Review

The Promise and Challenges of Developing miRNA-Based Therapeutics for Parkinson’s Disease

Simoneide S. Titze-de-Almeida, Cristina Soto-Sánchez, Eduardo Fernandez, James B. Koprich, Jonathan M. Brotchie and Ricardo Titze-de-Almeida

1. The Underlying Mechanisms of Parkinson’s Disease (PD)

Parkinson’s disease (PD) is a neurodegenerative disorder affecting over 6 million people globally and with rising incidence as the aged population continues to grow [1]. PD is typically classified as a movement disorder with a triad of bradykinesia, rigidity and tremor [2]. The major pathology contributing to the generation of these motor symptoms is degeneration of the dopaminergic nigrostriatal pathway [3]. Indeed, dopamine replacement therapy has provided the basis of symptomatic therapy for more than half a century. However, the benefits of symptomatic therapies are limited by the emergence of side effects and the loss of efficacy over time and the fact that they do not treat all symptoms of PD equally well [4]. These limitations emerge in part because of plastic changes in brain circuits in response to long term treatment, in part because of the insidious progression...
of the underlying disease and partly because not all symptoms of PD are driven by nigrostriatal dysfunction [5]. With respect to the latter, in addition to the above-mentioned motor symptoms, PD is also associated with a wide range of non-motor symptoms, and most of these remain significant unmet medical needs [6]. Such non-motor symptoms include cognitive and behavioral problems (executive dysfunction [7], psychosis [8], apathy [9], mood disorders [10], autonomic failure (e.g., urinary, bladder disturbances and constipation) [11], loss of sense of smell [12], sleep disturbance [13], and pain [14] and involve neural circuits in addition to the nigrostriatal pathway.

Irrespective of etiology, genetic or sporadic, most PD cases show alpha-synuclein (α-Syn)-related pathology marked by the presence of Lewy bodies and Lewy neurites [15]. While α-Syn accumulation is found within the nigrostriatal pathway and accompanies the degenerative process, this synucleinopathy is actually widespread throughout the brain and peripheral nervous system including in regions that likely contribute to non-motor symptoms, e.g., early in the disease in the enteric plexus, dorsal nucleus of the vagal nerve, locus coeruleus, dorsal raphe nucleus and, late in the disease, advancing into the cerebral cortex [5,16,17].

Thus, current symptomatic therapies do not provide effective long-term control of symptoms or underlying disease and do little to deal with problems emerging outside of the nigrostriatal pathway. There is a need for disease-modifying therapies for PD and, ideally, those that would slow neurodegeneration and functional pathology not only in the nigrostriatal pathway but also in other regions of the nervous system contributing to non-motor symptoms. This need is, so far, unmet. The finding that α-Syn pathology is a hallmark of PD, whatever the etiology, suggests that, despite likely divergence of upstream initiating events, there is a common downstream convergence leading to the involvement of α-Syn as a key player in disease pathogenesis. Many mechanisms of pathology have been proposed, and multiple targets identified to potentially manipulate these. These targets have, to varying degrees, been validated in cell-based systems, animal models and PD post-mortem material. Such targets include, of course, α-Syn, but extend to a panoply of mechanisms that might impact upon α-Syn, either in all or subsets of the PD population, these include oxidative stress [18], LRRK2 [19], apoptosis [20], neuroinflammation [21], misfolding proteins [22], age [2], melanin [23] and iron accumulation [24]. The most attractive of these targets also have support from genetic studies, and include LRRK2 [25], parkin, PINK1 [26] and GBA [27]. MicroRNAs are also emerging targets that have gained increased attention in neurodegenerative disorders, including PD [28]. These small RNAs play relevant roles in regulating cell transcriptome by RNA interference, but they can also contribute to disease pathogenesis if aberrantly expressed [29,30]. In such sense, a dysregulated expression of a specific microRNA can be set up to the physiological level by synthetic oligos for disrupting critical mechanisms of disease pathogenesis [31,32].

This review examines the role of microRNAs (miRNAs) in underlying mechanisms of PD and updates on advances in technologies that improve efficacy and delivery of small RNAs, paving the way toward a miRNA-based neuroprotective therapy.

2. MicroRNA Biogenesis

MicroRNAs are small RNA molecules that commonly originate from a ‘canonical’ pathway in which the biosynthesis begin in the cell nucleus and extend to the cytoplasm (Figure 1) [33]. Two other sources of miRNAs exist; these molecules can be formed in cells from a ‘noncanonical’ pathway, and, in addition, they can be chemically synthesized for experimental assays as oligonucleotides that imitate the endogenous molecules, the miRNA mimics [33,34]. The Box 1 of Supplementary Material presents brief definitions of miRNA mimics and AntimiRs.
The resulting miRNA precursor (pre-miRNA) moves to cytoplasm carried by Exportin 5. Dicer and TRBP enzymes execute the second round of processing, resulting in miRNA duplexes with 18–25 nucleotides. Sense strand is removed. The guide (or antisense) strand is the mature miRNA that will guide the RISC complex (miRISC) to target mRNAs bearing partially complementary sequences in 3'-UTR region. Silencing of miRNA-targeted mRNAs occurs through translational repression or degradation.AGO—Argonaute 2; CDS—Coding sequence region of mRNA; 3'-UTR-3' untranslated region; DGCR8—DiGeorge syndrome critical region gene 8; Dicer—a ribonuclease enzyme; Drosha—a ribonuclease enzyme; miRISC—RISC complex associated with a miRNA; Pol II—RNA polymerase II; pre-miRNA—miRNA precursor; pri-miRNA—primary miRNA; RISC—RNA-induced silencing complex; TRBP—HIV-1 Trans-activation response (TAR) RNA-binding protein. Reprinted with permission from Titze-de-Almeida and Titze-de-Almeida 2018, with modifications [34].

In the ‘canonical’ pathway, miRNA synthesis begins with a transcriptional process from the mammalian genome by RNA polymerase II (Pol II). A steam loop structure denominated primary miRNA (pri-miRNA) is formed in the cell nucleus. Two rounds of processing will subsequently occur. Drosha and DGCR8 enzymes cleave sequences outside the hairpin of the pri-miRNA, to form a precursor miRNA (pre-miRNA). Pre-miRNA are transported to the cytoplasm by exportin 5, and a
second round of processing takes place. Dicer enzyme removes the loop of pre-miRNA to generate a mature miRNA duplex, with ≈ 22 nucleotides [33,35]. The miRNA duplex is incorporated into the RNA-induced silencing complex (miRISC) and the passenger strand is removed. The remaining guide strand loaded in the miRISC will identify and bind to complementary sequences in the 3’-untranslated (3’-UTR) region of target mRNAs by Watson-Crick base-pairing, thus allowing the attached miRISC to exert a target-specific RNA interference. Base pairing between miRNA and mRNA contains mismatches that prevent AGO2 from cleaving the target mRNA. The RNAi process thus occurs in consequence of translational repression at ribosomes and further mRNA degradation [33–35].

While microRNAs play a critical role in regulating cell mRNA levels in a healthy brain, they can assume a pathological function if aberrantly expressed, thereby contributing to the underlying mechanisms of PD (Table 1) [28]. In terms of disease management, dysregulated miRNAs are candidate molecules for diagnosis and prognosis of disease progression, and promising targets for drug development as well, as shown in Figure 2.

Table 1. Selected microRNAs and related targets involved in PD pathogenesis.

| MicroRNA Target | Indirect Target | Reference |
|-----------------|-----------------|-----------|
| α-synuclein aggregation | | |
| miR-7, miR-153, miR34b/c, miR-214 | α-SYN | Jemm et al., 2009 [36]; Doxakis 2010 [37]; Choi et al., 2014 [38]; Fragkouli & Doxakis 2014 [39]; Kabaria et al., 2015 [40]; Wang et al., 2015 [41] |
| Let-7, miR-16-1 | HSP70 | Kumar et al., 2017 [42]; Zhang & Cheng 2014 [43] |
| Clearance of α-Syn | | |
| hsa-miR-21, hsa-miR-224; hsa-miR-373; and hsa-miR-379 | LAMP2a | Alvarez-Erviti et al., 2013 [44]; Alvarez-Erviti et al., 2013 [44] |
| hsa-miR-26b; hsa-miR-106a; and hsa-miR-301b | HSP70 | Deecossac et al., 2013 [45]; Chen et al., 2018 [46] |
| miR-128 | TFEB | |
| miR-13b-5p | SIAH1 | |
| Neuroinflammation | | |
| miR-17 | NLRP3 | Zhou et al., 2016 [47]; Ponomarev et al., 2013 [48] |
| miR-155 | FADD, SOC1, I KK, IL13Rα | |
| miR-27 | ATP5G3 | Prajapati et al., 2015 [49]; He et al., 2017 [50] |
| miR-716-5p | TNF-α | |
| Mitochondrial function and Oxidative stress | | |
| miR-7 | VDAC1; KEAP1 | Chaudhuri et al., 2016 [51]; Kabaria et al., 2015 [40] |
| Apoptosis | | |
| miR-7 | BAX; SIR2 | Li et al., 2016 [52]; Chaudhuri et al., 2015 [51] |
| Glycolytic pathway and gene expression | | |
| miR-7 | RELA; GLUT3; NFkB | Choi et al., 2014 [38]; Chaudhuri et al., 2015 [51] |
| Other pathways | | |
| miR-124 | calpain/ck5 pathway proteins | Kanagaraj et al., 2014 [53] |
| miR-126 | IGF-1/PI3K/AKT signaling | Kim et al., 2014 [54] |
| PRKN and PARK7-Oxidative stress in autosomal recessive parkinsonism | | |
| miR34b/c | PRKN and PARK7 | Mitones-Moyano et al., 2011 [55]; Xiong et al., 2014 [57]; Chen et al., 2017 [56] |
| miR-494; miR-4639-5p | | |
| LRRK2 related | | |
| miR-205; miR-138-2-3p | PARK8 | Cho et al., 2013 [57]; Cardo et al, 2014 [58] |
| Patogenic LRRK2 inhibit let-7 and miR-184; Let-7 and miR-184 regulate E2F1 and DP, respectively, protecting the cells. | | |
| miR-191-3p; miR-201-3p | E2F1; DP | Gehrke et al., 2010 [59] |
3. miRNAs and Alpha Synuclein (α-Syn) Accumulation

α-Syn is expressed abundantly in the brain of healthy individuals and has gained increased attention as a component of the Lewy bodies in PD and other synucleinopathies [60–63]. This small protein of 140-amino acids plays a critical role in clustering synaptic vesicles at the presynaptic terminals [64–66]. α-Syn also regulates protein ubiquitination, chaperone activity, kinase-dependent pathways, and, lastly, the metabolism of dopamine [67–70]. Indeed, genome-wide association studies (GWAS) identified that α-Syn expression, glycosphingolipid biosynthesis, and the protein ubiquitination are pathways affected by single nucleotide polymorphisms (SNP) and a set of microRNAs differentially expressed in PD [71–73].

In healthy individuals, α-Syn is a soluble protein arranged in a stably-folded tetramer [74]. In PD, α-Syn forms deposits of insoluble protofibrils and fibrils with an antiparallel β-sheet structure of monomers, dimers, and misfolded oligomers [75–77]. α-Syn aggregates are located inside the Lewy bodies, present widely in critical brain regions and in peripheral body tissues [5,78,79]. The mechanism of widespread distribution of Lewy bodies might be a progressive cell-to-cell propagation of α-Syn aggregates, consistent with the hypothesis that PD is a prionic disease [78,80,81], whereby α-Syn templates allow protein propagation that drives a complex, multifactorial, and yet poorly understood neuropathology [82].

Mounting evidence supports that miRNAs are critical players for α-Syn accumulation in apoptotic neurons [83,84]. First, functional studies reveal specific miRNAs that are downregulated in PD, which may contribute to α-Syn accumulation and the consequent loss of dopaminergic neurons [28,29]. Second, the 3′-UTR region of α-Syn mRNA contains functional miRNA binding sites that are highly conserved among different species [85]. At least four distinct microRNAs directly downregulate α-Syn, miR-7, miR-153, miR34b/c, and miR-214, meaning that a change in their levels may lead to α-Syn accumulation in the cells [36,37,39,40].

MiR-7 well represents how an alteration in microRNA content directly affects a targeted mRNA. Patients with PD present reduced levels of miR-7 in brain regions related to disease neuropathology, especially the substantia nigra, and depletion of this miRNA is functionally related with α-Syn accumulation and the further neuron loss [36,86]. In addition to miR-7, miR-153 also recognizes sequences in 3′-UTR region of α-Syn, thereby operating additively to downregulate the gene. MiR-7 causes a stronger down-regulation over α-Syn compared to miR-153 [37]. Increased expression of
miR-7 and miR-153 exert neuroprotective actions over dopaminergic cell types, as evidenced in the 1-methyl-4-phenylpyridinium (MPP+) in vitro model of PD [34,37,39].

Aberrant expression of miR-34c-5p also affects α-Syn deposition in PD brain tissues. This miRNA is depleted in amygdala, frontal cortex, substantia nigra and cerebellum of PD patients, as reported by Minones-Moyano and colleagues [55]. Depletion of miR-34c-5p begins in the premotor stages of the disease, thus we may speculate that a lack of this miRNA may have contributed to the slowly progressive loss of dopaminergic cells before the motor signs emerge. In addition, miR-34c-5p remains dysregulated later in the disease, thus having a potential role throughout disease progression [55]. A further report showed that miR-34b/c downregulates α-Syn by targeting to specific 3'-UTR sequence [40]. Conversely, the depletion of miR-34b/c leads to α-Syn accumulation with a subsequent reduction in viability of SH-SY5Y cells that also show oxidative stress, mitochondrial dysfunction, and reduction in DJ1 and Parkin expression [55].

Finally, three other microRNAs are involved in α-Syn regulation, exerting a direct regulation by binding to its 3'-UTR region (miR-214), or an indirect effect, as occurs for Let-7 and miR-16-1 [41,43,87].

Let-7 regulates α-Syn via the autophagy-lysosome network, a pathway that executes α-Syn clearance under physiological conditions [87–90]. In a Caenorhabditis elegans model of PD, depletion of let-7 led to α-Syn accumulation. The change was related to increased levels of lgg-1 and atg-13 and consequent degradation of cellular components and autophagy-related genes, respectively [87,91–93]. miR-16-1 has several binding sites in the 3'-UTR region of HSP70. Growing evidence supports the concept that dysregulated chaperones, especially Hsp70, are involved in PD pathogenesis [94,95]. In such a sense, Hsp70 plays a role in aggregation and cytotoxicity of α-Syn in PD, as confirmed by functional studies and miR-16-1 transfections in SH-SY5Y cells [43].

4. microRNAs and Neuroinflammation

MicroRNAs are important modulators of neuroinflammation, a process found in brain regions involved in PD pathogenesis [96,97]. First, the process of neuroinflammation develops in parallel to and contributes to the death of neuron cells [98,99]. Affected areas present a higher content of activated microglia and astrocytic cells, both findings reported in parkinsonian animals and PD patients [21,100–103]. In agreement with this role, pharmacological inhibitors of microglia activation prevent the loss of nigral neurons in animal models of PD [104–107].

Neuroinflammation is affected by, and affects, α-Syn accumulation in a bidirectional feedback loop. First, α-Syn aggregates activate microglial cells [108]. Activated microglia, in turn, lead to abnormal handling of α-Syn in neurons that further induces a pro-inflammatory process, with increased levels of cytokine, nitric oxide, and reactive oxygen species [109]. Accumulation of α-Syn in cerebral neurons correlates with the presence of HLA-DR (human homolog of MHCII) expressed by microglia and, in addition, with deposits of immunoglobulin G (IgG) in neuronal cells [110,111]. Moreover, the levels of pro-inflammatory cytokines IL1-β (interleukin-1 β), interleukin-2 (IL-2), interleukin-6 (IL-6), interferon-γ (IFN-γ) and tumor necrosis factor alpha (TNF-α) are increased in the nigrostriatal dopaminergic system and peripheral nerves of PD patients [112–117].

TNF-α, a critical regulator of inflammatory responses, was found elevated in the blood, CSF, and striatum of PD patients [112,114,115]. This cytokine can damage SH-SY5Y cells and, in addition, increase their vulnerability to 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP), 6-OHDA and rotenone [49]. TNF-α also regulates miRNAs that target proteins of the mitochondrial complex-I and V. Inhibition of miR-155—a microRNA that targets ATP5G3 (a subunit of F1-ATP synthase)—can attenuate the death of SH-SY5Y cells induced by TNF-α [49].

miR-7 has also been implicated in neuroinflammatory process related to the loss of nigrostriatal cells [36,47]. The gene NLRP3 (inflammasome nod-like receptor protein 3) expressed in microglial cells is down-regulated by miR-7 [47]. This work showed that injections of miR-7 mimics into mouse striatum can suppress NLRP3 inflammasome activation and reduce the loss of dopaminergic cells in MPTP-injured mice model of PD. Effects of miR-7 against neuroinflammation were corroborated by
other studies. Thus, miR-7 knocked-down RelA, a component of NF-κB—a transcriptional factor that regulates genes involved in inflammation and cellular death. Indeed, silencing of RelA induced by miR-7 attenuates the damage of MPP+ to SH-SY5Y cells [38,118–121].

5. Biotechnology for Moving microRNAs from the Bench to Clinics

5.1. Steps to Develop and Evaluate miRNA-Based Drugs

A biotechnological platform aimed to develop miRNA-based therapies broadly follows the stages applied to classic small molecule drug as shown (see Box 2 of the Supplementary Material); Step 1—Discovery and Development; Step 2—Preclinical Research; Step 3—Clinical Research (Phases 1–3 clinical trials); Step 4—FDA Review; Step 5—Post-Market Safety monitoring (Phase 4) [122].

Currently, ongoing clinical trials are employing miRNA- mimics or inhibitors that act on distinct targets and address diseases with different pathogenesis, revealing the flexibility of this biotechnology. Examples of microRNA-based therapeutics in clinical testing are: hepatitis C (AntimiR-122), type 2 diabetes and non-alcoholic fatty liver diseases (AntimiR-103/107), T-cell lymphoma and leukemias, mycosis fungoides (AntimiR-155), scleroderma (miR-29 mimic), mesothelioma and lung cancer (miR-16 mimic), wound healing and heart failure (miR-92), keloids and fibrous scar tissue formation (miR-92), and Alport syndrome (miR-21) [123–126].

5.2. Increased Duration of Effects and Site-Specific Delivery: Two Critical Issues for RNAi-Based Drugs for Brain Diseases

RNAi-based drugs have shown therapeutic benefits in chronic diseases outside the nervous system, as demonstrated by the first FDA-approved siRNA named patisiran [127]. At doses of 0.3 mg/Kg administered intravenously every three weeks, patisiran caused a gene knockdown stable for at least two years [128]. However, as previously discussed, PD patients develop a slowly progressive neuropathology that lasts for decades across the premotor and motor symptomatic phases [2,129]. The chronic and changing nature of brain diseases is a critical consideration for small RNA therapeutics, to provide an effective and durable concentration in brain areas may require dosing adjustments over time, and duration of treatment many times longer than those for which the existing technologies have been employed to date.

Chemical modification in RNAi oligonucleotides is a common strategy to produce more specific and sustainable effects [130–133]. Locked nucleic acids in ribose and phosphorothioate linkages between nucleotides have brought enhanced target specificity and higher nuclease resistance, respectively [134,135]. Both changes were incorporated in miravirsen, the first miRNA-based drug to enter clinical trials, and provided long-term and dose-dependent anti-HCV effects [136–139]. In addition to miravirsen, the first and second FDA-approved RNAi drugs, patisiran and givosiran, also contain chemically modified nucleotides. Patisiran incorporated oligonucleotides with a 2′-methoxy group on riboses to improve stability in serum, a change incorporated in other therapeutic siRNAs [130,133,135,140]. Givosiran has a modified ribose with either 2′-deoxy-2′-fluoro or 2′-O-methyl, like other siRNAs developed by Alnylam [141–145]. In addition, the drug presents phosphorothioate linkages at the 5′-end of both siRNA strands to prevent nuclease digestion after subcutaneous injection [146,147].

The delivery strategy of oligonucleotides used for brain diseases has been a matter of great importance in RNAi biotechnology. The blood–brain barrier represents a critical obstacle for systemically-injected drugs reaching areas inside the central nervous system [148–150]. Some strategies have improved brain penetration, such as conjugating liposomes with antibodies for delivering plasmid vectors [151]. Indeed, aptamers and monoclonal antibodies are viable molecules for association with RNAi duplexes for site-specific drug delivery [152], as revised elsewhere [153–159]

Therapeutic small RNAs can be administered locally into brain tissues by specific devices or stereotaxic surgery, thus circumventing the blood–brain barrier. Convection-enhanced delivery (CED)
has been experimentally used to inject siRNAs and microRNAs into cerebral areas where brain tumors grow [160–164]. Previous work showed that CED was viable for delivering nanoparticulated RNA duplexes against brain gliomas [161,165,166], a technology potentially exploitable to neurodegenerative diseases.

Stereotaxic surgery is another method to inject oligonucleotides into the brain, providing two main advantages: access to specific regions involved in disease pathogenesis and the bypass of blood–brain barrier. In such sense, new particles and methods have been developed to internalize oligonucleotides inside neurons [167,168].

Magnetic particles, along with magnetofection technology, has brought improvement in the delivery of oligonucleotides [169,170]. Our group recently tested a magnetic carrier, NeuroMag®, constituted by polystyrene copolymers coated with iron. This was first used in vivo for transfecting plasmids into pyramidal cells of the rat visual cortex [171]. This particle also shows ability to carry functional microRNA inhibitors to the rat striatal neurons when injected via intracerebroventricular route of administration, causing a significant change in miR-134 content in striatum [172]. In the following section, we will present some recently developed nanoparticles that provide remarkable inputs for miRNA-targeted therapeutics applied to neurodegenerative disorders.

5.3. Nanoparticles for microRNA Delivery

First, any miRNA-based therapy is aimed to return aberrantly expressed miRNAs to their physiological level, either by administering synthetic double-stranded RNAs that imitate them, the miRNA mimics, or those that reduce their content, the antisense oligos named anti-miRs [34,173]. To perform their function, these molecules must be first internalized into the cells, which requires some ability to penetrate cell membranes. Transfection is usually difficult due to the high molecular weight and high negative charge of oligonucleotides, leading them to be electrostatically repealed by anionic charges of the cell membrane [174]. Therefore, they commonly cross the lipid bilayer cloned using one of two main strategies, inside a virus genome or complexed with nanoparticles that facilitate their endocytosis [154].

Both viral and non-viral approaches have been widely used in the treatment of cancer [175,176], and many new applications have been reported for neurological diseases, including PD [176,177]. Actually, viral vectors are especially suitable for neurological diseases because they have high transfection efficiency in neurons. Different viral vectors have been used in animal models to deliver miRNAs that regulate genes involved in PD pathogenesis. To mention two examples, (i) a recombinant adeno-associated virus (rAAV) was injected in the mouse brain to overexpress miR-134, in order to provide evidence for a negative role of miR-134 in dendritic arborization of cortical layer V pyramidal neurons in vivo [178]; (ii) injections of lentivirus that overexpresses miR-126 were able to protect brain cells against 6-OHDA injury [54]. Lentivirus combined with siRNAs were effective to treat other cerebral pathology, focal cerebral ischemia. The study showed that microRNA-210 can increase microvascular density and the number of neural progenitors in ischemic brains in mice [179]. To date, several treatments based on viral vectors have entered pre-clinical and clinical trials, several of them to treat PD [180–182]. However, viral vectors still hold some risk factors, including immunogenicity, oncogenic transformation [183], off-target effects, as well as other biotechnological limitations like high costs for production, limiting gene packing capacity and difficulties for large scale production [184].

In order to avoid these limitations, research on non-viral vectors emerged as a safer and more affordable alternative. Nanoparticles have the ability to protect RNAi oligos against degradation by nucleases in the serum, and favor their penetration into target cells. Among the advantages of nanoparticles, are the ease of production, reproducibility, ease of storage, low toxicity and immunogenicity, and, in addition, most are biodegradable and biocompatible [142,185,186]. Numerous studies have demonstrated the efficiency of miRNA release through non-viral vectors, mainly in the cancer field. The most widely used transporting nanoparticles contain lipid compounds, polymers and inorganic molecules composed of gold, silica, calcium phosphate and iron oxides [187].
5.3.1. Lipid-Based Particles

Lipid particles containing cationic lipids have been extensively used in experimental research and biotechnological development. They are formed by one or several lipid bilayers between which the nucleic acid is trapped. Cationic lipids have a head with a positively charged amino group that interacts with the negative charges of nucleic acids for complexing and internalization. In addition, the positive charges help the particle to interact with the negatively charged membranes which aids cell transfection. Cationic liposomes can release both DNA [188,189] and RNA constructs [190,191]. In the case of miRNAs, once the nanoparticles loaded with miRNA are successfully internalized into the target cells, they need to avoid the endosomal pathway [192,193] for releasing viable miRNA into cytoplasm and avoid degradation of genetic material [194]. Many studies have generated lipid particles with improved capacity to deliver miRNAs, as reviewed elsewhere [195].

Finally, lipid exosomes can be employed for RNAi delivery [196]. Exosomes are small vesicles formed by lipid bilayers (40 to 100 nm) that carry biological molecules, including mRNAs and miRNAs, playing a role in cell signaling. These small lipid particles have already been used to release siRNAs as therapeutic molecules in numerous studies [196], which include pre-clinical applications for neurodegenerative diseases such as Alzheimer’s [197] and PD [198].

5.3.2. Polymeric Particles

Polymeric carriers are formed by synthetic or natural compounds. Several types of synthetic polymers have been successfully used for carrying miRNAs. Polymers based on poly(ethyleneamine)s (PEIs) provide efficient encapsulation and have demonstrated an ability to carry miR-145 and miR-33a [199,200]. Although a large number of positive charges and non-biodegradability of polymeric carriers can reduce their viability and use, several studies have shown efficacy in PD. Thus, an in vivo study in mice using PEI/siRNA complexes decreased the expression of α-synuclein mRNA and protein in the striatum by about 50% [201] without causing toxic signs. Likewise, several in vitro studies have demonstrated the efficiency of PEG/PEI complexes as vectors for delivering anti-α-Syn siRNAs to PC12 cells [202].

Poly(lactic-co-glycolic acid) (PLGA) polymers have advantages regarding biocompatibility and biodegradability. They are difficult to load with miRNAs due to their negative charge [203], but their combination with cationic polymers can overcome this limitation. PLGA nanoparticles coated with protamine sulfate are useful to carry miR-124; when injected in the mice brain, the treatment improved the motor signs in 6-OHDA-injured mice model [204]. PLGA particles can also be combined with a rabies virus glycoprotein RVG29 to enhance the transfection of miR-124 into cells of the substantia nigra, after injected by intraventricular route. This miRNA caused neuroprotection, by reducing pro-inflammatory cytokines and cell apoptosis [205].

Polyurethane based particles (PU-based NPs) are carriers with biocompatibility [206] and ability to inhibit brain tumor cells [207].

Natural polymers such as chitosan and hyaluronic acid, are highly biocompatible particles [208,209]. As these particles present a high positive charge that allows efficient RNA loading [210], they present, in contrast, difficulty to release the RNA oligos from the complex. Even so, some groups have managed to use hyaluronic acid-based nanoparticles combined with chitosan, for carrying miRNA for breast cancer therapy [211]. To improve the release, they have been complexed with negative polymers like PLGA [212–214].

5.3.3. Inorganic Particles

Inorganic particles possess several of the desirable characteristics for nucleic acid transportation, such as biocompatibility, ease of manufacture and storage, as well as ease of controlling their synthesis. To date, numerous studies have been carried out with this type of inorganic particles such as magnetic particles for miRNA release with different applications like cancer [215] and PD [172,216,217].
Other inorganic carriers, including silica nanoparticles and mesoporous silica have successfully release anti-miRs [218] or miRNA mimics [219] to treat brain cancers.

6. Neuroprotective miRNA-Based Approaches in Models of PD

Mounting evidence stands that changes in miRNA content by synthetic oligos could modify the underlying mechanisms of neurodegenerative diseases, as accessed by different experimental studies. The seminal work showing the value of microRNAs as a disease-modifying strategy used a model of temporal lobe epilepsy. Prof. Henshall and collaborators showed that antisense oligonucleotides (antimiRs) targeting miR-134 injected in mice brain ventricles could reduce the loss of hippocampal cells and the number of seizures induced by intra-amygdalar injection of kainic acid [220]. AntimiR-134 was also neuroprotective in multiple models of epilepsy, including pilocarpine, pentylenetetrazol, and stimulation of the perforant pathway [221,222]. Beyond evidencing that microRNAs play a role in the pathogenesis of a degenerative disorder, this ‘proof-of-concept’ work revealed that changing the content of specific microRNAs is a strategy to modify underlying mechanisms of neurodegeneration. Studies regarding PD also evidenced that microRNAs play a role in disease mechanisms and are promising targets for neuroprotection.

A significant decrease in miR-7 was found in the substantia nigra of patients with PD, and, therefore, this change was proposed to play a role in α-Syn accumulation [86]. The neuroprotective actions of miR-7 were first studied in cellular models of PD, which evidenced the ability of this miRNA to target and down-regulate α-Syn by binding to its 3’-UTR mRNA region [36,37]. In agreement, the expression of miR-7 in different tissues during neuronal development is inversely related to the α-Syn content, suggesting that this miRNA exerts a tuning role on α-Syn levels [37]. While the loss of miR-7 function contributes to α-Syn accumulation—which can be reversed by a microRNA replacement, this target also provides additional effects by regulating other genes related to PD pathogenesis [34,36,86]. miR-7 protects cells against oxidative stress by downregulating VDAC1 and Keap1. It also reduces the expression of genes related to the pro-inflammatory process (RelA and NRLP3) and cell glycolysis (RelA), among others, as discussed below [36,38,47,51].

The dysfunction of mitochondrial activity plays a critical role in underlying mechanisms of PD [223,224]. The membrane potential of this organelle is maintained by a complex of proteins named mitochondrial permeability transition pore (mPTP) [225]. However, an excessive opening of the mPTP pores leads to cell death [226,227]. A recent study showed that miR-7 regulates a component of mPTP, the voltage-dependent anion channel 1 (VDAC1) [51]. In MPP+ injured SH-SY5Y cells and primary cortical neurons, overexpression of miR-7 decreased the opening of mPTP channels, which reduced mitochondrial fragmentation/depolarization and mitigated oxidative stress and the release of pro-apoptotic proteins and calcium to the cytoplasm [51]. Also, a previous work using A53T α-Syn parkinsonian mice showed that VDAC1 affects the deposition of this mutant α-Syn in the brainstem, striatum, and cortex, and also induces the opening of mPTP pores [227]. By targeting distinct genes, VDAC1 and α-Syn, miR-7 causes a convergent effect against α-Syn accumulation [36,37,51].

MiR-7 also mitigates oxidative stress by silencing Kelch-like ECH-associated protein 1 (Keap1), a cytoplasmic inhibitory protein. In physiological conditions, Keap1 is found complexed with the erythroid 2–related factor 2 (Nrf2) [228], a nuclear factor that triggers the expression of antioxidant genes [40,229]. The movement of Nrf2 to the cell nucleus is inhibited by Keap1 [230]. In such sense, a miR-7 suppression over Keap1 will consequently allow Nrf2 to translocate to the nucleus for upregulating genes that alleviate oxidative stress. Indeed, overexpression of miR-7 increases glutathione levels causing a 50% decrease of hydroperoxides in MPP+-injured SH-SY5Y cells [40]. In summary, this work added valuable information regarding the cell pathways related to the ‘antioxidant’ effects of miR-7.

Regarding RelA gene, the protective role of miR-7 was also investigated in the MPP+ model [38]. First, miR-7 downregulates RelA, which is a component of the nuclear factor-κB (NF-κB). NF-κB is a pleiotropic transcriptional factor for signaling molecules such as chemokines, cytokines, pro-inflammatory enzymes, adhesion molecules, and pro-inflammatory transcription factors,
that ultimately protect neuron cells by orchestrating inflammatory reactions [231]. RelA plays a role in the death of neuron cells exposed to MPP+, and the respective knocking-down by miR-7 caused neuroprotection [38]. Cell glycolysis was also improved by miR-7-induced RelA depletion [51]. Replacement of miR-7 in MPP+-injured SH-SH5Y cells ameliorated glycolysis, with a positive increase in ATP/ADP ratio, glucose consumption, as well as lactic acid production. Indeed, the suppressive action of miR-7 on RelA upregulates the content of surface glucose transporter (Glut3) that also causes a glycolysis-promoting effect. These findings were confirmed by the silencing of the Glut3 gene or through inhibition of the hexokinase enzyme that catalyzes the first step in glycolysis [51].

Neuroinflammation is a component of the neurodegenerative process in PD that is affected by microRNAs [96,232]. The inflammasome protein NLRP3 is a message expressed in microglial cells directly regulated by miR-7. This protein was found activated in the serum of PD patients and also in midbrain regions of parkinsonian mice [47]. Intra-striatal injection of miR-7 mimics in MPTP-induced parkinsonian mice reduces the dopaminergic degeneration and inhibits microglial activation. In addition, the downregulation of NLRP3 inflammasome by miR-7 decreased the content of a pro-inflammatory cytokine, IL-1β. Injection of miR-7 mimics into the striatum also causes neuroprotective results in transgenic A53Ttg/tg parkinsonian mice, corroborating the results found in the MPTP model. Western blotting analysis confirmed that miR-7 down-regulates NLRP3 in mice midbrain, and also inhibits the activation of caspase-1 and decreases IL-1β production [47].

Finally, the microRNA Let-7 also plays a role in α-Syn regulation, affecting an autophagy-lysosome pathway that executes the clearance of α-Syn in healthy cells [87]. The role of let-7 in α-Syn accumulation was studied in a transgenic C. elegans model of PD that expresses human α-Syn. Inhibition of let-7 decreased α-Syn expression without affecting dopaminergic/acetylcholinergic neurons. The authors suggested this decrease in α-Syn was related to autophagy and induction of the daf-16 forkhead box O (FOXO) transcription factor that is involved in PD pathogenesis [233]. Igg-1 and atg were also increased in let-7 silenced cells. Finally, let-7 regulates genes that trigger cell apoptosis. Inhibition of let-7 protected cells from apoptosis by reducing two pro-apoptotic genes, ced-4 and jnk-1. Altogether, these results highlight let-7 as a promising miRNA target for PD therapeutics that act against multiple underlying mechanisms of the disease [87].

7. Summary and Perspectives

Translating a scientific discovery to the clinics represents a long time and challenging process that require huge efforts for planning and executing many assays, resilience, and money. The first FDA-approval of a small-interfering RNA (siRNA) drug was in 2018, twenty years after the seminal work of Fire & Mello has been published. When siRNA technology becomes available, the scientific community feels a disproportional excitement. Since any gene could be silenced, all diseases related to overexpressed genes would be treatable. However, limitations in siRNA technology rapidly appear, and the excitement gives place to disappointment. The siRNA’s effects lasted for a few days, some off-target effects emerged, and ribonuclease enzymes quickly cleaved RNA duplexes. The first lipid nanoparticles became widely available, the conventional liposomes that facilitated cell transfection. While this ‘first-generation’ liposomes offered protection for RNA molecules, they suffered rapid phagocytosis by circulating macrophages before reaching the target organs. So, only topic preparations were promising, and siRNAs for ophthalmic diseases were the leading drugs for an RNAi market yet in its infancy.

A massive investment of time and money, fortunately, has changed the scenario for the best. New nanoparticles were developed to escape phagocytosis in the blood and to deliver siRNAs safety inside the cell cytoplasm, either lipid or polymeric. Target delivery to specific organs is a reality today. The FDA-approved drugs, patisiran, and givosiran that are administered by intravenous or subcutaneous routes, are efficiently vectorized to the liver for transfecting hepatocytes with siRNAs. Furthermore, new chemical modifications in nucleic acid moieties have provided more specific and long-lasting effects.
Nowadays, several products with RNA interference applications are found in phase II-III clinical trials [130]. Although only a few gene therapy drugs are available in the market for human use, the number of products under clinical trials has increased in the last few years. This increment has augmented the interest of biotechnology companies in developing gene therapy-based drugs and, consequently, to obtain safer and more efficient genetic delivery systems, including miRNAs.

As already mentioned, miRNA mimics and AntimiRs have received biotechnological inputs to enhance the stability of RNA molecules and can be used in combination with nanoparticles with high transfection efficiencies [185]. This progress in nanotechnology aligned with improved changes in oligonucleotide chemistry has provided a notorious evolution in RNAi therapeutics [131,234,235]. Targeted delivery, reduced toxicity, and cost, are issues to be addressed in the future for even better use of short RNA nucleotides [134,236–239].

Finally, the treatment of brain diseases with small RNAs holds an additional challenge due to the blood–brain barrier. Stereotaxic surgery seems to be the most viable strategy to overcome this obstacle. Although laborious and requiring hospitalization, we could envisage benefits in this procedure, including a tissue-specific delivery that reduces the cost of treatment and avoids effects in other tissues. Moreover, as microRNAs act over different mRNAs, a more widespread effect is achieved and may counteract different underlying mechanisms of PD. In summary, we concluded that miRNAs are promising drug targets for modifying the progressive nature of PD, and future technological inputs will allow translating these small RNAs to the clinics in the coming future.

Supplementary Materials: The following are available online at http://www.mdpi.com/2073-4409/9/4/841/s1, Box 1: Basic definitions in RNA interference, Box 2. FDA guidelines for drug development process applied to miRNA-based drugs.

Author Contributions: S.S.T.-d.-A., C.S.-S., E.F., J.B.K., J.M.B., and R.T.-d.-A. contributed equally to this manuscript. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Acknowledgments: Simoneide S. Titze-de-Almeida and Ricardo Titze-de-Almeida are members of the Network for Translational Neuroscience-International Consortium for Academic Cooperation in Experimental and Clinical Studies Regarding Neurodegenerative Diseases (http://dgp.cnpq.br/dgp/espelhogrupo/5933421119277338).

Conflicts of Interest: The authors declare no conflict of interest.

References
1. Pringsheim, T.; Jette, N.; Frolkis, A.; Steeves, T.D. The prevalence of Parkinson’s disease: A systematic review and meta-analysis. Mov. Disord. 2014, 29, 1583–1590. [CrossRef]
2. Kalia, L.V.; Lang, A.E. Parkinson’s disease. Lancet 2015, 386, 896–912. [CrossRef]
3. Lees, A.J.; Hardy, J.; Revesz, T. Parkinson’s disease. Lancet 2009, 373, 2055–2066. [CrossRef]
4. Brotchie, J.M.; Lee, J.; Venderova, K. Levodopa-induced dyskinesia in Parkinson’s disease. J. Neural Transm. 2005, 112, 359–391. [CrossRef]
5. Beach, T.G.; Adler, C.H.; Sue, L.I.; Vedders, L.; Lue, L.; White Iii, C.L.; Akiyama, H.; Caviness, J.N.; Shill, H.A.; Sabbagh, M.N.; et al. Multi-organ distribution of phosphorylated alpha-synuclein histopathology in subjects with Lewy body disorders. Acta Neuropathol. 2010, 119, 689–702. [CrossRef]
6. Balestrino, R.; Schapira, A.H.V. Parkinson disease. Eur. J. Neurol. 2020, 27, 27–42. [CrossRef]
7. Smith, C.R.; Cullen, B.; Sheridan, M.P.; Cavanagh, J.; Grosset, K.A.; Grosset, D.G. Cognitive impairment in Parkinson’s disease is multifactorial: A neuropsychological study. Acta Neurol. Scand. 2020, in press. [CrossRef]
8. Ffytche, D.H.; Aarsland, D. Psychosis in Parkinson’s Disease. Int. Rev. Neurobiol. 2017, 133, 585–622. [CrossRef]
9. Pagonabarraga, J.; Kulisevsky, J. Apathy in Parkinson’s Disease. Int. Rev. Neurobiol. 2017, 133, 657–678. [CrossRef]
10. Schrag, A.; Taddei, R.N. Depression and Anxiety in Parkinson’s Disease. Int. Rev. Neurobiol. 2017, 133, 623–655. [CrossRef]
11. Palma, J.A.; Kaufmann, H. Treatment of autonomic dysfunction in Parkinson disease and other synucleinopathies. Mov. Disord. 2018, 33, 372–390. [CrossRef]
37. Doxakis, E. Post-transcriptional regulation of alpha-synuclein expression by miR-7 and miR-153. J. Biol. Chem. 2010, 285, 12726–12734. [CrossRef]

38. Choi, D.C.; Chae, Y.J.; Kabaria, S.; Chaudhuri, A.D.; Jain, M.R.; Li, H.; Mouradian, M.M.; Junn, E. MicroRNA-7 protects against 1-methyl-4-phenylpyridinium-induced cell death by targeting RelA. J. Neurosci. 2014, 34, 12725–12737. [CrossRef]

39. Fragkouli, A.; Doxakis, E. MiR-7 and miR-153 protect neurons against MPP(+) -induced cell death via upregulation of mTOR pathway. Front. Cell Neurosci. 2014, 8, 182. [CrossRef]

40. Kabaria, S.; Choi, D.C.; Chaudhuri, A.D.; Jain, M.R.; Li, H.; Junn, E. MicroRNA-7 activates Nrf2 pathway by targeting Keap1 expression. Free Radic. Biol. Med. 2015, 89, 548–556. [CrossRef]

41. Wang, Z.H.; Zhang, J.L.; Duan, Y.L.; Zhang, Q.S.; Li, G.F.; Zheng, D.L. MicroRNA-214 participates in the neuroprotective effect of Resveratrol via inhibiting alpha-synuclein expression in MPTP-induced Parkinson’s disease mouse. Biomed. Pharmacother. 2015, 74, 252–256. [CrossRef]

42. Kumar, M.; Sahu, S.K.; Kumar, R.; Subuddhi, A.; Maji, R.K.; Jana, K.; Gupta, P.; Raffetseder, J.; Lerm, M.; Ghosh, Z.; et al. MicroRNA let-7 modulates the immune response to Mycobacterium tuberculosis infection via control of A20, an inhibitor of the NF-kappaB pathway. Cell Host Microbe 2015, 17, 345–356. [CrossRef]

43. Zhang, Z.; Cheng, Y. MiR-16-1 promotes the aberrant alpha-synuclein accumulation in parkinson disease via targeting heat shock protein 70. Sci. World J. 2014, 2014, 938348. [CrossRef]

44. Alvarez-Erviti, L.; Seow, Y.; Schapira, A.H.; Rodriguez-Oroz, M.C.; Obeso, J.A.; Cooper, J.M. Influence of microRNA deregulation on chaperone-mediated autophagy and alpha-synuclein pathology in Parkinson’s disease. Cell Death Dis. 2013, 4, e545. [CrossRef]

45. Decressac, M.; Matsson, B.; Weikop, P.; Lundblad, M.; Jakobsson, J.; Bjorklund, A. TFEB-mediated autophagy rescues midbrain dopamine neurons from alpha-synuclein toxicity. Proc. Natl. Acad. Sci. USA 2013, 110, E1817–E1826. [CrossRef]

46. Chen, Y.; Lian, Y.J.; Ma, Y.Q.; Wu, C.J.; Zheng, Y.K.; Xie, N.C. LncRNA SNHG1 promotes alpha-synuclein aggregation and toxicity by targeting miR-15b-5p to activate SIAH1 in human neuroblastoma SH-SY5Y cells. Neurotoxicology 2018, 68, 212–221. [CrossRef]

47. Zhou, Y.; Lu, M.; Du, R.H.; Qiao, C.; Jiang, C.Y.; Zhang, K.Z.; Ding, J.H.; Hu, G. MicroRNA-7 targets Nod-like receptor protein 3 inflammasome to modulate neuroinflammation in the pathogenesis of Parkinson’s disease. Mol. Neurodegener. 2016, 11, 28. [CrossRef]

48. Ponomarev, E.D.; Veremeyko, T.; Weiner, H.L. MicroRNAs are universal regulators of differentiation, activation, and polarization of microglia and macrophages in normal and diseased CNS. Glia 2013, 61, 91–103. [CrossRef]

49. Prajapati, P.; Sripada, L.; Singh, K.; Bhateia, K.; Singh, R.; Singh, R. TFN-alpha regulates miRNA targeting mitochondrial complex-I and induces cell death in dopaminergic cells. Biochim. Biophys. Acta 2015, 1852, 451–461. [CrossRef]

50. He, Q.; Wang, Q.; Yuan, C.; Wang, Y. Downregulation of miR-7116-5p in microglia by MPP(+) sensitizes TNF-alpha production to induce dopaminergic neuron damage. Glia 2017, 65, 1251–1263. [CrossRef]

51. Chaudhuri, A.D.; Kabaria, S.; Choi, D.C.; Mouradian, M.M.; Junn, E. MicroRNA-7 Promotes Glycolysis to protect against 1-Methyl-4-phenylpyridinium-induced Cell Death. J. Biol. Chem. 2015, 290, 12425–12434. [PubMed]

52. Li, S.; Lv, X.; Zhai, K.; Xu, R.; Zhang, Y.; Zhao, S.; Qin, X.; Yin, L.; Lou, J. MicroRNA-7 inhibits neuronal apoptosis in a cellular Parkinson’s disease model by targeting Bax and Sirt2. Am. J. Transl. Res. 2016, 8, 993–1004. [PubMed]

53. Kanagaraj, N.; Beiping, H.; Dheen, S.T.; Tay, S.S. Downregulation of miR-124 in MPTP-treated mouse model of Parkinson’s disease and MPP iodide-treated MN9D cells modulates the expression of the calpain/cdk5 pathway proteins. Neuroscience 2014, 272, 167–179. [CrossRef] [PubMed]

54. Kim, W.; Lee, Y.; McKenna, N.D.; Yi, M.; Simunovic, F.; Wang, Y.; Kong, B.; Rooney, R.J.; See, H.; Stephens, R.M.; et al. miR-126 contributes to Parkinson’s disease by dysregulating the insulin-like growth factor/phosphoinositide 3-kinase signaling. Neurobiol. Aging 2014, 35, 1712–1721. [CrossRef] [PubMed]

55. Minones-Moyano, E.; Porta, S.; Escaramis, G.; Rabionet, R.; Iraola, S.; Kagerbauer, B.; Espinosa-Parrilla, Y.; Ferrer, I; Estivill, X.; Marti, E. MicroRNA profiling of Parkinson’s disease brains identifies early downregulation of miR-34b/c which modulates mitochondrial function. Hum. Mol. Genet. 2011, 20, 3067–3078. [CrossRef]
56. Chen, Y.; Gao, C.; Sun, Q.; Pan, H.; Huang, P.; Ding, J.; Chen, S. MicroRNA-4639 is a Regulator of DJ-1 Expression and a Potential Early Diagnostic Marker for Parkinson’s Disease. *Front Aging Neurosci.* 2017, 9, 232. [CrossRef]

57. Cho, H.J.; Liu, G.; Jin, S.M.; Parisiadou, L.; Xie, C.; Yu, J.; Sun, L.; Ma, B.; Ding, J.; Vancraenenbroeck, R.; et al. MicroRNA-205 regulates the expression of Parkinson’s disease-related leucine-rich repeat kinase 2 protein. *Hum. Mol. Genet.* 2013, 22, 608–620. [CrossRef]

58. Cardo, L.F.; Coto, E.; Ribacoba, R.; Mata, I.F.; Moris, G.; Menendez, M.; Alvarez, V. The screening of the 3’UTR sequence of LRRK2 identified an association between the rs66737902 polymorphism and Parkinson’s disease. *J. Hum. Genet.* 2014, 59, 346–348. [CrossRef]

59. Gehrke, S.; Imai, Y.; Sokol, N.; Lu, B. Pathogenic LRRK2 negatively regulates microRNA-mediated translational repression. *Nature* 2010, 466, 637–641. [CrossRef]

60. Boeve, B.F.; Silber, M.H.; Parisi, J.E.; Dickson, D.W.; Ferman, T.J.; Benarroch, E.E.; Schmeichel, A.M.; Smith, G.E.; Petersen, R.C.; Ahlskog, J.E.; et al. Synucleinopathy pathology and REM sleep behavior disorder plus dementia or parkinsonism. *Neurology* 2003, 61, 40–45. [CrossRef]

61. Rocha, E.M.; De Miranda, B.; Sanders, L.H. Alpha-synuclein: Pathology, mitochondrial dysfunction and neuroinflammation in Parkinson’s disease. *Neuropsychobiology* 2018, 79, 249–257. [CrossRef] [PubMed]

62. McCann, H.; Stevens, C.H.; Cartwright, H.; Halliday, G.M. alpha-Synucleinopathy phenotypes. *Park. Relat. Disord.* 2014, 20 (Suppl. 1), S62–S67. [CrossRef]

63. Goedert, M.; Jakes, R.; Spillantini, M.G. The Synucleinopathies: Twenty Years On. *J. Park. Dis.* 2017, 7, S51–S69. [CrossRef] [PubMed]

64. Alim, M.A.; Hossain, M.S.; Arima, K.; Takeda, K.; Izumiyama, Y.; Nakamura, M.; Kaji, H.; Shinoda, T.; Hisanaga, S.; Ueda, K. Tubulin seeds alpha-synuclein fibril formation. *J. Biol. Chem.* 2002, 277, 2112–2117. [CrossRef]

65. Diao, J.; Burre, J.; Vivona, S.; Cipriano, D.J.; Sharma, M.; Kyoung, M.; Sudhof, T.C.; Brunger, A.T. Native alpha-synuclein induces clustering of synaptic-vesicle mimics via binding to phospholipids and synaptobrevin-2/VAMP2. *Elife* 2013, 2, e00592. [CrossRef]

66. Totterdell, S.; Meredith, G.E. Localization of alpha-synuclein to identified fibers and synapses in the normal mouse brain. *Neuroscience* 2005, 135, 907–913. [CrossRef]

67. Bennett, J.P., Jr.; Piercey, M.F. Pramipexole—a new dopamine agonist for the treatment of Parkinson’s disease. *J. Neurol. Sci.* 1999, 163, 25–31. [CrossRef]

68. Jenco, J.M.; Rawlingson, A.; Daniels, B.; Morris, A.J. Regulation of phospholipase D2: Selective inhibition of mammalian phospholipase D isoenzymes by alpha- and beta-synucleins. *Biochemistry* 1998, 37, 4901–4909. [CrossRef]

69. Lotharius, J.; Brundin, P. Pathogenesis of Parkinson’s disease: Dopamine, vesicles and alpha-synuclein. *Nat. Rev. Neurosci.* 2002, 3, 932–942. [CrossRef]

70. Perez, R.G.; Waymire, J.C.; Lin, E.; Liu, J.J.; Guo, F.; Zigmond, M.J. A role for alpha-synuclein in the regulation of dopamine biosynthesis. *J. Neurosci.* 2002, 22, 3090–3099. [CrossRef]

71. Martin, E.R.; et al. Convergence of miRNA expression profiling, alpha-synuclein interacton and GWAS in Parkinson’s disease. *PLoS ONE* 2011, 6, e25443. [CrossRef] [PubMed]

72. Lees, A.J.; et al. The screening of the 3’UTR sequence of LRRK2 identified an association between the rs66737902 polymorphism and Parkinson’s disease. *J. Hum. Genet.* 2014, 59, 346–348. [CrossRef] [PubMed]

73. Visscher, P.M.; Wray, N.R.; Zhang, Q.; Sklar, P.; McCarthy, M.I.; Brown, M.A.; Yang, J. 10 Years of GWAS Discovery: Biology, Function, and Translation. *Am. J. Hum. Genet.* 2017, 101, 5–22. [CrossRef]

74. Dettmer, U.; Newman, A.J.; Soldner, F.; Luth, E.S.; Kim, N.C.; von Saucken, V.E.; Sanderson, J.B.; Jaenisch, R.; Bartels, T.; Selkoe, D. Parkinson-causing alpha-synuclein missense mutations shift native tetramers to monomers as a mechanism for disease initiation. *Nat. Commun.* 2015, 6, 7314. [CrossRef] [PubMed]

75. Rodriguez, J.A.; Ivanova, M.I.; Sawaya, M.R.; Cascio, D.; Reyes, F.E.; Shi, D.; Sangwan, S.; Guenther, E.L.; Johnson, L.M.; Zhang, M.; et al. Structure of the toxic core of alpha-synuclein from invisible crystals. *Nature* 2015, 525, 486–490. [CrossRef]
77. Roeters, S.J.; Iyer, A.; Pletikapic, G.; Kogan, V.; Subramaniam, V.; Woutersen, S. Evidence for Intramolecular Antiparallel Beta-Sheet Structure in Alpha-Synuclein Fibrils from a Combination of Two-Dimensional Infrared Spectroscopy and Atomic Force Microscopy. Sci. Rep. 2017, 7, 41051. [CrossRef]

78. Braak, H.; Del Tredici, K.; Rub, U.; de Vos, R.A.; Jansen Steur, E.N.; Braak, E. Staging of brain pathology related to sporadic Parkinson’s disease. Neurobiol Aging 2003, 24, 197–211. [CrossRef]

79. Kalia, L.V.; Lang, A.E. Parkinson disease in 2015: Evolving basic, pathological and clinical concepts in PD. Nat. Rev. Neuro. 2016, 12, 65–66. [CrossRef]

80. Goedert, M.; Spillantini, M.G.; Del Tredici, K.; Braak, H. 100 years of Lewy pathology. Nat. Rev. Neuro. 2013, 9, 13–24. [CrossRef]

81. Visanji, N.P.; Brooks, P.L.; Hazrati, L.N.; Lang, A.E. The prion hypothesis in Parkinson’s disease: Braak to the future. Acta Neuropathol. Commun. 2013, 1, 2. [CrossRef] [PubMed]

82. Surmeier, D.J.; Obeso, J.A.; Halliday, G.M. Parkinson’s Disease Is Not Simply a Prion Disorder. J. Neurosci. 2017, 37, 9799–9807. [CrossRef] [PubMed]

83. Jellinger, K.A. Neuropathology of sporadic Parkinson’s disease: Evaluation and changes of concepts. Mov. Disord. 2012, 27, 8–30. [CrossRef]

84. Spillantini, M.G.; Schmidt, M.L.; Lee, V.M.; Trojanowski, J.Q.; Jakes, R.; Goedert, M. Alpha-synuclein in Lewy bodies. Nature 1997, 398, 839–840. [CrossRef] [PubMed]

85. Sotiriou, S.; Gibney, G.; Baxevanis, A.D.; Nussbaum, R.L. A single nucleotide polymorphism in the 3′UTR of the SNCA gene encoding alpha-synuclein is a new potential susceptibility locus for Parkinson disease. Neurosci. Lett. 2009, 461, 196–201. [CrossRef] [PubMed]

86. McMillan, K.J.; Murray, T.K.; Bengoa-Vergniory, N.; Cordero-Llana, O.; Cooper, J.; Buckley, A.; Wade-Martins, R.; Uney, J.B.; O’Neill, M.J.; Wong, L.F.; et al. Loss of MicroRNA-7 Regulation Leads to alpha-Synuclein Accumulation and Dopaminergic Neuronal Loss In Vivo. Mol. Ther. 2017, 25, 2404–2414. [CrossRef] [PubMed]

87. Shamsuzzama; Kumar, L.; Nazir, A. Modulation of Alpha-synuclein Expression and Associated Effects by MicroRNA Let-7 in Transgenic C. elegans. Front. Mol. Neurosci. 2017, 10, 328. [CrossRef]

88. Boland, B.; Yu, W.H.; Corti, O.; Mollereau, B.; Henriques, A.; Bezd, E.; Pastores, G.M.; Rubinsztein, D.C.; Nixon, R.A.; Duchen, M.R.; et al. Promoting the clearance of neurotoxic proteins in neurodegenerative disorders of ageing. Nat. Rev. Drug Discov. 2018, 17, 660–688. [CrossRef]

89. Lee, H.J.; Khoshaghideh, F.; Patel, S.; Lee, S.J. Clearance of alpha-synuclein oligomeric intermediates via the lysosomal degradation pathway. J. Neurosci. 2004, 24, 1888–1896. [CrossRef]

90. Webb, J.L.; Ravikumar, B.; Atkins, J.; Skepper, J.N.; Rubinsztein, D.C. Alpha-Synuclein is degraded by both autophagy and the proteasome. J. Biol. Chem. 2003, 278, 25009–25013. [CrossRef]

91. Jenzer, C.; Manil-Segalen, M.; Lefebvre, C.; Largeau, C.; Glatigny, A.; Legouis, R. Human GABARAP can restore autophagosome biogenesis in a C. elegans lgg-1 mutant. Autophagy 2014, 10, 1868–1872. [CrossRef]

92. Keith, S.A.; Maddux, S.K.; Zhong, Y.; Chinchankar, M.N.; Ferguson, A.A.; Ghazi, A.; Fisher, A.L. Graded Proteasome Dysfunction in Caenorhabditis elegans Activates an Adaptive Response Involving the Conserved SKN-1 and ELT-2 Transcription Factors and the Autophagy-Lysosome Pathway. PLoS Genet. 2016, 12, e1005823. [CrossRef]

93. Manil-Segalen, M.; Lefebvre, C.; Jenzer, C.; Trichet, M.; Boulogne, C.; Satia-Jeunemaitre, B.; Legouis, R. The C. elegans LC3 acts downstream of GABARAP to degrade autophagosomes by interacting with the HOPS subunit VPS39. Dev. Cell 2014, 28, 43–55. [CrossRef] [PubMed]

94. Bandopadhyay, R.; de Belleruche, J. Pathogenesis of Parkinson’s disease: Emerging role of molecular chaperones. Trends Mol. Med. 2010, 16, 27–36. [CrossRef]

95. Witt, S.N. Hsp70 molecular chaperones and Parkinson’s disease. Biopolymers 2010, 93, 218–228. [CrossRef]

96. Slota, J.A.; Booth, S.A. MicroRNAs in Neuroinflammation: Implications in Disease Pathogenesis, Biomarker Discovery and Therapeutic Applications. Noncoding RNA 2019, 5, 35. [CrossRef] [PubMed]

97. Hirsch, E.C.; Hunot, S. Neuroinflammation in Parkinson’s disease: A target for neuroprotection? Lancet Neurol. 2009, 8, 382–397. [CrossRef]

98. Gao, H.M.; Hong, J.S. Why neurodegenerative diseases are progressive: Uncontrolled inflammation drives disease progression. Trends Immunol. 2008, 29, 357–365. [CrossRef] [PubMed]

99. Sherer, T.B.; Kim, J.H.; Betarbet, R.; Greenamyre, J.T. Subcutaneous rotenone exposure causes highly selective dopaminergic degeneration and alpha-synuclein aggregation. Exp. Neurol. 2003, 179, 9–16. [CrossRef]

100. Hirsch, E.C.; Jenner, P.; Przedborski, S. Pathogenesis of Parkinson’s disease. Mov. Disord. 2013, 28, 24–30. [CrossRef]
101. McGeer, P.L.; McGeer, E.G. Glial reactions in Parkinson’s disease. Mov. Disord. 2008, 23, 474–483. [CrossRef] [PubMed]

102. Przedborski, S. Inflammation and Parkinson’s disease pathogenesis. Mov. Disord. 2010, 25 (Suppl. 1), S55–S57. [CrossRef]

103. Teismann, P.; Tieu, K.; Cohen, O.; Choi, D.K.; Wu, D.C.; Marks, D.; Vila, M.; Jackson-Lewis, V.; Przedborski, S. Pathogenic role of glial cells in Parkinson’s disease. Mov. Disord. 2003, 18, 121–129. [CrossRef] [PubMed]

104. He, Y.; Appel, S.; Le, W. Minocycline inhibits microglial activation and protects nigral cells after 6-hydroxydopamine injection into mouse striatum. Brain Res. 2001, 909, 187–193. [CrossRef]

105. Du, Y.; Ma, Z.; Lin, S.; Dodel, R.C.; Gao, F.; Bales, K.R.; Triarhou, L.C.; Chernet, E.; Perry, K.W.; Nelson, D.L.; et al. Minocycline prevents nigrostriatal dopaminergic neurodegeneration in the MPTP model of Parkinson’s disease. Proc. Natl. Acad. Sci. USA 2001, 98, 14669–14674. [CrossRef] [PubMed]

106. He, Y.; Appel, S.; Le, W. Minocycline inhibits microglial activation and protects nigral cells after 6-hydroxydopamine injection into mouse striatum. Brain Res. 2001, 909, 187–193. [CrossRef]

107. Lazzarini, M.; Martin, S.; Mitkovski, M.; Vozari, R.R.; Stuhmer, W.; Bel, E.D. Doxycycline restraints glia and confers neuroprotection in a 6-OHDA Parkinson model. Glia 2013, 61, 1084–1100. [CrossRef]

108. Orr, C.F.; Rowe, D.B.; Mizuno, Y.; Mori, H.; Halliday, G.M. A possible role for humoral immunity in the pathogenesis of Parkinson’s disease. Brain 2005, 128, 2665–2674. [CrossRef]

109. Sanchez-Guajardo, V.; Tentillier, N.; Romero-Ramos, M. Microglia Response During Parkinson’s Disease: Alpha-Synuclein Intervention. Front. Cell Neurosci. 2018, 12, 247. [CrossRef]

110. Croisier, E.; Moran, L.B.; Dexter, D.T.; Pearce, R.K.; Graeber, M.B. Microglial inflammation in the parkinsonian substantia nigra: Relationship to alpha-synuclein deposition. J. Neuroinflamm. 2005, 2, 14. [CrossRef]

111. Reale, M.; Iarlori, C.; Thomas, A.; Gambi, D.; Perfetti, B.; Di Nicola, M.; Onofrj, M. Peripheral cytokines profile in Parkinson’s disease. Brain Behav. Immun. 2009, 23, 55–63. [CrossRef]

112. Stypula, G.; Kunert-Radek, J.; Stepien, H.; Zylinska, K.; Pawlikowski, M. Evaluation of interleukins, ACTH, cortisol and prolactin concentrations in the blood of patients with parkinson’s disease. Neuroimmunomodulation 1996, 3, 131–134. [CrossRef]

113. Blum-Degen, D.; Muller, T.; Kuhn, W.; Gerlach, M.; Przuntek, H.; Riederer, P. Interleukin-1 beta and interleukin-6 are elevated in the cerebrospinal fluid of Alzheimer’s and de novo Parkinson’s disease patients. Neurosci. Lett. 1995, 202, 17–20. [CrossRef]

114. Mogi, M.; Harada, M.; Kondo, T.; Riederer, P.; Inagaki, H.; Minami, M.; Nagatsu, T. Interleukin-1 beta, interleukin-6, epidermal growth factor and transforming growth factor-alpha are elevated in the brain from parkinsonian patients. Neurosci. Lett. 1994, 180, 147–150. [CrossRef]

115. Mogi, M.; Harada, M.; Riederer, P.; Narabayashi, H.; Fujita, K.; Nagatsu, T. Tumor necrosis factor-alpha (TNF-alpha) increases both in the brain and in the cerebrospinal fluid from parkinsonian patients. Neurosci. Lett. 1994, 165, 208–210. [CrossRef]

116. Stypula, G.; Kunert-Radek, J.; Stepien, H.; Zylinska, K.; Pawlikowski, M. Evaluation of interleukins, ACTH, cortisol and prolactin concentrations in the blood of patients with Parkinson’s disease. Neuroimmunomodulation 1996, 3, 131–134. [CrossRef]

117. Reale, M.; Iarlori, C.; Thomas, A.; Gambi, D.; Perfetti, B.; Di Nicola, M.; Onofrj, M. Peripheral cytokines profile in Parkinson’s disease. Brain Behav. Immun. 2009, 23, 55–63. [CrossRef]

118. Chen, L.F.; Greene, W.C. Shaping the nuclear action of NF-kappaB. Nat. Rev. Mol. Cell Biol. 2004, 5, 392–401. [CrossRef]

119. Oeckinghaus, A.; Ghosh, S. NF-kappaB family of transcription factors and its regulation. Cold Spring Harb. Perspect. Biol. 2009, 1, a00034. [CrossRef]

120. Verma, I.M.; Stevenson, J.K.; Schwarz, E.M.; Van Antwerp, D.; Miyamoto, S. Rel/NF-kappa B/I kappa B family: Intimate tales of association and dissociation. Genes Dev. 1995, 9, 2723–2735. [CrossRef]

121. Oeckinghaus, A.; Hayden, M.S.; Ghosh, S. Crosstalk in NF-kappaB signaling pathways. Nat. Immunol. 2011, 12, 695–708. [CrossRef] [PubMed]

122. FDA—U.S. Food and Drug Administration. The Drug Development Process. Available online: https://www.fda.gov/patients/learn-about-drug-and-device-approvals/drug-development-process (accessed on 2 October 2019).

123. Rupaimoole, R.; Slack, F.J. MicroRNA therapeutics: Towards a new era for the management of cancer and other diseases. Nat. Rev. Drug Discov. 2017, 16, 203–222. [CrossRef] [PubMed]
124. Setten, R.L.; Rossi, J.J.; Han, S.P. The current state and future directions of RNAi-based therapeutics. *Nat. Rev. Drug Discov.* **2019**, *18*, 421–446. [CrossRef] [PubMed]

125. Chakraborty, C.; Sharma, A.R.; Sharma, G.; Doss, C.G.P.; Lee, S.S. Therapeutic miRNA and siRNA: Moving from Bench to Clinic as Next Generation Medicine. *Mol. Ther. Nucleic Acids* **2017**, *8*, 132–143. [CrossRef] [PubMed]

126. Hanna, J.; Hossain, G.S.; Kocerha, J. The Potential for microRNA Therapeutics and Clinical Research. *Front. Genet.* **2019**, *10*, 478. [CrossRef]

127. Titze-de-Almeida, S.S.; Brandao, P.R.P.; Faber, I.; Titze-de-Almeida, R. Leading RNA Interference Therapeutics Part 1: Silencing Hereditary Transthyretin Amyloidosis, with a Focus on Patisiran. *Mol. Diagn. Ther.* **2020**, *24*, 49–59. [CrossRef]

128. Adams, D.; Gonzalez-Duarte, A.; O’Riordan, W.D.; Yang, C.C.; Ueda, M.; Kristen, A.V.; Tournev, I.; Schmidt, H.H.; Coelho, T.; Berk, J.L.; et al. Patisiran, an RNAi Therapeutic, for Hereditary Transthyretin Amyloidosis. *N. Engl. J. Med.* **2018**, *379*, 11–21. [CrossRef]

129. Poewe, W.; Seppi, K.; Tanner, C.M.; Halliday, G.M.; Brundin, P.; Schrag, A.E.; Lang, A.E. Parkinson disease. *Nat. Rev. Dis. Primers* **2017**, *3*, 17013. [CrossRef]

130. Titze-de-Almeida, R.; David, C.; Titze-de-Almeida, S.S. The Race of 10 Synthetic RNAi-Based Drugs to the Pharmaceutical Market. *Pharm. Res.* **2017**, *34*, 1339–1363. [CrossRef]

131. Haussecker, D. Current issues of RNAi therapeutics delivery and development. *J. Control. Release* **2014**, *195*, 49–54. [CrossRef]

132. Kim, D.H.; Rossi, J.J. Strategies for silencing human disease using RNA interference. *Nat. Rev. Genet.* **2007**, *8*, 173–184. [CrossRef] [PubMed]

133. Ku, S.H.; Jo, S.D.; Lee, Y.K.; Kim, K.; Kim, S.H. Chemical and structural modifications of RNAi therapeutics. *Adv. Drug Deliv. Rev.* **2016**, *104*, 16–28. [CrossRef] [PubMed]

134. Kanasty, R.; Dorkin, J.R.; Vegas, A.; Anderson, D. Delivery materials for siRNA therapeutics. *Nat. Mater.* **2013**, *12*, 967–977. [CrossRef] [PubMed]

135. Khvorova, A.; Watts, J.K. The chemical evolution of oligonucleotide therapies of clinical utility. *Nature* **2017**, *535*, 238–248. [CrossRef]

136. Lindow, M.; Kauppinen, S. Discovering the first microRNA-targeted drug. *J. Cell Biol.* **2012**, *199*, 407–412. [CrossRef] [PubMed]

137. Orom, U.A.; Kauppinen, S.; Lund, A.H. LNA-modified oligonucleotides mediate specific inhibition of microRNA function. *Gene* **2006**, *372*, 137–141. [CrossRef]

138. Elmen, J.; Lindow, M.; Schütz, S.; Lawrence, M.; Petri, A.; Obad, S.; Lindholm, M.; Hedjärn, M.; Hansen, H.F.; Berger, U.; et al. LNA-mediated microRNA silencing in non-human primates. *Nature* **2008**, *452*, 896–899. [CrossRef]

139. Janssen, H.L.; Reesink, H.W.; Lawitz, E.J.; Zeuzem, S.; Rodríguez-Torres, M.; Patel, K.; van der Meer, A.J.; Patrick, A.K.; Chen, A.; Zhou, Y.; et al. Treatment of HCV infection by targeting microRNA. *N. Engl. J. Med.* **2013**, *368*, 1685–1694. [CrossRef]

140. Czauderna, F.; Fechtner, M.; Dames, S.; Aygun, H.; Klippel, A.; Pronk, G.J.; Giese, K.; Kaufmann, J. Structural variations and stabilising modifications of synthetic siRNAs in mammalian cells. *Nucleic Acids Res.* **2005**, *33*, 2705–2716. [CrossRef]

141. Cummins, L.L.; Owens, S.R.; Risen, L.M.; Lesnik, E.A.; Freier, S.M.; McGee, D.; Guinossos, C.J.; Cook, P.D. Characterization of fully 2′-modified oligoribonucleotide hetero- and homoduplex hybridization and nuclease sensitivity. *Nucleic Acids Res.* **1995**, *23*, 2019–2024. [CrossRef]

142. Layzer, J.M.; McCaffrey, A.P.; Tanner, A.K.; Huang, Z.; Kay, M.A.; Sullenger, B.A. In vivo activity of nuclease-resistant siRNAs. *RNA* **2004**, *10*, 766–771. [CrossRef] [PubMed]

143. Takahashi, M.; Minakawa, N.; Matsuda, A. Synthesis and characterization of 2′-modified-4′-thioRNA: A comprehensive comparison of nuclease stability. *Nucleic Acids Res.* **2009**, *37*, 1353–1362. [CrossRef] [PubMed]

144. Allerston, C.R.; Sioufi, N.; Jarres, R.; Prakash, T.P.; Naik, N.; Berdeja, A.; Wanders, L.; Griffey, R.H.; Swayze, E.E.; Bhat, B. Fully 2′-modified oligonucleotide duplexes with improved in vitro potency and stability compared to unmodified small interfering RNA. *J. Med. Chem.* **2005**, *48*, 901–904. [CrossRef] [PubMed]

145. Prakash, T.P.; Kinberger, G.A.; Murray, H.M.; Chappell, A.; Riney, S.; Graham, M.J.; Lima, W.F.; Swayze, E.E.; Seth, P.P. Synergistic effect of phosphorothioate, 5′-vinylphosphonate and GalNAc modifications for enhancing activity of synthetic siRNA. *Bioorg. Med. Chem. Lett.* **2016**, *26*, 2817–2820. [CrossRef] [PubMed]
146. Nair, J.K.; Willoughby, J.L.; Chan, A.; Charissse, K.; Alam, M.R.; Wang, Q.; Hoekstra, M.; Kandasamy, P.; Ke’lin, A.V.; Milstein, S.; et al. Multivalent N-acetylglucosamine-conjugated siRNA localizes in hepatocytes and elicits robust RNAi-mediated gene silencing. *J. Am. Chem. Soc.* **2014**, *136*, 16958–16961. [CrossRef]

147. Nair, J.K.; Attarwala, H.; Sehgal, A.; Wang, Q.; Aluri, K.; Zhang, X.; Gao, M.; Liu, J.; Indrakanti, R.; Schofield, S.; et al. Impact of enhanced metabolic stability on pharmacokinetics and pharmacodynamics of GalNAc-siRNA conjugates. *Nucleic Acids Res.* **2017**, *45*, 10969–10977. [CrossRef]

148. Mathupala, S.P. Delivery of small-interfering RNA (siRNA) to the brain. *Expert Opin. Ther. Pat.* **2009**, *19*, 137–140. [CrossRef]

149. Pardridge, W.M. Blood-brain barrier delivery. *Drug Discov. Today* **2007**, *12*, 54–61. [CrossRef]

150. De Boer, A.G.; Gaillard, P. Drug targeting to the brain. *Annu. Rev. Pharmacol. Toxicol.* **2007**, *47*, 323–355. [CrossRef]

151. Boado, R.J. Blood-brain barrier transport of non-viral gene and RNAi therapeutics. [PubMed]

152. Juliano, R.L. The delivery of therapeutic oligonucleotides. *Nucleic Acids Res.* **2016**, *44*, 6518–6548. [CrossRef] [PubMed]

153. Lorenzer, C.; Dirin, M.; Winkler, A.M.; Baumann, V.; Winkler, J. Going beyond the liver: Progress and challenges of targeted delivery of siRNA therapeutics. *J. Control. Release* **2015**, *203*, 1–15. [CrossRef] [PubMed]

154. Wittrup, A.; Lieberman, J. Knocking down disease: A progress report on siRNA therapeutics. [PubMed]

155. Cuellar, T.L.; Barnes, D.; Nelson, C.; Tanguay, J.; Yu, S.F.; Wen, X.; Scales, S.J.; Gesch, J.; Davis, D.; van Brabant, K.; et al. Systematic evaluation of antibody-mediated siRNA delivery using an industrial platform of THIOMAB-siRNA conjugates. *Nucleic Acids Res.* **2015**, *43*, 1189–1203. [CrossRef]

156. McNamara, J.O., 2nd; Andrechek, E.R.; Wang, Y.; Viles, K.D.; Rempel, R.E.; Gilboa, E.; Sullenger, B.A.; Giangrande, P.H. Cell type-specific delivery of siRNAs with aptamer-siRNA chimeras. *Nat. Biotechnol.* **2006**, *24*, 1005–1015. [CrossRef]

157. Berezhnoy, A.; Castro, I.; Levay, A.; Malek, T.R.; Gilboa, E. Aptamer-targeted inhibition of mTOR in T cells enhances antitumor immunity. *J. Clin. Investig.* **2014**, *124*, 188–197. [CrossRef]

158. Zhou, G.; Wilson, G.; Hebbard, L.; Duan, W.; Liddle, C.; George, J.; Qiao, L. Aptamers: A promising chemical antibody for cancer therapy. *Oncotarget* **2016**, *7*, 13446–13463. [CrossRef]

159. Thiel, W.H.; Thiel, K.W.; Flecken, K.S.; Bair, T.; Dupuy, A.J.; McNamara, J.O., 2nd; Miller, F.J.; Giangrande, P.H. Cell-internalization SELEX: Method for identifying cell-internalizing RNA aptamers for delivering siRNAs to target cells. *Methods Mol. Biol.* **2015**, *1218*, 187–199. [CrossRef]

160. Lonser, R.R.; Sarntinoranont, M.; Morrison, P.F.; Oldfield, E.H. Convection-enhanced delivery to the central nervous system. *J. Neurosurg.* **2015**, *122*, 697–706. [CrossRef]

161. Cohen, Z.R.; Ramishetti, S.; Pesheh-Yaloz, N.; Goldsmith, M.; Wohl, A.; Zibly, Z.; Peer, D. Localized RNAi therapeutics of chemoresistant grade IV glioma using hyaluronan-grafted lipid-based nanoparticles. *ACS Nano* **2015**, *9*, 1581–1591. [CrossRef]

162. Querbes, W.; Ge, P.; Zhang, W.; Fan, Y.; Costigan, J.; Charissse, K.; Maier, M.; Nechev, L.; Manoharan, M.; Kotelianiski, V.; et al. Direct CNS delivery of siRNA mediates robust silencing in oligodendrocytes. *Oligonucleotides* **2009**, *19*, 23–29. [CrossRef] [PubMed]

163. Halle, B.; Marcusson, E.G.; Aaberg-Jesscen, C.; Jensen, S.S.; Meyer, M.; Schulz, M.K.; Andersson, C.; Kristensen, B.W. Convection-enhanced delivery of an anti-miR is well-tolerated, preserves anti-miR stability and causes efficient target de-repression: A proof of concept. *J. Neuroloncol.* **2016**, *126*, 47–55. [CrossRef] [PubMed]

164. Lewis, O.; Woolley, M.; Johnson, D.; Rosser, A.; Barua, N.U.; Bienemann, A.S.; Gill, S.S.; Evans, S. Chronic, intermittent convection-enhanced delivery devices. *J. Neurosci. Methods* **2016**, *259*, 47–56. [CrossRef] [PubMed]

165. Danhier, F.; Messaoudi, K.; Lemaire, L.; Benoit, J.P.; Lagarce, F. Combined anti-Galectin-1 and anti-EGFR siRNA-loaded chitosan-lipid nanocapsules decrease temozolomide resistance in glioblastoma: In vivo evaluation. *Int. J. Pharm.* **2015**, *481*, 154–161. [CrossRef] [PubMed]

166. Tsujuchi, T.; Natsume, A.; Motomura, K.; Kondo, G.; Ranjit, M.; Hachisu, R.; Sugimura, I.; Tomita, S.; Takehara, I.; Woolley, M.; et al. Preclinical evaluation of an O(6)-methylguanine-DNA methyltransferase-siRNA/liposome complex administered by convection-enhanced delivery to rat and porcine brains. *Am. J. Transl. Res.* **2014**, *6*, 169–178. [CrossRef]

167. Karra, D.; Dahm, R. Transfection techniques for neuronal cells. *J. Neurosci.* **2010**, *30*, 6171–6177. [CrossRef]

168. Sariyer, I.K. Transfection of neuronal cultures. *Methods Mol. Biol.* **2013**, *1078*, 133–139. [CrossRef]
190. Islam, M.A.; Reesor, E.K.; Xu, Y.; Zope, H.R.; Zetter, B.R.; Shi, J. Biomaterials for mRNA delivery. *Biomater. Sci.* 2015, 3, 1519–1533. [CrossRef]

191. Xia, Y.; Tian, J.; Chen, X. Effect of surface properties on liposomal siRNA delivery. *Biomaterials* 2016, 79, 56–68. [CrossRef]

192. Enlund, E.; Fischer, S.; Handrick, R.; Otte, K.; Debatin, K.M.; Wabitsch, M.; Fischer-Posovszky, P. Establishment of lipofection for studying miRNA function in human adipocytes. *PLoS ONE* 2014, 9, e98023. [CrossRef] [PubMed]

193. Malone, R.W.; Felgner, P.L.; Verma, I.M. Cationic liposome-mediated RNA transfection. *Proc. Natl. Acad. Sci. USA* 1989, 86, 6077–6081. [CrossRef] [PubMed]

194. Enlund, E.; Fischer, S.; Handrick, R.; Otte, K.; Debatin, K.M.; Wabitsch, M.; Fischer-Posovszky, P. Establishment of lipofection for studying miRNA function in human adipocytes. *PLoS ONE* 2014, 9, e98023. [CrossRef] [PubMed]

195. Campani, V.; De Rosa, G.; Misso, G.; Zarone, M.R.; Grimaldi, A. Lipid Nanoparticles to Deliver miRNA in Cancer. *Curr. Pharm. Biotechnol.* 2016, 17, 741–749. [CrossRef]

196. Maheshwari, R.; Tekade, M.; Gondaliya, P.; Kalia, K.; D’Emanuele, A.; Tekade, R.K. Recent advances in exosome-based nanovehicles as RNA interference therapeutic carriers. *Nanomedicine* 2017, 12, 2653–2675. [CrossRef]

197. Alvarez-Erviti, L.; Seow, Y.; Yin, H.; Betts, C.; Lakhal, S.; Wood, M.J. Delivery of siRNA to the mouse brain by systemic injection of targeted exosomes. *Nat. Biotechnol.* 2011, 29, 341–345. [CrossRef]

198. Cooper, J.M.; Wiklander, P.B.; Nordin, J.Z.; Al-Shawi, R.; Wood, M.J.; Vithlani, M.; Schapira, A.H.; Simons, J.P.; El-Andaloussi, S.; Alvarez-Erviti, L. Systemic exosomal siRNA delivery reduced alpha-synuclein aggregates in brains of transgenic mice. *Mov. Disord.* 2014, 29, 1476–1485. [CrossRef]

199. Ibrahim, A.F.; Weirauch, U.; Thomas, M.; Grunweller, A.; Hartmann, R.K.; Aigner, A. MicroRNA replacement therapy for miR-145 and miR-33a is efficacious in a model of colon carcinoma. *Cancer Res.* 2011, 71, 5214–5224. [CrossRef]

200. Liang, G.; Zhu, Y.; Jing, A.; Wang, J.; Hu, F.; Feng, W.; Xiao, Z.; Chen, B. Cationic microRNA-delivering nanocarriers for efficient treatment of colon carcinoma in xenograft model. *Gene Ther.* 2016, 23, 829–838. [CrossRef]

201. Helmschrodt, C.; Hobel, S.; Schoniger, S.; Bauer, A.; Bonicelli, J.; Gringmuth, M.; Fietz, S.A.; Aigner, A.; Richter, A.; Richter, F. Polyethyleneimine Nanoparticle-Mediated siRNA Delivery to Reduce alpha-Synuclein Expression in a Model of Parkinson’s Disease. *Mol. Ther. Nucleic Acids* 2017, 5, 57–68. [CrossRef]

202. Liu, Y.Y.; Yang, X.Y.; Li, Z.; Liu, Z.L.; Cheng, D.; Wang, Y.; Wen, X.J.; Hu, J.Y.; Liu, J.; Wang, L.M.; et al. Characterization of polyethylene glycol-polyethyleneimine as a vector for alpha-synuclein siRNA delivery to PC12 cells for Parkinson’s disease. *CNS Neurosci. Ther.* 2014, 20, 76–85. [CrossRef] [PubMed]

203. Yuan, X.; Shah, B.A.; Kotadia, N.K.; Li, J.; Gu, H.; Wu, Z. The development and mechanism studies of cationic chitosan-modified biodegradable PLGA nanoparticles for efficient siRNA drug delivery. *Pharm. Res.* 2010, 27, 1285–1295. [CrossRef] [PubMed]

204. Sariaga, C.; Paiva, J.; Santos, T.; Ferreira, L.; Bernardino, L. MicroRNA-124 loaded nanoparticles enhance brain repair in Parkinson’s disease. *J. Control. Release* 2016, 235, 291–305. [CrossRef] [PubMed]

205. Gan, L.; Li, Z.; Lv, Q.; Huang, W. Rabies virus glycoprotein (RVG29)-linked microRNA-124-loaded polymeric nanoparticles inhibit neuroinflammation in a Parkinson’s disease model. *Int. J. Pharm.* 2019, 567, 118449. [CrossRef]

206. Mattu, C.; Pabari, R.M.; Boffito, M.; Sartori, S.; Ciardelli, G.; Ramtoola, Z. Comparative evaluation of novel biodegradable nanoparticles for the drug targeting to breast cancer cells. *Eur. J. Pharm. Biopharm.* 2013, 85, 463–472. [CrossRef]

207. Yang, Y.P.; Chien, Y.; Chiou, G.Y.; Cherng, J.Y.; Wang, M.L.; Lo, W.L.; Chang, Y.L.; Huang, P.I.; Chen, Y.W.; Shih, Y.H.; et al. Inhibition of cancer stem cell-like properties and reduced chemoradioresistance of glioblastoma using microRNA145 with cationic polyurethane-short branch PEI. *Biomaterials* 2012, 33, 1462–1476. [CrossRef]

208. Bravo-Anaya, L.M.; Fernandez-Solis, K.G.; Rossegong, J.; Nano-Rodriguez, J.L.E.; Carvajal, F.; Rinaudo, M. Chitosan-DNA polyelectrolyte complex: Influence of chitosan characteristics and mechanism of complex formation. *Int. J. Biol. Macromol.* 2019, 126, 1037–1049. [CrossRef]

209. Costa, D.F.; Torchilin, V.P. Micelle-like nanoparticles as siRNA and miRNA carriers for cancer therapy. *Biomed. Microdevices* 2018, 20, 59. [CrossRef]
210. Santos-Carballal, B.; Aaldering, L.J.; Ritzefeld, M.; Pereira, S.; Sewald, N.; Moerschbacher, B.M.; Gotte, M.; Goycoolea, F.M. Physicochemical and biological characterization of chitosan-microRNA nanocomplexes for gene delivery to MCF-7 breast cancer cells. *Sci. Rep.* **2015**, *5*, 13567. [CrossRef]

211. Deng, X.; Cao, M.; Zhang, J.; Hu, K.; Yin, Z.; Zhou, Z.; Xiao, X.; Yang, Y.; Sheng, W.; Wu, Y.; et al. Hyaluronic acid-chitosan nanoparticles for co-delivery of MiR-34a and doxorubicin in therapy against triple negative breast cancer. *Biomaterials* **2014**, *35*, 4333–4344. [CrossRef]

212. Layek, B.; Singh, J. N-hexanoyl, N-octanoyl and N-decanoyl chitosans: Binding affinity, cell uptake, and transfection. *Carbohydr. Polym.* **2012**, *89*, 403–410. [CrossRef]

213. Bose, A.; Beal, M.F. Mitochondrial dysfunction in Parkinson's disease. *Transl. Neurodegener.* **2016**, *5*, 14. [CrossRef] [PubMed]

214. Niu, S.; Zhang, L.K.; Zhang, L.; Zhuang, S.; Zhan, X.; Chen, W.Y.; Du, S.; Yin, L.; You, R.; Li, C.H.; et al. Inhibition of multifunctional magnetic nanoparticles loaded with Alpha-Synuclein RNAi Plasmid in a Parkinson's Disease Model. *Theranostics* **2017**, *7*, 344–356. [CrossRef]

215. Yin, P.T.; Shah, B.P.; Lee, K.B. Combined magnetic nanoparticle-based microRNA and hyperthermia therapy to enhance apoptosis in brain cancer cells. *Small* **2014**, *10*, 4106–4112. [CrossRef] [PubMed]

216. Niu, S.; Zhang, L.K.; Zhang, L.; Zhuang, S.; Zhan, X.; Chen, W.Y.; Du, S.; Yin, L.; You, R.; Li, C.H.; et al. Inhibition by Multifunctional Magnetic Nanoparticles Loaded with Alpha-Synuclein RNAi Plasmid in a Parkinson's Disease Model. *Theranostics* **2017**, *7*, 344–356. [CrossRef]

217. Zhang, Y.; Li, H.; Sun, J.; Gao, J.; Liu, W.; Li, B.; Guo, Y.; Shen, J. DC-Chol/DOPE cationic liposomes: A comparative study of the influence factors on plasmid pDNA and siRNA gene delivery. *Int. J. Pharm.* **2010**, *390*, 198–207. [CrossRef]

218. Bertucci, A.; Prasetyanto, E.A.; Septiadi, D.; Manicardi, A.; Brogna, E.; Gambri, R.; Corradini, R.; De Cola, L. Combined Delivery of Temozolomide and Anti-miR221 PNA Using Mesoporous Silica Nanoparticles Induces Apoptosis in Resistant Glioma Cells. *Small* **2015**, *11*, 5687–5695. [CrossRef]

219. Tivnan, A.; Orr, WS.; Gubala, V.; Nooney, R.; Williams, D.E.; McDonagh, C.; Prenter, S.; Harvey, H.; Domingo-Fernandez, R.; Bray, I.M.; et al. Inhibition of neuroblastoma tumor growth by targeted delivery of microRNA-34a using anti-disialoganglioside GD2 coated nanoparticles. *PLoS ONE* **2012**, *7*, e38129. [CrossRef]

220. Jimenez-Mateos, E.M.; Engel, T.; Merino-Serrais, P.; McKiernan, R.C.; Tanaka, K.; Mouri, G.; Sano, T.; O’Tuathaigh, C.; Waddington, J.L.; Prenter, S.; et al. Silencing microRNA-134 produces neuroprotective and prolonged seizure-suppressive effects. *Nat. Med.* **2012**, *18*, 1087–1094. [CrossRef] [PubMed]

221. Jimenez-Mateos, E.M.; Engel, T.; Merino-Serrais, P.; Fernaud-Espinosa, I.; Rodriguez-Alvarez, N.; Reynolds, J.; Reschke, C.R.; Conroy, R.M.; McKiernan, R.C.; deFelipe, J.; et al. Antagomirs targeting microRNA-134 increase hippocampal pyramidal neuron spine volume in vivo and protect against pilocarpine-induced status epilepticus. *Brain Struct. Funct.* **2015**, *220*, 2387–2399. [CrossRef]

222. Reschke, C.R.; Silva, L.F.; Norwood, B.A.; Senthilkumar, K.; Morris, G.; Sanz-Rodriguez, A.; Conroy, R.M.; Costard, L.; Neubert, V.; Bauer, S.; et al. Potent Anti-seizure Effects of Locked Nucleic Acid Antagomirs Targeting miR-134 in Multiple Mouse and Rat Models of Epilepsy. *Mol. Ther. Nucleic Acids* **2017**, *6*, 45–56. [CrossRef]

223. Bose, A.; Beal, M.F. Mitochondrial dysfunction in Parkinson’s disease. *J. Neurochem.* **2016**, *139* (Suppl. 1), 216–231. [CrossRef] [PubMed]

224. Martin, L.J.; Semenkow, S.; Hanaford, A.; Wong, M. Mitochondrial permeability transition pore regulates Parkinson’s disease development in mutant alpha-synuclein transgenic mice. *Neurobiol. Aging* **2014**, *35*, 1132–1152. [CrossRef]
228. Itoh, K.; Wakabayashi, N.; Katoh, Y.; Ishii, T.; Igarashi, K.; Engel, J.D.; Yamamoto, M. Keap1 represses nuclear activation of antioxidant responsive elements by Nrf2 through binding to the amino-terminal Neh2 domain. Genes Dev. 1999, 13, 76–86. [CrossRef]  
229. Ma, Q. Role of nrf2 in oxidative stress and toxicity. Annu. Rev. Pharmacol. Toxicol. 2013, 53, 401–426. [CrossRef] [PubMed]  
230. Nguyen, T.; Nioi, P.; Pickett, C.B. The Nrf2-antioxidant response element signaling pathway and its activation by oxidative stress. J. Biol. Chem. 2009, 284, 13291–13295. [CrossRef]  
231. Shih, R.H.; Wang, C.Y.; Yang, C.M. NF-kappaB Signaling Pathways in Neurological Inflammation: A Mini Review. Front. Mol. Neurosci. 2015, 8, 77. [CrossRef]  
232. Wang, Q.; Liu, Y.; Zhou, J. Neuroinflammation in Parkinson’s disease and its potential as therapeutic target. Transl. Neurodegener. 2015, 4, 19. [CrossRef]  
233. Chen, P.; DeWitt, M.R.; Bornhorst, J.; Soares, F.A.; Mukhopadhyay, S.; Bowman, A.B.; Aschner, M. Age- and manganese-dependent modulation of dopaminergic phenotypes in a C. elegans DJ-1 genetic model of Parkinson’s disease. Metallomics 2015, 7, 289–298. [CrossRef]  
234. Haussecker, D. The Business of RNAi Therapeutics in 2012. Mol. Ther. Nucleic Acids 2012, 1, e8. [CrossRef]  
235. Whitehead, K.A.; Langer, R.; Anderson, D.G. Knocking down barriers: Advances in siRNA delivery. Nat. Rev. Drug Discov. 2009, 8, 129–138. [CrossRef]  
236. Yin, H.; Kanasty, R.L.; Eltoukhy, A.A.; Vegas, A.J.; Dorkin, J.R.; Anderson, D.G. Non-viral vectors for gene-based therapy. Nat. Rev. Genet. 2014, 15, 541–555. [CrossRef]  
237. Alabi, C.; Vegas, A.; Anderson, D. Attacking the genome: Emerging siRNA nanocarriers from concept to clinic. Curr. Opin. Pharmacol. 2012, 12, 427–433. [CrossRef] [PubMed]  
238. Zhao, J.; Feng, S.S. Nanocarriers for delivery of siRNA and co-delivery of siRNA and other therapeutic agents. Nanomedicine 2015, 10, 2199–2228. [CrossRef] [PubMed]  
239. Williford, J.M.; Wu, J.; Ren, Y.; Archang, M.M.; Leong, K.W.; Mao, H.Q. Recent advances in nanoparticle-mediated siRNA delivery. Annu. Rev. Biomed. Eng. 2014, 16, 347–370. [CrossRef] [PubMed]  

© 2020 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).