Introduction

Gangliosides are sialic acid containing glycosphingolipids widely expressed in vertebrate plasma membranes and intracellular compartments (Ledeen & Wu, 2011; Sandhoff & Harzer, 2013; Yu et al., 2011). Whilst ganglioside biosynthesis and distribution is widespread throughout the body, the nervous system in particular is very highly enriched in complex gangliosides relative to other organs and tissues. Gangliosides are present in both central and peripheral nervous system (CNS and PNS) grey and white matter, where they have roles in development and homeostasis, modulating cell-cell recognition, signal transduction, growth and motility. Gangliosides are key components of lipid raft membrane domains where they have been widely studied (Simons & Toomre, 2000; Sonnino et al., 2015). In this review, we will focus our attention on the role of gangliosides in the maintenance of the mammalian node of Ranvier (NoR), a specialised membrane domain formed by myelin-forming cells: Schwann cells in the PNS and oligodendrocytes in the CNS (Figure 1). The NoR is a critical relay station in the electrophysiological functioning of myelinating nerve fibres, essential for saltatory conduction, promoted by high density clustering of voltage-gated sodium (Nav) channels on the nodal axolemma. Anchoring of the myelin paranodal loops by adhesion molecules and their associated cytoskeletal linker proteins in both the glial and axonal membranes at the paranode segregate and condense the ion channels (Pinatel & Faivre-Sarrailh, 2021; Rasband & Peles, 2021). Of note to our review, disruption to the expression of the paranodal axo-glial membrane protein complex composed of glial Neurofascin 155 (NF155), axonal contactin-associated...
protein (Caspr) and Contactin markedly alters the normal conformation of the NoR (Bhat et al., 2001; Boyle et al., 2001; Pillai et al., 2009). A key readout of this includes a lengthening of nodal Nav channel domains and an invasion of juxtaparanodal voltage-gated potassium (Kv) channel domains into the paranode. Whilst the protein composition at the NoR has been the subject of a wealth of structural and functional studies, the glycolipid/ganglioside component and its effects on nodal integrity has been less extensively studied. The composition and physiology of PNS and CNS nodes have many similarities, but also differences; where relevant we will highlight distinctions between the two sites, but in general refer to the NoR as the generic term for both sites. All the information contained herein refers to the mouse NoR, except where specified otherwise. It has long been known that it cannot be assumed, and has never been systematically proven, that ganglioside distribution biology is identical in all species (Suzuki, 1965), or indeed at all NoR which also differ widely according to site and nerve fibre type.

2 | GANGLIOSIDE BIOSYNTHESIS

Gangliosides are synthesised in intracellular organelles through stepwise addition of donor sugars to an elongating glycan core, mediated by a family of developmentally and spatially regulated glycosyltransferases (Yu et al., 2011) (Figure 2a). They are subsequently trafficked to the plasma membrane, then recycled or degraded by lysosomal exoglycosidases. The ability
to inactivate glycosyltransferases, through genetic deletion in mice (Figure 2a) or through studying natural human mutations, has led to many of the recent advances in understanding ganglioside biology (Schnaar, 2016). GM3 is the first simple ganglioside synthesised through the addition of $\alpha$2,3-linked sialic acid to a lactosylceramide core. GM3 is then modified by the $\beta$1,4-N-acetylgalactosaminyltransferase 1 enzyme ($\text{B4galnt1}$ or $\text{GalNAc-T}$) enzyme to produce $a$-series gangliosides, respectively. It is GM1 and GD1a from the $a$-series, and GD1b and GT1b from the $b$-series that form the majority of the gangliosides expressed in the normal nervous system. The predominant gangliosides found in neural tissue are thus the complex gangliosides GM1, GD1a, GD1b and GT1b, although many minor gangliosides are also expressed, likely because of high activity of their synthesising enzyme, GalNAc-T in the nervous system (Dicesare & Dain, 1971). Humans with inherited ganglioside enzymatic deficiency syndromes develop complex neurodevelopmental and degenerative disorders (Boukhris et al., 2013; Simpson et al., 2004). Whilst these human disorders display a widespread phenotype, detailed morphological analysis of the NoR has not to date been conducted.

3 | GANGLIOSIDES AND OTHER GLYCOLIPIDS AT THE NODE OF RANVIER

Apart from their intrinsic functional roles, gangliosides also act as receptors for many microbial pathogens and toxins (Bullens et al., 2002; Ravindran et al., 2013) and peripheral neuropathy-associated autoantibodies (Kusunoki et al., 2008; Willison & Yuki, 2002). In these roles, the functional effects of ligand binding to gangliosides at NoRs have been determined not to induce pathophysiological perturbations, with the exception of fairly extensive studies on neuropathy-associated autoantibodies as described below. Knowledge of the anatomical localisation of gangliosides to the NoR first came from studies using ganglioside binding ligands (Ganser et al., 1983). Subsequently a wide range of immunohistological studies with mouse monoclonal antibodies (McGonigal et al., 2010; Paparounas et al., 1999; Susuki et al., 2007a, 2007b) and human antisera (Chiba et al., 1993; De Angelis et al., 2001; Illa et al., 1995; Lugaresi et al., 1997; O’Hanlon et al., 1998) have been conducted.
et al., 1998; Paparounas et al., 1999) demonstrated immunolocalisation of gangliosides to the nodal gap and paranodal structures, including GM1, GD1a and GQ1b. Much of this research was driven by investigators working on autoimmune neuropathy, in which anti-ganglioside antibody mediated injury at the node is believed to account in part for the paralytic features of disease (Griffin et al., 1996; Susuki et al., 2012). Since the small size of the nodal region prevents it from being spatially resolved either using mass-spectrometric imaging or biochemical isolation techniques, it has not been possible to unequivocally identify specific gangliosides in nodal regions, beyond extrapolation from immunolabelling methods. Transgenic mice with disruptions in specific glycosyltransferase genes, and thus deficient in the composition of all downstream gangliosides, have been critical tools in advancing understanding of their roles at NoRs, but since they are generally deleted ‘en bloc’ and subject to precursor build up according to the enzyme position in the biosynthetic pathway, attribution of specific functions to specific gangliosides has also not been possible.

3.1 | Galactocerebroside and sulphatide

The glycolipids galactocerebroside (GalC) and sulphatide, whilst not gangliosides (as they lack the defining feature, sialic acid), require special mention at the NoR where they have been first and extensively studied in transgenic mice (Coetzee et al., 1996; Dupree et al., 1998; Honke et al., 2002; Hoshi et al., 2007; Ishibashi et al., 2002). Through enzyme knockout technology, the myelin-localised glycosphingolipids, sulphatide and galactocerebroside were first shown to have a significant role in myelinated nerve maintenance. Their distinct or interdependent roles and interactions are unclear, particularly at the NoR, as GalC deficiency inevitably also leads to sulphatide deficiency which is downstream in the biosynthetic pathway (Figure 2a). Disruption of UDP-galactose:ceramide galactosyl transferase (CGT) results in mice that are incapable of synthesising either of these lipids; mice thus express a severe dysmyelinating phenotype including NoR disruption, which is more severe in the CNS than PNS (Coetzee et al., 1996; Dupree et al., 1998). Sulphatide, 3-O-sulphoglucosylceramide, synthesised from GalC through the addition of a 3-O-sulphate group by cerebroside sulphotransferase enzyme (CST), is enriched in the outer leaflet of the myelin and uncompacted glial membranes (Ishizuka, 1997) and small amounts are present in neurons and astrocytes (Eckhardt, 2008). Evidence from CST−/− mice indicates that sulphatides are crucial for the maintenance and stability of the NoR (Honke et al., 2002; Hoshi et al., 2007; Ishibashi et al., 2002). Indeed, in the absence of sulphatide, loss of NF155 immunostaining suggests a role in tethering this key paranodal axo-glial adhesion molecule to the glial membrane (Hoshi et al., 2007; McGonigal et al., 2019). However, only recently, owing to the development of new anti-sulphatide antibodies, have
FIGURE 4  Schematics depicting the likely impact of increasing glycolipid deficiency on paranodal arrangement at nodes of Ranvier based on known data from the PNS and CNS. Particular focus is placed on the paranodal axo-glial proteins Caspr, NF155 and MAG about which most is known. Lengthening of Nav channel domains are indicated by the pink arrow and invasion of juxtaparanodal Kv channels into the paranode suggestive of breakdown of the axo-glial junction at the paranodal/juxtaparanodal border, by the green arrow. The transgenic lines represented by each stage of nodal organisation are defined below the relevant schematic, and age is also defined. In ganglioside deficient states, there is nodal lengthening, followed by Caspr/NF155 disturbance and Kv1.1 invasion into the lateral paranode. Normally GD1a and GT1b in rafts will tether MAG but in their absence, MAG does not make the axo-glial connection. If NF155 is present, this protein can partner with Caspr/Contactin to make an axo-glial junction. However, when sulphatide is absent, NF155 is also lost from the paranode. In the absence of NF155, Caspr presence is also diminished, and this is exacerbated with the additional loss of complex ganglioside rafts. Under both ganglioside and sulphatide raft deficiency, we propose that MAG and NF155 are both absent from the paranodal axo-glial domain. This figure uses schematics modified from Figure 8, McGonigal et al., (2019) (https://doi.org/10.1523/JNEUROSCI.2095-18.2018) with permission under the Creative Commons CC-BY licence (http://creativecommons.org/licenses/by/4.0/)
immunolabelling studies shown that sulphatide appears highly enriched in paranodal loops, as well as being present along the Schwann cell membrane (Meehan et al., 2018) (Figure 3a).

3.2 | Complex gangliosides

The advent of glycosyltransferase transgenic mice lacking specific ganglioside species revealed both myelin and nodal disorganisation as a result of different deficiencies, thereby inferentially indicating a role for gangliosides in nodal organisation and maintenance (McGonigal et al., 2021; Susuki et al., 2007a; Yamashita et al., 2005).

The first mouse generated and relevant to the NoR had disruption in the B4galnt1 gene and thus deficiency in GalNAc-T (Takamiya et al., 1996). These mice are viable and appear grossly normal, indicating that complex ganglioside expression is not necessary for normal development, but later develop an age-dependent neurodegenerative phenotype characterised by weakness, ataxia, motor deficits, nerve degeneration and demyelination (Chiavegatto et al., 2000; Sheikh et al., 1999; Takamiya et al., 1996). In particular, loss of nodal axo–glial junction adhesion and integrity, disrupted paranodal loop attachment, invasion of voltage-gated potassium channels from the juxtaparanode to the paranode coinciding with attenuated paranodal Caspr and NF155 immunostaining at this border was observed (Susuki et al., 2007a). In these mice, overexpression of the precursor simple gangliosides GM3, GD3 and 9-O-Ac(etyl)-GD3 may play a compensatory developmental role that limits the severity of the phenotype (Furukawa et al., 2008; Ngamukote et al., 2007). Mice deficient in GD3 synthase and thus lacking b-series gangliosides whilst over-expressing a-series gangliosides (GD3s−/−) are grossly normal throughout life but repair peripheral nerve poorly (Okada et al., 2002). Nodal abnormalities have not been observed to date in GD3s−/− mice (McGonigal et al., 2010; Susuki et al., 2007a) which could indicate the particular importance of a-series gangliosides in maintaining the NoR. Mice whose ganglioside repertoire is restricted to GM3 (GalNAc-T and GD3s double knock-out) develop lethal audiogenic seizures (Kawai et al., 2001), age-dependent progressive motor and cognitive deficits (Tajima et al., 2009), and sensory loss (Inoue et al., 2002). The NoR in GM3 only mice was recently reported to show similar disruptions in nodal immunostaining to those found in GalNAc-T−/− mice (McGonigal et al., 2021), confirming the significance of complex gangliosides to this site as further loss of GD3 added minimal impact. Complete ganglioside ablation is not embryonic lethal; however, from 2 weeks of age, mice undergo progressive and severe neurodegeneration resulting in death at 2 months although the contribution of nodal dysfunction to this has not been assessed (Yamashita et al., 2005). Most ganglioside deficient mice display age-dependent abnormalities but are not as severely impaired as nodal protein null mice which have a marked reduction in life-expectancy. Additionally, CNS NoR are often more perturbed than PNS NoR, which could reflect the greater influence that the paranode bestows on CNS stability (Rasband & Peles, 2015). Together, these mouse data are suggestive of a more fundamental necessity for a-series gangliosides in age-related nervous system maintenance, although this is difficult to conclusively prove because a mouse with selective deficiency of a-series gangliosides has not been generated.

3.3 | Gangliosides, sulphatide and myelin-associated glycoprotein

A specific and interesting issue at the NoR relates to interactions between the glial myelin-associated glycoprotein (MAG) and gangliosides. The complex gangliosides GD1a and GT1b have been identified as axonal ligands for MAG (Collins et al., 1997; Vinson et al., 2001) (see Figure 2a), localised on the innermost wrap of the myelin sheath, paranodal loops and Schmidt-Lanterman incisures where glial cytoplasm is present (Bartsch et al., 1989; Trapp et al., 1989). The age-related degenerative phenotype, functional and morphological deficits displayed by GalNAc-T−/− mice paralleled those found in MAG−/− mice, both showing greater severity in the CNS than the PNS. Interestingly, generation of double deficient mice created from interbreeding GalNAc-T−/− and MAG−/− genotypes did not further exacerbate the phenotype of the single-null mice. This suggests a complementary and functional interaction between complex gangliosides and MAG in nerve integrity (Pan et al., 2005). Mice that lack key sialylation enzymes proposed to generate both GD1a and GT1b show a significant reduction in brain MAG immunostaining that coincides with abnormal nervous system development showing the specificity of this interaction (Sturgill et al., 2012). With the knowledge that both MAG and galactolipids modulate axo–glial stability, Marcus et al. (2002) investigated the significance of both molecules to axo–glial integrity by crossing MAG deficient mice with GaIC/sulphatide double-deficient mice, which individually have similar phenotypes. This genetic combination resulted in a lethal phenotype, with survival up to post-natal day 22 (P22). Again, the NoR appear to develop normally followed by subsequent generalised impairment of the paranodal axo–glial junction.

We advanced these studies to consider the compensatory roles of complex gangliosides and sulphatide in maintaining axo–glial stability at NoR by combining GalNAc-T−/− and CST−/− strains and examining the nodal phenotype (McGonigal et al., 2019). Depletion of both gangliosides and sulphatide exacerbated the phenotype, suggesting two independent roles. In these animals, a severe neurodevelopmental phenotype occurred with early death around P22. Degenerating axon number increased with diminishing lipid content and conduction became increasingly impaired. MAG expression in the myelin fraction from brain homogenates was significantly lower in the CST−/− x GalNAc-T−/− genotype compared with single deficiency genotypes and Nav channel cluster number, Caspr dimer number, and Nav channels flanked by NF155 dimers decreased with decreasing lipid expression. Once again the nodal phenotype was more severe in CNS compared to PNS nerve. Since MAG acts as a
myelin receptor for axonal gangliosides and also may be localised to the myelin membrane by sulphatide-rich lipid rafts, loss of both the axonal and glial membrane localisation domains for MAG could contribute to this severe phenotype. A progressive reduction in MAG and a significant reduction in NF155 (25% compared to wild type) in CST−/− mouse brains was recently reported (Palavicini et al., 2016), and extraction studies have suggested that sulphatide-containing lipid rafts could be anchors for MAG and NF155 (Pomicht et al., 2013). Taken together, it seems that glial sulphatide may act as a wide-ranging anchor for multiple myelin and glial membrane proteins in a complementary and interdependent relationship with axonal gangliosides.

4 | USE OF TISSUE SPECIFIC TRANSGENIC MICE TO STUDY GANGLIOSIDE FUNCTION AT THE NODE OF RANVIER

Whilst the above studies have indicated that complex gangliosides are required for nervous system maintenance, stability and repair including the NoR, it is unknown whether neuronal or glial ganglioside deficiency has the greater impact on the age-related phenotype and maintenance of the axon, myelin and axo-glial junction. Unlike GalC and sulphatide, which are almost exclusively localised to glial membranes, gangliosides are expressed in both glia and neurons (Gong et al., 2002; Ogawa-Goto & Abe, 1998), and thus function cannot be precisely attributed to tissue expression. An added complexity is that gangliosides can transfer between membranes by shedding and uptake (Heffer-Lauc et al., 2005; Olshelfski & Ladisch, 1996), potentially confounding the concept of discrete membrane localisation. Additionally there is likely to be heterogeneity in ganglioside lipid raft composition within a single membrane; for example distinct separation between GM1- and GD3-containing rafts has been reported (Vyas et al., 2001). To assess the relative significance and necessity of complex ganglioside expression in either neuronal or myelin-forming cells, we developed “rescue” mice. On a GalNAc-T−/− background, these rescue mice selectively express gangliosides either neuronally [GalNAc-T being driven by the neurofilament light (NFL) or Thy1 promoters: GalNAcT−/−-Tg(neuronal)] or in myelin [GalNAc-T being driven by the proteolipid protein (PLP) promoter: GalNAcT−/−-Tg(gial)] (McGonigal et al., 2016; Yao et al., 2014) (Figure 2b). Through this selective reintroduction of glycosyltransferase activity in a site-specific manner, we observed that neuronal, and not glial, rescue of complex gangliosides was both necessary and sufficient to prevent the age-dependent neurodegenerative phenotype seen in global GalNAcT−/− deficiency states (Yao et al., 2014). Despite the potential for ganglioside transfer between membranes as mentioned above, we detected no evidence of this transfer between axons and glia, or vice versa in our transgenic rescue mice. Additionally, there was a selective absence of ganglioside expression in primary cultured neurons or glial cells from our GalNAcT−/−-Tg(gial) or GalNAcT−/−-Tg(neuronal) transgenic lines, respectively, which validated the use of the cell-specific promoters. These findings clearly demonstrate the prime importance of neuronally expressed GalNAc-T in maintaining nervous system integrity throughout the lifespan.

With specific reference to the NoR, we considered that, because these glycolipids are differentially expressed in glial and axonal membranes (Figure 3b), they may act in partnership to retain clustered proteins in their respective domains. There is evidence that complex ganglioside deficiency can disturb lipid raft associated anchoring of membrane proteins as has been shown for complement regulators (Ohmi et al., 2011). Axonal Caspr and glial NF155 have been proposed to be in lipid rafts (Schafer et al., 2004), and ganglioside loss disrupts these raft-associations which coincides with loss of paranodal integrity (Susuki et al., 2007a). We sought to test the importance of specific ganglioside membrane expression by interbreeding strains of single-null mice and neuronal- and glial-specific rescue mice, thereby allowing us to assess interdependency and cooperativity in the role of axonal and glial glycolipids in paranodal organisation. The expression of a- and b-series gangliosides neuronally reversed the age-dependent breakdown of the axo-glial junction in both the PNS and CNS, demonstrated by invasion of Kv channels from the juxtaparanode to the paranode in GalNAc-T−/− mice (Yao et al., 2014). Glial ganglioside expression exacerbated rather than improved the phenotype. It is interesting to note that aged GalNAc-T−/−-Tg(neuronal) NoR did not completely return to a wild-type arrangement; the Nav channel clusters remained lengthened, and the Caspr domains were longer compared to all other genotypes. We further demonstrated that neuronal expression of a-series complex gangliosides could promote survival in the GalNAc-T x GD3s (i.e. GM3-only) double null mice, but complete restoration to normal would likely require b-series expression (McGonigal et al., 2021). In these mice, both CNS and PNS NoR showed only partial recovery to wild-type composition with the expression of a-series complex gangliosides. Shortening of NF155 and Caspr domains indicating loss of axo-glial integrity was more severe in the CNS and minimally improved. Notably, recovery of nodal protein organisation was dependent on the proteins localisation in either the neuronal or glial membrane. Glial neurofascin 155 was normalised with the expression of neuronal a-series gangliosides, likely owing to the contribution of glial sulphatide, but axonal juxtaparanodal Kv channels, Caspr domains and Nav channel clusters remained disrupted, suggesting the need for a contribution from b-series gangliosides. This suggests the requirement of specialised lipid raft associated anchoring domains that involve gangliosides. We feel this builds upon the work by Schafer et al. (2004) who proposed the formation of a paranodal lipid raft protein adhesion complex and strengthens the concept that Caspr requires two stabilising mechanisms: protein–protein and protein–lipid interactions that promote stability of the paranodal axo-glial junction (McGonigal et al., 2021).

Rescuing the neuronal, but not the glial, complex ganglioside expression reversed the lethality observed in the CST−/− x GalNAc-T−/− double null mice (McGonigal et al., 2019). Examining the relative importance of a- and b-series gangliosides, we observed that only modest improvement occurred with global a-series ganglioside expression on a sulphatide-null background. Similarly, normal CNS NoR composition in the absence of sulphatide and
complex gangliosides could be improved by a- and b-series gangliosides, and not with global a-series ganglioside expression. Collectively, these data indicate the importance of neuronal b-series gangliosides (e.g., GD1b and GT1b) in maintaining survival in the co-presence of sulphatide deficiency and indicate interdependency between the functions of these two groups of lipids.

These results suggest that the composition of the NoR is largely governed by neuronal a-series gangliosides, but that they are not completely sufficient for a normal phenotype. We thus propose that sulphatide and b-series ganglioside lipid domains on opposing membranes majorly contribute to a coordinated axo-glial adhesion and paranodal organisation, a combined loss of which leads to severe impairment of nerve integrity with a fatal outcome at an early age. Based on all of the studies discussed, we have created a series of schematics that represent the effect on nodal integrity of progressive glycolipid loss from neural membranes (Figure 4).

5 | USE OF TISSUE SPECIFIC TRANSGENIC MICE TO STUDY AUTOIMMUNE INJURY TO THE NODE OF RANVIER

As previously published data described in section 4 shows, generation of the rescue mice provides the previously unfeasible opportunity to characterise the contribution of membrane-specific ganglioside expression on nerve integrity and biological function. Additionally, they provide the ideal system for selectively targeting neural membranes at the NoR and determining the downstream consequences of site-specific autoimmune injury. Indeed, understanding the NoR pathophysiology of the human disease Guillain–Barré syndrome (GBS) has been the motivation for generating these mice and the subject of future publications. This anticipates a significant step forward as there exists a few enigmas within the field of inflammatory neuropathies associated with anti-ganglioside antibodies. Autoantibodies against gangliosides, expressed on both axonal and glial membranes, can be particularly associated with clinical syndromes dominated by specific cellular injury to only one. An example of this is the motor form of the peripheral neuropathy GBS, acute motor axonal neuropathy (AMAN), where motor axons appear to undergo selective injury in comparison with glial membranes (Feasby et al., 1986; Hafer-Macko et al., 1996). Traditionally, GBS cases are categorised as axonal or demyelinating variants depending on the presumed site of injury. However, it is becoming clear that this rigid classification system does not satisfactorily define patients and their symptoms. For example, patients with nodal dysfunction or disruption can have a spectrum of symptoms ranging from reversible conduction block to axon degeneration leading to quite different prognoses. There has been extensive discussions on the relative contribution nodal membrane injury can impart on nerve function in disease, and the concept of nodo-paranodopathy versus distinct axonal or demyelinating variants is gaining acceptance as a way to understand the pathophysiological continuum, especially pertaining to anti-ganglioside antibody driven disease at the NoR (Uncini et al., 2013).

Given the minor nodal abnormalities observed in aged rescue mice, it was essential to characterise the NoR at the age to be used in experimental studies. Herein, we also include data on a further neuronal and glial double rescue mouse. It has recently been reported that circulating anti-ganglioside antibodies can be sequestered by global ganglioside membrane expression in wild-type mice (Cunningham et al., 2016) rendering their use for passive immunisation injury studies limited. We observed this finding in wild-type mice on a C57BL/6 background and do not know if it can be generalised to other strains. Cunningham et al. (2016) showed that this phenomenon does not occur in single neuronal or glial rescue mice, and as such injury models using these mice are possible (McGonigal et al., 2016). It is our expectation that double rescue mice can substitute wild-type mice where we see no sequestration of anti-ganglioside antibody, but gangliosides are expressed on both neural membranes. Below we include the nodal characterisation of this newly generated double rescue line compared to the original transgenic mice.

At 4–6 weeks of age, there appears to be little detectable impact of membrane selective complex ganglioside expression on nodal organisation (Figure 5). There are no significant differences in the length of Nav channel clusters, Caspr domains, pan-neurofascin (pan-NFasc) domains or the gap between Kv1.1 channel dimers among the genotypes analysed in this study. Direct comparison between wild-type and GalNAc-T−/− nerve, showed subtle abnormalities in these parameters, also reported previously by Susuki et al. (2007a) at this age. Similar to GalNAc-T−/− mice, we did observe a slight preponderance of glial mice to exhibit mild Kv1.1 invasion into the juxtaparanodal border of the paranode (indicated in Figure 5), and would not recommend their use at older ages as this could indicate the early stages of the disrupted nodal phenotype.

Having established the normal nodal organisation, we assessed the potential for selective nodal targeting by pathogenic antibody. Using a single anti-GM1 antibody, we show the differential binding patterns at the NoR among wild-type, GalNAc-T−/−, neuronal, and glial ganglioside rescue mice. Figure 3b shows binding at both the nodal gap and paranodal loops in wild-type mice and an absence of any labelling in GalNAc-T−/− mice. Antibody binding is restricted to the nodal axolemma in neuronal rescue mice and to the paranodal loops in glial rescue mice. These results make these mice ideal for studying the downstream consequences of antibody binding to different neural membranes at the NoR. Knowing that the rescue mice do have an abnormal expression of gangliosides and are prone to show age-dependent decline in structure, it will be prudent to take this into consideration when interpreting results. However, the current immunohistological data demonstrate that the NoR are grossly normal in all of the lines studied at this age, thereby confirming their suitability for use in peripheral nerve injury models going forward.

6 | CONCLUSION

Transgenic manipulation of glycosyltransferases has provided a wealth of information on the roles of gangliosides and glycolipids in stabilising specialised membrane domains at the NoR. Whilst knockout studies on many nodal proteins has provided extensive and definitive information on nodal biology, the functional roles of glycolipids are much more complex to investigate and nuanced,
owing to redundancy and inability to knockout specific gangliosides. Nevertheless, it is clearly evident that the NoR is rich in glycolipids that function to maintain its integrity and can play a role in pathology. Our rescue mice provide an invaluable tool to explore this field further. It is clear that the NoR is enriched in glycolipids, and whilst this has been the focus of this review, it does not detract
from the important and specialised roles gangliosides have in other domains of the nervous system under investigation elsewhere. We anticipate that the legacy of our mice will be their use in exploring and understanding the emerging roles of gangliosides in the nervous system within selective membranes at the NoR and beyond.

7 | METHODS

7.1 | Mice

We used 4–6 week old mice, both male and female, from five mouse lines all on a C57BL/6J background: (a) wild-type; (b) GalNAc-T \(^{-/-}\); (c) GalNAc-T \(^{-/-}\)-Tg(neuronal); (d) GalNAc-T \(^{-/-}\)-Tg(glial); (e) GalNAc-T \(^{-/-}\)-Tg(neuronal/glial). GalNAc-T \(^{-/-}\) mice lack all complex gangliosides globally (Takamiya et al., 1996), while GalNAc-T \(^{-/-}\)-Tg(neuronal) and GalNAc-T \(^{-/-}\)-Tg(glial) mice have reconstituted site-specific expression of complex gangliosides on neuronal (McGonigal et al., 2016) or glial (Yao et al., 2014) membranes, respectively. GalNAc-T \(^{-/-}\)-Tg(neuronal/glial) were generated by interbreeding GalNAc-T \(^{-/-}\)-Tg(neuronal) and GalNAc-T \(^{-/-}\)-Tg(glial) mice. Mice were maintained under a 12h light/dark cycle at controlled temperature and humidity with ad libitum access to food and water. Mice were killed by a rising CO\(_2\) inhalation; all experiments complied with relevant guidelines on the care and use of animals outlined in the revised Animals (Scientific Procedures) Act of 1986.

7.2 | Anti-ganglioside antibody immunostaining

Sciatic nerves were rapidly dissected and desheathed in oxygenated (95% O\(_2\) and 5% CO\(_2\)) physiological Ringer’s solution containing the following (in mM): NaCl, 129; KCl, 3; NaH\(_2\)PO\(_4\), 1.2; CaCl\(_2\), 2.4; MgSO\(_4\), 1.3; HEPES, 3; NaHCO\(_3\), 20 and glucose, 10. Nerves were incubated for 1 h at 4\(^\circ\)C in 100µg/ml anti-GM1 IgG3 antibody that has been generated as previously described (Boffey et al., 2005; Townson et al., 2007). Nerves were washed and fixed in 4% PFA for 30 min at 4\(^\circ\)C, then washed in three 10 min changes of PBS, 0.1 M glycine and PBS. Sciatic nerves were gently teased out into single fibres onto APES coated slides. Slides were incubated overnight at 4\(^\circ\)C in blocking solution (0.3% Triton +3% normal goat serum) with rabbit anti-gliomedin antibody (Gldn, Abcam #ab24483, RRID:AB_2111616; 1:100). Nerves were washed with PBS followed by application of Alexa Fluor 488-conjugated goat anti-mouse IgG3 (Thermo Fisher Scientific Cat# A-21151, RRID:AB_2535784; 1:500) and Alexa Fluor 555-conjugated goat anti-rabbit IgG (Thermo Fisher Scientific Cat# A-21429, RRID:AB_2535850; 1:500) in PBS with 3% NGS for 2h at R.T. Slides were washed in PBS and mounted in Citifluor.

7.3 | Nodal Immunostaining

Sciatic nerves (n = 3/genotype) were rapidly dissected into 4% PFA and incubated for 30 min at 4\(^\circ\)C. The nerve was washed in three changes of PBS and moved to 30% sucrose for 1 h at 4\(^\circ\)C. Nerves were embedded in OCT mount medium and 10 µm longitudinal sections collected onto APES coated slides. Slides were pre-treated with 100% EtOH at -20\(^\circ\)C for 10 min, thoroughly washed in PBS, before application of a blocking solution (0.3% Triton +10% normal goat serum) for 1 h at 4\(^\circ\)C. Slides were then incubated overnight at 4\(^\circ\)C with one of the following combinations of primary antibodies: rabbit anti-voltage-gated potassium channel (Kv1.1, Alomone Laboratories #APC-009; RRID:AB_2040144; 1:200) plus mouse anti-Caspr (Antibodies incorporated #75-001; RRID:AB_2083496; 1:300); rabbit anti-pan neurofascin (anti-pan-NFasc; gifted from Professor Brophy, University of Edinburgh, UK; 1:1000) plus mouse anti-pan voltage-gated sodium channel (pNav; Sigma-Aldrich #8809; RRID:AB_477552; 1:100). Following washes in PBS, slides were incubated with secondary antibodies prepared in PBS plus 1% NGS for 2 h at R.T. as follows: Alexa Fluor 555-conjugated goat anti-rabbit IgG (Thermo Fisher Scientific Cat# A-21429, RRID:AB_2535850; 1:500); Alexa Fluor 647-conjugated goat anti-mouse IgG1 antibody (Thermo Fisher Scientific Cat# A-21240, RRID:AB_2535809; 1:500). After PBS washes, slides were mounted in Citifluor.

7.4 | Imaging and analysis

Images were captured at 40x or 63x magnification using a Zeiss Axiolamager Z1 with ApoTome attachment and processed with Zeiss Zen 2 blue edition software. For each staining combination, 5-10 z-stacks (step value 0.4 µm) were captured per mouse. For peripheral nerve node analysis, images were used to quantify the length of the Nav channel clusters, pan-NFasc and Caspr domains, and the distance between Kv1.1 dimers from 35 to 90 nodes of Ranvier per mouse. Results were plotted as the average length ±SEM. The tissue was coded upon collection, thereby the experimenter performing analysis was blinded and unaware of the genotype. Statistical differences among genotypes were determined by one-way ANOVA followed by Tukey’s post-hoc test using GraphPad Prism 6 software (RRID:SCR_002798). Differences were considered significant when p < 0.05.

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AUTHOR CONTRIBUTION
Both RM and HJW contributed equally to concept/design, drafting of the manuscript, critical revision and approval of the article. RM acquired the data and performed the data analysis.

ORCID
Rhona McGonigal https://orcid.org/0000-0001-9571-2526
Hugh J. Willison https://orcid.org/0000-0002-5997-1683

REFERENCES

Bartsch, U., Kirchhoff, F. & Schachner, M. (1989) Immunohistological localization of the adhesion molecules L1, N-CAM, and MAG in the developing and adult optic nerve of mice. The Journal of Comparative Neurology, 284, 451–462.

Bhat, M.A., Rios, J.C., Lu, Y., Garcia-Fresco, G.P., Ching, W., Martin, M.S. et al. (2001) Axon-glia interactions and the domain organization of myelinated axons requires neurexin IV/Caspr/Paranodin. Neuron, 30, 369–383.

Boffey, J., Odaka, M., Nicoll, D., Wagner, E.R., Townson, K., Bowes, T. et al. (2005) Characterization of the immunoglobulin variable region gene usage encoding the murine anti-ganglioside antibody repertoire. Journal of Neuroimmunology, 165, 92–103.

Boukhris, A., Schule, R., Loureiro, José L, Lourenço, C., Mundwiller, E., Gonzalez, M. et al. (2013) Alteration of ganglioside biosynthesis responsible for complex hereditary spastic paraplegia. American Journal of Human Genetics, 93, 118–123.

Boyle, M.E., Berglund, E.O., Murai, K.K., Weber, L., Peles, E. & Ranscht, B. (2001) Contactin orchestrates assembly of the septate-like junctions at the paranode in myelinated peripheral nerve. Neuron, 30, 385–397.

Bullens, R.W.M., O’Hanlon, G.M., Wagner, E., Molenaar, P.C., Furukawa, K., Furukawa, K. et al. (2002) Complex gangliosides at the neuro-muscular junction are membrane receptors for autoantibodies and botulinum neurotoxin but redundant for normal synaptic function. The Journal of Neuroscience, 22, 6876–6884.

Chiavegatto, S., Sun, J., Nelson, R.J. & Schnaar, R.L. (2000) A functional role for complex gangliosides: motor deficits in GM2/GD2 synthase knockout mice. Experimental Neurology, 166, 227–234.

Chiba, A., Kusunoki, S., Obata, H., Machinami, R. & Kanazawa, I. (1993) Serum anti-GQ1b IgG antibody is associated with ophthalmoplegia in Miller Fisher syndrome and Guillain-Barré syndrome: clinical and immunohistochemical studies. Neurology, 43, 1911–1917.

Coetzee, T., Fujita, N., Dupree, J., Shi, R., Blight, A., Suzuki, K. et al. (1996) Myelination in the absence of galactocerebroside and sulfatide: normal structure with abnormal function and regional instability. Cell, 86, 209–219.

Collins, B.E., Yang, L.S., Mukhopadhyay, G., Filbin, M.T., Kiso, M., Hasegawa, A. et al. (1997) Sialic acid specificity of myelin-associated glycoprotein binding. Journal of Biological Chemistry, 272, 1248–1255.

Cunningham, M.E., McGonigal, R., Meehan, G.R., Barrie, J.A., Yao, D., Halstead, S.K. et al. (2016) Anti-ganglioside antibodies are removed from circulation in mice by neuronal endocytosis. Brain, 139, 1657–1665.

De Angelis, M.V., Di Muzio, A., Lupo, S., Gambi, D., Uncini, A. & Lugaresi, A. (2001) Anti-GD1a antibodies from an acute motor axonal neuropathy patient selectively bind to motor nerve fiber nodes of Ranvier. Journal of Neuroimmunology, 121, 79–82.

Dicesare, J.L. & Dain, J.A. (1971) The enzymic synthesis of ganglioside. IV. UDP-N-acetylgalactosamine: (N-acetylneuraminyl)-galactosylglycosyl ceramide N-acetylgalactosaminyltransferase in rat brain. Biochimica Et Biophysica Acta, 231, 385–393.

Dupree, J.L., Coetzee, T., Blight, A., Suzuki, K. & Popko, B. (1998) Myelin galactolipids are essential for proper node of Ranvier formation in the CNS. Journal of Neuroscience, 18, 1642–1649.

Eckhardt, M. (2008) The role and metabolism of sulfatide in the nervous system. Molecular Neurobiology, 37, 93–103.

Feasby, T.E., Gilbert, J.J., Brown, W.F. et al. (1986) An acute axonal form of Guillain-Barre polyneuropathy. Brain, 109(PT 6), 1115–1126.

Furukawa, K., Aixinjueluo, W., Kasama, T., Ohkawa, Y., Yoshihara, M., Ohmi, Y. et al. (2008) Disruption of GM2/GD2 synthase gene resulted in overt expression of 9-O-acetyl GD3 irrespective of Tis21. Journal of Neurochemistry, 105, 1057–1066.

Ganser, A.L., Kirschnner, D.A. & Willinger, M. (1983) Ganglioside localization on myelinated nerve fibres by choleratoxin binding. Journal of Neurology, 12, 921–938.

Gong, Y., Tagaya, Y., Lunn, M.P., Laroy, W., Heffer-Lauc, M., Li, C.Y. et al. (2002) Localization of major gangliosides in the PNS: implications for immune neuropathies. Brain, 125, 2491–2506.

Griffin, J.W., Li, C.Y., Macko, C., Ho, T.W., Hsieh, S.-T., Xue, P. et al. (1996) Early nodal changes in the acute motor axonal neuropathy pattern of the Guillain-Barré syndrome. Journal of Neurology, 25, 33–51.

Hafer-Macko, C., Hsieh, S.-T., Ho, T.W., Sheik, K., Cornblath, D.R., Li, C.Y. et al. (1996) Acute motor axonal neuropathy: an antibody-mediated attack on axolemma. Annals of Neurology, 40, 635–644.

Heffer-Lauc, M., Lauc, G., Nimrichter, L., Fromholt, S.E. & Schnaar, R.L. (2005) Membrane redistribution of gangliosides and glycospiphosphatidylinositol-anchored proteins in brain tissue sections under conditions of lipid raft isolation. Biochimica Et Biophysica Acta, 1686, 200–208.

Honke, K., Hirahara, Y., Dupree, J., Suzuki, K., Popko, B., Fukushima, K. et al. (2002) Paranodal junction formation and spermatogenesis require sulfoglycolipids. Proceedings of the National Academy of Sciences of the United States of America, 99, 4227–4232.

Hoshi, T., Suzuki, A., Hayashi, S., Tohyama, K., Hayashi, A., Yamaguchi, Y. et al. (2007) Nodal protrusions, increased Schmidt-Lanterman incisures, and paranodal disorganization are characteristic features of sulfatide-deficient peripheral nerves. Glia, 55, 584–594.

Illa, I., Ortiz, N., Gallard, E., Juarez, C., Grau, J.M. & Dalakas, M.C. (1995) Acute axonal Guillain-Barré syndrome with IgG antibodies against motor axons following parenteral gangliosides. Annals of Neurology, 38, 218–224.

Inoue, M., Fujii, Y., Furukawa, K., Okada, M., Okumura, K., Hayakawa, T. et al. (2002) Refractory skin injury in complex knock-out mice expressing only the GM3 ganglioside. Journal of Biological Chemistry, 277, 29881–29888.

Ishibashi, T., Dupree, J.L., Ikenaka, K., Hirahara, Y., Honke, K., Peles, E. et al. (2002) A myelin galactolipid, sulfatide, is essential for maintenance of ion channels on myelinated axon but not essential for initial cluster formation. Journal of Neuroscience, 22, 6507–6514.

Ishizuka, I. (1997) Chemistry and functional distribution of sulfoglycolipids. Progress in Lipid Research, 36, 245–319.

Kawai, H., Allende, M.L., Wada, R., Kono, M., Sango, K., Deng, C. et al. (2001) Mice expressing only monosialoganglioside GM3 exhibit lethal audiogenic seizures. Journal of Biological Chemistry, 276, 6885–6888.

Kusunoki, S., Kaida, K. & Ueda, M. (2008) Antibodies against gangliosides and ganglioside complexes in Guillain-Barré syndrome: new aspects of research. Biochimica Et Biophysica Acta, 1780, 441–444.

Ledeen, R. & Wu, G. (2011) New findings on nuclear gangliosides: overview on metabolism and function. Journal of Neurochemistry, 116, 714–720.

Lugaresi, A., Ragni, M., Torrieri, F., Di Guglielmo, G., Fermani, P. & Uncini, A. (1997) Acute motor axonal neuropathy with high titer IgG and...
IgA anti-GD1a antibodies following Campylobacter enteritis. *Journal of the Neurological Sciences*, 147, 193–200.

Marcus, J., Dupree, J.L. & Popko, B. (2002) Myelin-associated glycoprotein and myelin galactolipids stabilize developing axo-glia interactions. *Journal of Cell Biology*, 156, 567–577.

McGonigal, R., Barrie, J.A., Yao, D., Black, L.E., McLaughlin, M. & Willison, H.J. (2021) Neuronally expressed α-series gangliosides are sufficient to prevent the lethal age-dependent phenotype in GM3-only expressing mice. *Journal of Neurochemistry*.

McGonigal, R., Barrie, J.A., Yao, D., McLaughlin, M., Cunningham, M.E., Rowan, E.G., Greenshields, K.N., Halstead, S.K., Ngamukote, S., Yanagisawa, M., Ariga, T., Ando, S. & Yu, R.K. (2007) *Neuronally expressed α-series gangliosides are sufficient to prevent the lethal age-dependent phenotype in GM3-only expressing mice. Journal of Cell Biology*, 133, 1944–1960.

Humphreys, P.D., Rother, R.P. et al. (2010) Anti-GD1a antibodies activate complement and calpain to injure distal motor nodes of Ranvier in mice. *Brain*, 133, 1944–1960.

Meehan, G.R., McGonigal, R., Cunningham, M.E., Wang, Y., Barrie, J.A., Okada, M., Itoh, M.-I., Haraguchi, M., Okajima, T., Inoue, M., Oishi, H. Olshefski, R. & Ladisch, S. (1996) Intercellular transfer of shed tumor cell gangliosides. *Acta Neuropathol Commun*, 4, 23.

Palavicini, J.P., Wang, C., Chen, L., Ahmar, S., Higuera, J.D., Dupree, J.L. et al. (2016) Novel molecular insights into the critical role of sulfatide antibodies to glial membranes. *Journal of Neuroimmunology*, 323, 28–35.

Ngamukote, S., Yanagisawa, M., Ariga, T., Ando, S. & Yu, R.K. (2007) Developmental changes of glycosphingolipids and expression of glycogens in mouse brains. *Journal of Neurochemistry*, 103, 2327–2341.

Ogawa-Goto, K. & Abe, T. (1998) Gangliosides and glycosphingolipids of peripheral nervous system myelins–a minireview. *Neurochemical Research*, 23, 305–310.

O’Hanlon, G.M., Paterson, G.J., Veitch, J., Wilson, G. & Willison, H.J. (1998) Mapping immunoreactive epitopes in the human peripheral nervous system using human monoclonal anti-GM1 ganglioside antibodies. *Acta Neuropathologica*, 95, 605–616.

Ohmi, Y., Tajima, O., Ohkawa, Y., Yamauchi, Y., Sugiura, Y., Furukawa, K. et al. (2011) Gangliosides are essential in the protection of inflammation and neurodegeneration via maintenance of lipid rafts: elucidation by a series of ganglioside-deficient mutant mice. *Journal of Neurochemistry*, 116, 926–935.

Okada, M., Itoh, M.-I., Haraguchi, M., Okajima, T., Inoue, M., Oishi, H. et al. (2002) b-series Ganglioside deficiency exhibits no definite changes in the neurogenesis and the sensitivity to Fas-mediated apoptosis but impairs regeneration of the lesioned hypoglossal nerve. *Journal of Biological Chemistry*, 277, 1633–1636.

Olsheski, R. & Ladisch, S. (1996) Intercellular transfer of shed tumor cell gangliosides. *FEBS Letters*, 386, 11–14.

Palavici, J.P., Wang, C., Chen, L., Ahmar, S., Higuera, J.D., Dupree, J.L. et al. (2016) Novel molecular insights into the critical role of sulfatide in myelin maintenance/function. *Journal of Neurochemistry*, 139, 40–54.

Pan, B., Fromholt, S.E., Hess, E.J., Crawford, T.O., Griffin, J.W., Sheikh, K.A. et al. (2005) Myelin-associated glycoprotein and complementary axonal ligands, gangliosides, mediate axon stability in the CNS and PNS: neuropathology and behavioral deficits in single- and double-null mice. *Experimental Neurology*, 195, 208–217.

Paparounas, K., O’Hanlon, G.M., O’Leary, C.P., Rowan, E.G. & Willison, H.J. (1999) Anti-ganglioside antibodies can bind peripheral nerve nodes of Ranvier and activate the complement cascade without inducing acute conduction block in vitro. *Brain*, 122(Pt 5), 807–816.

Pillai, A.M., Thaxton, C., Pribisko, A.L., Cheng, J.G., Dupree, J.L. & Bhat, M.A. (2009) Spatiotemporal ablation of myelinating glia-specific neurofascin (Nfasc NF155) in mice reveals gradual loss of parapomal axoglial junctions and concomitant disorganization of axonal domains. *Journal of Neuroscience Research*, 87, 1773–1793.
analytical studies on neuropathy-associated anti-GM1 antibodies. *Glycobiology*, 17, 294–303.

Trapp, B.D., Andrews, S.B., Wong, A., O’Connell, M. & Griffin, J.W. (1989) Co-localization of the myelin-associated glycoprotein and the microfilament components, F-actin and spectrin, in Schwann cells of myelinated nerve fibres. *Journal of Neurocytology*, 18, 47–60.

Uncini, A., Susuki, K. & Yuki, N. (2013) Nodo-paranodopathy: beyond the demyelinating and axonal classification in anti-ganglioside antibody-mediated neuropathies. *Clinical Neurophysiology*, 124, 1928–1934.

Vinson, M., Strijbos, P.J.L.M., Rowles, A., Facci, L., Moore, S.E., Simmons, D.L. et al. (2001) Myelin-associated glycoprotein interacts with ganglioside GT1b. A mechanism for neurite outgrowth inhibition. *Journal of Biological Chemistry*, 276, 20280–20285.

Vyas, K.A., Patel, H.V., Vyas, A.A. & Schnaar, R.L. (2001) Segregation of gangliosides GM1 and GD3 on cell membranes, isolated membrane rafts, and defined supported lipid monolayers. *Biological Chemistry*, 382, 241–250.

Willison, H.J. & Yuki, N. (2002) Peripheral neuropathies and anti-glycolipid antibodies. *Brain*, 125, 2591–2625.

Yamashita, T., Wu, Y.-P., Sandhoff, R., Werth, N., Mizukami, H., Ellis, J.M. et al. (2005) Interruption of ganglioside synthesis produces central nervous system degeneration and altered axon-glial interactions. *Proceedings of the National Academy of Sciences USA*, 102, 2725–2730.

Yao, D., McGonigal, R., Barrie, J.A., Cappell, J., Cunningham, M.E., Meehan, G.R. et al. (2014) Neuronal expression of GalNAc transferase is sufficient to prevent the age-related neurodegenerative phenotype of complex ganglioside-deficient mice. *Journal of Neuroscience*, 34, 880–891.

Yu, R.K., Tsai, Y.T., Ariga, T. & Yanagisawa, M. (2011) Structures, biosynthesis, and functions of gangliosides—an overview. *Journal of Oleo Science*, 60, 537–544.

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