

## A role for FNDC5/Irisin in the beneficial effects of exercise on the brain and in neurodegenerative diseases



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### ABSTRACT

The beneficial effects of exercise on the brain are well known. However, the underlying molecular mechanisms are much less well understood. Interestingly, myokines, hormones secreted by muscle in response to exercise, gained attention as such beneficial mediators. In this review, we will focus on FNDC5 and its secreted form, the newly discovered myokine “irisin”. We will discuss their role in the beneficial effects of exercise and its potential application in neurodegenerative disorders.

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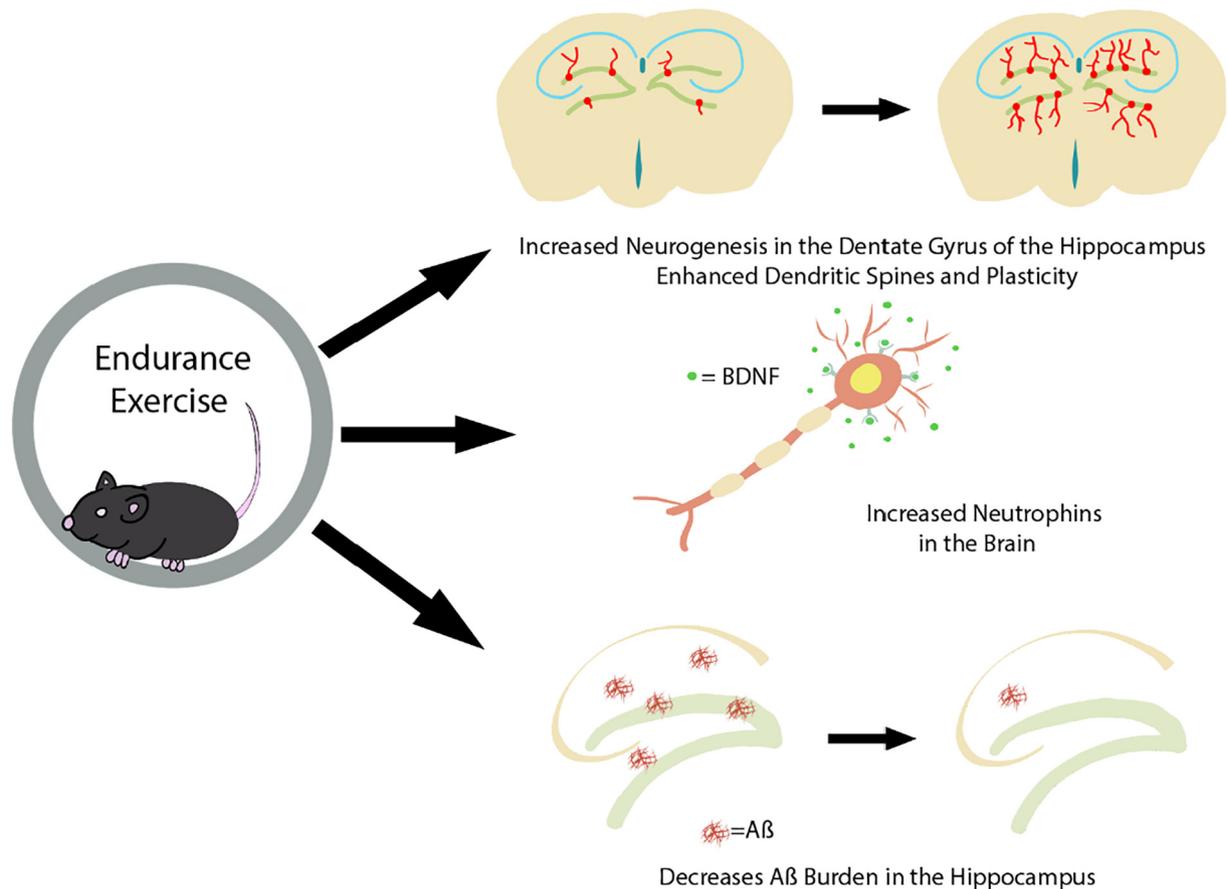
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**Abbreviations:** A $\beta$ , amyloid beta protein; AD, Alzheimer's disease; Akt/NF $\kappa$ B, protein kinase B/nuclear factor kappa-light-chain-enhancer of activated B cells; AMPK, 5' Adenosine monophosphate-activated protein kinase; BDNF, brain derived neurotrophic factor; BrdU, 5-bromo-2'-deoxyuridine; COX-2, cyclooxygenase 2; ELISA, enzyme-linked immunosorbent assay; FNDC5, fibronectin type III domains containing protein 5; GFAP, glial fibrillary acidic protein; HD, Huntington's disease; IL-1, interleukin 1; IL-6, interleukin 6; Map2, microtubule-associated protein 2; NADPH, nicotinamide adenine dinucleotide phosphate; PA, physical activity; PD, Parkinson's disease; PGC-1 $\alpha$ , peroxisome proliferator-activated receptor gamma coactivator 1-alpha; RNAi, RNA interference; TrkB, tropomyosin receptor kinase b.

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Physical activity (PA) has long been associated with tremendous benefits for the human body. In particular, increased PA has been correlated with improved health outcomes.<sup>1</sup> Furthermore, aerobic exercise, also known as endurance exercise, has been shown to have beneficial effects on cognitive function and overall brain plasticity.<sup>2–4</sup> This exercise-induced improvement in cognitive function has been particularly notable in older adults.<sup>2,5</sup> Exercise has also been shown to improve brain-related outcomes in particular neurodegenerative disorders, such as Parkinson's Disease (PD)<sup>6</sup> and Alzheimer's Diseases (AD)<sup>7</sup> as well neurovascular trauma such as stroke<sup>8</sup> (Fig 1). The hippocampus, an area of the brain associated with cognitive function and spatial awareness, is the main benefactor of the beneficial effects of exercise on the brain. Specific beneficial effects of exercise in the brain include



**Fig 1.** The beneficial effects of exercise on the brain. Exercise has many beneficial effects on the brain ranging from neurogenesis to improvements in memory function. These include a reduction in neuroinflammation, improvements in synaptic plasticity, increased adult hippocampal neurogenesis, and improvements in memory, mainly spatial learning and memory as well as pattern separation.

increases in the blood flow to the hippocampus and increases in its size in humans. In rodent models, exercise results in morphological changes in dendrites and dendritic spines, increased synapse plasticity, and importantly, enhanced de novo neurogenesis in the dentate gyrus.<sup>4,9</sup> Other beneficial effects of exercise include a reduction in neuroinflammation.<sup>10</sup> For a long time, the adult brain was thought to lack regenerative capabilities. Now it is known that two distinct regions of the brain seem to undergo de novo neurogenesis: the subgranular zone of the dentate gyrus of the hippocampus and a region connecting the rostral migratory stream to the olfactory bulb, the subventricular zone.<sup>11</sup> Moreover, exercise is one of the few stimuli known to provoke de novo neurogenesis in the adult mammalian brain as first shown by Van Praag et al.<sup>12</sup> In this study using BrdU to label dividing cells, the group demonstrated that voluntary free-wheel exercise in mice was sufficient to induce neurogenesis in the mouse dentate gyrus region of the hippocampus.<sup>12</sup> In this study, bromodeoxyuridine-positive cells, a marker for cell division, was used to quantify neurogenesis in the dentate gyrus. Later that year, van Praag also established that this exercise induced neurogenesis lead to both improvements in synaptic plasticity and spatial learning.<sup>13</sup> Interestingly, these effects were noted only with voluntary exercise and were not seen with an enriched environment lacking a running wheel.<sup>14,15</sup>

Although these beneficial effects of exercise on cognition are widely accepted, the actual mediator of these effects remains to be uncovered. Despite this, there has been considerable evidence regarding the gene FNDC5, along with its secreted form irisin, and its role in improving cognitive function in the brain following exercise. The goal of this review is to shed light on FNDC5/irisin and its role in the beneficial effects of exercise on cognitive function and its potential application in neurodegenerative disorders.

### Factors involved in brain health

Factors mediating the beneficial effects of PA/exercise on the brain can be broadly divided in two categories<sup>1</sup>: factors locally produced in the brain, so-called neurotrophins and<sup>2</sup> systemic factors circulating in the blood, for example, produced by skeletal muscle (myokines). During the last years, systemic factors released from skeletal muscle have gained a lot interest as regulators of the beneficial effects of exercise on the brain among them FNDC5 and its secreted form “irisin”. Other systemic factors are being discussed in detail in this issue by Tari et al. Neurotrophins play a major role in the growth and development of neurons, ranging from synaptic plasticity to survival. One of these neurotrophic factors is brain-derived neurotrophic factor (BDNF). Interestingly, as we will discuss later in this review, FNDC5 seems to be an important regulator of BDNF and therefore, we will briefly introduce BDNF. For an in-depth review on BDNF and exercise, please see.<sup>9,16,17</sup> BDNF has been shown to enhance neuronal survival, migration, and dendritic outgrowth.<sup>18,19</sup> In addition, it regulates synaptic plasticity and cognitive function.<sup>20–22</sup> BDNF has also been shown to be a crucial regulator of the beneficial effects of exercise in the brain. For example, Vaynman et al. has shown that blocking the tropomyosin receptor kinase B (TrkB), the BDNF receptor, with anti-TrkB antibodies leads to diminished expression of important synaptic proteins as well as decreased retention in spatial learning tasks, both of which have previously been seen to be part of the exercise-induced benefits on the brain.<sup>23,24</sup>

### Origins of FNDC5/Irisin

FNDC5 (fibronectin type III domains containing protein 5) was originally described in 2002 independently by two separate laboratories.

Teufel et al. found the FRCP2 gene while searching and characterizing different fibronectin type III domains,<sup>25</sup> and Ferrer-Martinez et al. discovered FRCP2, also known as the PeP protein, during peroxisomal studies.<sup>26</sup> FRCP2/PeP was later renamed FNDC5. The cleaved and secreted form of FNDC5 was discovered in 2012 and named “irisin” after the Greek goddess Iris, the messenger of the gods.<sup>27</sup>

FNDC5 is a glycosylated type I membrane protein. It contains a N-terminal signal peptide [amino acid (aa) 1–28], a FNIII domain (aa 33–124), a transmembrane domain (aa 150–170), and a cytoplasmic tail (aa 171–209) ([www.uniprot.org](http://www.uniprot.org)) (Fig 2). Irisin is generated via proteolytic cleavage of FNDC5 and contains 112 amino acids (aa 29–140). The enzyme that cleaves FNDC5 has yet to be uncovered. However, the crystal structure of irisin has been resolved.<sup>28</sup> Intriguingly, the FNIII-like domain shows an unusual confirmation, with a continuous intersubunit beta-sheet dimer, that has not been previously described for any other FNIII protein. Subsequent biochemical experiments confirmed the existence of irisin (bacterial recombinant) as a homodimer. In a recent paper investigating the receptor for irisin, biochemical and biophysical studies identified interactions between irisin and  $\alpha V/\beta 5$  integrin.<sup>29</sup> Chemical inhibition of the  $\alpha V$  integrin blocked irisin signaling in both osteocytes and fat cells.

### Quantification of irisin

Irisin is a unique polypeptide: the sequence is 100% conserved between humans and mice. It has an atypical start codon in humans (ATA not ATG). In addition, it circulates at very low levels in the blood.<sup>27,30</sup> These features of irisin led to a controversy whether irisin indeed was a circulating hormone. The atypical start codon of human FNDC5 sparked debate, despite previous studies establishing that a small percentage of functional eukaryotic mRNAs do indeed begin translation with this atypical ATA.<sup>31,32</sup> Some believed that the atypical start codon of human FNDC5 meant it was a “null mutation” or pseudogene and thus functional circulating irisin would not exist.<sup>33,34</sup> This was exacerbated by the lack of a defined antibody, ELISA, or other method of detecting or quantifying circulating irisin at the time.<sup>35,36</sup>

To address this issue, in 2015 we developed a method to quantify circulating irisin in human plasma via targeted mass spectrometry with control peptides enriched with stable heavy isotopes as internal standards.<sup>30</sup> This method was both state-of-the-art and precise; it demonstrated that the non-canonical ATA start codon of human irisin was in fact the main site of initiating translation. Further, we were able to quantify circulating irisin in sedentary individuals at a concentration of ~3.6 ng/ml. This circulating concentration was found to be significantly higher in individuals that participated in endurance exercise. This study determined that in fact human irisin does exist and can

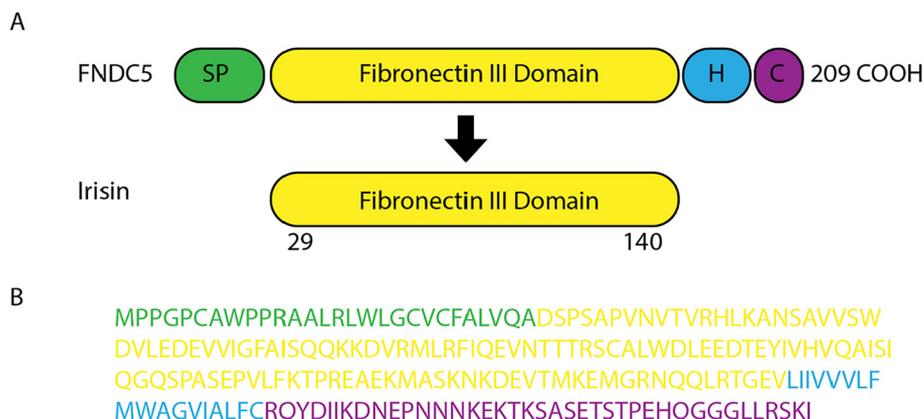
be regulated via aerobic exercise. Recently, another group using this targeted mass spectrometry approach with heavy internal control peptides found irisin to be present in human cerebrospinal fluid at approximately 0.26–1.86 ng/ml in men over 80 years of age with various diseases.<sup>37</sup> However, a knock-out validated antibody for irisin is still missing and would be a valuable resource for the scientific community.

### Irisin in the central nervous system-neural development effects

*Fndc5* is known to be profoundly expressed in the brain in many regions, including cerebellar Purkinje cells,<sup>25,26,38</sup> the hypothalamus,<sup>39</sup> as well as the hippocampus.<sup>40</sup> Further, the presence of irisin in the cerebrospinal fluid of humans was identified via Western blot and mass spectrometry.<sup>37,41</sup> During the maturation of primary mouse embryonic cortical neuron cultures or the differentiation of human embryonic stem cell-derived neural cells into neurons, FNDC5 levels are increasing.<sup>40,42</sup> High levels of FNDC5 are also found in the heart and oxidative skeletal muscle.<sup>40</sup> Forced overexpression of FNDC5 during the formation of neuronal precursors from a mouse embryonic stem cell population lead to increased BDNF, GFAP as well as Map2, b-tubulinIII, and Neurocan, all three being markers of neuronal maturation. However, when FNDC5 was overexpressed in an undifferentiated mouse embryonic stem cell population, the aforementioned effects were not detected.<sup>43</sup> Knockdown of FNDC5 in neuronal precursor cells has been shown to impair the maturation process of neurons and astrocytes.<sup>44</sup> Pharmacological dose of recombinant irisin in the mouse H19–7 hippocampal cell line led to increased cell proliferation.<sup>45</sup> We have previously shown that overexpression of FNDC5 in primary cortical neurons increased cell survival in culture whereas knockdown of FNDC5 reduces the cell survival.<sup>40</sup> Pharmacological dose of recombinant irisin in the mouse H19–7 hippocampal cell line led to increased cell proliferation.<sup>45</sup> Together, these data suggest that FNDC5/irisin play a developmental role in regulating the process of neuronal differentiation and maturation.

### The role of FNDC5/irisin in PA/exercise

Skeletal muscle composes 40% of the body mass in humans. Skeletal muscle is an endocrine organ and secretes myokines, muscle-derived secretory proteins with a wide array of biological functions.<sup>17</sup> Endurance exercise activates on a central transcriptional coactivator known as PGC-1 $\alpha$ . Irisin, the cleaved form of FNDC5 was shown to be a PGC-1 $\alpha$ -dependent myokine that was secreted from skeletal muscle during exercise by Bostrom et al. in 2012.<sup>27</sup> Some of the major metabolic benefits of PA/exercise were shown to be promoted by irisin; these include increased glucose tolerance as well as the “browning” of white



**Fig 2.** Analysis of Irisin Peptides by Mass Spectrometry. (a) Scheme of the murine FNDC5 protein structure (top) and murine irisin protein structure (bottom). SP signal peptide, H hydrophobic domain, C cytoplasmic domain. (b) Murine FNDC5 amino acid sequence with corresponding domains colored. The irisin sequence is in yellow.

adipose tissue.<sup>27,46</sup> Upregulation of circulating irisin by overexpression of FNDC5 in the liver, via adenoviral vector, led to an increase in both.<sup>27</sup> Many studies have confirmed the increase of *Fndc5* mRNA in skeletal muscle during exercise in mice<sup>40,47–49</sup> as well as humans<sup>34,36,50–52</sup> using RNA sequencing or qPCR. However, there are studies, which have not found an increase of *FNDC5* expression in muscle after exercise, suggesting that FNDC5 is not induced by any and all exercise interventions. Of note, several of these exercise regimens that did not see an upregulation of *FNDC5* also did not observe an increase in PGC1a, its transcriptional regulator<sup>52,53</sup> or did not assess changes of PGC-1a.<sup>54–56</sup> Other factors that could explain the differences between the different studies are: exercise regimen used, mode of exercise, time point of sampling, and study population.

In addition to the beneficial effects of irisin on metabolism, we later showed that endurance exercise induced irisin expression not only in skeletal muscle but also in the hippocampus, a region of the brain involved in memory and spatial awareness.<sup>40</sup> We demonstrated that neuronal *Fndc5* gene expression is also regulated by PGC-1 $\alpha$ . In addition, we identified FNDC5 as an important regulator of BDNF. Forced expression of FNDC5 in primary cortical neurons increased *Bdnf* expression and *Bdnf* expression was markedly reduced following RNAi-mediated knockdown of FNDC5. Similarly, the common human BDNF Val66Met polymorphism was shown to impair both BDNF and FNDC5 expression in the brain following exercise.<sup>49</sup> Likewise, Choi et al. showed that exercise-induced adult hippocampal neurogenesis was associated with increases in both BDNF and FNDC5, helping improve cognition in a mouse model of AD.<sup>57</sup> Importantly, peripheral delivery of FNDC5 to the liver via adenoviral vectors, which elevated blood irisin levels, induced the expression of *Bdnf* and other neuroprotective genes in the hippocampus. This finding implies that irisin, or factor induced by irisin, can cross the blood-brain barrier to affect gene expression in the brain. This discovery encouraged the idea that irisin has great therapeutic promise prompting further research on the role of irisin, the hippocampus, and improvements in cognitive function. Since then, there have been a few studies measuring the serum levels of irisin in relation to the cognitive function of either aging individuals,<sup>58</sup> young athletes<sup>59</sup> or obese and morbidly obese patients.<sup>60</sup> Two out of the three studies found that irisin positively correlates with better cognitive function,<sup>58,59</sup> particularly in the active individuals, whereas the other found a negative correlation of irisin with cognitive function.<sup>60</sup> In a

study investigating diabetic mild cognitive impairment, irisin overexpression showed elevated levels of BDNF in the serum and the hippocampus and lower levels of inflammatory advanced glycated end products and glycosylated hemoglobinA1c.<sup>61</sup> In primary hippocampal nerve cell culture from these diabetic rats, the overexpression of irisin produced higher levels of BDNF and cellular metabolic NAD(P)H-dependent activity following exposure to increasing levels of glucose. These results indicate that irisin may regulate the expression of BDNF and glycometabolism in diabetic rats. Taken together, these findings point toward the relationship of FNDC5 and BDNF expression in the brain that is linked to endurance exercise and subsequently the important metabolic mediators PGC-1 $\alpha$ . Further it indicates that irisin seems to be a key peptide at the interface between metabolism and brain function.

Of note, exercise can consist of either endurance or resistance exercise training, the former an aerobic, cardiovascular form of exercise and the latter focusing more muscle strength and hypertrophy. Currently, evidence suggest that irisin is not involved in resistance training. This is to be expected as endurance exercise activates PGC-1 $\alpha$ 1, an upstream transcriptional regulator of FNDC5, while resistance training utilizes the PGC-1 $\alpha$ 4 isoform, inducing hypertrophy in exercised muscle.<sup>62</sup> Similarly, a study found no differences in serum irisin levels between control and exercised individuals after high-intensity interval and resistance training.<sup>63</sup>

**Irisin and neuroprotection in neurodegeneration**

There is a vast body of literature demonstrating the beneficial effects of PA/exercise on neurodegenerative diseases (Fig 3), including AD (as reviewed by Tari et al. in this issue), PD,<sup>6</sup> and Huntington's disease (HD).<sup>64</sup> As discussed above, the neurotrophin BDNF plays an important role in the homeostatic function and maintenance of the neurons; particularly in synaptic plasticity and neurogenesis. Decreased levels of BDNF have been identified in serum, as well as in hippocampal and cortex samples of AD and PD patients.<sup>65–67</sup> To establish novel blood-based biomarkers of cognition and stress, Küster et al. investigated the serum levels of BDNF, irisin and molecules from the kynurenine pathway as potential biomarkers of cognitive decline and dementia in older adults at risk of dementia. Following PA/exercise or cognitive training compared to a control group, both BDNF and irisin serum levels positively correlated with global cognition scores and memory.<sup>58</sup>

<b>Neurodegenerative Diseases</b> 	<b>Alzheimer's</b> A $\beta$ plaques Neurofibrillary tangles Cognitive decline	<b>Parkinson's</b> Hypo/hyperkinesia Lewy bodies (a-synuclein plaques) Loss of dopaminergic neurons	<b>Huntington's</b> Hyperkinesia Atrophy Cognitive decline	<b>Depression</b> Atrophy/neuronal loss Decrease in BDNF in serum and the hippocampus
<b>Effects of Exercise/Irisin</b> 	<b>Alzheimer's</b> Improved BDNF and Irisin levels leading to improved cognition and AD pathogenesis (Choi et al. 2018)	<b>Alzheimer's</b> Irisin and A $\beta$ peptide conditioned astrocyte media led to a decrease in inflammatory factors (IL-1 $\beta$ , COX2, IL-6) and increase in overall neuronal viability (Wang et al. 2018)	<b>Huntington's</b> Increased systemic delivery of BDNF leads to neuroprotection and amelioration of neurological signs (Giampà et al. 2016)	<b>Depression</b> Recombinant irisin injection increased activity in mitochondrial complexes I, II, IV and creatine kinase in the prefrontal cortex of rats (Wang et al. 2016)

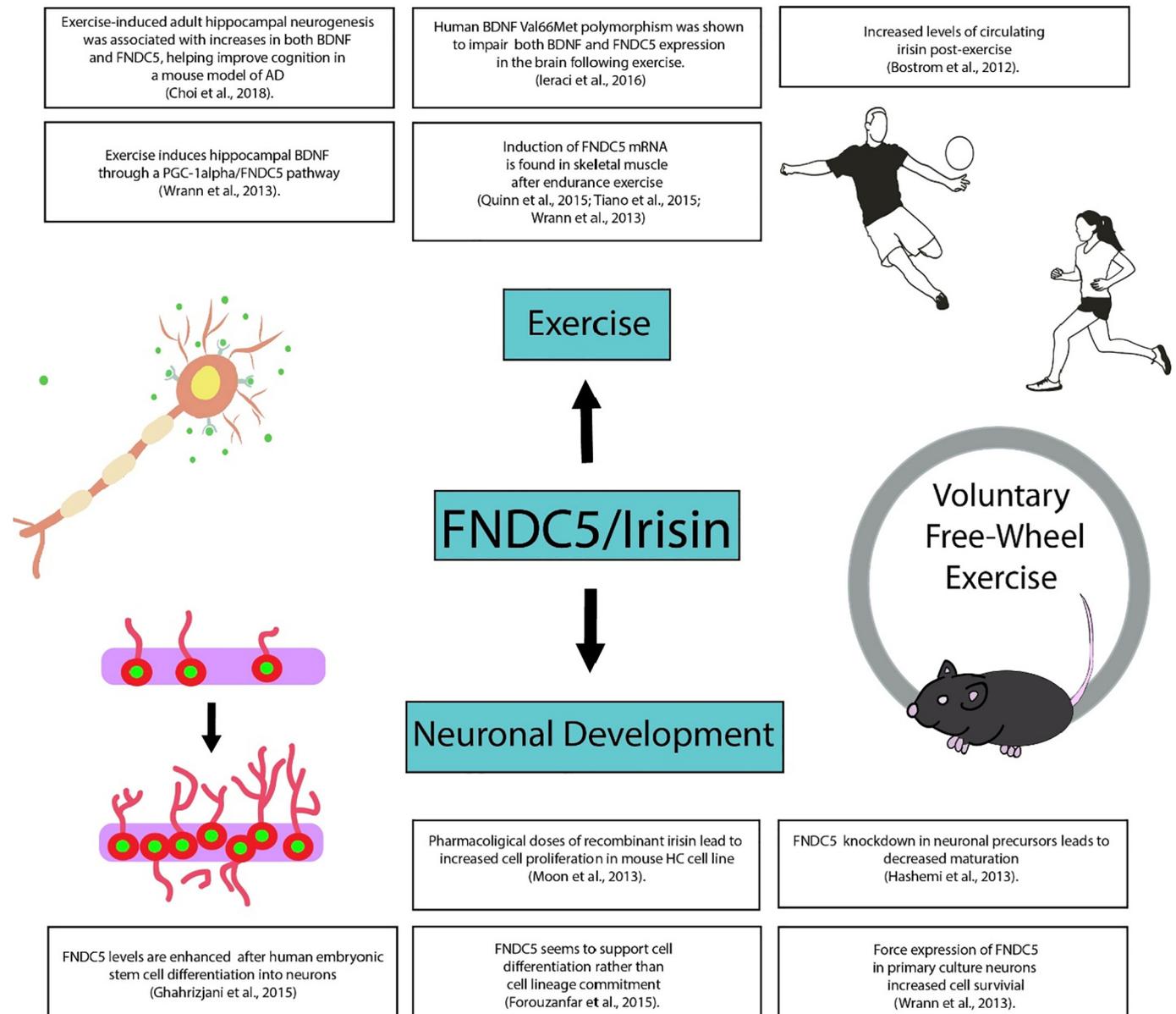
**Fig 3.** The effects of both irisin and exercise on neurodegenerative disease Irisin has been shown to have beneficial effects in a variety of neurodegenerative diseases.

Previous studies have shown that administration of recombinant BDNF in AD and HD animal models improves cognitive function and disease pathogenesis.<sup>66–68</sup> Choi et al. demonstrated that exercise provided cognitive benefit to 5 × FAD mice, a mouse model of AD, by inducing adult hippocampal neurogenesis (AHN) and elevating levels of BDNF and FNDC5. They successfully mimicked the beneficial effects of exercise on AD mice by genetically and pharmacologically inducing AHN in combination with elevating BDNF levels.<sup>57</sup> However, directly increasing levels of hippocampal IL-6 or FNDC5 via lentivirus failed to improve cognition or increase BDNF in 5 × FADProAHN mice (5x FAD injected with P7C3 and lentiviral Wnt3 to promote adult hippocampal neurogenesis). Recently, Lourenco et al. published that by delivering FNDC5 adenovirus either by intracerebroventricular injection or tail vein injection they were able to rescue memory impairments and synaptic plasticity in a mouse model of AD.<sup>69</sup> Interestingly, an in vitro study carried out by Wang et al. illustrated the potential neuroprotective effects of irisin in a primary neuron culture following a β-amyloid peptide (Aβ) insult. Notably, the effects of irisin were not observed by direct treatment of the neuronal cell culture with irisin and Aβ peptide<sup>25–35</sup>; but only

with the treatment of conditioned medium from astrocyte culture treated with irisin and Aβ peptide. Following treatment with astrocyte conditioned medium, there was significant improvement in neuronal cell viability after Aβ peptide exposure. The astrocyte cultures showed a reduction in inflammatory cytokine production of IL-6, IL-1β and COX-2 via the Akt/NFκB pathway upon Aβ peptide exposure. Another study has illustrated increased levels of glucose transport and phosphorylation by the AMPK pathway in astrocyte cell culture following treatment of recombinant irisin.<sup>70</sup> These findings indicate that irisin may be mediating neuroprotective effects via astrocytes. Mitochondrial dysfunction has been implicated with multiple neurodegenerative diseases such as PD.<sup>71</sup> This link between the metabolic effects of irisin and its effects on the brain warrant further investigation for potential therapeutic applications of irisin in the future.

**The future of FNDC5/irisin, PA/exercise, and brain health**

A large amount of interest has been generated by the myokine irisin in recent years; a simple PubMed database search of the keyword



**Fig 4.** The role of FNDC5/irisin in exercise and the brain. Exercise has been shown to be a strong stimulus for the upregulation of FNDC5/irisin. In addition to the metabolic benefits of FNDC5/irisin, it has also been shown to play a role in neuronal differentiation and neuroprotection in the brain.

“irisin” yields over 700 peer-reviewed articles. Although irisin is a myokine secreted from skeletal muscle, it has a large impact on varying regions of the body ranging from lipid metabolism, thermogenesis and browning of white adipose tissue to bone resorption as well as neuronal differentiation and neuroprotection in the brain (Fig 4). As previously discussed, aerobic exercise has been shown to have the greatest impact upon cognitive function in the aging population and in neurodegenerative diseases. Although more research is necessary to determine whether the FNDC5/irisin protein can improve cognitive function in animals, our prior studies suggest that a hormone administered peripherally could induce some of the effects of endurance exercise on the brain. The future for FNDC5/irisin is very exciting, and we look forward to the potential therapeutic uses of this myokine and exercise in ameliorating cognitive deficits associated with neurodegenerative disease.

### Statement of conflict of interest

C.D.W. is a holder of US Patent Application, WO 2015051007 A1.

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