting that Cav_{2.1} channels mediate Ca^{2+} influx into the oligodendroglial cells upon neuronal activity in vivo. In this way, neuronal activity may trigger and/or modulate Ca^{2+}-dependent signaling in oligodendroglial cells. RNA-seq detected CACNA1A gene in microglia (Hammond et al., 2019). Cav_{2.1} channels may contribute to glioblastoma progression because their inhibition reduced proliferation of glioblastoma cells, although to a lesser extent than blockade of N-type channels (Nicoletti et al., 2017).

Expression and Function in MS

Bulk RNA-seq found CACNA1A upregulation in CA lesions (Elkjær et al., 2019; Frisch et al., 2020). The snRNA-seq revealed significant expression of CACNA1A transcripts in neuronal and OPCs clusters (Jakel et al., 2019; Tables 1, 2). CACNA1A upregulation in MS may reflect the necessity to build new nodes of Ranvier on demyelinated axons within the CA lesions. In oligodendroglial cells, Ca^{2+} entry through Cav_{2.1} channels may be required for activation of intracellular signaling pathways necessary for differentiation of OPCs and pre-myelinating oligodendrocytes.

Cav_{2.3} (CACNA1E), R-Type VGCCs

Neurons

Cav_{2.3} channels are localized to the dendritic spines and pre-synaptically (Parajuli et al., 2012). Cav_{2.3}-mediated Ca^{2+} currents activate upon strong membrane depolarization and are distinguished by sensitivity to low NiCl_{2} concentrations and SNX-482 toxin. Presynaptic R-type channels play a role in neurotransmitter release (Wu et al., 1999; Gasparini et al., 2001) and synaptic plasticity (Dietrich et al., 2003; Yasuda et al., 2003; Takahashi and Magee, 2009), but their efficiency in triggering neurotransmitter release may be lower compared to P/Q- or N-type VGCCs if they are placed distant from vesicle release sites (Wu et al., 1999). Dendritic R-type channels are coupled to SK channels and provide Ca^{2+} influx for their activation during excitatory postsynaptic potentials and back-propagating APs (Bloodgood and Sabatini, 2008; Jones and Stuart, 2013). The capacity of dendritic SK channels to promote generation of dendritic Ca^{2+} spikes also depends on Cav_{2.3} activation (Bock et al., 2019). Besides, Ca^{2+} influx via Cav_{2.3} channels may be necessary for activation of K_{4.2} channels (Wang et al., 2014). The Cav_{2.3} channels also form complexes with BK channels, and this functional interaction modulates AP properties and short-term plasticity in hippocampal neurons (Gutzmann et al., 2019). Studies in KO mice revealed that Cav_{2.3} channels are involved in the mechanisms of sleep modulation, fear response, pain, and seizures (Saegusa et al., 2000; Lee et al., 2002; Weiergraber et al., 2007; Siwek et al., 2014; Zamponi et al., 2015; Wormuth et al., 2016). Deletion of Cav_{2.3} channels in mice resulted in larger infarct size after middle cerebral artery occlusion in vivo and larger Ca^{2+} entry into the cells upon oxygen-glucose deprivation in slices, suggesting that Cav_{2.3} channels are protective during ischemic tissue damage (Toriyama et al., 2002).

Glia

In primary astrocyte cultures, mRNA and Cav_{2.3} proteins were detected using RT-PCR, Western blotting, immunohistochemistry, and electrophysiological recordings (Latour et al., 2003; D’Asenzo et al., 2004). During myelinosogenesis, oligodendrocytes within WM of the brainstem,
cerebellum, and telencephalon transiently express Cav2.3 channels, but their expression strongly decreases into adulthood (Chen et al., 2000). Ultrastructural analysis demonstrated Cav2.3 immunoreactivity in soma and processes of oligodendrocytes, paranodal loops, and loose myelin sheaths (Chen et al., 2000). RNA-seq detected only negligible CACNA1 expression in microglia (Hammond et al., 2019).

Expression and Function in MS
Bulk RNA-seq showed CACNA1E upregulation in CA lesions (Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq found significant expression of CACNA1E transcripts in neuronal clusters (Jakel et al., 2019; Tables 1, 2). The functional role of Cav2.3 channels in MS is unknown.

T-Type VGCCs
The T-type channels (Cav3) are low-voltage activated Ca\(^{2+}\) channels with \(\alpha 1\)-subunit being encoded by CACNA1G (Cav3.1), CACNA1H (Cav3.2), or CACNA1I (Cav3.3) gene. They are widely distributed in the brain, spinal cord, and DRGs. Cav3 channels activate around \(V_{\text{rest}}\), show fast inactivation kinetics (Cav3.1 > Cav3.2 > Cav3.3), and mediate tiny Ca\(^{2+}\) currents (Perez-Reyes, 2003; Weiss and Zamponi, 2019). Cav3 channels regulate neuronal excitability and play a role during rhythmic AP bursts of thalamic relay neurons, which underlie generation of neuronal oscillations under physiological (sleep) and pathophysiologial (epilepsy) conditions (Suzuki and Rogawski, 1989; Astori et al., 2011). Cav3 channels are involved in regulation of nociceptive pathways, sensory processing, hormone, and neurotransmitter release (Weiss and Zamponi, 2019). Mutations in Cav3 genes are linked to neurodevelopmental, neurological, and psychiatric diseases (Lory et al., 2020). Pharmacological non-selective T-type channel blockers are clinically used as antiepileptic drugs and also show anti-nociceptive effects (Zamponi et al., 2015).

Cav3.1, Cav3.2, and Cav3.3 (CACNA1G, CACNA1H, and CACNA1I)

Neurons
Cav3 isoforms display distinct distribution patterns with prominent somatodendritic expression in thalamic and hippocampal neurons (McKay et al., 2006). Cav2\(^{2+}\) imaging and pharmacological experiments showed that Cav3.2 and Cav3.3 subtypes located in the AIS influence the generation and the timing of APs (Bender and Trussell, 2009; Kole and Stuart, 2012). In rodent WM, Cav3 transcripts were detected at low level (Agudo et al., 2016), and information on cellular distribution is lacking.

Glia
Some studies detected Cav3.1 transcripts and proteins in rat cortical astrocytic cultures (Latour et al., 2003), while others found only scarce Cav3.1 expression in cultured astrocytes (Cheli et al., 2016b; Kim et al., 2018). Divergent findings showed that Cav3.2 immunoreactivity was absent (Chen et al., 2015) or present (Li et al., 2017) in rat spinal cord astrocytes. Cav3.1 and Cav3.2 transcripts were detected in clonal oligodendroglial CG4 cell line (Rui et al., 2020) and in OPCs isolated from mouse cortex (Zhang et al., 2014) or hippocampal slices (Haberlandt et al., 2011). In microglia, RNA-seq did not detect the Cav3 isoforms (Hammond et al., 2019).

Expression and Function in MS
Bulk RNA-seq revealed upregulation of Cav3.2 and Cav3.3 genes in CA lesions and upregulation of Cav3.1 in ILs (Elkjaer et al., 2019; Frisch et al., 2020; Table 1). The snRNA-seq detected Cav3.1 and Cav3.3 transcripts in neuronal clusters, while it did not detect the Cav3.2 (Jakel et al., 2019; Tables 1, 2). Genome-wide sequencing identified significant association of a Cav3.2 mutation (CACNA1H_R1871Q) with patients suffering relapsing-remitting MS (Sadovnick et al., 2017). Cav3 upregulation in MS lesions may be triggered by inflammatory mediators and may contribute to axonal dysfunction. Indeed, prostanoids and hydrogen sulfide modulate Cav3.2 expression and function, and increased Cav3.2 channel activity and axonal accumulation is associated with inflammation and pain (Sadovnick et al., 2017; Chen et al., 2018). T-type currents contribute to Ca\(^{2+}\)-mediated injury of spinal cord WM axons triggered by anoxia (Imaizumi et al., 1999) and to peripheral nerve injury (Watanabe et al., 2015). L-type T-type VGCC blocker lomerizine prevents retinal ganglion cell death after diffuse axonal injury (Karim et al., 2006).

Animal studies suggest that Cav3.1 upregulation in IL, a lesion type with complete demyelination and substantial axonal loss, may play a detrimental role. Specifically, the Cav3.1-deficient mice are markedly resistant to EAE induction, and this effect may be mediated by lower production of granulocyte–macrophage colony-stimulating factor (a cytokine implicated in EAE susceptibility) by CNS-infiltrating Th1 and Th17 cells (Wang et al., 2016). The Cav3.1 subunit is a functionally predominant T-type channel in CD4\(^{+}\) T cells (Trebak and Kinet, 2019). The Cav3.1-mediated Ca\(^{2+}\) increase is critical for calcineurin-NFAT activation driving transcription of cytokines in T cells, and T cells from Cav3.1-deficient mice show decreased IL-17A, IL-17F, and IL-21 production. The development of isoform-specific modulators should help in establishing the differential role of Cav3 subtypes in MS lesions.

Ryanodine Receptors
RyRs encompass three mammalian isoforms, RyR1–3, which form homo-tetrameric channels on the ER. RyRs are highly conductive Ca\(^{2+}\) channels: they get activated by Ca\(^{2+}\) influx upon plasma membrane depolarization mediating CICR from the ER (Fill and Copello, 2002; Lanner et al., 2010). In the brain, the RyR2s show predominant expression, followed by RyR3s, and then RyR1s (McPherson and Campbell, 1990; Giannini et al., 1995).

RyR2 and RyR3

Neurons
RyRs localize along ER of neurons, including WM axons (Giannini et al., 1995). They play a role in vesicle fusion, neurotransmitter release, synaptic plasticity, and growth cone dynamics (Giannini et al., 1995; Kushnir et al., 2018). RyRs form
complexes with L-type Ca\(^{2+}\) channels: RyR1-Ca\(_v\), 1.2 and RyR2-Ca\(_v\), 1.3 (Ouardouz et al., 2003). WM axons transduce membrane depolarization to Ca\(^{2+}\) release from ER, whereby L-type VGCCs gate RyRs, analogous to “excitation–contraction coupling” in muscles (Ouardouz et al., 2003; Stirling and Stys, 2010). Genetic mutations or oxidative stress can render RyRs leaky to Ca\(^{2+}\) and promote defective signals as observed in neurodegenerative disorders, heart failure, and muscular dystrophy (Kushnir et al., 2018).

**Glia**

**RYR2** and **RYR3** transcripts, but only RyR3 protein, were found in cultured astrocytes from mouse brain (Matyash et al., 2002; Keshwerani and Agrawal, 2012). *RyR2* transcripts and proteins were upregulated in spinal WM astrocytes after hypoxic injury (Keshwerani and Agrawal, 2012) and SCI (Liao et al., 2016; Pelisch et al., 2017). All **RyRs** subunits were found in rat optic nerve oligodendrocyte cultures (Ruiz et al., 2010), but **RyR3** was selectively expressed in rat cortical OPCs (Haak et al., 2001; Li T. et al., 2018). RyR3s amplify small inward Ca\(^{2+}\) currents in astrocytes and OPC, regulating behavior of these cells (Simpson et al., 1998; Matyash et al., 2002; Haberlandt et al., 2011). RyRs mediate stress response in oligodendrocytes, and RyR inhibition attenuated intracellular Ca\(^{2+}\) overload following AMPA excitotoxicity (Ruiz et al., 2010). RyR1 and RyR2 mRNAs were detected in adult human microglia, whereas only RyR3 was found in fetal microglia (Klegeris et al., 2007). RNA-seq did not detect **RYR2** and **RYR3** in mouse microglia (Hammond et al., 2019).

**Expression and Function in MS**

Bulk RNA-seq found upregulation of RyR2 transcripts in CA lesions and downregulation of RyR3 in ILs (Table 1; Elkjaer et al., 2019; Frisch et al., 2020). The snRNA-seq revealed significant expression of RyR2 in neuronal clusters and of RyR3 in the astrocyte1 cluster (Table 1; Jäkel et al., 2019). RyR subunits probably play a differential role in perturbed intracellular Ca\(^{2+}\) homeostasis in WM cells of SPMS brain. RyR2 in CA lesions may contribute to axonal dysfunction because intraxonial Ca\(^{2+}\) overload mediated by RyRs and IP3Rs activates the mitochondrial permeability transition pore and contributes to axonal dieback and degeneration following WM ischemic injury (Ouardouz et al., 2003; Stirling and Stys, 2010; Keshwerani and Agrawal, 2012) and SCI (Stirling et al., 2014; Liao et al., 2016). The RyRs inhibitor ryanodine significantly attenuates mitochondrial dysfunction (Villegas et al., 2014), axonal dieback, and secondary axonal degeneration in injured WM (Thorell et al., 2002; Stirling et al., 2014; Orem et al., 2017). In line, mice with RyR2 gain-of-function mutation exhibit more axonal damage than wild-type controls following SCI (Stirling et al., 2014), while RyR2 knockdown attenuates mitochondrial dysfunction and ER stress and improves functional recovery (Liao et al., 2016).

Functional RyR3s may contribute to astrocyte migration in response to injury, which is important for tissue remodeling and wound healing. In fact, RyR3s control astrocyte motility because astrocytes from RyR3 KO mice display reduced migratory activity (Matyash et al., 2002). Conversely, RyR3 downregulation in ILS may influence the formation of dense astrocytic scar imposing a major barrier to axonal and myelin regeneration. RyR3s also contribute to intracellular Ca\(^{2+}\) transients during OPCs differentiation, while RyR3 inhibition prevents OPCs development (Li T. et al., 2018). Interaction between RyRs and NCX in oligodendrocyte processes may represent an amplification mechanism to generate Ca\(^{2+}\) transients required for oligodendrocyte differentiation in vitro (Casamassa et al., 2016; Hammann et al., 2018; de Rosa et al., 2019; Boscia et al., 2020). However, it remains unclear whether these mechanisms play a role in human MS. The development of selective modulators will help to establish function of RyRs in MS.

**TRP Channels**

Transient receptor potential (TRP) channels are tetrameric non-selective cation channels which encompass 30 different types (Nilius and Owsianik, 2011). Upon TRP channel activation, the membrane potential depolarizes, leading to activation or inactivation of voltage-gated ion channels and regulation of Ca\(^{2+}\) signaling (Gees et al., 2010). Various intracellular or extracellular stimuli, including chemical and osmotic stress, can trigger activation of TRP channels (Clapham, 2003). TRP channels are involved in pain, regulation of neurotransmitter release, and immune functions. Vanilloid TRP channels (TRPV), melastatin TRP channels (TRPM), and polycystin TRP channels (TRPP) have been detected in WM lesions of patients with progressive MS.

**TRPV1**

**Neurons**

In the CNS, TRPV1 channels are mainly localized on cell bodies and dendritic spines, but also in synaptic vesicles (Goswami et al., 2010). TRPV1 channels are activated by exogenous (i.e., capsaicin) or endogenous (i.e., high temperatures, acid pH, anandamide, 2-arachidonoylglycerol, and lipid metabolites) stimuli (Van Der Stelt and Di Marzo, 2004). They play a role in weight, appetite, and energy homeostasis (Derbenev and Zsombok, 2016; Christie et al., 2018); synaptic plasticity (Gibson et al., 2008; Wang et al., 2020); neuropathic pain (Rivat et al., 2018); and regulation of inflammatory response (Kong et al., 2017).

**Glia**

TRPV1 channels are expressed in astrocytes (Ho et al., 2014), microglia (Sappington and Calkins, 2008), and, to a lesser extent, oligodendrocytes (Gonzalez-Reyes et al., 2013; Marques et al., 2016).

**Expression and Function in MS**

Bulk RNA-seq showed significant TRPV1 downregulation in CA lesions (Figure 2, Table 1; Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq barely detected TRPV1 (Table 2; Jäkel and Williams, 2020). The downregulated TRPV1 in CA lesions may influence neural plasticity and glia response both in the hypocellular inactive demyelinated core and in the hypercellular rim filled with activated glia. However, it is unclear whether dysfunctional TRPV1 has pro- and anti-inflammatory roles,
and whether it favors or prevents CA lesion expansion and progression, because experimental findings are inconsistent. In rodents, administration of TRPV1 agonists reduced EAE severity (Tsujii et al., 2010), while the TRPV1 antagonist capsazepine, although ineffective for EAE severity (Paltsar et al., 2013), reversed the beneficial effects of the endocannabinoid uptake inhibitor (Cabranez et al., 2005). Beneficial effects of TRPV1 may be mediated by its ability to promote micro-vesicle release from microglia, which enhances glutamatergic transmission in neurons (Marrone et al., 2017). However, on the other hand, TRPV1 stimulation induces the pro-inflammatory phenotype of microglia while downregulation promotes the anti-inflammatory phenotype (Hassan et al., 2014; Marrone et al., 2017). TRPV1 also regulates microglia migration, cytokine production, ROS generation, phagocytosis, and death (Kim et al., 2006; Schilling and Eder, 2009; Miyake et al., 2015). Furthermore, TRPV1 mediates migration and chemotaxis of astrocytes, their activation during stress and injury (Ho et al., 2014), and inflammasome activation. The picture becomes even more complex because TRPV1-KO mice show higher lethality during EAE peak but better recovery in the chronic stage (Musumeci et al., 2011). In addition, genetic deletion of TRPV1 in mice resulted in significant protection in the MOG-EAE model, and less severe breakdown of BBB (Paltsar et al., 2013). Interestingly, patients with severe MS progression show over-representation of single-nucleotide polymorphisms (SNPs) in the TRPV1 gene (Paltsar et al., 2013) that can affect the expression and activity of the channel and cortical excitability, and modulate pain (Xu H. et al., 2007; Mori et al., 2012; Stampanoni Bassi et al., 2019).

**TRPV6**

TRPV6 channels are distinguished by high Ca\(^{2+}\) selectivity (van de Graaf et al., 2006) and constitutive activity at low intracellular Ca\(^{2+}\) levels and \(V_{\text{rest}}\) (Vennekens et al., 2000). TRPV6 channels can form homo- or hetero-tetramers. TRPV5–6 are mainly expressed in epithelial and bone cells (Hoenderop et al., 2003).

**Neurons and Glia**

In the mouse brain, TRPV6 channels are expressed in neurons, while transcripts were detected in astrocytes by RNA-seq (Riccio et al., 2002; Nijenhuis et al., 2003; Batiuk et al., 2020).

**Expression and Function in MS**

Bulk RNA-seq found TRPV6 downregulation in all MS lesion types and in NAWM (Figure 2, Table 1: Elkjaer et al., 2019; Frisch et al., 2020), but snRNA-seq failed to detect TRPV6 transcripts (Table 2: Jakel et al., 2019). Little is known about the functional role of TRPV6 in brain cells. However, TRPV6 deletion in trophoblasts correlates with altered extracellular matrix (ECM) formation in the labyrinth during pregnancy (Winter et al., 2020). Hence, it will be important to investigate whether TRPV6 downregulation contributes to ECM alterations observed in SPMS lesions and believed to be a key remyelination-inhibiting factor.

**TRPM2**

**Neurons**

TRPM2 channels are found in cell bodies and neurites (Nagamine et al., 1998; Olah et al., 2009) and often co-localize with a marker of dopaminergic neurons (Bai and Lipski, 2010). They are Ca\(^{2+}\)-permeable sensors of various stimuli (Huang et al., 2020), contribute to synaptic plasticity, and inhibit neurite outgrowth (Sita et al., 2018).

**Glia**

TRPM2 transcripts are intensely expressed in mouse microglia (Malko et al., 2019), but only at lower levels in astrocytes and oligodendrocytes (Choi et al., 2015; Marques et al., 2016; Falcao et al., 2018; Batiuk et al., 2020; Table 3). TRPM2 plays a critical role in microglia activation and generation of pro-inflammatory mediators, thus contributing to neuropathic pain, brain damage due to chronic hypo-perfusion, neonatal hypoxia–ischemia, and amyloid-beta (Malko et al., 2019).

**Expression and Function in MS**

Bulk RNA-seq showed increased TRPM2 expression in the ILs (Table 1: Elkjaer et al., 2019; Frisch et al., 2020). SnRNA-seq found TRPM2 in neuronal, microglia, and ImOLG clusters. The functional role of TRPM2 channels in ILs, lesions that display reduced microglia density, axonal loss, and upregulation of stress response genes (Elkjaer et al., 2019; Frisch et al., 2020), may be related to neuronal and microglia damage. Indeed, TRPM2 channel is upregulated by diverse pathological stimuli (Malko et al., 2019) and is an important element during oxidative stress, mitochondrial dysfunction (Freestone et al., 2009), and neurodegenerative disorders (Chung et al., 2011). Constitutive TRPM2 activation is triggered by ROS and leads to pathological Ca\(^{2+}\) signaling and cell death (Eisfeld and Luckhoff, 2007; Naziroglu and Luckhoff, 2008). Knockout of TRPM2 gene in mice, or blocking the channels with miconazole, improves pathological outcome in EAE and attenuates painful behavior (Melzer et al., 2012; So et al., 2015; Tsutsui et al., 2018). TRPM2-KO mice show reduction of CXCL2 chemokine production by CNS-infiltrating macrophages and suppressed neutrophil infiltration of the brain tissue (Tsutsui et al., 2018). These findings suggest that TRPM2 may represent a promising target in SPMS.

**TRPP1 and TRPP3 (PKD2 and PKD2L2)**

The TRPP(PKD2) channels are encoded by TRPP1(PKD2), TRPP2(PKD2L1), and TRPP3(PKD2L2) genes (www.guidetopharmacology.org) and form Ca\(^{2+}\)-permeable non-selective cation channels. In the mouse brain, PKD2 and PKD2L2 transcripts are detected in neurons and glia (Table 3). TRPP1 is present on the ER, primary cilia, and plasma membrane, and TRPP3 is widely expressed in fetal tissues (Guo et al., 2000).

**Expression and Function in MS**

Bulk RNA-seq detected significant downregulation of PKD2 or PKD2L2 in ILs and CA lesions, respectively (Elkjaer et al., 2019; Frisch et al., 2020) (Figure 2, Table 1). SnRNA-seq detected PKD2 transcripts in neuronal and glia clusters but did not detect...
PKD2L2 (Table 2). It is unclear whether TRPP downregulation in MS lesions is beneficial or detrimental. On one hand, it may be detrimental because TRPP1 and TRPP3 channels are important for maintaining Ca\(^{2+}\) homeostasis and contribute to cell proliferation (Xiao and Quarles, 2010; Xiao et al., 2010), while TRPP1 knockdown results in increased susceptibility to stress-induced cell death in kidney epithelial cells (Brill et al., 2020). On the other hand, overexpression of TRPP contributes to apoptosis (Xiao and Quarles, 2010; Xiao et al., 2010), and TRPP1 is upregulated as a direct consequence of ER and oxidative stress during pathological conditions.

Chloride Channels
CIC channels mediate voltage-dependent transmembrane transport of Cl\(^{-}\). They are expressed in plasmalemma and intracellular membranes forming transmembrane dimers (Weinreich and Jentsch, 2001). CIC proteins can function as Cl\(^{-}\) channels or as Cl\(^{-}\)/H\(^{+}\) exchangers. CICs regulate \(V_{\text{rest}}\) in skeletal muscle, trans-epithelial Cl\(^{-}\) reabsorption in kidneys, and intracellular pH and Cl\(^{-}\) concentration through coupled Cl\(^{-}\)/H\(^{+}\) exchange in several cell types including brain cells.

CIC-2 (CLCN2)
The CLCN2 gene encodes a voltage- and volume-regulated CIC-2 channel (Chu et al., 1996), essential for efflux of accumulated Cl\(^{-}\) and control of cell volume homeostasis. CIC-2 is expressed in neurons and glia (Jentsch et al., 2005) and is upregulated at low osmolarity, cell swelling, and membrane hyperpolarization (Grunder et al., 1992; Clark et al., 1998).

Neurons
CIC-2 localizes on inhibitory interneurons and regulates GABA\(_{A}\) receptor-mediated synaptic inputs from basket cells (Foldy et al., 2010). Cl\(^{-}\) extrusion by CIC-2 following hyperpolarization ensures the maintenance of low intracellular Cl\(^{-}\) concentration following synaptic inhibition (Foldy et al., 2010). The link of CIC-2 mutations with generalized epilepsies in humans suggests an important role of CIC-2 in regulating neuronal excitability (Kleeu↵-Lie et al., 2009).

Glia
Astrocytes express CIC-2 that interacts with AQP4 to regulate Cl\(^{-}\) influx and efflux (Benfenati et al., 2007). CIC-2 is expressed in microglia and may regulate cell volume and phagocytosis (Ducharme et al., 2007). In oligodendrocyte lineage cells, CIC-2 positively regulates OPCs differentiation (Jentsch and Pusch, 2018) and transcription factors for myelin genes, thus contributing to myelin formation and WM integrity (Hou et al., 2018).

Expression and Function in MS
Bulk RNA-seq showed significant CLCN2 downregulation in CA lesions (Elkjaer et al., 2019; Frisch et al., 2020; Figure 2, Table 1). SnRNA-seq detected CLCN2 transcripts in oligodendrocyte clusters, while they were only faintly observed or absent in other clusters (Table 2). Several findings suggest that CLCN2 downregulation in MS may reflect altered WM integrity and/or contribute to the mechanisms of myelin destruction: first, CLCN2\(^{-/-}\) mice exhibit abnormal WM morphology (Blanz et al., 2007); second, loss-of-function CLCN2 mutations lead to leukodystrophy; third, loss of cell adhesion molecule GlialCAM, which binds to CIC-2 in glia, is associated with leukodystrophy (Jeworutzki et al., 2012; Hoegg-Beiler et al., 2014). Of note, though, is a recent report showing that leukodystrophy fully develops only when CIC-2 is disrupted in both astrocytes and oligodendrocytes (Goppner et al., 2020). It remains to be investigated whether CLC-2 loss in glia contributes to the failure of myelin repair in human CA lesions.

CIC-7 (CLCN7)
The CLCN7 gene encodes for the chloride-proton antiporter CIC-7 localized to lysosomes and crucial for function of osteoclasts and brain cells (Kornak et al., 2001; Jentsch and Pusch, 2018).

Neurons and Glia
In mice, neurons and microglia express CIC-7 protein (Kasper et al., 2005; Majumdar et al., 2011; Weinert et al., 2014), while transcripts were found in astrocytes and oligodendrocyte lineage (Falcao et al., 2018; Batiuk et al., 2020). Mutations in the human CLCN7 gene are associated with osteopetrosis and neurodegeneration (Kornak et al., 2001).

Expression and Function in MS
Bulk RNA-seq detected significant CLCN7 downregulation in CA lesions (Elkjaer et al., 2019; Frisch et al., 2020; Figure 2, Table 1). SnRNA-seq found CLCN7 transcripts in neuronal and all glia clusters (Table 2). Functional role of CIC-7 under demyelinating conditions is unknown. In neurons, CIC-7 on lysosomes contributes to the function of the endosomal–lysosomal pathway (Poet et al., 2006; Bose et al., 2021). Lysosomal localization of CIC-7 increases during microglia activation, leading to increased lysosomal acidification and A\(_{\beta}\) degradation (Majumdar et al., 2011). CICN7-deficient mice display widespread WM atrophy, neuronal loss, microglia activation, astrocytosis, and accumulations of storage material in lysosomes (Kornak et al., 2001; Kasper et al., 2005; Pressey et al., 2010). In SPMS lesions, dysfunctional CIC-7 activity may directly affect the luminal pH and Cl\(^{-}\) concentrations and lysosomal protein degradation (Wartosch et al., 2009), which, in turn, may lead to neuronal and glial degeneration in the WM.

Connexins
Connexins (Cxs) are transmembrane proteins with channel and non-channel functions. Channel functions include the formation of gap junctions (GJs) and hemichannels (HCs) (Saez et al., 2003; Wang et al., 2013; Gajardo-Gomez et al., 2016), while non-channel functions involve adhesion properties and intracellular signaling (Zhou and Jiang, 2014; Leithe et al., 2018). More than 20 Cxs genes have been described in humans, and 11 of them are expressed in the brain (Willecke et al., 2002; Theis et al., 2005). Cxs are essential players in ionic homeostasis, intercellular Ca\(^{2+}\) signaling and Ca\(^{2+}\) waves propagation, glialtransmission, synaptic transmission and plasticity, brain metabolism, brain-blood barrier development and integrity, and myelination.
(Takeuchi and Suzumura, 2014). In the WM, GJs are essential for K+ buffering in response to neuronal activity, they facilitate transport of nutrients and ions from oligodendrocyte soma to myelin layers and from astrocytes to oligodendrocytes (Bradl and Lassmann, 2010). In the WM, HCs are involved in metabolic coupling and energy supply to neurons, and provide a major pathway for glucose entry into OPCs and oligodendrocytes (Niu et al., 2016).

Cx37 (GJA4)
Cx37, encoded by GJA4 gene, predominantly builds heterotypic GJs with Cx40 and Cx43 in vascular cells and plays an essential role in vasomotor activity, endothelial permeability, and maintenance of body fluid balance (Falcao et al., 2018; Li et al., 2018).

Expression and Function in MS
Bulk RNA-seq revealed significant GJA4 upregulation in ILs (Elkjaer et al., 2019; Frisch et al., 2020), while snRNA-seq showed high GJA4 expression in pericyte cluster (Jakel et al., 2019; Tables 1, 2). In chronically demyelinated axons, as those within ILs, hypoxia due to imbalance between increased energy demand and reduced ATP production because of mitochondrial dysfunction may drive angiogenesis. However, while providing trophic factors for tissue remodeling, angiogenesis may contribute to hyperperfusion and neurovascular uncoupling (Girolamo et al., 2014). Interestingly, Cx37 knockdown with siRNA in human umbilical vein endothelial cells diminishes capillary branching (Gartner et al., 2012), but Cx37−/− mice develop a more extensive vasculature under ischemic conditions and show enhanced recovery after hind limb ischemia (Fang et al., 2011). In the future, it will be important to investigate whether Cx37 protein contributes to aberrant cerebrovascular and angiogenic responses in human ILs during MS.

Pannexins
The Pannexin (Px) family consists of three members, encoded by Panx1, Panx2, and Panx3 genes. Pannexins do not form GJ in vivo but operate as plasma membrane channels (pannexons) and participate in paracrine and autocrine signaling in brain GM and WM (Sosinsky et al., 2011; Sahu et al., 2014; Dahl, 2015).

Px1 (PANX1)
Px1 is permeable to anions, some negatively charged molecules (glutamate, aspartate, and ATP), and fluorescent dyes (Ma et al., 2012; Yeung et al., 2020). Opening of Px1 may be promoted by voltage, increased intracellular Ca2+, mechanical stress, extracellular K+, oxygen deprivation, caspases cleavage, ATP binding to P2Y or P2X7 receptors, activation of α1-adrenergic, NMDA, and thromboxane receptors (Chiu et al., 2018; Dahl, 2018; Whyte-Fagundes and Zoidl, 2018).

Neurons and Glia
Px1 is distributed in GM and WM regions, including cerebellum, corpus callosum, and fimbria fornix of mice (Bruzzone et al., 2003) and rats (Vogt et al., 2005). Px1 is expressed in neurons, astrocytes, microglia, oligodendrocytes, vascular cells, and peripheral immune cells (Iglesias et al., 2009; Swayne et al., 2010; Orellana et al., 2013; Good et al., 2018; Lapato and Tiwari-Woodruff, 2018). In neurons, Px1 may be co-expressed with Px2 and is found in cell soma, dendrites, and axons (Cone et al., 2013).

Interaction between Px1 and purinergic signaling deserves special attention because Px1 forms complexes with P2X7Rs (Taruno, 2018). Binding of ATP to P2X7-R triggers opening of Px1 channels with subsequent ATP release (Locovei et al., 2007; Iglesias et al., 2008; Pelegrin et al., 2008; Chiu et al., 2018). ATP signaling involving Px1 channels regulates neurite outgrowth and synaptic plasticity in neurons, while in glia, it underlies intercellular propagation of Ca2+ waves, cell differentiation, and migration (Giaume et al., 2021).

Expression and Function in MS
Bulk RNA-seq showed significant PANX1 upregulation in ILs (Table 1; Elkjaer et al., 2019; Frisch et al., 2020), but snRNA-seq did not detect PANX1 transcripts (Jakel et al., 2019). ILs are lesions with little/no inflammatory activity but with sharply demarcated hypocellular areas of demyelination and axonal degeneration. Px1 activation is known to enable ATP release, and ATP is a “find me” signal promoting chemotaxis of microglia/macrophages to the injury site for fast clearance of dead cells and a molecule important for myelination (Chekeni et al., 2010; Gajardo-Gomez et al., 2016). Hence, PANX1 upregulation in ILs may be a compensatory mechanism that stimulates glial activity. On the other hand, upregulated Px1 mRNA expression in cerebellum and spinal cord in chronic EAE contributes to WM damage (Lutz et al., 2013). Uncontrolled opening of P2X7-R-Px1 complex in response to demyelination triggers excessive glutamate and ATP release, altered Ca2+ dynamics, excitotoxicity, damage of axons, and myelin (Orellana et al., 2011; Crespo Yanguas et al., 2017). Knockout or blockade of Px1 with probenecid in rodents restrains EAE symptoms and results in reduced inflammation and decreased oligodendrocyte damage (Hainz et al., 2017), suggesting that Px1 activity supports damage during MS. More studies are required to establish how Px1 should be modulated in order to halt neurodegeneration during MS.

CatSper Channels
CatSperg and Caspere
Cation channel of spermatozoa (CatSper) is a highly complex multi-subunit voltage-gated Ca2+-permeable ion channel. Four distinct α-subunits (CatSper1–4) and several accessory subunits are encoded by CATSPERE genes (Qi et al., 2007). The CatSper channel is essential for the activity of sperm flagellum and sperm fertility (Lishko and Mannowetz, 2018). RNA-seq detected only CATSPERG transcripts in mouse neurons, oligodendrocytes, and microglia (Marques et al., 2016; Hammond et al., 2019; Jakel et al., 2019).

Expression and Function in MS
Bulk RNA-seq found downregulation of the auxiliary subunit gamma (CASPERG) and epsilon (CATSPERE) in CA lesions and ILs, respectively (Table 1; Elkjaer et al., 2019; Frisch et al., 2020). SnRNA-seq did not find CATSPERE transcripts and barely detected CATSPERG in neuronal and glia clusters. It is difficult to
speculate on the role of CatSper channels in MS lesions because characterization of these subunits is limited to sperm cells, and no data on CatSper protein expression or function in the brain are available.

CONCLUSIONS

Understanding how distinct ion channels regulate CNS ionic homeostasis in WM neurons, axons, glia, and vascular cells under chronic demyelinating conditions is of critical importance for the development of novel therapeutic strategies to prevent neurodegeneration and disability progression and improve functional recovery and repair in MS. Recent Bulk RNA-seq (Elkjaer et al., 2019; Frisch et al., 2020) revealed a considerable number of ion channel genes that are altered in different types of WM lesions of the SPMS brain, particularly in WM CA lesions, a type of lesion that develops in MS patients despite disease-modifying therapy and predicts a more aggressive disease course (Absinta et al., 2019; Elliott et al., 2019). SnRNA-seq found that transcripts for dysregulated ion channels belong to the clusters of neurons, astrocytes, oligodendrocyte lineage, microglia/macrophages, and pericytes (Jäkel et al., 2019). The dysregulation of ion channel genes in MS may be detrimental or beneficial for functions of neurons, including interstitial neurons. Intense upregulation of genes encoding voltage-gated Na$^+$ channels in CA lesions may reflect the imbalance of Na$^+$ homeostasis observed in SPMS brain (Inglese et al., 2010). Conversely, the upregulation of a large number of voltage-gated K$^+$ channel genes may be linked to a protective response to limit neuronal excitability. The altered Cl$^-$ homeostasis, revealed by the significant downregulation of voltage-gated Cl$^-$ channels in MS lesions, may contribute to an altered inhibitory neurotransmission and increased excitability. Depending on the type of alterations, dysregulated ion channels in MS may favor AP propagation and dampen neuronal hyperexcitability or, on the contrary, may contribute to axonal dysfunction and cell death. Altered expression and/or function of ion channels may also influence key properties of glia including proliferation, migration, spatial buffering, cytokine release, cell metabolism, myelin repair, angiogenesis, BBB permeability, and several other important functions.

We described the importance of uniquely dysregulated genes well-known to play a role in WM dysfunction in the MS brain (KCNA1, KCNA2, SCNA2A, and SCNA8), or in experimental models of MS (KCN3, KCNQ3, KCNK2, CACNA1C, CACNA1G, TRPV1, TRPM2, and PANX1). Furthermore, we highlighted the importance of ion channel genes that are uniquely dysregulated in SPMS lesions but have never been previously explored in MS brain. Those genes are expressed in OPC (KCND2, SCNA1, SCNA3, and CACNA1A), ImOLG (KCNQ3), mature oligodendrocyte (KCNH8), microglia (KCNQ3), astrocyte (KCNN3 and RYR3), and pericyte (GJA4 and CACNA1C) clusters of healthy and SPMS brain. It remains to be investigated whether and how the ionic imbalance in different glial cells, particularly oligodendroglia, contributes to impaired recovery and failure of myelin repair.

Several genes, including KCNA1, SCNA8, SCNA11A, CACNA1H, PKD2L2, TRPV6, PANX1, and CATEPHERE transcripts, were detected in bulk transcriptome (Elkjaer et al., 2019; Frisch et al., 2020), but were not found by snRNA-seq (Jäkel et al., 2019). This discrepancy may be explained by several observations: (1) the transcriptional profiling may vary when lesions analyzed by different studies come from different WM regions (Jäkel and Williams, 2020); (2) snRNA-seq analysis lacks information on gene expression in WM axons that may also contain ion channel transcripts; and (3) snRNA-seq only includes RNA transcripts from the nucleus and may therefore lack RNA transcripts from cytoplasm.

Future experiments on dysregulated ion channels predicted by transcriptomic analysis are expected to provide a better understanding of the molecular mechanism of MS progression and may pave the way for the identification of new therapeutic targets to limit lesion expansion, reduce neurological impairment, and stimulate functional recovery.

AUTHOR CONTRIBUTIONS

FB, MK, ZI, and ME: writing-original draft preparation. FB and MK: writing-review and editing. FB, MK, and ZI: funding acquisition. All authors have read and agreed to the final version of the manuscript.

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